Modular Physiologic Monitoring Systems, Kits, and Methods

CROSS REFERENCE TO RELATED APPLICATION

[0001] The present application is an international application which claims benefit of and priority to U.S. Provisional Application Ser. No. 62/032,515, filed on August 1, 2014 and U.S. Provisional Application Ser. No. 62/032,565, filed on August 2, 2014, each entitled "Modular Physiological Monitoring Systems, Kits, and Methods," by Landy Toth et al., the entire contents of both of which are incorporated by reference herein for all purposes.

BACKGROUND

Technical Field

[0002] The present disclosure relates to the field of physiologic monitoring. The disclosure relates to systems and methods for reliable measurement of one or more physiologic parameters of a subject. In particular, the disclosure relates to aspects of systems and methods for unobtrusively monitoring electrophysiological activity and/or related information from an ambulatory subject in an uncontrolled setting.

Background

[0003] As chronic diseases continue to proliferate throughout the world, there is a heightened need to treat such conditions in a cost effective manner. Remote monitoring of patients with cardiovascular diseases (heart failure, post stroke, etc.), diabetes, kidney failure, COPD, obesity, neurological disorders (depression, Alzheimer's disease, migraines, stress disorders, etc.), arthritis, among other ailments, for purposes of treatment or prevention of such diseases may substantially improve patient outcomes.

[0004] Although physiologic monitoring is performed today for a range of purposes, existing technologies are not without shortcomings.

[0005] There is a need to measure physiologic parameters of subjects, reliably, simply, and without cables. As the proliferation of mobile and remote medicine

increases, simplified and unobtrusive means for monitoring the physiologic parameters of a patient become more important.

[0006] Patient compliance is critical to the success of such systems and is often directly correlated to the ease of use and unobtrusiveness of the monitoring solution used.

[0007] Existing monitoring systems are often prone to false alarms, usage related failures, unreliable user interfaces, cumbersome interfaces, artifact or electromagnetic interference (EMI) related interference, etc. Such problems decrease productivity of using these systems, can result in lost data, and lead to dissatisfaction on the part of both the subject being monitored and the practitioners monitoring the subject. In the case of a hospital setting, the continual drone of alarms can lead to alarm fatigue and decreased productivity.

[0008] Long term compliance of subjects may suffer due to uncomfortable interfaces with monitoring devices, involved maintenance or change-over of disposables, painful or itchy reactions to materials in the devices, and the like.

[0009] More reliable, redundant, and user friendly systems are needed that can provide valuable patient data even when operating with limited supervision, expert input, or user manipulation.

SUMMARY

[0010] One illustrative, non-limiting objective of this disclosure is to provide systems, devices, methods, and kits for monitoring physiologic and/or physical signals from a subject. Another illustrative, non-limiting objective is to provide simplified systems for monitoring subjects. Another illustrative, non-limiting objective is to provide comfortable long term wearable systems for monitoring subjects. Yet another illustrative, non-limiting objective is to provide systems for facilitating interaction between a user and a subject with regard to physiologic monitoring of the subject.

[0011] The above illustrative, non-limiting objectives are wholly or partially met by devices, systems, and methods according to the appended claims in accordance with the present disclosure. Features and aspects are set forth in the appended claims, in the following description, and in the annexed drawings in accordance with the present disclosure.

[0012] According to a first aspect there is provided a system for monitoring one or more physiologic and/or physical signals from a subject including one or more patches each in accordance with the present disclosure configured for attachment to the subject, and one or more modules each in accordance with the present disclosure configured and dimensioned to mate with a corresponding patch, and to interface with the subject there through. In aspects, one or more of the modules may be configured to convey and/or store one or more physiologic and/or physical signals, a signal derived therefrom, and/or a metric derived therefrom obtained via the interface with the subject.

[0013] In aspects, the system may include or interface with a host device in accordance with the present disclosure coupled in wireless communication with one or more of the modules configured to receive one or more of the signals and/or metrics therefrom. In aspects, the host device may include features for recharging and/or performing diagnostic tests on one or more of the modules.

[0014] According to aspects there is provided, use of a system in accordance with the present disclosure to monitor a subject, to monitor an electrocardiogram of a subject, to perform one or more tasks in accordance with the present disclosure, etc.

[0015] According to aspects there is provided an interface (i.e. a patch in accordance with the present disclosure) for monitoring a physiologic and/or physical signal from a subject, including a substrate, an adhesive coupled to the substrate formulated for attachment to the skin of a subject, and one or more sensors and/or electrodes each in accordance with the present disclosure coupled to the substrate, arranged, configured, and dimensioned to interface with the subject.

[0016] In aspects, the substrate may be formed from an elastic or polymeric material and the patch is configured to maintain operation when stretched to more than 25%, more than 50%, or more than 80%.

[0017] In aspects, the interface (i.e. the patch) may be configured with a moisture vapor transmission rate of between 200 grams/square meter (g/m^2)/24 hours (hrs) and 20,000g/m^2/24hrs, between 500g/m^2/24hrs and 12,000g/m^2/24hrs, or between 2,000g/m^2/24hrs and 8,000g/m^2/24hrs, etc.

[0018] In aspects, the interface may be used for a range of applications, some non-limiting examples of which include electrocardiography, sleep assessment,

bruxism assessment, sleep apnea, traumatic brain injury, black box event based monitoring (e.g. for syncope, atrial fibrillation, etc.), biofeedback, stress management, relaxation, physiotherapy, stroke or surgical recovery, or the like. Additional uses and details thereof are described throughout the present disclosure.

[0019] According to aspects there is provided a device (i.e. a module in accordance with the present disclosure) for monitoring a physiologic, physical, and/or electrophysiological signal from a subject including, a housing, a printed circuit board (PCB) including one or more microcircuits, and an interconnect configured for placement of the device onto a subject interface (i.e. a patch in accordance with the present disclosure).

[0020] In aspects, the printed circuit board may constitute at least a portion of the housing.

[0021] In aspects, the device may include a three dimensional antenna coupled to the microcircuits (i.e. coupled with a transceiver, transmitter, radio, etc. included within the microcircuits). In aspects, the antenna may be printed onto or embedded into the housing.

[0022] According to aspects there is provided a kit for monitoring a physiologic, physical, and/or electrophysiological signal from a subject, including one or more patches in accordance with the present disclosure, one or more modules in accordance with the present disclosure; a recharging bay in accordance with the present disclosure, and one or more accessories in accordance with the present disclosure.

[0023] In aspects, one or more of the accessories may include an adhesive removing agent configured to facilitate substantially pain free removal of one or more of the patches from a subject.

[0024] According to aspects there is provided, a service system for managing the collection of physiologic data from a customer, including a customer data management service, configure to generate and/or store the customer profile referencing customer preferences, data sets, and/or monitoring sessions, an automated product delivery service configured to provide the customer with one or more monitoring products or supplies in accordance with the present disclosure, and a datacenter configured to store, analyze, and/or manage the data obtained from the customer during one or more monitoring sessions.

[0025] In aspects, the service system may include a report generating service configured to generate one or more monitoring reports based upon the data obtained during one or more monitoring sessions, a report generating service coupled to the datacenter configured to generate one or more monitoring reports based upon the data obtained during one or more monitoring sessions, and/or a recurrent billing system configured to bill the customer based upon the number or patches consumed, the data stored, and/or the reports generated throughout the course of one or more monitoring session.

[0026] According to aspects there is provided a method for monitoring a physiologic, physical, and/or electrophysiological signal from a subject, including one or more steps in accordance with the present disclosure.

[0027] In aspects, one or more of the steps may be performed at least in part by a system in accordance with the present disclosure.

[0028] According to aspects there is provided, an isolating patch for providing a barrier between a handheld monitoring device with a plurality of contact pads and a subject, including a flexible substrate with two surfaces, a patient facing surface and an opposing surface, and an electrically and/or ionically conducting adhesive coupled to at least a portion of the patient facing surface configured so as to electrically and mechanically couple with the subject when placed thereupon, wherein the conducting adhesive is exposed within one or more regions of the opposing surface of the substrate, the regions patterned so as to substantially match the dimensions and layout of the contact pads.

[0029] In aspects, the conducting adhesive may include an anisotropically conducting adhesive, with the direction of conduction oriented substantially normal to the surfaces of the substrate.

[0030] According to aspects there is provided, a patch interface (i.e. a patch in accordance with the present disclosure) for monitoring one or more physiologic and/or electrophysiological signals from a subject, including a substrate, an adhesive coupled to the substrate formulated for attachment to the skin of a subject, an interconnect embedded into the substrate for attachment of the patch to a microcircuit, and one or more sensors and/or electrodes attached to or embedded onto the surface of

the substrate, the sensors and/or electrodes arranged, configured, and dimensioned to interface with the subject when the adhesive is attached thereto.

[0031] In aspects, the adhesive may be patterned onto the substrate so as to form one or more exposed regions of the substrate, one or more of the sensors and/or electrodes arranged within the exposed regions. One or more of the electrodes may include an inherently or ionically conducting gel adhesive.

[0032] In aspects, one or more of the electrode may include an electrode feature arranged so as to improve the electrical connection between the electrode and the skin upon placement on a subject. In aspects, the improved electrical connection may be achieved after pressure is applied to the electrode (i.e. after the patch is secured to the subject and then a pressure is applied to the electrode). The electrode feature may include one or more microfibers, barbs, microneedles, or spikes to penetrate into a stratum corneum of the skin. The electrode feature may be configured to penetrate less than 2mm into the skin, less than 1mm, less than 0.5mm, less than 0.2mm, or the like during engagement therewith. In aspects, a gel adhesive in accordance with the present disclosure located adjacent to the electrode features (i.e. between the features and the skin) may be configured to maintain the improved electrical connection to the skin for more than 1hr, more than 1day, or more than 3days after the electrode contacts the skin or pressure is applied to the electrode.

[0033] In aspects, a patch interface in accordance with the present disclosure may include one or more stretchable electrically conducting traces attached to the substrate, arranged so as to coupled one or more of the sensors and/or electrodes with one or more of the interconnects.

[0034] In aspects, the interconnect may include a plurality of connectors, the connectors physically connected to each other through the substrate. The patch may include an isolating region arranged so as to isolate one or more of the connectors from the skin while the patch is engaged therewith.

[0035] In aspects, the patch interface may be sufficiently physically frail such that it cannot retain a predetermined shape in a free standing state. The patch interface may include a temporary stiffening member attached to the substrate, the temporary stiffening member configured to provide retention of the shape of the patch interface prior to attachment to the subject, the stiffening member being removable from the

substrate after attachment to the subject. In aspects, after removal of the stiffening member, the retention of the shape of the patch interface may be provided by the skin of the subject. Removal of the patch interface from the skin of the subject may result in a permanent loss in shape of the patch interface without tearing of the patch interface. In aspects, the interconnect may be sufficiently frail such that removal of the patch interface from the skin of the subject may result in a permanent loss of shape of the interconnect.

[0036] In aspects, an adhesive in accordance with the present disclosure may have a peel tack to mammalian skin of greater than 0.02 Newton/millimeter (N/mm), greater than 0.1N/mm, greater than 0.25N/mm, greater than 0.50N/mm, greater than 0.75N/mm, or the like. The patch interface may have a tear strength of greater than 0.5N/mm, greater than 1N/mm, greater than 2N/mm, greater than 8N/mm, or the like.

[0037] In aspects, a patch interface in accordance with the present disclosure may have a ratio between the tear strength of the patch and the peel tack of the adhesive to mammalian skin is greater than 8:1, greater than 4:1, greater than 2:1, or the like. In aspects, the substrate may be formed from a soft pseudo-elastic material and the patch interface may be configured to maintain operation when stretched to more than 25%, more than 50%, more than 80%, or the like. In aspects, the patch interface may be configured with a moisture vapor transmission rate of between 200g/m^2/24hrs and 20,000g/m^2/24hrs, between 500g/m^2/24hrs and 12,000g/m^2/24hrs, between 1,000g/m^2/24hrs and 8,000g/m^2/24hrs, or the like.

[0038] According to aspects, there is provided a module for monitoring one or more physiologic and/or electrophysiological signals from a subject, including a housing, a circuit board including one or more microcircuits, and a module interconnect coupled to one or more of the microcircuits configured for placement and coupling of the device onto a patch interface in accordance with the present disclosure.

[0039] In aspects, the module interconnect may be embedded into the circuit board, and/or the circuit board may constitute at least a portion of the housing. The module may include a three dimensional antenna in accordance with the present disclosure, the antenna coupled to one or more of the microcircuits, the microcircuits including a transceiver or transmitter coupled to the antenna. In aspects, the antenna

may be printed on an interior wall of or embedded into the housing, the circuit board providing a ground plane for the antenna. In aspects, the housing may be shaped like a dome and the antenna may be patterned into a spiraling helix centered within the dome.

[0040] In aspects, a module in accordance with the present disclosure may include a sensor coupled with one or more of the microcircuits, the sensor configured to interface with the subject upon attachment of the module to the patch interface. The module may include a sensor and/or microelectronics configured to interface with a sensor included on a corresponding patch interface. In aspects, one or more of the sensors may include an electrophysiologic sensor, a temperature sensor, a thermal gradient sensor, a barometer, an altimeter, an accelerometer, a gyroscope, a humidity sensor, a magnetometer, an inclinometer, an oximeter, a colorimetric monitor, a sweat analyte sensor, a galvanic skin response sensor, an interfacial pressure sensor, a flow sensor, a stretch sensor, a microphone, a combination thereof, or the like.

[0041] In aspects, the module may be hermetically sealed. The module and/or patch interface may include a gasket coupled to the circuit board or the substrate, the gasket formed so as to isolate the region formed by the module interconnect and the patch from a surrounding environment, when the module is coupled with the patch.

[0042] According to aspects there is provided, a device for monitoring one or more physiologic and/or electrophysiologic signals from a subject including a patch interface in accordance with the present disclosure; and a module in accordance with the present disclosure. In aspects, the module interconnect included within the module may be sized and dimensioned to interface with a corresponding interconnect included within the patch interface, wherein to form an operable interconnection between the patch interface and the module, the patch interface may first be coupled to the subject (i.e. so as to maintain the shape thereof during the process of coupling the patch interface to a corresponding module).

[0043] In aspects, the module interconnect ay include an electrically conducting magnetic element, and the patch interface may include one or more ferromagnetic regions coupled to the substrate, the magnetic elements arranged so as to physically and/or electrically couple the module to the patch interface when the magnetic elements are aligned with the ferromagnetic regions. In aspects, the ferromagnetic

regions may be formed from stretchable pseudo elastic material and/or may be printed onto the substrate. In aspects, the module and/or the patch interface may include one or more fiducial markings to visually assist with the alignment of the module to the patch during coupling thereof.

[0044] According to aspects there is provided, a system for monitoring one or more physiologic and/or electrophysiological signals from a subject including a patch interface in accordance with the present disclosure configured for attachment to the subject, and a module in accordance with the present disclosure configured and dimensioned to mate with the patch, and to interface with the subject there through, the module configured to convey and/or store one or more physiologic, electrophysiological, and/or physical signals, a signal derived therefrom, and/or a metric derived therefrom obtained via the interface with the subject.

[0045] In aspects, the system may include a host device coupled in wireless communication or physical communication with the module, configured to receive one or more of the signals and/or metrics therefrom. In aspects, the host device may include one or more features for recharging and/or performing diagnostic tests on one or more of the modules. In aspects, the system may include a plurality of modules, the modules being hot swappable with the patch interface, so as to maintain a nearly continuous or continuous operation thereof.

[0046] In aspects, the system may include a plurality of modules and associated patch interfaces for placement onto a signal subject, the host device, and/or one or more of the modules configured to coordinate synchronous monitoring of the signals amongst the modules on the subject. In aspects, a host device in accordance with the present disclosure may be integrated into a bedside alarm clock, housed in an accessory, within a purse, a backpack, a wallet, is or is included in a mobile computing device, a smartphone, a tablet computer, a pager, a laptop, a local router, a data recorder, a network hub, a server, a secondary mobile computing device, a repeater, a combination thereof, or the like.

[0047] According to aspects there is provided, use of a device, a module, a patch, and/or a system each in accordance with the present disclosure to monitor an electrocardiogram of a subject.

[0048] According to aspects there is provided, a method for monitoring one or more physiologic and/or electrophysiological signals from a subject, including attaching one or more soft breathable and hypoallergenic devices to one or more sites on the subject, obtaining one or more local physiologic and/or electrophysiological signals each of the devices, and analyzing the signals obtained from each of the devices to generate a metric, diagnostic, report, and/or additional signals therefrom.

[0049] In aspects, the method may include hot swapping one or more of the devices without interrupting the step of obtaining, and/or calibrating one or more of the devices while on the subject. In aspects, the step of calibrating may be performed with an additional medical device (e.g. a blood pressure cuff, a thermometer, a pulse oximeter, a cardiopulmonary assessment system, a clinical grade electrocardiogram (EKG) diagnostic system, etc.).

[0050] In aspects, the method may include determining the position and/or orientation of one or more of the devices on the subject, and/or determining the position and/or orientation from a photograph, a video, or a surveillance video.

[0051] In aspects, one or more steps of a method in accordance with the present disclosure may be performed at least in part by a device, patch interface, module, and/or system each in accordance with the present disclosure.

[0052] According to aspects there is provided, an isolating patch for providing a barrier between a handheld monitoring device with a plurality of contact pads and a subject, including a flexible substrate with two surfaces, a patient facing surface and an opposing surface, and an electrically and/or ionically conducting adhesive coupled to at least a portion of the patient facing surface configured so as to electrically and mechanically couple with the subject when placed thereupon, wherein the conducting adhesive is exposed within one or more regions of the opposing surface of the substrate, the regions patterned so as to substantially match the dimensions and layout of the contact pads.

[0053] In aspects, the conducting adhesive may include an anisotropically conducting adhesive, with the direction of conduction oriented substantially normal to the surfaces of the substrate.

[0054] According to aspects there is provided, a system for measuring blood pressure of a subject in an ambulatory setting including an EKG device in accordance

with the present disclosure (i.e. a patch/module pair in accordance with the present disclosure configured to measure local electrophysiological signals in adjacent tissues), configured for placement onto a torso of the subject, the EKG device configured to measure an electrocardiographic signal from the torso of the subject so as to produce an EKG signal, one or more pulse devices (i.e. patch/module pairs in accordance with the present disclosure configured to measure local blood flow in adjacent tissues) each in accordance with the present disclosure, configured for placement onto one or more sites on one or more extremities of the subject, each of the pulse devices configured to measure a local pulse at the placement site so as to produce one or more pulse signals; and a processor included in or coupled to one or more of the EKG device and the pulse devices, the processor configured to receive the EKG signal, the pulse signals, and/or signals generated therefrom, the processor including an algorithm, the algorithm configured to analyze one or more temporal metrics from the signals in combination with one or more calibration parameters, to determine the blood pressure of the subject.

[0055] In aspects, the system for monitoring blood pressure of a subject may include a blood pressure cuff configured to produce a calibration signal, the processor configured to generate one or more of the calibration parameters, from the calibration signal in combination with the EKG signal, and pulse signals.

[0056] In aspects, one or more of the devices may include an orientation sensor, the orientation sensor configured to obtain an orientation signal, the processor configured to receive the orientation signal or a signal generated therefrom, and to incorporate the orientation signal into the analysis. Some non-limiting examples of orientation sensors include one or more of an altimeter, a barometer, a tilt sensor, a gyroscope, combinations thereof, or the like.

[0057] A system for measuring the effect of an impact on physiologic state of a subject including an electroencephalogram (EEG) device (i.e. a patch/module pair in accordance with the present disclosure configured to measure local electrophysiological signals associated with brain activity in adjacent tissues) in accordance with the present disclosure, configured for placement behind an ear, on the forehead, near a temple, onto the neck of the subject, or the like, the EEG device configured to measure an electroencephalographic signal from the head of the subject so as to produce an EEG signal, and configured to measure one or more kinetic and/or

kinematic signals from the head of the subject so as to produce an impact signal, and a processor included in or coupled to the EEG device, the processor configured to receive the EEG signal, the impact signals, and/or signals generated therefrom, the processor including an algorithm, the algorithm configured to analyze the impact signals to determine if the subject has suffered an impact, to separate the signals into pre impact and post impact portions and to compare the pre and post impact portions of the EEG signal, to determine the effect of the impact on the subject.

[0058] In aspects, the EEG device may include additional sensors such as a temperature sensor configured to generate a temperature signal from the subject or a signal generated therefrom, the processor configured to receive the temperature signal and to assess a thermal state of the subject therefrom. In aspects, the EEG device may include a hydration sensor configured to generate a fluid level signal from the subject, the processor configured to receive the fluid level signal or a signal generated therefrom, and to assess the hydration state of the subject therefrom.

[0059] In aspects, the EEG device and/or the processor may include or be coupled to a memory element, the memory element including sufficiently large space to store the signals for a period of 3 minutes, 10 minutes, 30 minutes, or 1 hr.

[0060] In aspects, the system for measuring the effect of an impact on physiologic state of a subject may include an EKG device (i.e. a patch/module pair in accordance with the present disclosure configured to measure local electrophysiological signals in adjacent tissues) in accordance with the present disclosure, the EKG device configured for placement onto the torso or neck of the subject, the EKG device configured to measure an electrophysiological signal pertaining to cardiac function of the subject so as to produce an EKG signal, the processor configured to receive the EKG signal or a signal generated therefrom, the algorithm configured so as to incorporate the EKG signal into the assessment. In aspects, the processor may be configured to extract a heart rate variability (HRV) signal from the EKG signal, a pre impact and post impact portion of the HRV signal compared to determine at least a portion of the effect of the impact.

[0061] According to aspects there is provided, a system for assessing a sleep state of a subject including an electromyography (EMG)/ electrooculography (EOG) device (i.e. a patch/module pair in accordance with the present disclosure configured to

measure local electromyographic and/or electrooculographic signals from adjacent tissues), in accordance with the present disclosure, configured for placement behind an ear, on a forehead, substantially around an eye, near a temple, or onto a neck of the subject, the EMG/EOG device configured to measure one or more electromyographic and/or electrooculographic signals from the head or neck of the subject so as to produce an EMG/EOG signal, and a processor included in or coupled to the EMG/EOG device, the processor configured to receive the EMG/EOG signal, and/or signals generated therefrom, the processor including an algorithm, the algorithm configured to analyze EMG/EOG signal, to determine the sleep state of the subject.

[0062] In aspects, the EMG/EOG device may include a microphone, the microphone configured to obtain an acoustic signal from the subject, the processor configured to receive the acoustic signal or a signal generated therefrom, the algorithm configured so as to incorporate the acoustic signal into the assessment.

[0063] In aspects, the system may include a sensor for evaluating oxygen saturation (sPO2) at one or more sites on the subject to obtain an oxygen saturation signal from the subject, the processor configured to receive the oxygen saturation signal or a signal generated therefrom, the algorithm configured so as to incorporate the oxygen saturation signal into the assessment.

[0064] In aspects, the processor may include a signal analysis function, the signal analysis function configured to analyze the EMG/EOG signals, the acoustic signal, and/or the oxygen saturation signal to determine the sleep state of the subject, identify snoring, identify a sleep apnea event, identify a bruxism event, identify a rapid eye movement (REM) sleep state, identify a sleep walking state, a sleep talking state, a nightmare, or identify a waking event. In aspects, the system may include a feedback mechanism, configured to interact with the subject, a user, a doctor, a nurse, a partner, a combination thereof, or the like. The processor may be configured to provide a feedback signal to the feedback mechanism based upon the analysis of the sleep state of the subject. The feedback mechanism may include a transducer, a loudspeaker, tactile actuator, a visual feedback means, a light source, a buzzer, a combination thereof, or the like to interact with the subject, the user, the doctor, the nurse, the partner, or the like.

[0065] According to aspects there is provided, a system for assessing a gait and/or a muscle movement of a subject including an EMG device (i.e. a patch/module pair in accordance with the present disclosure configured to measure local electromyographic signals from adjacent tissues) in accordance with the present disclosure, configured for placement over a muscle group on the subject, the EMG device configured to measure one or more electromyographic signals from the muscle group of the subject so as to produce an EMG signal, and a processor included in or coupled to the EMG device, the processor configured to receive the EMG signal, and/or signals generated therefrom, the processor including an algorithm, the algorithm configured to analyze EMG signal to determine at least a portion of the gait and/or the muscle movement of the subject.

[0066] In aspects, the system may include a plurality of EMG devices, each EMG device configured to monitor a separate muscle group on the subject, the processor configured to synchronize and analyze the EMG signals received from each EMG device to determine at least a portion of the gait and/or the muscle movement of the subject.

[0067] In aspects, one or more of the EMG devices may include an orientation, kinetic, kinematic, and/or proprioception sensor each in accordance with the present disclosure, configured so as to generate a kinematic signal, the processor configured to incorporate the kinematic signal into the analysis. In aspects, the processor may be configured to analyze one or more of the EMG signals to generate a muscle exertion metric.

[0068] According to aspects there is provided, a patch interface for monitoring neural activity from the skin of a subject, including a substrate with a surface; an adhesive coupled to the substrate formulated for attachment to the skin of a subject; an interconnect embedded into or coupled to the substrate for attachment of the patch to a microcircuit; and a plurality microelectrodes attached to or embedded into the surface of the substrate, the microelectrodes arranged, configured, and dimensioned to interface with the subject when the adhesive is attached thereto, the microelectrodes electrically coupled with the interconnect and/or the microcircuit.

[0069] In aspects, the adhesive may be patterned onto the substrate so as to form one or more exposed regions of the substrate, one or more of the microelectrodes

arranged within the exposed regions, the adhesive may include an electrically or ionically conducting component, and/or the adhesive may provide a macroelectrode function for interfacing with the subject when attached thereto, the macroelectrode electrically coupled to the interconnect and/or the microcircuit. Such a macroelectrode may provide a guard, reference, or counter electrode function or the like for one or more of the microelectrodes.

[0070] In aspects, one or more of the microelectrodes may be configured to penetrate into the skin when the adhesive is attached to the subject. One or more of the microelectrodes may have a needle-like shape, the microelectrode including an insulated region and an exposed region (so as to interface with local tissues). The microelectrode may have a length such that the exposed region is positioned within the epidermal layer, the dermal layer, and/or the hypodermal layer of the skin upon attachment of the patch to the surface thereof. In aspects, one or more of the microelectrodes may be shaped so as to penetrate down a root of a follicle, to reach a sweat gland, a sebaceous gland, a muscle, a vessel, an arrector pili follicular muscle, or the like upon attachment of the patch interface to the skin.

[0071] In aspects, one or more of the microelectrodes may include an electrode feature in accordance with the present disclosure arranged so as to improve the electrical connection between the microelectrode or an exposed region thereof and the skin upon placement on a subject. In aspects, the improved electrical connection may be achieved after pressure is applied to the microelectrode.

[0072] In aspects, the microcircuit may include a switch bank and an amplifier with two or more inputs, the switch bank adaptably configurable to couple one or more of the microelectrodes with an input of the amplifier, and one or more alternative microelectrodes and/or the macroelectrode to another input of the amplifier. The switch bank may be adaptively configurable to connect one or more clusters of microelectrodes to a single input of the amplifier.

[0073] In aspects, one or more of the microelectrodes may be sized, dimensioned, oriented, configured and shaped so as to penetrate less than 3mm, less than 2mm into the skin, less than 1 millimeter (mm), less than 0.5mm, less than 0.2mm, or the like into the skin during engagement therewith. One or more of the microelectrodes and/or exposed regions thereof may have an active surface area of less than $4mm^2$,

less than 1mm^2, less than 100,000um^2, less than 10,000um^2, less than 1,000um^2, less than 100um^2, or the like.

[0074] In aspects, one or more of the macroelectrodes may have an active surface area of greater 5mm^2, greater than 50mm^2, greater than 100mm^2, greater than 500mm^2, or the like.

[0075] In aspects, the macroelectrode may be shaped and/or patterned on the substrate so as to reduce movement artifacts, monitor a macroscopic electrophysiologic field, provide a reference electrode function, provide a guard electrode function, provide a drive electrode function, a combination thereof, or the like during use.

[0076] In aspects, the patch interface may include a sensor coupled with the substrate, and electrically coupled with the microcircuit, the sensor configured to monitor one or more physiologic parameters of the subject when coupled thereto. Some non-limiting sensor components include an electrophysiologic sensor, a temperature sensor, a thermal gradient sensor, a barometer, an altimeter, an accelerometer, a gyroscope, a humidity sensor, a magnetometer, an inclinometer, an oximeter, a colorimetric monitor, a sweat analyte sensor, a perfusion sensor, a galvanic skin response sensor, a hydration sensor, an interfacial pressure sensor, a flow sensor, a stretch sensor, a microphone, a combination thereof, and the like.

[0077] In aspects, the patch interface may include a stimulating device selected from an electrical stimulator, a thermoregulating device, a heating coil, a thermoelectric device, a Peltier device, a tactile stimulating component, a vibratory stimulating element, a combination thereof, or the like arranged along the substrate so as to interface with the skin of the subject when the adhesive is attached thereto. In aspects, the patch interface and/or stimulating device may include a feedback sensor coupled with the stimulating device, configured and arranged so as to monitor the interaction of the stimulating device with the skin during activation thereof.

[0078] In aspects, the patch interface may include one or more stretchable electrically conducting traces attached to the substrate, arranged so as to couple one or more of the sensors, microelectrodes, and/or macroelectrodes with the interconnect or microcircuit. In aspects, the patch interface may be sufficiently physically frail such that it cannot retain a predetermined shape in a free standing state.

[0079] In aspects, the patch interface may include a temporary stiffening member attached to the substrate, the temporary stiffening member configured to provide retention of the shape of the patch interface prior to attachment to the subject, the stiffening member being removable from the substrate after attachment to the subject. In aspects, after removal of the stiffening member, the retention of the shape of the patch interface may be provided by the skin of the subject, and/or removal of the patch interface from the skin of the subject may result in a permanent loss in the shape of the patch interface without tearing of the patch interface.

[0080] In aspects, the patch interface may be constructed it is sufficiently frail and that removal of the patch interface from the skin of the subject results in a permanent loss of shape of the interconnect. Such a configuration may be advantageous in balancing frailty of the freely supported patch interface against substantially maximizing comfort for the subject during use. In aspects, the adhesive may have a peel tack to mammalian skin of greater than 0.02N/mm, greater than 0.1N/mm, greater than 0.25N/mm, greater than 0.50N/mm, greater than 0.75N/mm, or the like. In aspects, the patch may have a tear strength of greater than 0.5N/mm, greater than 1N/mm, greater than 2N/mm, greater than 8N/mm, or the like. In aspects, the ratio between the tear strength of the patch and the peel tack of the adhesive to mammalian skin may be greater than 8:1, greater than 4:1, greater than 2:1, or the like. In aspects, the substrate may be formed from a soft pseudo-elastic material and the patch interface may be configured to maintain operation when stretched to more than 25%, more than 50%, more than 80%, or the like. In aspects, the patch interface may be configured with a <through thickness> moisture vapor transmission rate of between 200g/m^2/24hrs and 20,000g/m^2/24hrs, between 500g/m^2/24hrs and 12,000g/m^2/24hrs, between 1,000g/m^2/24hrs and 8,000g/m^2/24hrs, or the like.

[0081] In aspects, the adhesive may include anendothermic-reactive specie, an exothermic-reactive specie, a local analgesic, a neuro-blocker, a neurotoxin, a neural antagonist, and/or a neural agonist. Such components may be advantageous for affected local neural traffic at a monitoring site on a body with a high degree of specificity.

[0082] According to aspects there is provided, a module for monitoring one or more physiologic, physical, skin based neural activity, and/or electrophysiological signals from a subject, including a housing; a circuit board including one or more

microcircuits, the circuit board coupled to the housing; and a module interconnect coupled to one or more of the microcircuits configured for placement and coupling of the module onto a patch interface in accordance with the present disclosure.

[0083] In aspects, the module interconnect may be embedded into the circuit board, and/or the circuit board may constitute at least a portion of the housing. In aspects, a three dimensional antenna may be coupled to one or more of the microcircuits, the microcircuits including a transceiver or transmitter coupled to the antenna, and/or the antenna may be printed on an interior wall of or embedded into the housing, the circuit board providing a ground plane for the antenna.

[0084] In aspects, the housing may be shaped like a dome and the antenna may be patterned into a spiraling helix centered within the dome.

[0085] In aspects, the module may include a sensor coupled with one or more of the microcircuits, the sensor configured to interface with the subject upon attachment of the module to the patch interface.

[0086] In aspects, the module may include a sensor and/or microelectronics configured to interface with a sensor included on the patch interface, one or more of the sensors including an electrophysiologic sensor, a temperature sensor, a thermal gradient sensor, a barometer, an altimeter, an accelerometer, a gyroscope, a humidity sensor, a magnetometer, an inclinometer, an oximeter, a colorimetric monitor, a perfusion sensor, a sweat analyte sensor, a galvanic skin response sensor, an interfacial pressure sensor, a flow sensor, a stretch sensor, a microphone, a combination thereof, or the like.

[0087] In aspects, the module may include a stimulating device selected from an electrical stimulator, a thermoregulating device, a heating coil, a thermoelectric device, a Peltier device, a tactile stimulating component, a vibratory stimulating element, a combination thereof, or the like arranged along the substrate so as to interface with the skin of the subject when coupled to the patch interface.

[0088] In aspects, the module may be hermetically sealed and/or the module may include a gasket coupled to the circuit board, the gasket shaped so as to isolate the region formed by the module interconnect and the patch from a surrounding environment, when the module is coupled with the patch.

[0089] According to aspects there is provided, a device for monitoring one or more physiologic, physical, and/or electrophysiologic signals from a subject including a patch interface in accordance with the present disclosure; and a module in accordance with the present disclosure, the module interconnect included within the module, sized and dimensioned to interface with the interconnect included within the patch interface, wherein to form an operable interconnection between the patch interface and the module, the patch interface must first be coupled to the subject.

[0090] In aspects, the module interconnect may include an electrically conducting magnetic element, and the patch interface may include one or more ferromagnetic regions coupled to the substrate, the magnetic elements arranged so as to physically and electrically couple the module to the patch interface when the magnetic elements are aligned with the ferromagnetic regions.

[0091] In aspects, one or more of the ferromagnetic regions may be formed from a stretchable pseudo elastic material and may be printed onto the substrate. In aspects, the module and/or the patch interface may include one or more fiducial markings to visually assist with the alignment of the module to the patch during coupling thereof.

[0092] According to aspects there is provided, a system for monitoring one or more physiologic, physical, and/or electrophysiological signals from a subject including a patch interface in accordance with the present disclosure configured for attachment to the subject; and a module in accordance with the present disclosure configured and dimensioned to mate with the patch, and to interface with the subject there through, the module configured to convey and/or store one or more physiologic, neural activity, and/or physical signals, a signal derived therefrom, and/or a metric derived therefrom obtained via the interface with the subject.

[0093] In aspects, the system may include a host device coupled in wireless communication or physical communication with the module, configured to receive one or more of the signals and/or metrics therefrom.

[0094] In aspects, the host device may include one or more features for recharging and/or performing diagnostic tests on one or more of the modules. In aspects, the system may include a plurality of modules, the modules being hot swappable with the patch interface, so as to maintain a nearly continuous operation thereof.

[0095] In aspects, the system may include a plurality of modules and associated patch interfaces for placement onto a single subject, the host device, and/or one or more of the modules configured to coordinate synchronous monitoring of the signals amongst the modules on the subject. In aspects, the host device may be integrated into a bedside alarm clock, housed in an accessory, within a purse, a backpack, a wallet, is or is included in a mobile computing device, a smartphone, a tablet computer, a pager, a laptop, a local router, a data recorder, a network hub, a server, a secondary mobile computing device, a repeater, a combination thereof, or the like.

[0096] According to aspects there is provided, use of a device in accordance with the present disclosure, and/or a system in accordance with the present disclosure, to confirm completion of, follow up on, confirm partial completion of, monitor a patient response to, or patient selection in connection with, a denervation procedure, a renal denervation procedure, ablation of a renal nerve, ablation of renal artery, or ablation of an accessory renal artery.

[0097] According to aspects there is provided, a method for assessing an autonomic nervous system of a subject, including monitoring neural activity at one or more sites on the skin of the subject to generate one or more signals; performing a stress test on the subject; and analyzing the signals obtained before, during, and/or after the stress test to generate a metric, diagnostic, report, and/or additional signals therefrom relating to the autonomic nervous system of the subject.

[0098] In aspects, the stress test may include administration of a chemical, a drug, medicament, a hormone, an enzyme, a diuretic, a solution, electrolytes, a peptide, steroid, saline, a hypotonic solution, a hypertonic solution, a combination thereof, or the like to the subject. In aspects, the administration may be topical, systemic, intravenous, intra-arterial, intra-parenchymal, sub-dermal delivery, transdermal delivery, rectal, via vaginal suppositories, via urethral suppositories, via nasal suppositories, via rectal suppositories, inhaled, a combination thereof, or the like.

[0099] In aspects, the stress test may include delivery of energy, stimulation of, electrical stimulation of, presenting an audio field to, application of thermal stress to, presenting a light field to, presenting an image to, asking a question to, or playing music for, the subject. In aspects, the stress test may include providing a tactile input to one or more sites on the subject.

[00100] In aspects, the stress test may include stimulating one or more of a carotid sinus, a carotid body, a baroreceptor, a vagus nerve receptor, an erogenous zone, or the like of the subject.

[00101] In aspects, the stress test may include applying an electromagnetic field to, injecting a current into, applying pressure to, applying stroking to, or applying a change in barometric pressure surrounding the subject.

[00102] In aspects, the stress test may include having the subject sleep, cry, laugh, lie down, jump, walk, run, change posture, exercise, perform a breath holding exercise, climb stairs, or the like.

[00103] In aspects, one or more of the sites may be coupled to the perineal region, the perianal region, the pubic region, the inner thigh region, the posterior knee region, the neck, the ear, the ocular region, the breast, the axilla, the elbow, the wrist, the palm, the foot, the lips, an erogenous zone of the subject, a combination thereof, or the like.

[00104] In aspects, one or more of the steps of a method in accordance with the present disclosure may be performed at least in part by a device in accordance with the present disclosure, a system in accordance with the present disclosure, a module in accordance with the present disclosure, and/or a patch interface in accordance with the present disclosure.

[00105] According to aspects there is provided, a system for performing a neuromodulation and/or ablation procedure on a target site within a subject including a treatment system for delivering energy or a chemical to the target site; one or more monitoring devices configured to interface with the skin of the subject at one or more monitoring sites on the subject, each of the monitoring devices configured to measure one or more electrophysiological signals, neural traffic signals, and/or physiologic parameters from the monitoring sites so as to produce an activity signal; and a processor included in or coupled to one or more of the monitoring devices, the processor configured to receive the activity signal(s), and/or one or more signals generated therefrom, the processor including an algorithm, the algorithm configured to analyze the activity signal(s) to determine the effect of the treatment system on the target site.

[00106] In aspects, the algorithm may be configured to compare the activity signal(s) generated by a plurality of monitoring devices against each other to determine the effect of the treatment system on the target site.

[00107] In aspects, the treatment system may include one or more of an ablation system, a neuromodulation device, a neuromodulation implant, an ablation catheter, a focused energy delivery device, a radio frequency ablation system or catheter, a microwave ablation system or catheter, an ultrasound energy delivery system, a high intensity focused ultrasound [HIFU] delivery system or catheter, a cryoablation system or catheter, a chemical ablation system or catheter, a radiosurgical system, an optical ablation system, an infrared ablation system, a laser ablation system, an magnetic resonance (MR) guided HIFU system, or the like.

[00108] In aspects, the algorithm may be configured to indicate when only a partial neuromodulation and/or ablation procedure has been performed on a target site, and/or when a complete procedure has been performed on the target site.

[00109] In aspects, one or more of the monitoring devices may include a chemical agent selected from an endothermic reactive specie, an exothermic reactive specie, a local analgesic, a neuro-blocker, a neurotoxin, or a neural agonist for delivery to the associated monitoring site when coupled thereto, the algorithm configured to compensate for the presence of the chemical agent in the analysis of the activity signal(s).

[00110] In aspects, the system may include a stimulating device selected from an electrical stimulator, a thermoregulating device, a heating coil, a thermoelectric device, a Peltier device, a tactile stimulating component, a vibratory stimulating element, a combination thereof, or the like, the stimulating device configured to stimulate the subject at one or more stimulation sites, the algorithm configured to compensate for the stimulation in the analysis.

[00111] In aspects, one or more of the monitoring devices and/or one or more of the stimulating devices may be embedded into or included in the treatment system.

[00112] In aspects, the treatment system may be configured to deliver a stimulating energy or chemical agent to the target site, and/or one or more stimulatory sites within the subject.

BRIEF DESCRIPTION OF THE DRAWINGS

[00113] Several aspects of the disclosure can be better understood with reference to the following drawings. In the drawings, like reference numerals designate corresponding parts throughout the several views.

[00114] Figs. 1a-d show aspects of modular physiologic monitoring systems in accordance with the present disclosure.

[00115] Figs. 2a-d show aspects of a patch and a corresponding module in accordance with the present disclosure.

[00116] Figs. 3a-f show aspects of patches in accordance with the present disclosure.

[00117] Figs. 4a-f show top, side, and isometric views of a mated patch and module in accordance with the present disclosure.

[00118] Fig. 5 shows a perspective view of aspects of a mated patch and module in accordance with the present disclosure.

[00119] Fig. 6 shows a schematic of aspects of a module in accordance with the present disclosure.

[00120] Fig. 7 shows a schematic of a patch/module pair attached to a subject in accordance with the present disclosure.

[00121] Figs. 8a-e show aspects of patch layouts in accordance with the present disclosure.

[00122] Fig. 9 shows aspects of an impact sensing patch and a feedback component in accordance with the present disclosure.

[00123] Figs. 10a-e show aspects of a patch in accordance with the present disclosure.

[00124] Figs. 11a-b show aspects of patches and modules in accordance with the present disclosure.

[00125] Figs. 12a-c show aspects of patches and modules in accordance with the present disclosure.

[00126] Fig. 13 shows a module configured to apply energy to a subject in accordance with the present disclosure.

[00127] Fig. 14 shows a module for interrogating a subject in accordance with the present disclosure.

[00128] Fig. 15 shows a vibrating module configured to apply a tactile input to a subject in accordance with the present disclosure.

[00129] Figs. 16a-c show arrangements of patches on a subject for generating an EKG in accordance with the present disclosure.

[00130] Figs. 17a-c show aspects of electrode features and methods for engaging such features with skin in accordance with the present disclosure.

[00131] Fig. 18 illustrates an isolation patch in accordance with the present disclosure.

[00132] Fig. 19 shows aspects of a modular physiologic monitoring kit in accordance with the present disclosure.

[00133] Figs. 20a – l illustrate aspects of a multi-sensor patch including a plurality of microelectrodes in accordance with the present disclosure.

[00134] Fig. 21 shows aspects of a circuit schematic to interface with a plurality of microelectrodes in accordance with the present disclosure.

[00135] Figs. 22a – c show aspects of multi-site monitoring, stimulation, stress application, and/or treatments applied to a subject each in accordance with the present disclosure.

[00136] Fig. 23 illustrates aspects of a range of different types of skin, innervation, vasculature, and hair follicles thereof and approaches for interaction therewith in accordance with the present disclosure.

[00137] Figs. 24a – c illustrate perineal region, cutaneous innervation of the perineal skin, and associated patches in accordance with the present disclosure for interaction therewith.

[00138] Figs. 25a – b illustrate cranial and ear cutaneous innervation, and associated patches in accordance with the present disclosure for interaction therewith.

[00139] Figs. 26a – b illustrate lower lumbar extremity and plantar cutaneous innervation, and associated patches in accordance with the present disclosure for interaction therewith.

[00140] Figs. 27a – b illustrate arm and hand cutaneous innervation, and associated patches in accordance with the present disclosure for interaction therewith.

[00141] Figs. 28a – b illustrate torso cutaneous innervation, and associated patches in accordance with the present disclosure for interaction therewith.

[00142] Figs 29 – d, illustrate temporal readings of metrics associated with stress testing and procedures in accordance with the present disclosure.

[00143] Fig. 30 illustrates spatial readings of skin neural activity with a mult-sensor patch in accordance with the present disclosure.

[00144] Figs. 31a – c illustrate aspects of methods for monitoring, stressing, and/or treating one or more regions of a subject each in accordance with the present disclosure.

DETAILED DESCRIPTION

[00145] Particular embodiments of the present disclosure are described herein below with reference to the accompanying drawings; however, the disclosed embodiments are merely examples of the disclosure and may be embodied in various forms. Therefore, specific structural and functional details disclosed herein are not to be interpreted as limiting, but merely as a basis for the claims and as a representative basis for teaching one skilled in the art to variously employ the present disclosure in virtually any appropriately detailed structure. Like reference numerals may refer to similar or identical elements throughout the description of the figures.

[00146] A modular physiologic monitoring system in accordance with the present disclosure for assessing one or more physiologic parameters of a subject (e.g. a human subject, a patient, an athlete, a trainer, an animal, such as equine, canine, porcine, bovine, etc.) with a body may include one or more patches, each patch adapted for attachment to the body of the subject (e.g. attachable to the skin thereof, reversibly attachable, adhesively attachable, with a disposable interface and a reusable module, etc.). In aspects, the physiologic monitoring system may include one or more modules, each module may include a power source (e.g. a battery, a rechargeable battery, an energy harvesting transducer, microcircuit, and an energy reservoir, a thermal gradient harvesting transducer, a kinetic energy harvesting transducer, a radio frequency energy harvesting transducer, a fuel cell, a biofuel cell, etc.), signal

conditioning circuitry, communication circuitry, one or more sensors, or the like, configured to generate one or more signals (i.e. physiologic and/or physical signals).

[00147] One or more of the patches may include one or more interconnects, configured and dimensioned so as to couple with one or more of the modules, said modules including a complimentary interconnect configured and dimensioned to couple with the corresponding patch. The patch may include a bioadhesive interface for attachment to the subject, the module retainable against the subject via interconnection with the patch.

[00148] In aspects, the patch may be configured so as to be single use (i.e. disposable). The patch may include a thin, breathable, stretchable laminate. In aspects, the laminate may include a substrate, a bioadhesive, one or more sensing elements in accordance with the present disclosure, and one or more interconnects for coupling one or more of the sensing elements with a corresponding module.

[00149] In aspects, to retain a high degree of comfort and long term wear-ability of the patch on a subject, to limit interference with normal body function, to limit interference with joint movement, or the like, the patch may be sufficiently thin and frail, such that it may not substantially retain a predetermined shape while free standing. Such a definition is described in further detail below. The patch may be provided with a temporary stiffening film to retain the shape thereof prior to placement of the patch onto the body of a subject. Once adhered to the subject, the temporary stiffening film may be removed from the patch. While the patch is adhered to the subject, the shape and functionality of the patch may be substantially retained. Upon removal of the patch from the subject, the, now freestanding patch is sufficiently frail such that the patch can no-longer substantially retain the predetermined shape (i.e. sufficiently frail such that the patch will not survive in a free standing state). In aspects, stretch applied to the patch while removing the patch from the subject may result in snap back once the patch is in a freestanding state that renders such a patch to crumple into a ball and no longer function.

[00150] In aspects, the patch may include a film (e.g. a substrate), with sufficiently high tear strength, such that, as the patch is peeled from the skin of a subject, the patch does not tear. In aspects, the ratio between the tear strength of the patch and the peel adhesion strength of the patch to skin (i.e. tear strength: peel adhesion strength),

is greater than 8:1, greater than 4:1, greater than 2:1, or the like. Such a configuration may be advantageous so as to ensure the patch may be easily and reliably removed from the subject after use without tearing.

[00151] In aspects, the patch may include a bioadhesive with peel tack to mammalian skin of greater than 0.02N/mm, greater than 0.1N/mm, greater than 0.25N/mm, greater than 0.50N/mm, greater than 0.75N/mm, or the like. Such peel tack may be approximately determined using an American Society for Testing and Materials (ASTM) standard test, ASTM D3330: Standard test method for peel adhesion of pressure-sensitive tape.

[00152] In aspects, the patch may exhibit a tear strength of greater than 0.5N/mm, greater than 1N/mm, greater than 2N/mm, greater than 8N/mm, or the like. Such tear strength may be approximately determined using an ASTM standard test, ASTM D624: Standard test method for tear strength of conventional vulcanized rubber and thermoplastic elastomers.

[00153] In aspects, the patch may be provided with a characteristic thickness, of less than 50 micrometer (um), less than 25um, less than 12um, less than 8um, less than 4um, or the like. Yet, in aspects, a balance between the thickness, stiffness, and tear strength may be obtained so as to maintain sufficiently high comfort levels for a subject, minimizing skin stresses during use (i.e. minimizing skin stretch related discomfort and extraneous signals as the body moves locally around the patch during use), minimizing impact on skin health, minimizing risk of rucking during use, and minimizing risk of maceration to the skin of a subject, while limiting risk of tearing of the patch during removal from a subject, etc.

[00154] In aspects, the properties of the patch may be further altered so as to balance the hydration levels of one or more hydrophilic or amphiphilic components of the patch while attached to a subject. Such adjustment may be advantageous to prevent over hydration or drying of an ionically conducting component of the patch, to manage heat transfer coefficients within one or more elements of the patch, to manage salt retention into a reservoir in accordance with the present disclosure, and/or migration during exercise, to prevent pooling of exudates, sweat, or the like into a fluid measuring sensor incorporated into the patch or associated module, etc. In aspects, the patch or a rate determining component thereof may be configured with a

moisture vapor transmission rate of between 200g/m^2/24hrs and 20,000g/m^2/24hrs, between 500g/m^2/24hrs and 12,000g/m^2/24hrs, between 2,000g/m^2/24hrs and 8,000g/m^2/24hrs, or the like.

[00155] Such a configuration may be advantageous for providing a comfortable wearable physiologic monitor for a subject, while reducing on material waste, cost of goods, preventing contamination or disease spread through uncontrolled re-use, and the like.

[00156] In aspects, one or more patches and/or modules may be configured for electrically conducting interconnection, inductively coupled interconnection, capacitively coupled interconnection, with each other. In the case of an electrically conducting interconnect, each patch and module interconnect may include complimentary electrically conducting connectors, configured and dimensioned so as to mate together upon attachment. In the case of an inductively or capacitively coupled interconnect, the patch and module may include complimentary coils or electrodes respectively, configured and dimensioned so as to mate together upon attachment.

[00157] Each patch or patch/module pair may be configured to monitor one or more local physiologic and/or physical parameters of the attached subject (e.g. local to the site of attachment, etc.), local environment, combinations thereof, or the like, and to relay such information in the form of signals to a host device (e.g. via a wireless connection, via a body area network connection, or the like), one or more patches or modules on the subject, or the like.

[00158] In aspects, the host device may be configured to coordinate information exchange to/from each module and/or patch, and to generate one or more physiologic signals, physical signals, environmental signals, kinetic signals, diagnostic signals, alerts, reports, recommendation signals, commands, combinations thereof, or the like for the subject , a user, a network, an electronic health record (EHR), a database (e.g. as part of a data management center, an EHR, a social network, etc.), a processor, combinations thereof, or the like.

[00159] In aspects, a system in accordance with the present disclosure may include a plurality of substantially similar modules (i.e. generally interchangeable modules, but with unique identifiers), for coupling with a plurality of patches, each patch,

optionally different from the other patches in the system (e.g. potentially including alternative sensors, sensor types, sensor configurations, electrodes, electrode configurations, etc.). Each patch may include an interconnect suitable for attachment to an associated module. Upon attachment of a module to a corresponding patch, the module may validate the type and operation of the patch to which it has been mated. In aspects, the module may then initiate monitoring operations on the subject via the attached patch, communicate with one or more patches on the subject, a hub, etc. The data collection from each module may be coordinated through one or more modules and/or with a host device in accordance with the present disclosure. The modules may report a time stamp along with the data in order to synchronize data collection across multiple patch/module pairs on the subject, between subjects, etc. Thus, if a module is to be replaced, a hot swappable replacement (i.e. replacement during a monitoring procedure) can be carried out easily by the subject, a caregiver, practitioner, etc. during the monitoring process. Such a configuration may be advantageous for performing redundant, continuous monitoring of a subject, and/or to obtain spatially relevant information from a plurality of locations on the subject during use.

[00160] In aspects, the modules and/or patches may include corresponding interconnects for coupling with each other during use. The interconnects may include one or more connectors, configured such that the modules and patches may only couple in a single unique orientation with respect to each other. In aspects, the modules may be color coded by function. A temporary stiffening element attached to a patch may include instructions, corresponding color coding, etc. so as to assist a user or subject with simplifying the process of monitoring.

[00161] The following overviews aspects described herein which may be generally applied to the appended Figures where applicable.

MODULAR MONITORING ASPECTS:

[00162] In aspects, a system in accordance with the present disclosure may include a plurality of patches or patch/module pairs to generate higher level function and/or increased clinically relevant data than may be obtained from a single site on the body. Some non-limiting examples of such applications include multi-electrode electrocardiograms, traumatic brain injury assessment, touch restoration, gait analysis,

cardiorespiratory assessment, metabolic assessment, breath/gait synchronization, blood pressure monitoring, combined reading analysis (i.e. combining a first function such as EEG, with another such as HRV to elucidate deeper understanding of the state of a subject or the extend of a condition or disease state).

[00163] In one non-limiting example, a high level of cardiovascular information may be evaluated from a multi-site EKG diagnostic system (i.e. a 12 lead EKG collected from several individual patches or patch/module pairs). In an application relating to the capture of diagnostic grade EKG data, a plurality of patch/modules each in accordance with the present disclosure may be attached to a subject. A calibration step may be performed to determine the general location of the patch/modules on the body or the relationship between patch/modules on the body during the usage case. In one non-limiting example, an image of the arrangement may be acquired (e.g. by a smartphone camera, a host device, etc.) and analyzed so as to establish the physical layout of patch/modules on the subject. Such an analysis may be performed by comparing the location of each patch/module to each other, and/or to one or more body features (e.g. head, chest, shoulders, waist, etc.), collected from the acquired image.

[00164] In aspects wherein one or more patch/module pairs are equipped with a pulse generator and one or more electrodes suitable for emitting one or more pulses into the subject, a calibration step may include emitting one or more pulses from one or more patch/module pairs and monitoring for evoked potentials at one or more of the additional patch/module pairs. A combination of the timing delay, polarity, and/or amplitude of the received pulses as recorded collectively by the additional patch/module pairs may be used to generate a location metric. The location metric may be used to estimate the location of one or more patch/module pairs on the body of the subject.

[00165] In aspects wherein one or more patch/module pairs are equipped with a plurality of electrophysiological sensing electrodes, a plurality of patch/module pairs may collect electrophysiological information synchronously. A plurality of metrics as collected from the different sites: amplitude, time delay, polarity, ratio between wave components of the signal, movement artifacts, breathing artifacts, etc. may be used to generate a series of location metrics. Such information may be compared against previously collected maps (e.g. generated from studies with correlated camera images

and electrophysiologically collected signals, etc.) and compared against the data collected during a calibration test to determine the location of one or more patch/module pairs.

[00166] In aspects, additional kinematic information may be used to determine and/or refine the location determining aspects of a calibration procedure. One or more patch/module pairs may be equipped with one or more orientation determining sensors, such as one or more accelerometers, barometers, tilt sensors, gyroscopes, combinations thereof, etc. Information gleaned from one or more of such orientation determining sensors may be used in combination with one or more methods in accordance with the present disclosure to determine, enhance, confirm, etc. placement of the patch/module pairs on the subject.

[00167] The analysis may provide suitable information from which relationships between EKG data collected from each patch/module may be coordinated to form a higher level diagnostic function (e.g. such as to reconstruct a 12-lead EKG, etc.). Such a configuration may be advantageous for providing detailed diagnostic information from a subject without requiring precise electrode layout, application of a wired Holter monitor, etc.

[00168] In aspects, a system in accordance with the present disclosure may include a plurality of patches configured to coordinate simultaneous multi-site electrocardiographic signal capture on a subject. Such signal capture may include redundant monitoring of heart rate, regularity of heartbeats, synchronization of heart rhythm with other phenomena, detection of myocardial contraction, detection of P-waves, QRS complexes, ST-segments, and T-wave configurations, calculating/showing/displaying/analyzing the standard electrocardiogram (ECG) lead configurations (Limb leads I,II,III,AVR, AVL, AVF) and precordial leads (V1-V6), combinations thereof, and the like.

[00169] In aspects, a system in accordance with the present disclosure may include a plurality of patches, each patch including one or more electrodes. In aspects, such electrode arrangements may be bipolar, tripolar, quadripolar, or otherwise multi-polar. In one non-limiting example, a single patch may contain one or more electrodes, the one or more electrodes arranged so as to establish a (virtual) reference for the system. During operation, the system may simultaneously monitor signals from the plurality

of electrodes (i.e. via each patch/module pair), possibly in conjunction with the virtual reference. Within a patch/module pair a plurality of such electrodes may be coupled directly through a corresponding bioamplifier (i.e. such as may be located onboard the corresponding module). During a monitoring session, the patch/module pair, a hub, a coordinating module, or the like may be configured to extract multipolar signals, to detect, to amplify, and/or to algebraically combine such signals with one or more other multipolar signals monitored at sites located elsewhere on the body of the subject. In this manner, standard and/or higher level EKG lead configurations may be derived from multipolar signals obtained from a plurality of patch/module pairs on a subject.

[00170] In another non-limiting example, patch/module pairs including bipolar or multipolar electrode configurations may be placed on a body at standard EKG locations (limb or precordial sites), but not physically connected to one another. One or more electrode sites within the arrangement may be automatically designated as 'references' for the purposes of intra patch signal comparison. In aspects, the system may be configured to designate creation of a 'virtual' reference, and 'virtual' (i.e. calculated) standard leads (limb and precordial) equivalent recordings, performed using mathematics/signal processing techniques (e.g. algebraic transformations based upon one or more of the monitored signals, an image of the patch arrangement on the subject, a network topology of the patches, etc.).

[00171] Such a configuration may be advantageous for partially or completely eliminating the necessity of physical interconnection of electrodes via direct wire leads. Thus, a system in accordance with the present disclosure may include a function for the mathematical combination of signals and/or supplemental data (e.g. orientation based images, orientation based sensor data, etc.) to derive a clinically recognizable EKG signal (e.g. a standard EKG signal that would include lead I, II, III, aVR, aVL, aVF, or the epicardial leads V1, V2, V3, V4, V5 or V6, etc.).

[00172] In aspects, one or more parameters for a signal transformation between patches, may be calculated from an image of the patches on the subject. Each patch may be given a coordinate vector determine from the position and orientation of the patch on the subject (i.e. optionally with respect to one or more standard lead application sites). The coordinate vector may be used in such calculations to calculate a standard lead configuration from a collection of patch/module pairs on the subject.

[00173] In aspects, an ad hoc arrangement of patch/module pairs may be used in harmony to provide redundant monitoring of physiologic parameters from the subject. Sensor fusion of such redundant signals may be used to substantially remove movement artifacts, reduce movement noise, determine a faulty connection on one or more modules, eliminate false alarms caused by movement or other physiologic processes (e.g. brushing teeth, eating, a physiological event, a seizure, an epileptic seizure, an asthma attack, a pulmonary event, wheezing, or the like).

[00174] In aspects, one or more signals, with the assistance of one or more parameters, maybe transformed from the signal as monitored, to a clinically recognizable signal. Such transforms may include a linear algebraic operation, a sum of waveforms, a difference of waveforms, and other more sophisticated signal processing methods such as frequency domain analysis, complex vector representation (amplitude and phase at the multiple sites), vector transformations, convolution or another signal processing technique.

[00175] In aspects, vector (complex real and imaginary components) combination of lead signals may be performed by an operably system in the time domain, or frequency domain for the purpose of transforming one or more signals into a clinically relevant equivalent, etc.

[00176] In aspects, a system in accordance with the present disclosure may include a plurality of patches or patch/module pairs to form a redundant (i.e. for reliable recording, to extract higher level coordination of data from various sites on a subject) physiologic monitor. Such a system may be configured to implement one or more algorithms to coordinate information from each pair to determine higher level functions, maintain operation when a component fails, maintain operation when one or more signals are corrupted (e.g. by movement, stretch artifacts, a poor body connection, etc.), identify components in the network, identify and/or indicate when a component needs to be swapped (i.e. hot swapping for continuous monitoring from the subject), combinations thereof, or the like.

[00177] In aspects, such modular monitoring solutions may be applied to a wide range of monitoring situations. Some non-limiting examples of such applications include hospital based monitoring of patients, remote monitoring of patients, heart-rate monitoring, electrocardiographic monitoring of fitness, athletic, aerobic activities,

yoga, stress management, biomechanics and biometric monitoring systems (e.g. so as to monitor EMG, proprioceptive inputs, etc.), heart-rate variability training, heart-rate variability assessment, traumatic brain injury assessment, muscle tension assessment, tissue assessment (e.g. determination of fat content in tissues around the body, changes in fat content during workout, etc.), sleep studies, sleep monitoring, sleep apnea assessment, physiologic assessment of sleep state, sleep biofeedback, snoring analysis, bruxism monitoring, physiotherapy, event response (e.g. stroke capture and response, heart attack, heart attack prediction, atrial fibrillation, syncope, ST-segment depression or elevation, onset of myocardial ischemia, p-wave analysis, onset of snoring, night terrors, sleep walking, etc.), hydration and fluid management, long-term monitoring, gaming or computer input devices, product testing, marketing analysis, virtualization of emotional experiences, physiotherapy, combinations thereof, or the like.

[00178] In aspects, a system in accordance with the present disclosure may include one or more feedback components (e.g. a device with audible feedback, tactile feedback, visual feedback, combinations thereof, etc.), to provide a subject, coach, practitioner, caregiver, partner, or the like with information, commands, or prompts, pertaining to the physiologic and/or physical signals captured by one or more patch/modules arranged upon the subject. In aspects, such feedback may be used to enhance the sleep state of a subject, interrupt a sleep event to return a subject to a safe or comfortable sleeping state (e.g. interrupt a sleep walking event, a snoring event, a sleep apnea event, night terrors, nightmares, etc.). In aspects, such feedback may be analyzed in combination with the electrophysiological and/or physiologic signals to alter the state of the subject (e.g. the mood, the sleep pattern, the state of sleep, to prevent wake-up, to initiate wake-up, etc.).

[00179] In aspects, a feedback component in accordance with the present disclosure may include or be included in a wristwatch (e.g. a biometric watch, a smart watch, etc.). Such a wristwatch may include a display, a touch screen, or user input device, a tactile (i.e. vibrating) aspect, an audible feedback aspect or the like. Such feedback components may be used to convey signals, or metrics relating to the physiologic and/or physical signals to the wearer (e.g. the subject, a coach, a physician, a caregiver, a partner, etc.).

[00180] In aspects, a feedback component in accordance with the present disclosure may include a heads-up-display (HUD), such as may be provided by a pair of HUD ready glasses, Google Glass™, or the like. In aspects, the HUD may include visual representation of the physiologic and/or physical signals for a wearer (e.g. the subject, a coach, a caregiver, etc.), and/or signals or metrics related thereto or derived therefrom. In aspects, a plurality of such feedback mechanisms may be used to enhance the user experience, such as a combination of audible feedback (i.e. via a loudspeaker), and visual feedback (e.g. on a HUD, via a light emitting diode (LED), etc.).

[00181] In aspects, an augmented reality application may be envisaged using a pair of HUD ready glasses, or via a handheld device with both display and camera functionality (e.g. a tablet, etc.). In aspects, aspects associated with muscle exertion, electrocardiographic data, etc. may be superimposed onto movements associated with the monitoring site so as to highlight such activities to an observer. In one non-limiting example, heart-rate data may be translated into an amplitude parameter for pixel movements and overlaid onto the display or HUD over top of the torso of the subject as displayed in the image. In such an example, a physiotherapist may be able to visualize "exertion" of a muscle group of a subject as it is overlaid onto that particular muscle group during a monitoring session. The exertion may be compared against previous bests, in the context of physiotherapy, may be compared against capabilities (i.e. from previously collected history) and compared against maximal exertion levels, etc. so as to avert injury, optimize an exercise for a subject, maximize the exertion of a local muscle group within a safety window, monitor muscle fatigue during exercise, or the like. Such a system may be advantageous for allowing a user (e.g. the subject, a physiotherapist, a physician, a nurse, etc.) to assess one or more physiologic parameters of the subject while observing the subject or aspects thereof in a display (i.e. without taking attention away from the subject).

[00182] In aspects, one or patch/modules may include a vibrating actuator (e.g. an eccentric motor, an electroactive material actuator, etc.) configured so as to provide a local tactile sensation to the subject. The tactile sensation may be driven by one or more of the physiologic and/or physical signals, by an input from a coach, a caregiver, or the like. In aspects, a system in accordance with the present disclosure may be used to transfer touch sensation from a site without adequate feedback (e.g. a foot, a

shin, a knee, a site of neuropathy, an injured region of the body, etc.), to an alternative site on the body, which still has functioning touch feedback. In aspects, a system in accordance with the present disclosure may be used to convey touch sensation between remotely located subjects, to convey haptic touch information from an object (e.g. a portion of a wheelchair, a bumper, etc.) to a site on the body of the subject.

[00183] In aspects, a patch/module worn by an alternative subject (i.e. a second subject) may be configured so as to provide tactile feedback related to the actions of the first subject (i.e. feedback based upon the physiologic and/or physical signals), so as to convey a sense to the alternative subject of such signals.

[00184] In aspects, a physiotherapist may wear one or more patch/modules corresponding to patches worn by the subject (i.e. a patient). The system may be configured such that physiologic signals and/or physical signals measured on the patient (e.g. electromyographic signals relating to muscle activity, kinetic data, respiration rate, edema level, exertion parameters, etc.) may be "felt" by the physiotherapist via one or more patch/modules worn thereupon. Such a configuration may be advantageous for improving the data available to a physiotherapist during a training session with a patient.

[00185] In aspects, a physiotherapist may wear a HUD ready glasses, Google Glass™, or the like, to which information relating to the physiologic and/or physical signals measured on the patient may be conveyed (i.e. in the form of audio and/or visual feedback). In aspects, the HUD ready glasses may include one or more tactile feedback elements, so as to provide the wearer with further sensations related to the signals.

[00186] In aspects, an application linking an instructor to a student is described herein. In aspects, the instructor may interact with one or more feedback devices (e.g. visual display, HUD ready glasses, tactile feedback device, audio feedback device, etc.) and the student may be coupled to one or more patch/modules in accordance with the present disclosure. In aspects, the instructor may obtain feedback from one or more of the patch/modules on the student pertaining to the physiologic and/or physical signals measured thereby. Such a configuration may provide the instructor with more detailed information that may be unavailable otherwise (such as being able to quantify exertion levels of a student, visualize one or more physiologic parameters

of the student, capture information relating to fatigue, cardiopulmonary changes in the student, etc.).

[00187] In aspects, an application linking two or more partners is envisaged. In aspects, one or more partners may be fashioned with one or more patch/modules in accordance with the present disclosure and one or more feedback devices in accordance with the present disclosure. In aspects, the exchange of physiologic data from patch/module to feedback device may be used to enhance interactions between the partners, remotely link the partners (perhaps in real-time, pseudo real-time, etc.).

[00188] In aspects, a system in accordance with the present disclosure may include a plurality of patch/modules each in accordance with the present disclosure. The plurality of patch/modules may be configured to form at least part of a body area network (BAN). In aspects, the patch/modules may be wirelessly connected to a host device and/or to each other for purpose of communicating physiologic and/or physical signals, network configuration data, time stamps, patch/module configuration data, patch/module IDs, etc. In aspects, the patch/modules may form at least part of a star, line, mesh, tree, spanning tree, network topology to provide such communication.

[00189] In aspects, component operation on and data communication over the network may be coordinated through addition of a time stamp. The time stamp may be used by an associated processor to temporally compare data collected by a plurality of patches on a subject, between a patch on a subject and another recording device located in the environment, etc. One or more patch/module pairs may include a clock, an ultra-low power clock, for generation of the time stamp.

[00190] In aspects, each patch/module may be allocated a temporal window (i.e. temporally multiplexed) within which to broadcast a signal to the host and/or a commanding patch/module in the network. Such a configuration may be advantageous to coordinate significant amounts of data on a network within a limited number of channels.

[00191] In aspects, a system in accordance with the present disclosure may be applied to a stress monitoring application. Such a system may include one or more patches or patch/modules each in accordance with the present disclosure, attached to a subject. The system may be configured to measure one or more physiologic parameters from the subject (e.g. heart rate variability, sympathetic tone, muscular

sympathetic nerve activity, galvanic skin response, skin sympathetic tone, electromyographic activity, respiration rate, etc.). Such information may be combined to form a metric relating to the stress state of the subject. Such a stress state may be represented by a feedback component in accordance with the present disclosure, as part of a biofeedback loop (e.g. a centering algorithm, a calming algorithm, etc.), provided in conjunction with a light and sound show, provided as an "emotional" input to a light and sound show, etc.

[00192] In aspects, the stress state may be used in conjunction with a biofeedback algorithm to help a subject lower the stress state during a monitoring session. Such a system may be advantageous for helping subjects reduce anxiety, reach a meditative state, realize when their stress state is elevated, help adjust respiratory rate, enter into a meditative state, etc.

[00193] In aspects, one or more patches and/or patch/module pairs may include one or more sensors configured to monitor one or more physiologic, environmental, and/or physical parameter locally on the subject. Some non-limiting examples of such sensors include electrophysiologic sensors (e.g. EKG, EMG, EEG, electroretinogram (ERG), EOG, respiration, bioimpedance, activity, etc.), temperature sensor (e.g. near to the skin, within a module, ambient [environmental], etc.), thermal gradient sensor (i.e. so as to calculate a heat transfer vector locally on the body of the subject, to estimate a core temperature), barometer, altimeter, accelerometer, gyroscope, humidity sensor, magnetometer, inclinometer, oximeter, colorimetric monitor (e.g. color change analysis of underlying tissue, for respiration, blood flow, pulse, etc.), sweat analyte sensor (e.g. so as to measure sweat constituents, salt content, etc.), galvanic skin response, neural activity (e.g. skin sympathetic activity), interfacial pressure sensors (e.g. for contact assessment, compliance measurement, blood pressure, etc.), flow sensor (e.g. airflow over a module, or the like), surface strain sensor (e.g. via integration of stretch sensors into the patch, evaluation of stretch along one or more electrical interconnect within the patch, integrated capacitive stretch sensors, etc.), a microphone, combinations thereof, and the like.

[00194] One or more patches and/or patch/module pairs may be configured so as to harvest power from a nearby power source (e.g. via inductive coupling, optical radiation, radio frequency waves, thermal gradients, kinetic energy, or the like). In aspects, one or more patches and/or patch/module pairs may include an antenna

configured to communicate inductively with a nearby radio frequency power source. Such a configuration may be advantageous to work towards minimization of size and weight of the patch or patch/module pair, to reduce costs, recharge the modules without the need for physical connections, etc.

[00195] In aspects, an application for providing a physiologic input tool is envisaged. A system in accordance with the present disclosure may include one or more patches and/or patch/module pairs configured and arranged upon a subject so as to monitor one or more proprioceptive and/or electromyographic signals therefrom (e.g. such as from a muscle group, a bicep, a forearm, a finger, etc.). Such information may be coupled to a software program, such as a music generation program, a toy, a training program, etc. In aspects, the effort with which a muscle group, and/or appendage is moved may be used as an input to the software. In aspects, such input may be attributed to a musical note, an instrument, an avatar of the subject, etc. In aspects, such notes, instruments, etc. may be attributed to a particular patch and/or module, thus an orchestrated score may be generated from measurements made synchronously by a plurality of such patch/modules (e.g. with instruments, tones, etc. being emulated by a particular muscle group, etc.). Such a system may be advantageous for physiologic training for accident or stroke victims, for assisting with gait correction, as a means for treating depression, for entertainment, etc.

[00196] In aspects, such inputs may be coordinated across multiple subjects to orchestrate a coordinated event, collaborate on a musical or dance piece, etc. Particular movements may be used to adjust the tones of a musical stream (i.e. running in parallel with the activity of the subject), etc. Such tuning may be coupled with a gait analysis package so as to provide an elegant biofeedback system for the coupling of music with movement (i.e. such that the music is in tune when form of the movements matches an ideally determined structure).

[00197] Such movement may be coupled with a musical stream. In such cases, the movements may be used to reinforce aspects of the music (e.g. to reinforce beats of the music, etc.), but also to decrease particular beats, as the movements fall out of sync with the music.

[00198] In aspects, the physiologic tool may be used to determine an effort related to a given task, to map a particular movement, to a task space, etc. Such information

may be useful for use in a training program (e.g. a running program to assist with training a student the biomechanics of the sport, etc.). In aspects, strategically placed patches may be used to capture electromyographic information from muscle groups during movement. In aspects, such information may be coupled to a biofeedback system to assist with the correction of movements made by the subject.

[00199] In aspects, such tools may be used to improve the gait of a subject, such as during physiotherapy, during long distance running, during athletic training, during a fitness routine, etc. In aspects, a system in accordance with the present disclosure may include a patch configured for placement over the gluteal muscle (e.g. gluteus maximus, gluteous medius, gluteus minimus) and monitoring during movement (e.g. during a running practice, during a fitness routine, etc.). The system may also include means for monitoring impact of the foot of the subject against the ground, means for determining the basic state of the gait of the subject, etc. (e.g. with a heel-strike sensor, with sensors maintained in one or more patches placed on the body of the subject, etc.). The relationship between the EMG monitored near to a gluteus muscle may be analyzed with respect to other aspects of the gait (such as timing in relation to a heel-strike event, a knee bent, a foot extension, etc.). The relationship may be used to help the subject correct their gait, so as to alleviate pain, reduce injury, recover from injury, improve performance, etc.

[00200] In aspects, a system in accordance with the present disclosure may be used to analyze, assess the quality of, and/or improve the physical output of a performance animal (e.g. a racehorse, a dog, a camel, etc.). Such a system may be advantageous for performing gait analysis, cardiopulmonary assessment, endurance assessment, etc. of the animal adapted from one or more methods in accordance with the present disclosure.

[00201] In aspects, a system in accordance with the present disclosure may be used to assess heart function (e.g. heart-rate, heart-rate variability, electrocardiographic information, respiration, etc.) of an animal, such as in the context of veterinary medicine, preventative veterinary medicine, home pet care, etc.

[00202] In aspects, one or more patches and/or patch/module pairs may include a stretch sensor, for evaluating a state of movement in the muscles underlying a patch on the subject (e.g. or across a joint, along an abdomen, along a spine, etc.). Such

monitoring may be combined with EMG monitoring to give a more thorough session with a subject (e.g. so as to combine exertion with extension of muscular groups, joints, etc.). Such information may be advantageous for evaluating the extent of movement of a subject during an exercise. Feedback derived therefrom may be suitable for correcting gait, limiting such movement, thus reducing injury, as part of a treatment routine for Achilles tendonitis, etc. Such stretch sensing may be combined with general electrophysiological monitoring to remove a stretch related artifact from the signal. Such stretch related artifact may be removed by adaptively subtracting the stretch signal from the electrophysiological signal, so as to minimize the coherence between the two signals.

[00203] In aspects, a stretch sensor in accordance with the present disclosure may be embedded into a patch in accordance with the present disclosure. In aspects, a pair of interconnects may be coupled with a stretchable electrically conducting trace (optionally meandering in nature) on the patch. The stretchable electrically conducting trace may change impedance in accordance with stretch of the patch during use. Such stretch related changes in impedance may be used to determine the overall stretch to which the patch is subjected during use. In aspects, an alternative stretch sensor may embedded into a patch in accordance with the present disclosure. The alternative stretch sensor may include a pair of interconnects for communication with a corresponding module, the alternative stretch sensor including a first stretchy electrode (sized, shaped, and dimensioned to provide a stretchy electrode that responds to stretch preferably in a first direction), a soft dielectric film (e.g. a region of ink coating the first electrode), and a second stretchy electrode placed so as to sandwich the soft dielectric film between the first and second stretchy electrodes. The impedance (e.g. capacitance, ac impedance, etc.), as measured between the coupled interconnects, will change with the stretch of the patch. In aspects, the concepts provided herein for the integration of a conductivity based and impedance based stretch sensor may be combined to give a combination device, which may result in a higher sensitivity to stretch. In aspects, the baseline impedance of the stretch sensor may be measured by the module and used as a method for assessing the interconnection between a module and a correspondingly equipped patch. Such a configuration may be advantageous for monitoring one or more of tissue strain, posture, respiration, respiration depth, movement of digits, movement of joints, or the

like. Such functionality may be integrated into a patch and a corresponding module for use a range of applications requiring stretch related information.

[00204] In aspects, such monitoring may be used to assist with one or more forms of exercise (e.g. any exercise that works and/or stretches the buttocks, for example). Such monitoring may be suitable for evaluating exertion, stretch, etc. during lunges, hip thrusts, climbing stairs, fencing, bicycling, rowing, squats, tennis, arabesque, aerobics, and various specific exercises for the buttock and lower limbs. The information obtained from one or more muscle groups may be used as feedback for training, biomechanics, exertions, etc. as relevant to the sport or exercise under study.

[00205] In aspects, such monitoring may be used to assist with powerlifting exercises which are known to significantly strengthen the gluteal muscles include the squat, deadlift, leg press, feet in squats, and good mornings (bend over with a bar on the shoulders with a light amount of weight). Such monitored information may be used to assist with exertion feedback, limiting movement, and the like, so as to provide a subject or trainer with information during the exercise, limit the risk of damage, optimize training, maximize performance gains, or the like.

[00206] In aspects, such monitoring may be applied to virtually any muscle group or groups on the subject. Such approaches as described herein may be adapted to serve a range of exercises and sports, so as to assist subjects, trainers, physiotherapists, and the like with improving performance, adjusting gait, monitoring progression of an injury, comparing capabilities to previous efforts, assess or assist with sexual dysfunction, etc.

[00207] In aspects, a system in accordance with the present disclosure may be configured for long-term wear by a subject. The system may include one or more patches and/or patch/modules in accordance with the present disclosure, configured to comfortably attach to the subject and remain attached for a prolonged period of time (e.g. days, weeks, greater than 3 weeks, etc.). In aspects, the patch may include a bioadhesive with suitable breathability for long term wear (i.e. with moisture vapor transmission rate (MVTR) in a range such that tissues underlying the patch do not undergo maceration but also do not sufficiently dry out so as to impede electrophysiological measurement thereupon). In aspects, the module may include a

gasket arranged so as to bias towards a mating patch when the two are assembled (i.e. so as to form a substantially water tight structure).

[00208] In aspects, the module may include a moisture sensor and/or humidity sensor positioned so as to monitor the environment between the module and the patch during use. Such information may be advantageous for determining if the patch needs to be swapped out during use.

[00209] In aspects, such a system may be suitable for use as a heart-rate variability (HRV) monitor. In aspects, such a module may include circuitry or be programmed with a function to calculate HRV from an EKG or extracted heart-rate metric. Thus, physiologic signals pertaining to HRV, heart rate (HR), etc. may be conveyed from the module to the host during a monitoring session (i.e. as a means for reducing the amount of raw data to be sent from the module to the host during a session). HRV monitoring may be advantageous for gauging training recovery, for evaluating a subject who has experienced a traumatic brain injury (e.g. via a sudden impact, repeated impacts, etc.), to grossly interpret activity of the sympathetic nervous system, to evaluate an emotional state or response of a subject, or the like.

[00210] In aspects, a system in accordance with the present disclosure may be adapted to monitor impact to a subject during a monitoring session. In aspects, the system may include a patch and/or patch/module pair including an accelerometer, a gyroscope, an altimeter, a barometer, stretch sensor, and/or a magnetometer configured so as to measure a sudden impact on the patch during a monitoring session. In aspects, the module may include circuitry or be programmed to monitor data derived from the sensors so as to determine if an impact has occurred (e.g. a rapid change one or more of the sensor data streams, a change of sufficient magnitude, etc.). Such information may be directed to a trainer, a doctor, a coach, a parent, or the like to alert them to the impact. In aspects, such information may be useful to determine if an impact requires further investigation, etc.

[00211] A system in accordance with the present disclosure may be configured for analyzing traumatic brain injury of a subject. The system may include one or more patch/module pairs in accordance with the present disclosure sized, dimensioned, and configured for placement at one or more sites on the head (e.g. sites 50, 131, 137, 140 as indicated in Fig. 1a, along a neck, along a forehead, along a temple, behind an ear,

etc.) forming a head mounted patch/module pair, and optionally elsewhere on the body (e.g. on the chest, over a muscle group, etc.) forming a second patch/module pair.

[00212] In aspects, the head mounted patch/module pair may include one or more electrodes for monitoring local physiologic activity (e.g. EEG, EOG, EMG, etc.), near the brain of the subject, one or more kinetic or kinematic sensors (e.g. accelerometer, gyroscope, barometer), one or more additional sensors (e.g. local bioimpedance, hydration, temperature), and one or more acoustic sensors (e.g. a down facing microphone). The head mounted patch/module pair, or a processor coupled thereto may be configured to capture signals related to the above sensors in a black box, virtual black box, rolling first in first out (FIFO) buffer, or the like. In addition a processor on or coupled with the head mounted patch/module pair may be configured to analyze the signals to determine the timing of an impact, the kinematics of an impact, etc. In aspects, the second patch/module pair may collect heart rate information, heat rate variability, further body based temperature, hydration, and/or kinematic information. Upon detection of an impact, the system may save the physiologic information captured prior to the impact so as to form a pre-impact dataset, and continue to collect physiologic and/or physical data so as to form a post-impact dataset. The system may include an algorithm configured to automatically compare the pre-impact and post-impact datasets to assess the response of the subject to the impact. In aspects, an apparent change in heart rate variability, a change in the EEG, or EMG collected by the head mounted patch/module pair, or the like may be a strong indication that the subject was adversely affected by the impact. In aspects, the system may be configured to generate an alert, a report, an alarm, etc. upon determination of such a response.

[00213] The system may include an algorithm (e.g. either incorporated into a processor on a patch/module pair, in a processor coupled thereto, etc.), configured to analyze temperature, heat transfer, hydration level, or the like from the subject. Such information may be combined to form a metric relating to dehydration, an over temperature, and/or exhaustion state of the subject from which further action may be taken (e.g. generate an alert, an alarm, a report, feedback to the subject, to a coach, to a parent, etc.).

[00214] In aspects, a system in accordance with the present disclosure may be adapted to monitor thermal stress of a subject during a monitoring session. In aspects, the system may include one or more patches and/or patch/module pairs including one or more temperature sensors, thermal gradient sensors, hydration sensors, core temperature estimating sensors or algorithms, etc. so as to capture a relevant temperature of the subject during a monitoring session. In aspects, the system may include a feedback component in accordance with the present disclosure to convey such thermal information to a coach, a trainer, a physician, a parent, a race coordinator, etc.

[00215] In aspects, one approach for forming a core temperature estimating sensor in accordance with the present disclosure includes providing a patch in accordance with the present disclosure, the patch including at least one temperature sensor (e.g. a microcircuit based temperature sensor, a thermocouple, a bimetal strip, etc.), the patch configured so as to mate with a corresponding module, the module including one or more additional temperature sensors. The combination patch/module pair thus includes a plurality of temperature sensors (e.g. included in the patch, included in the module, etc.). Heat transfer from the adjacent tissues past the plurality of temperature sensors and into the surrounding environment will follow different pathways. The patch and/or module may include a plurality of controlled pathways with known heat transfer coefficients (e.g. such that the ratio of heat transfer coefficients between the pathways can be reasonably predicted). In aspects, the heat transfer coefficient along the pathway between the skin and each sensor may be known, and the heat transfer coefficient between each sensor and the surrounding environment may be known (e.g. to within a ratio). In aspects, the patch/module pair may include an additional temperature sensor, positioned so as to monitor a temperature near that of the surrounding environment (e.g. such as exposed on the surface of the module, on the surface of the patch pointing way from the body, collected from a local hub, from a smartphone, from a local weather report, or the like).

[00216] The temperature at a site remote from the patch (e.g. deeper into the tissue), may be estimated from the plurality of temperature readings in combination with thermal pathway coefficients. In aspects, temperature measurement may be improved by single point calibration with a traditional core temperature sensor. After calibration, the temperature measurements may be sufficient to estimate changes in

core temperature over time while the subject wears the patch/module pair. In aspects, a plurality of patches worn by a subject may be used to improve an estimate of core temperature thereof.

[00217] In aspects, one approach for forming an embedded hydration sensor into a patch/module pair may include, embedding a plurality of electrodes into a patch in accordance with the present disclosure (e.g. 2 or more electrodes, 4 electrodes, etc.). The corresponding module may include a signal generator and one or more bioamps (or multiplexing circuits in combination with a reduced number of bioamps, etc.), the signal generator configured so as to provide a signal between two or more of the electrodes (e.g. in a frequency range of generally between 1hertz (Hz) and 10GHz, 1kHz and 10MHz, 5kHz and 1MHz, or the like, at multiple frequencies, swept over a range of frequencies, etc.), while the bioamps are configured to capture one or more signals from two or more of the electrodes. The processor, gate array, digital signal processor, or an associated microcircuit, configured to analyze the captured signals to determine a bioimpedance of the nearby tissues. Such a bioimpedance may be advantageous for monitoring a change in fluid level in the adjacent tissues. Such a configuration may be advantageous for assessing the fluid level of a subject. In aspects, two or more of the electrodes may be coupled to a second bioamp (an additional amplifier, the same amplifier, etc.), the second bioamp configured to capture a biosignal (e.g. EKG, HR, EMG, EOG, EEG, ERG, or the like) from the electrodes. Assessment of the signals at a plurality of frequencies may provide sufficient data to extract electrode impedance, from the reading, to estimate tissue impedance at different frequencies, and to extract the extracellular (ECW), intracellular (ICW), and total body water (TBW) content from the tissues under analysis by the patch/module pair.

[00218] In aspects, the patch/module pair includes an antenna, the antenna coupled to convey one or more signals to an external hub, additional patch/module pair, etc. The antenna may be coupled to the adjacent body, such that during signal transfer, the impedance of the antenna varies with water concentration of the adjacent tissues. During operation, changes in the S11 parameter for the antenna may be monitored, so as to determine local fluid levels in the adjacent tissues, while also transferring data back and forth for completing other functions.

[00219] In aspects, a second approach for forming an embedded hydration sensor into a patch/module pair may include, embedding a visible, near infrared, or infrared emitter into the module such that, upon coupling of the module with a corresponding patch, the emitter is arranged such that radiation emitted therefrom is directed into the tissues of the subject, the module including a photodetector (e.g. a narrow band detector, centered generally about 510 nanometer (nm), 578nm, 630nm, 750nm, 1000nm, 1180nm, 1040nm, 1210nm, 1300nm, 1500nm, a multi-band detector <combinations thereof>, a broad band detector, multiple detectors, etc.) to capture reflected or back scattered radiation from the skin from the emitter. In aspects, the corresponding patch may be configured with a window transparent to, or polarized so as to exclude light, such that when the module is coupled to the corresponding patch, the emitted and/or detected light pass through the window. In aspects, the module may include a corresponding cross polarized window, such that the two windows may be used in unison to exclude light from the surroundings, eliminate incident light from the emitter reaching the detector, etc. Such a configuration may be advantageous to improve signal to noise ratio in such readings.

[00220] The collected signals may be used to assess the tissues adjacent to the patch/module pair. In aspects, the patch/module pair may include or be coupled to a processor equipped with an algorithm, the algorithm configured to interpret the collected signals and determine the state, analyte concentration (e.g. oxygen, water, lipid, melanin, myoglobin, collagen, elastin, etc.), and/or composition of a fluid and/or tissue adjacent to the patch/module pair. In aspects, the algorithm may be configured to extract a metric related to water content of the tissues, the metric suitable for use as a surrogate for hydration of the subject. In aspects, the collected data may be analyzed so as to extract oxygen blood concentration, water, lipid level, or the like. Such a configuration may be advantageous for providing a consistently oriented measurement configuration, with substantially minimized external influences, highly controlled pressure on the tissues (e.g. near zero contact pressures, through to consistently controllable pressures with inclusion of a feature on the patch), exclusion of ambient light, and cost reduced measurement configuration. When placed near to an artery, the configuration may be suitable for obtaining a local pulse signal (i.e. for combination with a plurality of patch/module pairs to determine a time-of flight based blood pressure reading on the subject).

[00221] In aspects, the emitter may be pulsed with a duty cycle, and the sensory circuits may be configured so as to operate with a duty cycle. Such a configuration may be used so as to substantially conserve power during operation.

[00222] Such a configuration may be combined with one or more sensors in accordance with the present disclosure (e.g. such as electrodes into the patch, etc.), so as to provide a higher level of diagnostic information.

[00223] Such a system may be combined with one or more features in accordance with the present disclosure to provide diagnostic input relating to assessment and capture of a sleep apnea event, choking, limb status, post-operative tissue flap status, or the like (i.e. assess local blood flow to a tissue site covered by the patch/module pair).

[00224] According to aspects there is provided, a patch/module pair in accordance with the present disclosure, the patch configured for placement onto a body part after a surgical operation (e.g. after a plastic surgery, a tummy tuck, a breast augmentation, a stoma formation, a fistula repair, a wound closure, a tissue transplant, etc.), the patch configured so as to shape the tissue to which it has been attached (e.g. provide a restraining shape, so as to enhance the healing of the procedure, minimize scaring, heal in a particular configuration, under a particular stress state, etc.), the module and patch including one or more sensors in accordance with the present disclosure, the sensors arranged and oriented so as to assess the health of the tissue. In aspects, the patch/module pair may include a hydration sensor in accordance with the present disclosure, to assess the hydration state of the tissues, a change in color of the tissues, a change in oxygen saturation level, or the like. The module may include a processor equipped with an algorithm to assess changes in the tissue state and generate an alert, an early warning, etc. if the changes are not conducive to a successful surgical outcome for the patient.

[00225] In aspects, the system may include a feedback component in accordance with the present disclosure, such as a pair of HUD ready glasses, such as Google glass™ configured to be worn by a data recipient (e.g. a trainer, a coach, the subject, a parent, a physician, a nurse, a caregiver, etc.). The feedback component may be configured to alert the wearer of the thermal state of the subject (e.g. so as to warn of heat exhaustion, dehydration, etc.). In aspects, a feedback component on a single data

recipient (e.g. a trainer, a coach, a race coordinator, etc.) may be configured to display data corresponding to an entire team, a class of students, a group of competitors, etc.

[00226] In aspects, thermal monitoring, impact monitoring, and HRV monitoring may be combined within a single system to provide a heath monitor for use during physical training, sports, fitness events, etc.

[00227] In aspects, a system in accordance with the present disclosure may be configured to provide one or more of a cuff-less blood pressure monitor (e.g. via inclusion of one or more patch/module pairs equipped with an interfacial pressure sensor, local pulse assessment sensor, a compliance sensor, or the like), a modular EKG monitor (i.e. via inclusion of one or more patches or patch/module pairs configured to monitor local electrophysiological information from the torso of a subject), a modular EMG monitor (i.e. via inclusion of one or more patches or patch/module pairs configured to monitor electrophysiological information in the vicinity of functional muscle groups on the subject), an edema assessment system (i.e. via inclusion of a patch or patch/module pair configured to measure water content of tissues in the vicinity of the patch as placed onto the subject), a blood clot detection system (i.e. via inclusion of two or more patches to monitor changes in blood flow dynamics between the patches as placed along one or more appendages of a subject, e.g. flow dynamics as measured along the leg), a peripheral vasculature diagnostic system (e.g. via inclusion of one or more patches to monitor arterial flow, arterial brachial index, etc.), or utility in the monitoring of heart failure patients, patients that are short of breath, chronic obstructive pulmonary disease (COPD) patients, patients suffering or suspected of suffering from sleep apnea, valvular disease, metabolic syndrome, etc. to track cardiopulmonary parameters as indicators of disease state/progression, diagnostics, and/or therapeutic progress with the disease.

[00228] According to aspects a system for monitoring blood pressure of a subject is provided, the system including a plurality of patch/module pairs, at least one patch module pair configured to measure an electrophysiological signal in accordance with the present disclosure (i.e. an EKG patch), and at least one or more patch module pairs configured to measure a local signal relating to blood oxygen level and/or pulse in a tissue volume in the vicinity thereof (e.g. via optical methods, bioimpedance, spectroscopic, local counter pressure assessment, or the like in accordance with the present disclosure) (i.e. a local pulse patch). The EKG patch configured for

placement on the torso of the subject, and one or more of the local pulse patches configured for placement at least on an extremity of the subject (e.g. on the torso and on an extremity, near a wrist, on an arm, on a leg, etc.). The system may include a time stamp in accordance with the present disclosure such that the timing of the electrophysiological signal obtained by the EKG patch may be compared to the pulse response signal(s) obtained by the local pulse patch(es). A time delay(s) between a reference wave on the electrophysiological signal (e.g. onset, peak, etc. of a QRS complex), and the pulse signal (e.g. pulse peak, valley, etc.) of the local pulse patch(es) may be generated to assess blood pressure. The time delay(s) may be combined to generate a blood pressure metric for the subject. In aspects, one or more of the patches may include a kinematic sensor, positioning sensor (e.g. a barometer, a tilt sensor, etc.), to determine a positioning between the patches in accordance with a local gravitational field (so as to calculate a head loss, etc. between the patches). The positioning between the patches may be assessed by an algorithm in combination with the time delay(s) to improve the estimate of the blood pressure of the subject. In aspects, the system may include or interface with a blood pressure measurement device (e.g. a blood pressure cuff), the measurement taken by the blood pressure measurement device used to calibrate the algorithm, so as to further improve the blood pressure estimate. In a usage case, the blood pressure reading may be calibrated daily, weekly, etc. so as to maintain an acceptable accuracy thereof. Such a configuration may be advantageous to provide a non-invasive, comfortable ambulatory blood pressure monitor for assessing real-time changes in blood pressure of a subject during a prolonged monitoring period.

[00229] In aspects, one or more patches or modules may include a microphone, optionally directed so as to collect an audio signal from an adjacent skin surface. The microphone may be used to measure an acoustic signal from the subject in the vicinity of the patch or module. In aspects, the patch or module may include acoustic isolation such that the microphone doesn't pick up substantial ambient noise. In aspects, the microphone may be suitable for capturing data relating to heart murmur, a sleep apnea event, an airway obstruction, wheezing, an asthmatic event, weeping, joint noise, or the like.

[00230] In aspects, a system in accordance with the present disclosure for assessing traumatic brain injury may include a microphone within a module and/or patch, the

microphone configured to monitor an acoustic signal related to bone movement, echo, and bone on bone impact associated with a potentially concussive impact applied to the subject. Such information may be combined with electrophysiological monitoring, or the like in accordance with the present disclosure to assess the influence of an impact on a subject.

[00231] In aspects, a system in accordance with the present disclosure may be adapted for follow-up of a joint replacement surgery. In aspects, the system may include one or more patches and/or patch/module pairs configured for placement in the vicinity of the joint, or arranged so as to monitor muscle movement or exertion associated with the joint. In addition the to the physiotherapy related aspects discussed throughout the present disclosure, the patch and or module may be configured to monitor one or more aspects of the joint (e.g. joint acoustic signature, thermal changes in the joint during use, joint impacts, etc.). The system may be configured so as to analyze such data to predict a lifetime of the joint, predict patient outcome, detect early signs of grinding, or the like.

[00232] In aspects, one or more patches and/or modules in accordance with the present disclosure may include a micro compliance sensor and/or an interfacial pressure sensing element so as to assess local changes in blood pressure, pulse, arterial pressure, or the like. The microcompliance sensor can be used to assess local tissue stiffness, or placed in close proximity to vascular structures in order to measure vessel stiffness and/or pulse pressure. Arrangements of such sensors along the extremities of a subject may be advantageous for assessing blood flow (i.e. via pulse time of flight measurements), blood pressure, pulse, or the like. A microcompliance sensor may include a deformable part, and a Micro-Electro-Mechanical system (MEMs), or optoelectronic sensing element to determine the state of deformation of the deformable part. In one non-limiting example, a microcompliance sensor may include an elastomeric deformable part, configured so as to move when a pressure is applied thereto, and an optical source and detector, configured to deliver a light source into the deformable part and monitor a return signal therefrom. Changes in the return signal may be correlated to the deformation state of the deformable part, thus providing the necessary feedback. Such a patch may be advantageously applied to a blood pressure monitoring system as a local pulse patch in accordance with the present disclosure.

[00233] In aspects, a system in accordance with the present disclosure may be configured for simplified use by a subject. One or more patches or modules may include an interfacial monitor to determine if the system is attached to the subject. Upon attachment, the module may be configured to determine the quality of the attachment, and connect to the BAN to start a monitoring session. In aspects, one or more patches or modules may include a battery monitor, interface monitor, etc. to determine if the patch should be swapped out or not during a monitoring session. In aspects, a plurality of patches or patch/module pairs may be attached to the subject such that a level of redundancy in the monitoring process may be achieved. In such situations, the system may continue to effectively monitor the subject even when one or more patch/module pairs is not properly attached, low on power, etc. Such a configuration may be advantageous for reducing the number of false alarms that may be detrimental to the monitoring process in a hospital, for example.

[00234] In aspects, to further simplify use, one or more modules in accordance with the present disclosure may include one or more orientation sensor, motion sensor, barometer, or the like and/or means for determining the network topology (ex. wireless signal strength between modules in the network, user method for identifying the module, etc.). Such information may be used to automatically or at least partially determine the placement of the module on the subject.

[00235] From such an assessment, the relative positions and orientations of modules in the network and on the subject can be determined and used to improve the clinical quality of a multi-module monitoring session (i.e. by generating a standardized EKG lead configuration from the available configuration), improving the accuracy of a multi-lead EKG, identifying a heart location in comparison with the modules, and/or determining the orientation of the subject (e.g. standing, sitting, supine, etc.).

[00236] In aspects, one or more of the modules, a host, or a system coupled thereto may be programmed with a function to determine the effectiveness of the capture of the intended data by one or more of the patch/module pairs (i.e. the quality of the collected data) and to determine whether such data should be trusted in the collected data stream or not. In aspects, the data may be analyzed to determine if a particular data stream has been corrupted by movement (e.g. due to EMG interference, relative movement at the site of the patch, stretch based artifacts, etc.), by water ingress (e.g.

due to moisture entrainment into the module, interface, etc.), poor connection to the subject (e.g. via determination of high electrode impedance, etc.), or the like. Upon detection of an issue, the algorithm may be configured to dismiss data collected form that patch/module pair, de-emphasis such data, etc. until the issue is resolved. The algorithm may be configured to assess whether the data collected from the remaining patches is sufficient to capture the sought after information (e.g. sufficient data to rule out a heart attack, to assess atrial fibrillation, to assess syncope, to determine if a syncope event is cardiogenic, reflex, and/or orthostatic hypertension, etc.), and continuing with monitoring if this is the case, while raising an alarm, alert, etc. if the quality of recording cannot be maintained in light of the issue. Such algorithms may be advantageously implemented to assist with managing a system in accordance with the present disclosure.

[00237] In aspects, one or more systems in accordance with the present disclosure may be coupled to a control console (e.g. a computer terminal, a system management software front end, a server, a virtual server, a cloud based service, etc.) whereby aspects of the system(s) may be assessed and altered rapidly to improve workflow therewith, or the like.

[00238] In aspects, a system in accordance with the present disclosure may be coupled with a patient management system, configured to quantitatively manage customer/patient progress, improvement, and engagement with a therapy, exercise routine, local community event, etc.

[00239] In aspects, a system in accordance with the present disclosure may be adapted for use within a home care setting. In such settings, data collected by the host (e.g. a smartphone, a WiFi® hub, a Bluetooth low energy® hub, etc.) may be sent onto a data center for further analysis. Such information may be collected efficiently without interfering with the subject's daily routine, etc.

[00240] In aspects, a system in accordance with the present disclosure may be configured for entertainment purposes. Such a system may include one or more functions to report (e.g. notify, Tweet™, m2m text message, post, communicate, etc.) one or more aspects of a subject's physiologic and/or physical response to a peer group. In aspects, such a system may include connections to a theme park customer management system, a product evaluation feedback system, etc. In one non-limiting

example, a system in accordance with the present disclosure may be configured to monitor and report the heart-rate of a subject during an amusement park ride (e.g. during a roller coaster, haunted house, etc.), during an extreme sport (e.g. sky diving, water skiing, hang gliding, etc.), or the like, and to report such metrics to a peer group associated with the subject, optionally along with one or more contextual data points (e.g. roller coaster name, subject location, etc.). Such information may be reported during peak physiologic events (e.g. during peak heart rate, during peak respiratory rate, etc.). Such information may be used to quantitatively track customer response to a product, process, to track subject "activities", or the like.

[00241] In aspects, a system in accordance with the present disclosure may be configured to communicate one or more aspects of the collected data, or signals/metrics derived therefrom, goals achieved, or the like to a social forum associated with the subject (e.g. a social network, Facebook™, Instagram™, Google+™, Patient's Like Me™, or the like). Such information may be included in a feedback loop (e.g. so as to encourage a patient, congratulate a subject on an outcome, etc.). In aspects, one or more processors integrated with the social forum may be configured to automatically analyze the collected data and produce one or more metrics relating to disease progression, health state, performance, events (e.g. excitement, amusement park reporting, product usage feedback, intimacy assessment, stroke, physiotherapy progress, etc.).

BODY INTERFACE ASPECTS:

[00242] A patch in accordance with the present disclosure may include a substrate, one or more interconnects fastened to the substrate, each interconnect including one or more connectors, configured, and dimensioned for interfacing with a module in accordance with the present disclosure. The patch may include one or more sensors in accordance with present disclosure coupled with the substrate, arranged so as to interface with an adjacent skin surface after placement on a subject, the sensors electrically, mechanically, and/or optically coupled with one or more interconnects. In aspects, one or more of the sensors may be an electrode. Optionally the patch may include a microcircuit embedded in or attached to the substrate or one or more interconnects, configured to convey information between the interconnect and one or more associated sensors included in the patch. In aspects, the patch may include an adhesive or attachment layer coupled with or integrated into the substrate, configured

such that the patch may be applied to a surface of a subject (i.e. the skin of the subject).

[00243] In aspects, the patch may be configured such that it is sufficiently breathable to water vapor and/or oxygen that it can be worn comfortably by a subject for a prolonged period of time. In aspects, one or more regions of the patch may be configured with a moisture vapor transmission rate of greater than 200g/m^2/24hrs, greater than 500g/m^2/24hrs, greater than 2,000g/m^2/24hrs, etc. In aspects, one or more regions of the patch may be configured with a moisture vapor transmission rate of less than 20,000g/m^2/24hrs, less than 12,000g/m^2/24hrs, less than 8,000g/m^2/24hrs, or the like. Such moisture vapor transmission rate (MVTR) may be approximately determined using ASTM standard ASTM E96: Standard test methods for water vapor transmission of materials.

[00244] In aspects, the substrate or the patch in accordance with the present disclosure on the whole may be stretchable, so as to maintain operation while stretching in conjunction with the surface of the subject where it is attached (e.g. on the torso, across a joint, along a muscle, etc.). In aspects, the substrate or the patch on the whole may be highly elastic so as to easily stretch with the skin without rucking, curling, lifting from the skin surface, etc. Such a soft and thin patch may be advantageous for substantially maximizing comfort of the subject, reducing skin irritation, etc. thus improving subject compliance for long term monitoring.

[00245] In aspects, an adhesive layer in accordance with the present disclosure may include a dielectric or an electrically conducting adhesive, a biocompatible pressure sensitive adhesive, a gel layer, a hydrogel layer, or the like. In aspects, the adhesive layer may include one or more patterned features, configured so as to provide regional variation in adhesion, to isolate regions of the skin surface in one region from adhesive located in other regions, and the like. In aspects, the adhesive layer may include a biodegradable polymer.

[00246] In aspects, one or more regions of an adhesive layer in accordance with the present disclosure may be electrically and/or ionically conducting and patterned onto the substrate so as to provide adhesion thereof to a skin surface. The substrate may be patterned with one or more electrical traces configured to connect one or more electrodes with one or more connectors in an interconnect each in accordance with the

present disclosure. In aspects, the electrical traces may be isolated from one or more regions of the adhesive layer (e.g. with a dielectric overcoat, via a passivation layer, etc.) so as to form regions of substantial isolation and regions of substantial connection between an attached skin surface and the electrodes and/or electrical traces in the patch.

[00247] In aspects, the patch may include an adhesive layer coupled with the substrate for making contact with the subject. To maintain a breathable, flexible interface, the thin adhesive layer may have a thickness of less than 50um, less than 25um, less than 12um, less than 6um, less than 4um, etc. The thin adhesive layer may be formed from a pressure sensitive adhesive (e.g. an acrylic adhesive, a silicone adhesive, a hot melt pressure sensitive adhesive, a thin hydrogel adhesive, etc.), a silicone gel adhesive, a hydrogel, a biopolymer, a hot melt adhesive, combinations thereof, or the like. The thin adhesive layer may be formulated in combination with one or more salts, so as to impart suitable ionic/electrical conductivity to communicate electrically between one or more aspects of the patch and the skin surface of the subject.

[00248] In aspects, the adhesive layer may be formulated with one or more soothing constituents, such a menthol, mint, honey extract, aloe, essential oils (e.g. lavender oil, vanilla oil, camphor, cinnamon extract, orange extract, etc.). Such soothing constituents may be added to the adhesive formulation during fabrication, misted onto the patches, and/or added to the packaging of the product, etc.

[00249] In aspects, the adhesive layer may be patterned so as to form a heterogeneously distributed adhesive pattern to an adjacent skin surface, to form regions of varied MVTR, as a means for reducing the amount of adhesive required to interface with the subject, etc.

[00250] In aspects, one more elements within the adhesive layer, the substrate, or the patch may include a hydrogel, including a biocompatible click chemistry based gel (i.e. for improved strength and minimization of gel residual transfer to the subject during a monitoring session), a radiation cured gel, a high tear strength gel, an ionically conducting gel, an intrinsically conducting (e.g. combined with conjugated polymers, in situ polymerized conjugated polymers, etc.).

[00251] In aspects, the patch may include a ferromagnetic layer or patterned region, ferromagnetic layer with adhesion (i.e. for integration into the adhesive layer), ferromagnetic material with combined electrical conductivity (i.e. for combined attachment and function as an electrical interconnect), etc.

[00252] In aspects, a patch in accordance with the present disclosure may include a substrate, coupled to the thin adhesive layer. In aspects, the substrate may be formed from a sufficiently thin polymeric material (e.g. a polyester, a polyurethane, ethyl vinyl acetate, polyether block amides, an elastomer, polyisobutylene, polyisoprene, a thermoplastic elastomer, butadiene block copolymer, styrene butadiene carboxy block copolymer, vinyl methyl silicone, polysiloxane, styrene ethylene butylene styrene copolymer, styrene butadiene, polyurethane, acrylonitrile butadiene, isobutylene isoprene butyl, hydrogenated nitrile butadiene, fluorocarbon rubber, fluoro silicone, fluoronated hydrocarbon, polybutadiene, ethylene-acrylate rubber, polyester urethane, etc.). To maintain a sufficiently flexible, comfortable and/or breathable interface for a subject, the substrate may be formed from a thin polymeric material with thickness less than 50um, less than 25um, less than 12um, less than 8um, less than 4um, etc. In aspects, the substrate may be formed at least partially from a thin polyurethane substrate with thickness less than 13um. In aspects, the polyurethane film may exhibit an MVTR of greater than 200g/m^2/24hrs, greater than 400g/m^2/24hrs, greater than 1,000g/m^2/24hrs, greater than 4,000g/m^2/24hrs. Such a configuration may be advantageous for providing a low profile, soft, conforming disposable adhesive part that may stretch and move with the adjacent tissues of the subject during an ambulatory monitoring session.

[00253] In aspects, one or more regions of the substrate may be coated with a lubricous or otherwise low friction material so as to provide a desirable surface texture to a subject when placed during a monitoring session (so as to limit rucking of the patch during a monitoring session, etc.). In aspects, the substrate may be coated with a thin lubricious coating of a silicone and/or silicon micro/nano bead layer. Such a coating may provide a soft, lubricious feel for the substrate while maintaining a hypoallergenic quality and further limit rucking during a monitoring session.

[00254] Alternatively or in combination, the patch may include a thin ionically and/or electrically conducting adhesive, coupled to the substrate and/or the thin adhesive layer so as to make physical contact with a subject during placement on the

body of the subject. The ionically and/or electrically conducting adhesive may be one or more adhesive known in the art. In aspects, the ionically and/or electrically conducting adhesive may include a hydrogel.

[00255] In aspects, the patch may include an electrically conducting member (e.g. a stud, an electrical trace, an electrical patch, a wire, a coil, a thin film conducting trace, a printed electrical trace, etc.) for interfacing an operably connected module to one or more sensors (e.g. sensors in accordance with the present disclosure, electrodes, etc.), the thin adhesive layer, and/or the ionically and/or electrically conducting adhesive. The electrically conducting member may include a ferromagnetic material (e.g. iron, nickel, etc.) so as to provide a bias force for holding the patch to an associated module including one or more mating magnetic elements. In aspects, a ferromagnetic material region may be added to the patch such that the conducting layer would be sandwiched between the ferromagnetic material region and a magnetic connector on a corresponding module. Such a configuration may be advantageous to maintain a conductive interface between the module and the patch when mated together, without the need for a large mechanically interlocking connection between the patch and the module.

[00256] In aspects, a patch in accordance with the present disclosure may include a module-patterned adhesive structure oriented so as to interface with an operably placed module in accordance with the present disclosure during placement onto the body of a subject. In aspects, the module-patterned adhesive may be formulated so as to provide a reversibly attachable adhesive option for a corresponding module. In aspects, the module-patterned adhesive may be formed from a silicone adhesive, a low tack pressure sensitive adhesive, or the like.

[00257] In aspects, the patch may include a substrate formulated so as to provide both support (e.g. to one or more interconnects, sensors, microcircuits, etc.) and means for securing the patch to a subject (i.e. attachment means to a skin surface). In aspects, the substrate may be formed from a suitably porous, hydrophilic material. The porous hydrophilic material may be configured to wick fluids from the surface of the skin during attachment, so as to form intimate contact therewith and to bond to the skin during the period of intimate contact (i.e. to establish Van der Waal bonds thereto).

[00258] In aspects, a patch in accordance with the present disclosure may include two or more electrodes, patterned so as to measure a local electric field (e.g. a bipolar, multipolar reading, a field vector, etc.) on the surface of the subject during a monitoring session. The local electric field may be used in part to construct one or more electrocardiographic signals from the subject during a monitoring session. In aspects, one or more patches may include an increased number of electrodes, and such electrodes may be monitored to glean further information related to electric field orientation and propagation over the body of the subject during the monitoring session. Such information may be advantageous in so far as obtaining diagnostic information from a subject during a monitoring session as well as to assist in constructing higher lead count EKGs from an array of patches (e.g. extracting a 12-lead EKG equivalent from an array of patches attached to the subject, etc.).

[00259] In aspects, an electrode in accordance with the present disclosure may include one or more electrode features (e.g. microfibers, barbs, microneedles, spikes, or the like), arranged and dimensioned so as to penetrate into the stratum cornium of the skin during an attachment and/or during an engagement procedure. In aspects, the electrodes may be made to puncture and/or penetrate through the stratum corneum during placement, an attachment procedure, and engagement procedure, etc. The electrodes may be forced to cause penetration of one or more of the electrodes or electrode features into the skin wall so as to enhance the electrical connection thereto during a monitoring session. In aspects, one or more electrodes may be configured for microscopic or macroscopic spatial recording. In aspects, one or more of the electrodes may include a barb to retain the electrode into the skin of the subject after engagement. Such a configuration may be advantageous to obtain a low impedance, reliable interface with the subject during a monitoring session, to assist in decreasing movement artifacts, and/or help in decreasing stretch based artifacts in an acquired electrophysiological signal.

[00260] In aspects, one or more electrode features in accordance with the present disclosure may be spring mounted, such that they do not normally engage directly with the skin surface after attachment of the corresponding patch thereto. Upon pressure application to the electrode by an external entity (e.g. a thumb, an applicator, etc.), the electrode features may be biased towards the skin, thus penetrating the stratum corneum and enhancing the electrical connection thereto during the

monitoring session. In aspects, once engaged with the skin, the electrode features may remain in place for the duration of the monitoring procedure. In aspects, after removal of the biasing force (i.e. the force provided by the external entity) the electrode features may withdraw from the skin, leaving an array of micropunctures through the stratum corneum for enhanced electrical contact therewith.

[00261] In aspects, the electrode features may be coupled to a region or layer of an electrically and/or ionically conducting substance (e.g. a gel, a hydrogel, a salt laden hydrogel, an intrinsically conducting hydrogel, etc.), and penetrate through the substance when engaging with the underlying skin. In aspects, the substance region or layer may provide a biasing force to disengage the electrode features from the skin layer after a brief engagement with the skin. Such a configuration may be advantageous to provide a high quality interconnect between the patch electrodes and the skin of the subject, while maintaining a comfortable environmentally controlled contact with the skin and protecting (and optionally maintaining) any micropunctures in the stratum corneum during the monitoring session. The initial engagement of the electrode features with the skin may assist in lowering the local impedance of the stratum corneum, so as to improve contact between the electrode and the body, and reduce stretch and movement based artifacts, but a gel based intermediary may maintain the lowered impedance throughout a monitoring session, or over a prolonged period of time without causing undue irritation or discomfort for the subject.

[00262] In aspects, the electrode features may be configured and dimensioned so as to penetrate less than 2mm into the skin, less than 1mm, less than 0.5mm, less than 0.2mm, or the like during engagement. In aspects, the electrode features may include one or more microneedles with a length of greater than 0.1mm, greater than 0.25mm, greater than 0.5mm, greater than 1mm, or the like.

[00263] In aspects, one or more electrode features may be coated or impregnated with a medication (e.g. an anesthetic, a steroid, a skin care product, aloe, a therapeutic agent, a pain killer, etc.) configured to attach to or diffuse into the adjacent tissues during engagement thereof. In aspects, the electrode features may include one or more medicated tips (e.g. tips formed from a medication filled biodegradable polymer, sugar, etc.) in the form of an arrow structure, a barb, etc. so as to maintain the electrode feature in place after an initial engagement with an adjacent tissue (i.e. so as to form an indwelling electrode – tissue arrangement). Such a configuration

may improve retention of the tips within the tissues. Biodegradable or soluble tips may be advantageous to provide a slowly disengaging interconnection between the electrode features and the tissues over a desired time frame.

[00264] In aspects, a patch in accordance with the present disclosure may include one or more electrically conducting and/or one or more electrically insulating features (e.g. microfibers, barbs, microneedles, spikes, or the like) arranged so as to intimately interconnect the patch to the patient during a monitoring session. Such a configuration may be advantageous for reducing and/or eliminating the need for an adhesive layer in such a patch configuration.

[00265] In aspects, an electrode including one or more features in accordance with the present disclosure may be dimensioned with an area of less than 4 centimeter (cm)^2, less than 1cm^2, less than 0.5cm^2, less than 0.25cm^2, or the like for monitoring a macro signal such as is related to an EKG, an EEG, a surface EMG, etc. One or more of the electrodes (or one or more microneedle electrode features) may be dimensioned with an exposed electrode area of less than 1000um^2, less than 200um^2, less than 100um^2, less than 50um^2, less than 5um^2, or the like, in the case of the microneedle electrode features optionally exposed at the tip thereof, so as to measure one or more micro-electrophysiological signals, a multi-unit nerve study, or the like. In aspects, a plurality of microneedle electrode elements may be electrically isolated from the others such that a plurality of micro electrophysiological measurements may be made therefrom. In aspects, the array may be coupled with a plurality of preamplifiers, multiplexers, analog to digital converters, comparators, or the like, so as to collectively capture micro-electrophysiological signals therefrom. In aspects, such a configuration may be advantageous for monitoring neural traffic at the surface of the skin, measuring skin sympathetic neural activity, ocular nerve activity, touch analysis, etc.

[00266] In aspects, the microneedles may be configured for providing a stimulating current into the adjacent tissues, the patch and/or module including one or more pulse generators configured to provide the stimulating current. Such a configuration may be advantageous for providing an easy to apply acupuncture system for treating a subject.

[00267] In aspects, a system in accordance with the present disclosure may include one or more patches for attachment to a subject at alternative, seemingly arbitrary locations. The patches may be attached over the body of the subject to investigate physiologic parameters related to specific regions of the body, to determine blood flow propagation, electric field propagation, local muscle response, etc. as discussed throughout the present disclosure.

[00268] In aspects, one or more sensors may be formed from one or more organic electronic elements, patterned versions thereof, semiconducting versions thereof, etc.

[00269] In aspects, other methods for rendering the skin conductive and/or for generating conductive portals on the body include subdermally implantable studs, infusions of conducting materials, infusion of hyaluronic acid and optional in situ gelation thereof in the tissues, in situ polymerization of an intrinsically conducting polymer, application of electroporation (e.g. optionally via a patch/module pair, a handheld device, etc.), abrasion, or the like.

[00270] In aspects, a wide range of functions for microcircuits in accordance with the present disclosure are envisaged, including storing identification information, configuring an inductive interface (i.e. with an antenna embedded into a patch) to provide inductive coupling of one or more patch elements to a corresponding module, etc. In aspects, one or more microcircuit may be embedded into a patch in accordance with the present disclosure, embedded into the housing of a module, etc.

[00271] In aspects, a microstud in accordance with the present disclosure may be configured to form an interconnect between a patch and a mating structure on a module as well as to provide a medium for conducting electricity between the patch and the module during use.

[00272] In aspects, a patch and/or module in accordance with the present disclosure may include means for biasing a sensor included therein towards the subject so as to maintain more intimate contact between the sensor and the subject during a monitoring session.

[00273] In aspects, a system in accordance with the present disclosure may include an isolating patch for interfacing a handheld unit monitoring device with a subject. In aspects, the isolating patch may include one or more subject side electrodes and one or more device side electrodes. Each subject side electrode coupled to one or more of

the device side electrodes so as to provide a means for communicating between the skin of the subject and the handheld unit (i.e. without the handheld unit coming into direct contact with the subject). Such an isolating patch may be advantageous for interfacing a handheld EKG monitoring apparatus (e.g. a hand held unit with contact electrodes, an AliveCor™ module, etc.).

[00274] In aspects, such an isolating patch may include a z-axis conducting adhesive, formed so as to provide a conductive medium through the thickness of the isolating patch without allowing for transfer of biological species there between.

[00275] In aspects, the z-axis conductive adhesive may be patterned such that an operator of the handheld unit may observe one or more visual cues when aligning the handheld unit with the isolating patch in order to interface with the subject.

[00276] In aspects, a patch in accordance with the present disclosure may be configured and dimensioned for attachment to the bridge of a subject's nose (i.e a nasal patch configuration). Such a nasal patch may include an optical sensor and optionally a light source each in accordance with the present disclosure. The optical sensor may be configured to capture light from an adjacent skin surface (i.e. nasal tissues) for purposes of measuring local blood flow, a heart-rate, etc. therefrom. In aspects, the nasal patch may include a flexing member (e.g. a stiffened polymer element, etc.) optionally integrated into, or representing the substrate, the flexing member configured so as to pull upon the sides of the nose after placement, so as to improve airflow there through. Such a configuration may be advantageous for monitoring a subject during exercise while improving performance of the subject (i.e. via improving breathe exchange).

[00277] In aspects, the optical sensor may be arranged so as to capture blood flow readings off of the septum, tissue near the nose (philtrum, at the anterior nares), or along an intranasal wall (vestibule, alar of the nose, alar lobule, etc.). The patch may include the optical sensor coupled with a microcircuit in accordance with the present disclosure into a septum clip, an alar clip (e.g. for placement along the outside or inside of the nose, etc.)

[00278] In aspects, the optical sensor may be configured to measure sPO2, or a signal related thereto near to a line of contact between the nose and the cheek or alternatively at the root of the nose (near the bridge of the nose). In aspects, the

flexing member may be configured so as to bias the sensor towards the tissue during a monitoring session. The sensor may be biased towards the tissues with a pressure of greater than 5mmHg, greater than 15mmHg, greater than 40mmHg, etc. In aspects, the sensor may be biased towards the tissue with a pressure of approximately 5mmHg, approximately 10mmHg, approximately 25mmHg, approximately 35mmHg, or the like.

MODULE FEATURES AND PERFORMANCE ENHANCING ASPECTS:

[00279] A system in accordance with the present disclosure may include one or more modules each in accordance with the present disclosure. The modules may be configured to interface with a patch in accordance with the present disclosure. In aspects, a module may include a power source, a housing, one or more interconnects, signal conditioning circuitry, communication circuitry, a processor, a transceiver, a transducer, one or more sensors, an antenna, a buzzer, a button, a light source, and/or the like, configured to generate one or more signals (e.g. physiologic, electrophysiologic, and/or physical signals) or a feedback signal in accordance with the present disclosure. The signal conditioning circuitry may be configured to amplify, de-noise, pre-filter, generate a trigger, analyze aspects, extract a metric, etc. from one or more physiologic and/or physical signals during a monitoring session, a calibration session, an attachment event, etc.

[00280] In aspects, one or more interconnects may be configured for electrically conducting interconnection, inductively coupled interconnection, capacitively coupled interconnection, between the module and a corresponding patch. In the case of an electrically conducting interconnect, each patch and module interconnect may include complimentary electrically conducting connectors, configured and dimensioned so as to conduct current there between upon attachment.

[00281] In aspects, the module may include a housing, the housing including one or more aspects of the interconnect, optionally providing a sealed cavity in which other components of the module may reside. One or more of the circuits may be integrated into the housing. In aspects, the interconnect may be integrated into one or more circuits included in the housing. The antenna may be integrated into the housing (e.g. printed onto the housing, embedded into the housing, sealed within the housing, etc.).

[00282] In aspects, an interconnect in accordance with the present disclosure may be embedded into one or more of the circuits. The interconnect may be at least partially formed from a through-hole within the circuit (i.e. formed in conjunction with a through hole interconnect on a circuit board of the module). The interconnect may include a cap configured as a surface mount component to form a hermetic seal around the interconnect itself (i.e. so as to isolate the interior of the module from the interconnect).

[00283] In aspects, an interconnect, circuit, circuit board, high density interconnect (HDI) board, or the like may include a magnetic. In aspects, the magnet may be embedded into a circuit of the module (e.g. embedded into a region, attached to, or embedded into a through-hole of the circuit). Such a configuration may be advantageous to simplify assembly of the module and reduce the form factor taken up by the interconnects within the module.

[00284] In aspects, an interconnect and/or circuit board including a magnet may be configured such that the magnet forms the electrical connection to the patch as well as maintains a bias force between the module and the patch during the attachment period.

[00285] In aspects, the module may include one or more sensors each in accordance with the present disclosure. A system in accordance with the present disclosure may include a plurality of module types (i.e. modules with differing sensing aspects). In one non-limiting example, the system includes a kinetic module including an accelerometer, a gyroscope, and/or a magnetometer, a thermal module including one or more temperature sensors, an airflow sensor, and/or a moisture sensor, and an optical module including an optical sensor configured to monitor one or more optical parameter in an associated patch or skin surface. Such module functionality may be included in addition to the communication and power aspects provided to interface with a corresponding patch (which may include one or more sensors, electrodes, etc.).

[00286] In aspects, the interconnect may include one or more snap connectors (i.e. so as to interface with a corresponding microstud based patch, etc.).

[00287] In aspects, one or more modules in accordance with the present disclosure may include one or more sensors and/or microcircuitry configured to interface with

the sensor(s), the sensors configured to monitor one or more physiologic, environmental, and/or physical parameters locally on the subject. Some non-limiting examples of such sensors include electrophysiologic sensors (e.g. EKG, EMG, EEG, ERG, EOG, respiration, bioimpedance, activity, etc.), temperature sensor (e.g. near to the skin, within a module, ambient [environmental], etc.), thermal gradient sensor (e.g. so as to calculate a heat transfer vector locally on the body of the subject, to estimate a core temperature, etc.), barometer, altimeter, accelerometer, gyroscope, humidity sensor, magnetometer, inclinometer, oximeter, colorimetric monitor (e.g. color change analysis of underlying tissue, for respiration, blood flow, pulse, etc.), sweat analyte sensor (e.g. so as to measure sweat constituents, salt content, etc.), galvanic skin response, neural activity (e.g. skin sympathetic activity), interfacial pressure sensors (e.g. for contact assessment, compliance measurement, blood pressure, etc.), flow sensor (e.g. airflow over a module, etc.), surface strain sensor (e.g. via integration of stretch sensors into the patch, evaluation of stretch along one or more electrical interconnect within the patch, integrated capacitive stretch sensors, etc.), a microphone, combinations thereof, and the like.

[00288] In aspects, a module in accordance with the present disclosure may include a three dimensional antenna. The antenna may be arranged along, or embedded within a wall of the module housing (e.g. printed along the wall of the housing, embedded within the outer wall of the housing, particularly in a dome shaped, or pseudo hemi-spherically shaped module, etc.). In aspects, the antenna may be formed from a freestanding conducting structure, suspended above a ground plane situated within the module (e.g. incorporated into a circuit thereof), etc. Such a configuration may be advantageous for boosting a communication range, providing more reliable wireless connections, or the like in the context of monitoring physiologic signals on a subject. Such a configuration may be advantageous for reducing power consumption during wireless communications performed therewith.

[00289] In aspects, the three dimensional antenna may be printed along the inner wall of a suitable shaped housing (i.e. a pseudo hemispherical housing) with one or more pickups arranged along the base thereof. A module in accordance with the present disclosure may include a circuit configured and dimensioned so as to form a wall of the housing, including one or more electrical connections configured to interface with the pickups during assembly. Such a configuration may be

advantageous for providing an improved wireless performance while minimizing the form factor thereof.

[00290] The module may include one or more lights, buttons, buzzers, etc. in order to interact with a user (e.g. a subject, a practitioner, a caregiver, etc.).

[00291] In aspects, a module may include a three dimensional circuit, a stacked circuit, or the like. One or more of the circuit elements may be interconnected with a high density interconnect (HDI) printed circuit board, the HDI printed circuit board optionally configured to function as part of the housing of a module in accordance with the present disclosure.

[00292] In aspects, one or more elements within the module may be encapsulated in a potting material to provide a hermetically sealed element, etc.

[00293] In aspects, a module in accordance with the present disclosure may include a power source (e.g. a battery, a rechargeable battery, an energy harvesting system, etc.). As such, each module may be a self-powered device. In aspects, a module may include a processor and an internal power source.

[00294] The microcircuit may include one or more of signal conditioning circuitry, a system on chip, a processor, a radio, a power management system, an energy harvesting system, a memory module, etc.

[00295] In aspects, the processor may be programmed to operate in a range of power states (e.g. a low power state, a diagnostic state, a monitoring state, a subject detected state, a synchronization state, a calibration state, a communicating state, a recharging state, an alert state, a troubleshooting state, etc.). The processor may operably remain in a low power state so as to improve the lifetime of the power source. The processor may switch between states based on conditions determined via the sensors, a recharge unit, a calibration unit, a host device, etc.

[00296] Each module may be configured to communicate with one or more patches, additional modules, an analysis device, and/or a host device, etc. Such communication may be performed wirelessly (e.g. acoustically, via infrared, via radio frequency communication, etc.) through the environment surrounding the subject, through the body of the subject (e.g. acoustically, optically, capacitively, resistively, and/or inductively coupled signal transmission, etc.). In aspects, one or more patches may relay a combination of an energy signal (e.g. to determine a physiologic

parameter) as well as to communicate an information signal to one or more patches, modules, a host device, etc.

[00297] The processor may be programmed and configured via connection with one or more sensors to determine when a module and/or patch/module pair has been placed onto a subject. The processor via data collected from one or more sensors may be configured to determine the quality of the interface with the subject. In aspects, the patch may include two or more electrode elements to be placed into electrical contact with the subject during a monitoring session. The processor may, via the electrode elements and/or signal conditioning or test electronics attached thereto, estimate the impedance between the electrodes and the body of the subject. If the impedance levels are within acceptable ranges, the processor may initiate collection of bioelectrical information from the subject during a monitoring session. If impedance levels are deemed outside acceptable ranges, the processor may opt not to monitor the subject during the monitoring session. In this case, the module may communicate a "bad connection" signal to one or more modules, patches, an analysis device and/or a host device during a monitoring session. The module may alternatively or in combination send a compromised signal, one or more modules, patches, an analysis device and/or a host device may be used to determine as much information as possible from the signal (e.g. in relation to an EKG example, the signal measured may not be of diagnostic quality, yet detection of the QRS pulse may be adequate for timing blood flow events between patches, determining heart-rate, etc.). As such, analysis of degraded signals may be advantageous for completing a monitoring session with at least a minimum quantity of viable signal information.

[00298] In aspects, the module may include a microcircuit configured to periodically monitor an interconnect to the patch so as to determine if the module has been attached to a corresponding patch or not. Upon identification of attachment to the patch, the microcircuit may perform a wakeup function, a test function, a network communication function, etc. Such detection may be advantageous for hot swapping of monitoring units on a subject to ensure substantially continuous and/or continuous monitoring thereof during a usage scenario.

[00299] In aspects, a module in accordance with the present disclosure may be formed as a watertight and sterilizable component so as to be reused across multiple subjects (e.g. washed and/or sterilized and used across multiple patients, multiple gym

members, multiple students, etc.). In such usage cases, the module may be hermetically sealed, such that fluid ingress into the module is limited. In aspects, the module may be filled with a low dielectric permittivity potting material, so as to limit the potential for fluid ingress during use.

[00300] In aspects, one or more of the patch and/or the module may include registration and/or alignment features to assist a user (e.g. the subject, a physician, a trainer, a caregiver, an emergency medical technician, etc.), with rapidly and/or reliably attaching a module to a patch to initiate a monitoring procedure (or to swap out a module during a monitoring procedure, etc.). Such registration marks and/or alignment features may be utilized by one or more multi-patch coordinating algorithms in accordance with the present disclosure (e.g. so as to assist with more accurately/precisely identifying module/patch placement on the body, identifying particular patches and modules on the subject, for identification and/or orientation purposes if cataloging placement/orientation or determining placement/orientation of modules, via photograph, etc.).

[00301] One or more modules may include electronics and/or software algorithms for detecting one or more fault conditions related to contact with the body of the subject. In aspects, the module may be configured to detect when the impedance between an electrode and the subject is within an acceptable range for measurements (e.g. less than 2Mohm, less than 200kohm, less than 20kohm, less than 2kohm, or the like). In aspects, the electronics may be configured to glean such information by measuring or estimating the impedance between two or more electrode pickups on a coupled patch. In aspects, an impedance estimate may be determined by applying a brief voltage or current pulse to a first electrode, applying a load to a second electrode, and monitoring the temporal response of the second electrode against the first. The rise time of the temporal response compared against the load may be used to indicate the collective impedance of the electrodes and tissues there between.

[00302] In aspects, one or more modules may include a signal source for imparting an energy signal (e.g. electrostatic, electromagnetic, magnetic, vibrational, thermal, optical, etc.) into the body of the subject. The energy signal may be used to communicate to the user, as a form of alert, for diagnostic purposes, to determine a physiologic and/or physical parameter, to configure an array of patches, provide sensation to the subject, etc. In aspects, the energy signal may be emitted into the

body of the subject by a first patch and simultaneously monitored by one or more patches to determine a physiologic parameter of the subject, location based calibration of the patches on the body, etc. In aspects, an identification signal may be merged with the energy signal in order to identify the patch from which the signal was emitted (e.g. so as to automatically configure a patch network, to determine the health of the patch network, to determine a location parameter of a patch with respect to other patches on the body, etc.).

[00303] In aspects, one or more modules and/or patches may include an optical sensor for measuring colorimetric changes in the adjacent tissues during the monitoring process. Such information may be used, optionally in combination with an energy signal to determine one or more optically variable physical parameters and/or one or more optically variable physiologic parameters of the subject, local to the associated patch. In aspects, the optical sensor may be used in combination with one or more optical emitters (e.g. light emitting diodes, laser diodes, bulbs, etc.) to monitor a physiologic signal related to local blood perfusion on the body of the subject. A plurality of such patches may simultaneously monitor such physiologic signals at discrete locations on the body of the subject and relay such information to one or more patches, modules, a host device, and/or an analysis device. The combination of information from such patches or modules may be used to determine blood flow dynamics throughout one or more regions of the body of the subject, to characterize the underlying vasculature in one or more regions of the body of the subject, etc. In aspects, blood perfusion related signals are simultaneously measured at multiple locations on the body of the subject (e.g. the chest, arm[s], leg[s]) and the phase and/or time delays between such signals, as well as the shapes and characteristics of the signals may be used to determine an arterial brachial index of the subject. Such techniques may also be used to determine one or more regions of the subject that may suffer from arterial or venous insufficiency. In aspects, such techniques may be used to estimate the location and/or presence of a blood clot in an extremity of the subject.

[00304] In aspects, one or more modules may include a barometer and/or an altimeter to measure a local environmental parameter (e.g. local pressure, temperature, etc.) during a monitoring session. In aspects, such information may be used to determine the posture of the subject, determine if the subject has fallen, etc.

In aspects, the posture of the subject may be used to determine and/or improve such physiologic measurements as those relating to blood pressure of the subject, correcting EKG data, determining positional relationships between a plurality of patches positioned on the body of the subject, etc.

[00305] In aspects, one or more modules may include an activity sensor (e.g. an accelerometer, a gyroscope, a pedometer, etc.) to measure one or more inertial parameter (e.g. local acceleration, rotation, vibration, etc.) at a location on the body of the subject during a monitoring session. In aspects, information obtained from one or more activity sensors may be used to remove movement artifacts from a physiologic signal, calculate a trajectory, determine a gravitational reference frame, orientation of the module and/or accompanying patch, etc. In aspects, one or more modules may include a tri-axis accelerometer for characterizing the local inertial vector of the body of the subject to which the module is attached. In aspects, one or more modules may include a tri-axis accelerometer, a gyroscope, and optionally a magnetometer. Information from one or more such sensors may be used to calculate an improved local trajectory of the body part of the subject during a monitoring session.

[00306] In aspects, an optical sensor housed within a patch or module may be tailored to monitor a blood flow parameter. In aspects, the optical sensor may be used to monitor a real-time blood perfusion parameter in a tissue of the subject, in the immediate vicinity of the optical sensor. The blood pressure measurement device may include a plurality of such sensors, each sensor configured to monitor a local blood perfusion parameter in the tissue of the subject. Such information may be collected from each sensor in real-time. Correlation of delays, waveform changes, and the like over the body of the subject may be used to generate a correlated signal. In aspects, the correlated signal may be used to create a diagnostic signal (e.g. blood flow volume, blood ejection rate, peripheral vascular parameter, blood oxygen saturation, blood oxygen partial pressure, blood carbon dioxide partial pressure, blood pressure, etc.).

[00307] In aspects, a barometer and/or altimeter housed within a module may be configured with an absolute pressure sensor and optionally a temperature sensor. The altimeter may be sufficiently sensitive to determine local changes in altitude, preferably in the range of decimeters, centimeters, millimeters, etc. The blood pressure measurement device may include a plurality of modules and corresponding

patches, one or more modules including a barometer and/or altimeter, the differences between which may be used to algorithmically coordinate the location of each patch for further measurement analysis. In aspects, information from each altimeter may be used to determine the vertical distance between altimeters on the subject. Such information may be used to determine the posture of the subject, the heart location relative to the sensors, etc.

[00308] In aspects, a module in accordance with the present disclosure may include a recharging connector (or series of contacts), a wireless recharging antenna, or the like for communicating with a recharging bay in accordance with the present disclosure. The recharging connector or antenna may be used to communicate with the recharging bay for purposes of recharging a power source on the module, applying firmware updates, performing diagnostics, or the like.

ALGORITHM ASPECTS:

[00309] A method for monitoring one or more physiologic and/or physical signals from the body of a subject in accordance with the present disclosure includes applying one or more patches each in accordance with the present disclosure to the body of the subject, and attaching a corresponding number of modules each in accordance with the present disclosure to the patches (i.e. so as to form one or more patch/module pairs in accordance with the present disclosure), establishing a body area network among the modules, and collecting physiologic and/or physical signals from the subject using the patches and modules during a monitoring session (i.e. for a period of time suitable for the desired purpose of the method, e.g. 10seconds, 1 minute (min), 1hr, 8hrs, 24hrs, 1week, 1month, 3 months, chronically, etc.).

[00310] In aspects, the method may include storing the collected signals on a memory device (e.g. a memory location on the patches, the modules, a host device, a user device, a datacenter, etc.). In aspects, the body area network may be extended to include a host device in accordance with the present disclosure. The method may include transferring the signals and/or one or more signals and/or metrics derived therefrom from the patches and/or modules to the host device, in real-time, intermittently, in a time synchronous fashion, or the like, during and/or after the monitoring session. In a range of applications, the system may be configured to

monitor for an event (e.g. a change in heart function, a change in EMG, a change in posture, an impact, a change in breathing rate, etc.).

[00311] In aspects, there may be applications where real-time or even pseudo real-time data collection is not necessary (i.e. during aspects of a home sleep study, etc.). In such scenarios, a module in accordance with the present disclosure may be configured to store the collected data locally on a memory device. The module may be configured to download the data to a recharging bay in accordance with the present disclosure at the conclusion of the monitoring session, periodically throughout the monitoring session, or the like in order to transfer the data to a processor for analysis, review, etc.

[00312] In aspects, a method for interacting with a subject with one or more patch/module pairs in accordance with the present disclosure may include measuring one or more physiologic signals therefrom. The method may include deriving a feedback signal, a command, an alert, a metric, a diagnostic value, a schedule, an augmented reality overlay, etc. from one or more of the signals. The method may include identifying when one of the modules requires attention (e.g. the battery is low, a poor interconnection has been made with a corresponding patch, or between a corresponding patch and the subject, a malfunction has occurred, a poor signal quality is being obtained therefrom, etc.). Attention may include swapping the module with a new module, swapping the module out without interrupting the monitoring procedure, removing the module and corresponding patch from the subject, etc.

[00313] In aspects, the method may include providing feedback to a user (e.g. the subject, a physician, a therapist, an officer, a soldier, a group leader, a teacher, a student, an emergency medical technician (EMT), a coach, a trainer, a partner, etc.) relating to the physiologic and/or physical signals. The method may include representing a signal, value, metric, graphic, etc. related to the signals on a feedback component in accordance with the present disclosure (e.g. on a display, a HUD, a wristwatch, an earpiece, a loudspeaker, a tactile display, etc.).

[00314] In aspects, the method may include coordinating the monitoring session across multiple subjects, and optionally synchronizing data collection across the subjects for purposes of calibration, comparative analysis, etc.

[00315] In aspects, a method for identifying a patch/module pair participating in a body area network (BAN) on a subject in accordance with the present disclosure includes placing an object (e.g. a hand, a phone, etc.), over a patch/module pair on the subject while listening to the wireless signal strength of all patches within range, and identifying the patch/module pair by a change in the wireless signal strength measured at a receiving hub (e.g. the phone, a hub, one or more modules, etc.), witnessed during the placing process. Such a configuration may be advantageous for establishing a communication link (i.e. pairing) one or more patch/module pairs on a subject with a host device within an environment with a large number of wireless devices or significant wireless signal clutter (e.g. a gym, a spin class, a conference, an airport, etc.).

[00316] In aspects, one or more steps in the method may be included as part of a BAN initiation routine, to identify placement of one or more modules on a subject, etc. The method may include prompting a user to place a hand over each module in order to identify it on the subject. The method may include prompting the subject with an image, showing where to place the patch/module pairs. In aspects, the method may include assessing the accuracy of the placement of a patch on the subject (e.g. by assessing the physiologic signal obtained therefrom, comparing such readings against a reference, prompting the subject to perform a test routine such as moving an arm, transitioning the subject from a sitting to a standing position, etc.).

[00317] In aspects, a method for identifying a patch/module pair on a subject in accordance with the present disclosure irrespective of the number of patches placed in the vicinity thereof may include, applying one or more patches to the subject (e.g. in a predetermined pattern, substantially random pattern, etc.). Once the system is in communication with and optionally sending data to a host device (e.g. via a wireless link), the host device or a feedback component may prompt the user (e.g. the subject, the care giver, the nurse, the physician, etc.) to place a hand over a patch, a preselected patch, the patch generally nearest to the heart. Upon placement of the hand over the patch, the radio frequency (RF) signal power strengths from the patch in question may change appreciably compared with other RF signal power strengths within communication range and thus a positive identification (ID) can be attained without additional hardware requirements.

[00318] Such a method may be adapted for simplifying the pairing process between a host device (e.g. a smartphone, a wireless local area network (WLAN), etc.), and one or more patches arranged on the subject. In one non-limiting example, such a method may be used to determine which module a user's hand is placed over during a pairing procedure (e.g. perhaps within range of 100s of other wireless devices, modules, etc.). Such a method may be advantageous for use within a gym setting, a fitness class setting, etc.

[00319] In aspects, a system may, at least roughly, determine the general location of one or more patch/module pairs with respect to a host device, by assessing the RF signal strength as received by the host device from one or more of the patch/module pairs in comparison with those received from the other patches in the system.

[00320] In aspects, a modular physiologic monitoring system in accordance with the present disclosure may include a plurality of patches (e.g. patches, patch/module pairs, etc.). In a method of monitoring a subject with such a system, the positioning of the patch/module pairs onto a subject may be visually assessed during placement. One or more patches, modules, or both may include an orientation marker and/or an identifying marker that may be visually assessed from a local observer after placement on the body of the subject. In aspects, the precise placement of the patch/module pairs on the subject may be calculated post attachment by taking an image of the subject after the patches have been placed on the subject. The image may be taken with a coordination device (e.g. a smartphone, a camera, a Kinect™ camera, a HUD ready pair of glasses, Google Glass™, etc.), a host device, etc. In aspects, the orientation markers may be segmented, identified, and extracted from the images to calculate one or more calibration parameters from the orientation of the patches over the body of the subject. In aspects, one or more features associated with the subject (e.g. neck, shoulders, arms, legs, torso, etc.) may be detected and categorized, so as to be incorporated into a patch placement calculation or assessment algorithm.

[00321] In aspects, the coordination device may be used by a user (e.g. the subject, a practitioner, a clinician, a trainer, a coach, a friend, etc.) to take an image of the subject or a portion thereof after placement of the patches. Patch locations and orientations on the subject (e.g. position vectors, positions with respect to anatomical features on the subject, etc.) may be calculated from the image and used to produce a

corrected or standard EKG output, calibrate an EMG based physiotherapy assessment system, automatically assign muscular group behavior to corresponding patches, etc. The system, the host device, the coordination device, etc. may alert a user as to the adverse placement of a patch, the need for more patches, etc. in order to determine a particular cardiovascular function. In aspects, the user may be directed to place one or more additional patches and/or adjust the position of an already placed patch in order to favorably adjust the physiologic data obtained therefrom.

[00322] In aspects, the coordination device may also be used to direct the user to properly place patches on the subject dependent upon the goal of the particular monitoring session. In aspects, an augmented reality display may be employed to direct a clinician to properly place electrodes on the body given the goal of the particular monitoring session (e.g. to assist with placement for EKG, EMG, to match placements from previous sessions, etc.). The augmented reality display may overlay orientation markings onto a camera generated display, highlighting where on the subject the user should place one or more patches in order to better achieve the goals of the indicated monitoring session.

[00323] In aspects, a method for removing a physiologic monitoring system in accordance with the present disclosure may include spraying or wiping an adhesive removing solution onto one or more of the patches, waiting for the adhesive in the patches to disengage from the subject and to remove the patches.

[00324] In aspects, a method in accordance with the present disclosure may include generating one or more metrics from a captured signal (e.g. within a patch, a module, or a patch/module pair in accordance with the present disclosure), prior to wirelessly sending information to another entity in an associated BAN.

[00325] In aspects, data transfer from a module may occur only when a valid physiologic and/or physical signal has been captured from the subject. A module and/or processor coupled thereto may include a function configured to analyze the captured signal in order to determine if a valid reading has been obtained (e.g. such as via detection of a heart-beat, via validation of such signals with those obtained from one or more alternative patches in the system, via detection of substantial movement artifacts, via detection of a low quality electrical interface with the body).

[00326] In aspects, a module in accordance with the present disclosure may include one or more algorithms (e.g. implemented on a processor, SoC, etc.) configured to analyze the signals obtained from the subject. In aspects, an algorithm in accordance with the present disclosure may be configured to extract a metric from the signal including a heart-beat, time-stamping of a QRS complex, etc. (or other metrics as described herein). In aspects, to save on wireless bandwidth and associated power consumption, a module may include an algorithm to efficiently extract such metrics from the raw data and send the metrics rather than the raw data. In aspects, the modules may include multiple modes of operation (e.g. a low priority mode, a high priority mode). Some modes may be configured so as to send small amounts of data (i.e. such as when a heart-rate or monitored function of a subject is within a 'normal' range), or metrics extracted from the raw data (e.g. a simple heart-rate, etc.) so as to maintain a low wireless bandwidth. Some modes may be configured so as to send all available data (i.e. such as when an 'event' is occurring, when a previously 'normal' signal changes) so as to provide a user with as much information as possible during the 'event'. Such a configuration may be advantageous to balance power consumption of hardware within the modules with the depth of the monitored signal.

[00327] In aspects, when configured for monitoring of an electrocardiographic signal, a module may be configured to adjust the sampling rate thereof with an adaptive sampling algorithm (e.g. a local adaptive sampling algorithm, etc.). In aspects, an adaptive sampling algorithm may be constructed to use a timestamp associated with the QRS complex. In aspects, the sampling algorithm may be configured to increase sampling rate to just beyond the frequency determined by the Nyquist-Shannon sampling theorem for the given heart-rate of the subject (i.e. adjustable with heart-rate), and the other samples obtained within a single heart-beat, may be taken at a much lower rate (i.e. as the signal generally does not include the same high frequency content in those regions of the beat). Such a configuration may be advantageous to reduce power consumption within a module, decrease the amount of data sent wirelessly from the module, etc.

[00328] In aspects, the BAN may implicitly extend to a recognized user device (e.g. a HUD ready pair of glasses worn by a user, a coach's wrist watch, etc.) when that entity enters the range of the BAN. Thus one or more signals, metrics, identification records, etc. relating to the subject or monitoring session may

automatically start streaming to the user device upon recognition. In aspects, such a configuration may be advantageous for seamlessly brining new users into a monitoring session without requiring extensive effort on the part of the user.

[00329] In aspects, a method in accordance with the present disclosure may include determining a priority metric for one or more signals captured by a module in the system (e.g. via assigning a priority level, determining a degree of redundancy, etc.). Such a priority metric may be used in an algorithm to determine the type and urgency of an "alert" generated by a failure on one or more modules in the system. In one non-limiting example, a system including 5 modules is deployed onto a subject to monitor a 3 lead equivalent EKG in accordance with the present disclosure. A priority metric for the system is determined based on the number of modules that would have to fail in order to risk obtaining a low quality EKG from the subject. In aspects, the priority may be more or less affected by the removal of one or more modules in the system (i.e. based upon the location of the module on the subject), etc. If a module on the subject fails or indicates that it is about to fail (i.e. a battery low alert). The system may be configured to assess how such a failure will affect the priority metric, thus adjusting a potential alert accordingly (i.e. from "do nothing" through to "needs immediate attention"). Such a configuration may be advantageous for reducing false alarms within a hospital setting, thus reducing alarm fatigue, or the like while providing more robust monitoring of EKG.

[00330] A method for managing a monitoring session with a system in accordance with the present disclosure may include, mounting one or more modules on a recharge bay. Diagnosing function of and/or recharging one or more modules with the recharge bay. Indicating to a user when a module in the recharge bay is ready for use. Swapping a module from the body to the recharge bay and vice versa. Integrating a newly placed module into the BAN and initiating the streaming of signals therefrom.

[00331] In aspects, a method in accordance with the present disclosure may include generating a local field vector from a patch/module pair including three or more electrodes. The method may include calculating a vector electrocardiogram from a collection of local field vectors, or the like. The method may include displaying one or more field vectors over an image of the subject to assist with the assessment thereof.

AESTHETIC DESIGN ASPECTS:

[00332] In aspects, the patch may be shaped so as to contour a specific region of the body so as to better capture signals there from during a monitoring session. In aspects, the modules may be designed so as to maintain a low profile and with no sharp edges, etc. in order to better hide the modules and/or reduce snags during a procedure.

[00333] In aspects, the modules may be formed as hemispheres or pseudo hemispheres for purposes disclosed herein. In aspects, the modules may include one or more low profile interconnects arranged along an underside thereof, to interface with a corresponding patch for purposes of monitoring.

[00334] The modules may include one or more orientation markings to assist with alignment of the modules onto one or more patches on the body, automatically determine orientation and positioning of the modules one the subject (e.g. with a camera, a video system, etc.).

MONITORING KIT ASPECTS:

[00335] A system in accordance with the present disclosure may be provided as part of a monitoring kit. In aspects, a monitoring kit in accordance with the present disclosure may include one or more modules, a recharging bay, one or more patches, or set of patches (i.e. a series of patches configured and dimensioned to perform a particular type of monitoring on a subject), and (optionally) one or more accessories such as an adhesive removal wipe/spray, skin preparation tools, instructions, software access, etc.

[00336] In aspects, the recharging bay may be configured to hold one or more modules each in accordance with the present disclosure. The recharging bay may be configured so as to act as a host device (e.g. as a wireless hub, etc.) so as to provide multiple functions for a user.

[00337] The recharging bay may include one or more connectors to interface with each module. The connectors may be configured to provide recharging services for the modules, to interface with a processor on the module, to transfer data (e.g. stored session data, etc.), to perform firmware updates, to assess functionality of the module (i.e. to test interconnects of the module, onboard circuitry, etc.). The recharging bay

may include one or more status or diagnostic indicators there upon, to identify if a held module is ready for use, recharging, if there is an error with the module, etc.

[00338] The recharging bay may be configured for mounting to a wall, such as near to the bed of a subject, etc. so as to facilitate simplified hot swapping of modules to the subject, etc.

[00339] The recharging bay may be configured to interface with a host device, a user device, etc. for purposes of monitoring the subject, and/or providing alerts, assisting with configuration of a system in accordance with the present disclosure, etc.

[00340] In aspects, the kit may include a patch set configured for providing specific functionality on a subject, or sized to fit the subject (e.g. small, medium, large, custom, etc.). In aspects, the patch set may be configured to perform a 3 lead EKG, a redundant 12 lead EKG, a biomechanic assessment (e.g. for a leg, a hand, an arm, a foot, etc.), a sleep lab assessment, an impact study, a sport specific application, a heart-rate monitoring session, a thermal overload monitoring session (e.g. for monitoring for heat exhaustion, etc.), a weight lifting workout, etc.

[00341] In aspects, there is provided, an adhesive test kit for determining preferences of a user prior to performing a monitoring session in accordance with the present disclosure. The adhesive test kit may include a series of patches, each patch with identification markings and an associated adhesive (i.e. each with an alternative type of adhesive). The adhesive test kit may be provided to a perspective customer, so as to choose the adhesive type that best suits their needs (e.g. the adhesive with the best long-term wear ability for their skin type, for their diet, adhesive with the lowest irritation for that person, etc.).

[00342] In aspects, the adhesive test kit may include a plurality of test patches with a range of adhesive options for potential customers to test during a workout, in a shower, during rest, etc. in order to determine which best suits their needs, best limit skin irritation, best adhere to their skin, etc. In aspects, a customer to a service plan in accordance with the present disclosure may receive periodic adhesive test kits to determine if alternative adhesives would better serve their changing needs, etc.

[00343] In aspects, one or more adhesive samples in the kit may include polyacrylate, polyisobutylene, polysiloxane, polyurethane, pressure sensitive adhesive, amphiphilic adhesive, polyvinylpyrrolidone (PVP), polyvinyl Alcohol

(PVA), polyethylene glycol (PEG), or poly (2-acrylamido-2-methyl propyl sulfonic acid (PAMPS) based hydrogel adhesives, combinations thereof, or the like.

[00344] As part of a service plan, an adhesive test kit may be provided to a customer. The customer may wear the samples provided in the kit for a suitable period of time depending on the intended application, to determine which adhesive is best for them (e.g. most compatible with their skin, wearable, reliable, etc.).

[00345] The test patches may be configured with additional weight, etc. to simulate the attachment of a module thereto. One or more test patches may include an impedance monitoring circuit to assess the reliability of a monitoring process provided therewith.

[00346] The recharging bay may be integrated into a display (e.g. into the chassis of a signal display system, etc.). Such a display may provide a means for a user to interact with the recharging bay, provide data, metrics, or visualization of the signals, etc. Such a display may be advantageous for use within a hospital setting, as a monitor and/or analyzer in an intensive care unit (ICU), an operating room (OR), etc. with a cluster of modules, patient interfaces and hand-held coordination devices.

BUSINESS METHOD ASPECTS:

[00347] According to aspects, there is provided a service (i.e. optionally embodied in the form of a service system) for managing the collection of physiologic data from a customer, including a subscription service in which a customer is assigned a profile including usage related information (e.g. purposes for monitoring, billing, etc.), an automated patch delivery system configured to provide a customer with a timely series of patches for use in one or more associated monitoring sessions, a datacenter configured to store, analyze, etc. the data obtained from the customer during one or more monitoring sessions, a report generating service configured to generate one or more monitoring reports based upon the data obtained during one or more monitoring sessions, and a recurrent billing system configured to bill the customer based upon the number or patches consumed, the data stored, the reports generated, or the like.

[00348] The service may include a module replacement system configured to identify from the data when a customer may require additional modules and to include such modules along with a replacement patch set delivery, for immediate delivery, etc. depending on the particular needs of the customer.

[00349] The service may include one or more application programming interfaces (APIs) whereby one or more 3rd parties (e.g. a data repositories, government agencies, pharmaceutical companies, health data providers, EHRs, a social network, an employer, a healthcare provider, etc.) may access the data, reports generated therefrom, etc.

[00350] In aspects, the service may include an automated adhesive selection algorithm configured to assign a customer preference for adhesives in a patch set based upon the outcome of an adhesive test (i.e. performed via an adhesive test kit in accordance with the present disclosure).

[00351] In aspects, the service may include an automated algorithm for assessing the performance of a patch set on a customer from the data. Such an algorithm may be configured to identify if a customer is experiencing irregular, erratic, problematic, shortened, wear times, etc. and may issue an alert to contact the customer, issue an alternative adhesive preference, issue for delivery of a complimentary adhesive test kit to the customer, etc.

[00352] In aspects, the service may include one or more algorithms for providing remote coaching, stress management, physiotherapy, and/or related services to a customer, hospital, clinic, etc.

GENERAL DISCUSSION ON HOST DEVICES:

[00353] In aspects, the host device may be operably worn/held by the subject, located near to the subject, integrated into a bedside alarm clock, or housed in an accessory (e.g. a purse, a backpack, a wallet, etc.). In aspects, the host device may be a mobile computing device (e.g. a smartphone, a tablet computer, a pager, etc.). In aspects, the host device may be a local router, a data recorder, a network hub, a server, a secondary mobile computing device, a router, a repeater, etc.

[00354] In aspects, the host device may be a dongle or accessory for a mobile computing device. In such aspects, the host device may be configured to coordinate communication with one or more patches/modules, analyze incoming patch data, fuse sensor information from one or more patches, condition and/or de-noise information signals obtained from one or more patches, correlate connectivity of one or more patches, to reconstruct signals from parameters sent by one or more patches/modules,

or the like. In aspects, the host device may be configured to generate one or more physiologic signals, alerts, etc. therefrom.

[00355] In aspects, one or more patches, modules, a host device, user device, and/or an analysis device may fuse sensory information from one or more patches during a monitoring session. If sensory information is missing from a particular patch, module, etc. or if it is in some way compromised, etc. the one or more patches, modules, the host device and/or the analysis device may ignore, remove, de-emphasize, etc. the information. As such, the system may be advantageous for providing a robust, fault tolerance means for monitoring one or more physiologic parameters of a subject.

[00356] In aspects, one or more patches, modules, a host device, a user device, and/or an analysis device may generate various levels of alerts for maintaining the monitoring session during a long-term monitoring session on a subject. Such alerts may be related to a subject emergency (e.g. a fall, a heart arrhythmia, a neurological arrhythmia [e.g. due to a seizure], an elevated heart-rate, syncope, an accident, an impact, a sleep apnea event, respiratory arrhythmia, choking, a drop in arterial carbon dioxide (CO2), hypercapnia, a missing heart-rate, etc.), a moderate priority maintenance need (e.g. a high number of compromised signals, a high number of low or depleted power sources, etc.), a low priority maintenance need (e.g. a limited number of compromised signals, one or more low battery indications, etc.).

[00357] In aspects, one or more patches, modules, a host device and/or an analysis device may generate an "information quality" signal related to the overall quality of one or more signals (e.g. individual information signals, a collective signal, a physiologic parameter, etc.) related to one or more patches on the subject, and/or the overall system. Such an "information quality" signal may be used to determine and/or convey the degree of confidence that the system has in the physiologic parameters of a subject being measured during a monitoring session. The signal may be good, average, compromised, poor, unacceptable and/or the like. An alert may be advantageously constructed from the information quality signal so as to optimally compromise between functionality (e.g. basic quality of the monitoring session) and productivity (e.g. number of alerts requiring attention) during a monitoring session.

[00358] A modular physiologic monitoring system in accordance with the present disclosure may include a host device in communication with one or more patches. The host device may coordinate the monitoring network, etc. One or more patches, and/or the host device may be configured in operable electrical communication with each other. In aspects, the patches and/or the host may be configured to communicate with each other during a calibration procedure (e.g. so as to calibrate the altimeters). Such calibration may be advantageous for improving the quality of the monitoring session.

[00359] The host device may communicate with an external network (e.g. a WiFi network, a cellular network, a LAN, etc.). In aspects, the host device may be a stationary device wired to a local area network (LAN), configured to communicate one or more signals and/or information derived therefrom to another device, via the LAN.

[00360] In aspects, the host device may be configured to monitor the wireless signal strength received from one or more patch/module pairs in an associated BAN and coordinate signal transfer from patch/module pairs to the hub, or between each other, so as to maximize the strength of the BAN + hub network (i.e. in the case that the hub is not situated on the body of the subject). In aspects, the hub may coordinate that a module with a poor signal transfer data to high signal strength module on the subject, and the high signal strength module relay the data to the hub (if the hub communicates that the data was not received, if a signal could not be obtained directly from the module, etc.).

ASSESSMENT OF NEURAL ACTIVITY

[00361] In aspects, a system, device, method, and/or component in accordance with the present disclosure may be used for assessing the neural tone of a subject (e.g. neural tone associated with a region of skin of a subject, tone associated with an organ of a subject, tone associated with neuroendocrine function in a body, signals, traffic, etc. associated with central and/or peripheral neural traffic between a brain, ganglion, neural structure, and an organ, etc.). The discussion now turns to discussion of non-invasive systems and methods for determining a state of one or more aspects of an autonomic neural system (ANS) of a subject, determining a relationship between the state of one or more aspects of an ANS and a stress test, determining the outcome of a

neural traffic modifying procedure, determining if a subject is a suitable candidate for a procedure, medical treatment, combinations thereof, and the like.

[00362] Signals traveling through the autonomic nerves system of a subject include bidirectional signals: afferent, efferent traffic. Efferent traffic can trigger changes in different parts of the body simultaneously. Relating to some non-limiting examples, the sympathetic nervous system can accelerate heart rate; widen bronchial passages, decrease motility (movement) of the large intestine; constrict blood vessels, increase peristalsis in the esophagus, cause pupillary dilation, piloerection (goose bumps) and perspiration (sweating), raise blood pressure, etc. The parasympathetic system can affect various systems and bodily functions as well, generally in an approach opposing the action of the sympathetic system. The differential traffic between the sympathetic nervous system (SNS) and peripheral nervous system (PNS) innervating a particular organ may be as important to the overall function of that organ, as the individual afferent/efferent traffic of each neural network.

[00363] Relating to aspects, one or more synapses in the skin (preganglionic neuron to postganglionic neuron) may be mediated by nicotinic receptors activated by acetylcholine (a neurotransmitter), one or more synapses of the postganglionic neuron may be mediated by adrenergic receptors and may be activated by either noradrenaline (norepinephrine) or adrenaline (epinephrine). Sweat glands receive sympathetic innervation but include muscarinic acetylcholine receptors, which are normally characteristic of the parasympathetic nervous system. Other exceptions exist, such as with certain deep muscle blood vessels, which dilate (rather than constrict) with an increase in sympathetic tone. This is because of the presence of more beta2 receptors, rather than alpha1, which are more frequently found on other vessels of the body. Traffic associated with such nerves may be monitored, blocked, stimulated, and or assessed with a system, device, patch, patch/module pair, and/or method each in accordance with the present disclosure.

[00364] Such systems may be advantageous for assessing a disease state of a subject, an autonomic neural disorder, a peripheral neuropathy, the extent of a neural block (e.g. such as via a local analgesic, application of a neuro-blocker, a neurotoxin, etc.), the state of a neural block, a neuroendocrine relationship, a state of a sympathetic neural branch, a state of parasympathetic neural branch, etc.

[00365] In aspects, a patch and/or module in accordance with the present disclosure may include a plurality of electrodes, microelectrodes, or the like for assessing a skin neural activity, skin sympathetic neural activity, skin somatosensory neural activity, skin parasympathetic neural activity, combinations thereof, or the like. Additionally, alternatively, or in combination a patch and/or module may include a sensor for assessing local hydration, galvanic skin response, or the like in accordance with the present disclosure, an optical sensor for assessing a blood perfusion and/or oxygenation, etc., combinations thereof, or the like. Such combinations may be advantageous to assess differing aspects of a local neural response to a stimulus, a stressor, a procedure, etc.

[00366] In aspects, a patch and/or module may be configured to assess skin neural tone (autonomic, somatosensory, sympathetic, parasympathetic, follicular erection, smooth muscle neural activity, vascular contraction/dilation, etc.), in combination with blood perfusion, and/or local hydration (e.g. due to sweating, exudate migration, etc.), or the like. In aspects, a system in accordance with the present disclosure may include one or more patch module pairs configured to monitor skin neural tone at one or more sites on a body, in combination with one or more patch module pairs configured to monitor EKG, heart rate, heart rate variability, breathing rate, breathing effort, muscle tone, tissue hydration, sweating, blood perfusion to tissues, combinations thereof, or the like.

[00367] Additionally, alternatively, or in combination a neural tone may be extracted from an EKG recording (such as separated from the noise floor thereof), so as to further assess an autonomic neural state from a subject.

[00368] Such systems may be used to determine the autonomic neural state of a subject (i.e. a state of the autonomic nervous system [ANS]), to determine the sympathetic and/or parasympathetic component of the autonomic state of a subject, to determine the state of a branch of the ANS, to determine the relationship between the ANS state, a branch thereof, and/or a change in state thereof and a change in organ function, to determine a contribution of afferent traffic from an organ in a subject to the autonomic neural state of the subject or to changes thereof, to determine a relationship (e.g. a qualitative relationship, a causal relationship, a quantitative relationship, a transfer function, etc.) between an ANS state of a subject and an input parameter, a stress test, a medical procedure, delivery of a medication, a change in

state of an organ in the body, the outcome of an interventional procedure, a neural ablation, a combination thereof, or the like.

[00369] In aspects, such systems may be configured to assess changes an overall autonomic neural state of a subject as influenced by a subsystem of the body, a change in neural traffic along a nerve, nerve plexus, ganglion, receptors, efferent/afferent traffic, and/or sensory traffic from one or more sites in a subject (i.e. so as to establish a cause – effect relationship between the target and the overall ANS state). Alternatively, additionally, or in combination, such systems may be configured to actively influence the autonomic neural state of a subject while monitoring changes in one or more organ states, organ functions, or the like (i.e. so as to establish an ANS – organ functional influence). In aspects, a patch module pair in accordance with the present disclosure may be configured to apply a stress state to a first tissue site, one or more additional patch module pairs configured to monitor one or more forms of neural traffic, and/or surrogates thereof at one or more additional sites on the body of the subject.

[00370] Alternatively, additionally, or in combination a system in accordance with the present disclosure may include and/or may be configured to work in conjunction with a therapeutic system in accordance with the present disclosure. Such a therapeutic system may include an ablation system, a neuromodulation device/implant, an ablation catheter, a focused energy delivery device, a radio frequency ablation system or catheter, a microwave ablation system or catheter, an ultrasound energy delivery system (e.g. a high intensity focused ultrasound [HIFU] system, catheter, or the like) or catheter, a cryoablation system or catheter, a chemical ablation system or catheter, a radiosurgical system, an optical ablation system (e.g. an infrared ablation system, a laser ablation system, etc.), a magnetic resonance (MR) guided HIFU system, a combination thereof or the like. In aspects, the system may be configured to temporarily and/or substantially permanently alter the neurological state of one or more nerves in a subject through a procedure (e.g. delivery of energy, delivery of a chemical, stimulation, etc.), the system configured to monitor one or more aspects of a neural state in the subject to determine the completion of such a procedure. Such monitoring may be performed so as to measure activity of a related branch of the ANS, a surrogate physiologic parameter associated therewith, tone associated with at least a portion of the sympathetic or parasympathetic nervous

system, a relationship between a cursory stress test and the ANS (e.g. a change in the results of a stress test applied to a subject pre and post procedure), or the like.

[00371] In aspects, a system in accordance with the present disclosure may be used to determine if a subject is a suitable candidate for a procedure, such as a neuromodulation procedure, a neural ablation procedure, a sympathectomy, a peripheral neural block, or the like. Such an assessment may be determined by comparing the functional relationship between the ANS or an aspect thereof, with one or more stress states, over a range of stress stimulating inputs, as assessed during a stress test, combinations thereof, or the like. Subject inclusion / exclusion criteria may be developed around one or more metrics generated from one or more stress tests completed on the subject in accordance with the present disclosure, a baseline autonomic, sympathetic, and/or parasympathetic tone of a subject as measured with a system, device, or method in accordance with the present disclosure, etc.

[00372] Some non-limiting examples of stress tests in accordance with the present disclosure include administration of a chemical, a drug, medicament, a hormone, an enzyme, a diuretic, a solution, electrolytes, a peptide, steroid, a combination thereof, or the like to a subject (e.g. topical, systemic, intravenous, intra-arterial, intra-parenchymal, sub-dermal delivery, transdermal delivery, rectal, vaginal, urethral, or nasal suppositories, in-hailed, or the like), delivery of energy, stimulation, electrical stimulation, presenting an audio field to a subject, application of thermal stress, a light field, an image, asking the subject a question, playing music, generating an audible signal for the subject, a change in humidity, a tactile input (e.g. to one or more sites on the body, to a region of skin, to a carotid sinus, to a carotid body, to a baroreceptor, to a vagus nerve receptor, to an erogenous zone on the skin, etc.), application of an electromagnetic field, injection of a current, application of pressure, application of stroking to a region of skin, a change in barometric pressure, a change in posture, an exercise, a breath holding exercise, a stair climbing exercise, to evoke an emotional response therefrom, to alter an environmental state thereabout, and/or combinations thereof to a subject or one or more sites there upon or therein. The stress tests may be devised and implemented so as to cause a differential response between the parasympathetic state and the sympathetic state of a subject (e.g. overall, of a branch of the ANS, as relating to afferent traffic of one or more parts of the ANS, or the like).

[00373] The stress test may include having the subject perform a Valsalva maneuver, a tilt table test, elevating one or more legs, transient siting to standing exercises, execute a change in posture, move from a prone position to a sitting or standing position, a breath hold technique, or combinations thereof. In aspects, the stress test may include injection or infusion of a vasodilator (e.g. endothelium-derived hyperpolarizing factor (EDHF), potassium, nitric oxide, β-2 adrenergic receptors, histamine, prostacyclin, prostaglandin, vasoactive intestinal peptides, adenosine, adenosine triphosphate (ATP), adenosine diphosphate (ADP), L-arginine, bradykinin, substance P, niacin, CO2, etc.), or a vasoconstrictor (e.g. ATP, muscarinic agents, acetylcholine, neuropeptide Y (NPY), adrenergic agonists, epinephrine, norepinephrine, dopamine, thromboxane, endothelin, angiotensin II, asymmetric dimethylarginine, antidiuretic hormone, vasopressin, etc.), a neuroblocker, a neurostimulant, a diuretic, insulin, glucose, beta-adrenergic receptor antagonist, angiotensin-II converting enzyme inhibitor, calcium channel blocker, an 3-hydroxy-3-methylglutaryl-coenzyme A (HMG-CoA) reductase inhibitor, digoxin, anticoagulants, beta blockers, angiotensin-converting-enzyme (ACE) inhibitors, one or more steroids (e.g. diflorasone, betamethasone, dexamethasone, clobetasol, prednisolone, mometasone, methylprednisolone, Deprodone, difluprednate, fluocinonide, amcinonide, triamcinolone, difluprednate, hydrocortisone, etc.), testosterone, or the like, into the body of the subject, into an organ of the subject, into a lumen of the subject, into an artery, a vein, a renal artery, into one or more of the monitoring sites, etc.

[00374] In aspects, a stress test may include having a subject breath air that is a different temperature than that of the skin of the subject and/or have air of a different temperature blown onto the skin thereof, so as to elicit a strong ANS response. In aspects, a warm air gust may substantially stimulate more of the parasympathetic system of the subject and a cold air gust may stimulate the SNS of the subject.

[00375] In aspects, the stress test may be devised so as to elucidate a differential response between aspects of the sympathetic state of a subject, a branch of the sympathetic nervous system, and a parasympathetic state of a subject, an initial neural state, etc. One non-limiting example may include assessment of acetylcholine nerve states separately from that of epinephrine nerve states, or that of substance P, cholinergic, and/or vesicular acetylcholinetransporter (VAChT) nerve states. In

aspects, such differentiation may be determined by assessing neural traffic at a plurality of sites on the body such as near a particularly sweaty region of the skin to assess acetylcholine nerve states, or near a muscular structure for assessment of epinephrine related nerve states, simultaneously so as to assess relationships there between, near an erogenous zone for assessment of cholinergic or vesicular acetylcholinetransporter related nerves, and the like. Comparison between a plurality of responses recorded over alternative regions of skin of a subject may be advantageous for extracting independent measures of the functionality of one or more branches of the autonomic nervous system, sympathetic nerves, parasympathetic nerves, somatosensory nerves, or the like.

[00376] In aspects, a method may include delivery of an alpha or beta agonist intravenously or intra arterially to a subject and following the response of one or more nerves with a system, device, or method in accordance with the present disclosure to determine the state, response, suitability for treatment, thereof, etc. Such a stress test may include monitoring the activity for a period of greater than 30 seconds, greater than 1 minute, greater than 5 minutes, greater than 15 minutes after initiation of the stress test, etc. The analysis may include comparing the neural response associated with one or more tissue sites of the subject, generating an overall assessment of traffic (e.g. an assessment of tone, afferent traffic, efferent traffic, etc.), for use in the comparison, etc. The analysis may include mapping the input strength – output intensity of the relationship between the stress test input and the monitored neural response, or surrogate thereof.

[00377] In aspects, such an approach may be advantageous to assess and/or troubleshoot a disease state, treatment, medication effect, etc. on a subject, troubleshooting and/or state characterization of various autonomic functions of a subject, individual assessment of sympathetic and parasympathetic state of a subject, or the like.

[00378] Some non-limiting examples of uses for such a system include, assessment of autonomic function of a subject, CNS disorders, assessing impact of a medication on the ANS of a subject, assessing medication dosage parameters (e.g. personalizing medication for a subject, personalizing dosage for a subject, timing dosage delivery for a subject, assessing periods of activity for a medication on a subject, assessing pharmacokinetics of a substance on a subject, assessing differences between

bioavailability of substances, assessing effectiveness of a generic medication on a subject, assessing a difference in delivery rate between medications, etc.), hypohidrosis, hyperhidrosis, neuroendocrine function, suitability for a denervation procedure, suitability for a renal or carotid body denervation procedure, diabetic neuropathy assessment, peripheral neuropathy, analgesic feedback, neural block feedback, in-procedure feedback, procedural follow up, cardiac conditions, lie detection, assessment of sexual dysfunction, psychiatric assessment, urinary/fecal incontinence, combinations thereof, and the like.

[00379] In aspects, the posture of the subject may be altered during the monitoring (such as via a tilt table), alteration of a peripheral resistance (e.g. such as via application / removal of a tourniquet, etc.), a sudden rise or fall in a peripheral resistance, a change in local barometric pressure, a sudden decompression, a sudden compression, a sitting to standing movement, a Valsalva maneuver, an exercise, or the like may be performed during monitoring to further assess aspects of the ANS of the subject.

[00380] In aspects, a stress test may include applying a visual, olfactory, gustatory "taste", and/or audible field or experience to a subject, while monitoring a neural response or surrogate thereof. Such sensation of sight, smell, sound, taste, or the like may evoke and ANS response from individuals. SNS or PNS outflow associated with such a response may be monitored with a system, device, patch, and/or method in accordance with the present disclosure. Abnormal responses may be determined during such tests, the abnormal response being an indication of a disease state, suitability, or unsuitability for a treatment, etc.

[00381] In aspects, a stress test in accordance with the present disclosure may include application of a thermal input (such as heating, cooling, or regulating the response of), a region of skin of a subject, airflow around a subject, air breathed by a subject, or the like. In aspects, such a stress test may be performed by a system, or device in accordance with the present disclosure, the device including a thermal regulating unit (e.g. a thermoelectric device, a Peltier device, an endothermic reactive specie, an exothermic reactive specie, a temperature regulating gel, a fluid cooling/heating system, etc.). In aspects, the device and/or system may be configured to monitor the generated afferent traffic associated with the thermal stress that may adjust the SNS/PNS differential relationship in the body, an SNS/PNS output or

surrogate thereof associated with the changes in ANS caused by application of the thermal stress state etc. Such outflow may be monitored with one or more devices in accordance with the present disclosure (at an alternative site on the body, at the same site on the body as the thermal stress application, etc.). An associated system may be configured so as to monitor changes in PNS related outflow, SNS related outflow, somatosensory response, etc. and determine the relationships associated therewith. In aspects, the system may include one or more sensors located within or near to the thermal regulating site, the sensors configured to monitor afferent traffic generated in the skin by the change in thermal load thereupon. Simulatanously, additionally, alternatively, or in combination, the system may include one or more sensors each in accordance with the present disclosure, each sensor located at one or more alternative sites on the body, configured to monitor corresponding outflow from the ANS caused, at least in part by the stimulus. Such a system may be advantageous to map correlating ANS inputs (e.g. afferent traffic from various ANS coupled organs, eye, skin, audio, scent, taste, thermal, tactile skin response, etc.), to changes in the ANS outflows (e.g. SNS outflow, PNS outflow, changes in branches, changes in physiologic parameters associated with the ANS outflow, such as changes in HR, HRV, BP, etc.). By varying the stimulatory loads during stress tests, a processor programmed with an appropriate stress evaluation algorithm may be suitably configured to generate a transfer function for the subject in this regard (e.g. a transfer function, a single-input multi-output (SIMO) transfer function, a multi-input multi-output (MIMO) transfer function, or the like relating the stress inputs to the monitored afferent, efferent, and outflows). Such a transfer function may be suitably derived and/or compared between tests performed before/during/after a medical procedure, etc. Changes in the test observed after a procedure may be used to determine the extent of the procedure, etc.

[00382] In aspects, a stress test may include a subsystem in accordance with the present disclosure for the application of a tactile input to one or more regions of the skin of a subject. Such tactile input may include penetration, penetration-like movements, vibratory movement, stroking, lateral movements, rotary vibrations, or the like. In aspects, the site of the excitation may be selected so as to excite particular regions of the ANS (e.g. such as exciting the parasympathetic nervous system, via tactile input to one or more parasympathetic receptors located near to the ear, etc.).

[00383] In aspects, energy, tactile input, or the like may be applied to a carotid sinus, carotid body, etc. In aspects, energy delivery may be provided by a device in accordance with the present disclosure. In aspects, a systemic variable, such as a change in blood pressure, heart rate, heart rate variability, neural activity, skin SNA, etc. (such as may be monitored by a device in accordance with the present disclosure) may provide a global feedback that the stimulus, energy input, etc. was delivered to the carotid sinus, and reflect the degree of excitation thereof for the subject, one or more devices in accordance with the present disclosure may be arranged on the body so as to monitor an associated change in the ANS, the PNS, the SNS, an overall change, in a branch of the ANS, such as a skin branch, may be representative of the ANS, SNS, and/or PNS outflow generated therefrom. In aspects, such a stress test may be used to determine the suitability of a subject for a procedure, a sympathectomy, a neuromodulation implant, a neural ablation procedure, a renal denervation procedure, etc.

[00384] In aspects, the amount of change in one or more of the monitored signals, whether the signals recovered near to the original values after the test, how quickly the signals changed, recovered, differential changes between signals, etc. may be considered in deciding if the subject is a suitable candidate for a procedure, a therapy, an implant, or the like.

[00385] In aspects, a stress test may include altering an environment around a subject, such as changing the temperature, ambient light levels, humidity, airflow, etc. In aspects, such ambient changes may be monitored with a sensor in accordance with the present disclosure, the magnitudes of the changes compared against the response in the generation of a transfer function, a response map, etc.

[00386] In aspects, a system in accordance with the present disclosure may include one or more components, sensors, and/or subsystems for assessing a change in a state of the autonomic neural system (ANS) and/or a relationship between a component of the ANS and a stress test at a site that is non-invasively accessible (inter aural, intra nasally, salivary, skin sites <groin, armpit, neck, anal-rectal region, palm of hand, sole of foot> eye, pupil, iris, iris EMG, EOG, ERG, etc.). In aspects, one or more neural activity sensing elements may be configured and arranged near the trachea of a subject so as to assess the larynx tone during application of a stress test there upon. Such an assessment may be a suitable surrogate for PNS activity along the nearby

vagus nerve plexus. In aspects, such activity may be monitored with one or more multi-sensor patches in accordance with the present disclosure.

[00387] In aspects, the system may include one or more devices to monitor one or more of ocular neural tone, facial muscular tone, electroretinography, nasalis muscular tone, temporalis tone, zygonaticus tone, orbicularis tone, occipitofrontalis tone, etc. Such tone may be assessed and change as the relationship between the overall SNS and PNS of a subject change, during the stress state, during a procedure, after completion of a procedure, etc.

[00388] In aspects, the system may include one or more devices configured to monitor one or more physiologic signals including but not limited to heart rate, heart rate variability, heart murmur, electrophysiologic signals associated with low level autonomic activity (e.g. as extractable from an EKG signal, etc.), perfusion, sweating, hydration, or the like. The physiologic signals may be compared, analyzed, etc. before, during, and/or after one or more stress tests, procedures, etc. to determine the extent thereof, the body response thereto, etc.

[00389] In aspects, a device in accordance with the present disclosure may be configured to monitor one or more aspects of neural traffic, and/or surrogates thereof within one or more regions of the skin of a subject. Such neural traffic may include somatosensory traffic, receptor response, sympathetic outflow, parasympathetic outflow, muscular response (i.e. to SNS, PNS, etc.), smooth muscle electrophysiological response, etc. In aspects, such neural traffic may be monitored by a plurality of microelectrodes, one or more of the microelectrodes electrically isolated from the others such that the local electrophysiological signals can be teased out from the overall macro electrophysiological traffic in the vicinity of the region.

[00390] According to aspects there is provided a method for assessing the sympathetic neural state of a subject including, interfacing a system in accordance with the present disclosure with a subject, and monitoring one or more of neural traffic, a physiologic parameter, or a surrogate of neural traffic at one or more sites on the body of the subject.

[00391] In aspects, the method may include monitoring neural traffic at two or more locations (e.g. such as a PNS innervated site, a primarily SNS innervated site, a sweat gland heavy site, a somatosensory dense site, etc.). Such monitoring at

morphologically different skin regions on a body may be advantageous to extract different sub system responses from the overall traffic.

[00392] In aspects, a device in accordance with the present disclosure may include a local analgesic, a local neural blocker, a local neural agonist, or the like, integrated into the bioadhesive thereof, configured so as to be transdermally delivered to the tissues when the device is placed thereupon. Such a configuration may be advantageous for selectively blocking / discriminating between different types of neural traffic near to one or more sensors in the device. In aspects, a system in accordance with the present disclosure may include a plurality of such patches, each patch configured so as to selectively block one or more types of neural traffic in the nearby tissues (e.g. such that, through use of multiple patches, sensors, etc. different types of nerve traffic to/from the skin may be differentiated from the other types). In aspects, such a configuration may be advantageous to selectively highlight one or more types of autonomic outflow to a region of skin, or to selectively block one or more types of afferent traffic, from a stress site back to the CNS of the subject.

[00393] The method may include, performing a stress test in accordance with the present disclosure on the subject and monitoring the response thereto in accordance with the present disclosure

[00394] The method may include, assessing one or more changes in traffic associated with a neural state, parasympathetic neural state, sympathetic neural state, or the like, and determining the response of the sympathetic system of the subject to the stress test.

[00395] The method may include, performing a procedure in accordance with the present disclosure on the subject and monitoring the response, monitoring completion thereof, monitoring follow up thereof, etc. in accordance with the present disclosure.

[00396] According to aspects there is provided, a method for quantifying the contribution of a neurological state to a disease state of a subject including, non-invasively monitoring the neural state or a surrogate of the neural state of the subject to generate data, performing a stress test on the subject while monitoring, and analyzing the data to determine the change in neurological state of the subject, and/or to determine the relationship between the stress test and the neurological state of the subject, the analysis relating to the contribution.

[00397] According to aspects there is provided, a system for assessing the autonomic neural state of a subject and/or the relationship between sympathetic and parasympathetic autonomic neural state of a subject including a sensor (such as but not limited to an EMG, micro electrode array on a contact, ERG, etc.) for monitoring a physiologic state of an ocular element of a subject (e.g. state of a pupil, iris, ocular muscle, retinal state, retinal traffic, the tone of an ocular muscle, etc.), and a light source and/or display for providing one or more visual cues, optical stresses, incident light profiles, light scans, optical stress tests, etc. into the eye or eyes of the subject.

[00398] According to aspects there is provided a system for assessing autonomic neural state of a subject, and/or the response thereof to a stress state applied to the subject, configured to interface with the cutaneous innervation of one or more regions of skin of the subject.

[00399] The system may include a patch and/or an associated module for placement onto a site on the skin of a subject. The patch may include an array of microelectrodes, the patch and/or module including signal conditioning circuitry coupled with the microelectrodes, and a processor programmed with machine readable instructions coupled to the signal conditioning circuitry, the signal conditioning circuitry and/or the processor configured to extract one or more neural signals from one or more of the microelectrodes.

[00400] In aspects, the patch may include one or more of the microelectrodes includes a microneedle, the microneedle configured so as to penetrate a structure nearby the patch upon engagement therewith. In aspects, the microneedle may be sized with a predetermined length, the length thereof arranged so as to position an electrode arranged thereupon near to a characteristic depth of a nerve structure of interest within a region of skin on a body (e.g. a hairy skin surface, a glabrous skin surface, a mucosal surface, near a sweat gland, near a bulb of a hair follicle, near an arrector pili follicular muscle, near a sebaceous gland, near an erogenous zone, etc.).

[00401] In aspects, the patch/module may be arranged such that the processor and/or signal conditioning circuitry is/are configured to extract the neural signal amid one or more electromyographic signals associated with a nearby striated or smooth muscle structure, stretch based surface potential changes, movement, combinations thereof, or the like. In aspects, EMG artifacts may be algorithmically removed from

microelectrode signals, stretch based surface potential changes may be removed by subtracting a stretch surrogate signal (e.g. as obtained from a stretch sensor, a perfusion sensor, etc. in accordance with the present disclosure), movement may be removed by subtracting a movement signal (e.g. as measured by kinematic or kinetic sensor in accordance with the present disclosure), or the like.

[00402] In aspects, the patch may be configured such that one or more of the microelectrodes may be substantially electrically isolated from each other prior to placement onto the skin of the subject.

[00403] In aspects, the patch may include an engagement mechanism in accordance with the present disclosure, the patch including one or more of the microneedles sized, dimensioned, and arranged such that upon placement of the patch and/or initiation of the engagement mechanism, one or more of the microneedles may penetrate into the surface of the skin.

[00404] In aspects, the patch may include a gel layer arranged amid the microelectrodes, the gel layer including an adhesive configured so as to adhere the patch to the skin. In aspects, the gel layer being ionically and/or electrically conductive such that it may provide a macro-electrode useful for providing a reference electrode to the microelectrodes, an electrode for measuring an ECG, EMG, ERG, signal, etc. In aspects, the microelectrodes may be electrically isolated from the macro-electrode (e.g. such as via patterned regions, via an isolating region on microneedles, etc.).

[00405] In aspects, the adhesive or gel layer may be configured with an electrical and/or ionic conductivity such that the gel layer provides essentially an anisotropic electrical interconnection between the microelectrodes and the skin during engagement therewith.

[00406] In aspects, the adhesive or gel layer may be configured such that the anisotropic electrical interconnection is arranged so as to decrease an effective impedance between one or more of the microelectrodes and the skin upon engagement, while maintaining a relatively high impedance between adjacent microelectrodes in the patch.

[00407] In aspects, the adhesive or gel layer may be sized such that a thickness of the gel layer is sufficiently thing such that the microneedles may penetrate through the gel layer upon engagement with a region of tissue.

[00408] In aspects, the gel layer may be configured so as to have a moisture vapor transfer rate in the range of 100 – 4000 g/m^2/24hrs, in the range 200 – 1000 g/ m^2/24hrs, etc. so as to maintain a relatively high humidity environment around the tissues of the skin upon engagement.

[00409] In aspects, the signal conditioning circuitry and/or processor may be configured to assess the bioimpedance between two or more microelectrodes upon engagement with the skin so as to determine a hydration state thereof, to determine a fluid content thereof, or the like.

[00410] In aspects, the patch may include a plurality of macroelectrodes (2, 3, 4, greater than 4, etc.), each macroelectrode configured to interface with the skin upon placement thereupon, the signal conditioning circuitry and/or processor configured to assess the bioimpedance between the macroelectrodes (e.g. in a two point measurement configuration, 3 point, 4 point measurement configuration, etc.), upon engagement with the skin so as to determine a hydration state thereof, to determine a fluid content thereof, or the like.

[00411] In aspects, the device, patch, and/or module may include one or more temperature sensors configured to assess a thermal state of an adjacent tissue upon engagement therewith.

[00412] In aspects, the patch may include a thermal stimulatory component arranged in the patch so as to control a thermal state of an adjacent tissue upon engagement.

[00413] The thermal stimulatory component may include a thermoelectric component, an RF heating circuit, an ultrasound source, a component for providing a Seebeck effect, Peltier effect, and/or Thomson effect in the adjacent tissues upon engagement. The patch/device/module may include an insulating layer so as to limit heat transfer between the skin site and the surrounding environment after placement of the patch/module/device thereupon. A Peltier device may be incorporated into the device so as to heat and/or cool the adjacent tissues as part of a stress test. The processor may be programmed so as to drive the Peltier device over a predetermined

temperature range (e.g. to perform one or more temperature sweeps, to maintain a particular temperature, etc.), an embedded temperature sensor arranged so as to provide feedback of the adjacent tissue temperature. Such a configuration may be advantageous for generating a transfer function between thermal stress and afferent traffic, efferent outflow, etc. associated with one or more neural structures of the subject.

[00414] In aspects, the patch may include one or more photodetectors and optionally optical sources, the photodetectors configured to determine one or more of absorption spectroscopy, laser speckle analysis, narrow band absorption, and/or the like of an adjacent tissue, so as to determine perfusion, composition, and/or hydration of the tissue or fluid contained therein upon engagement therewith.

[00415] In aspects, the patch may include a sonography component, configured to provide an ultrasonic signal to and/or receive a sonographic signal from an adjacent tissue upon engagement with the skin. The sonography component may be configured so as to image, and/or capture a metric from an adjacent tissue upon engagement with the skin. In aspects, the metric may include a perfusion parameter, a tissue stiffness parameter, a hydration level, a temperature rise, a vessel diameter, a combination thereof, or the like.

[00416] In aspects, the patch may include an electromagnetic field generator configured to form an electromagnetic field in the vicinity of the patch, the signal(s) measured by the photodetectors related to the electromagnetic field.

[00417] The electromagnetic field generator may be adjustable dependent upon the absorption, reflection, speckle, or the like may be assessed under altered field conditions. The patch may include an electromagnetic field detector configured to assess the state of an electromagnetic field in the vicinity of one or more microelectrodes included in the patch. Together, an assessment of the changes in electromagnetic field and absorption, reflection, speckle, etc. may be advantageous for improving the number of analytes, or type of metrics which may be extracted from the measurements (e.g. such as measurement of challenging analytes such as glucose, measurement of hydration in addition to blood oxygen saturation, etc.).

[00418] In aspects, the device/patch/module may include a vibratory stimulatory component, configured so as to provide a tactile stress state to an adjacent region of

tissue. Such stimulation may be advantageous to controllably fire one or more types of somatosensory nerves in the tissue region (e.g. such as excitation of Pacinian corpuscles in the tissue, Merkel cell excitation, etc.). In aspects, the stimulatory component may be arranged to provide one or more tactile forms of stimulation to the tissues (e.g. indentation like tactile stimulus, stretch like stimulus, hair follicle deflection, skin shear, vibration, or pain <noxious mechanical stimulation>). Such tactile inputs may be coordinated so as to selectively stimulate one or more neural structures in the skin including Merkel cells, Ruffini cells, Meissner corpuscles, longitudinal lanceolate endings, Pacinian corpuscles, free nerve endings, combinations thereof, or the like.

[00419] In aspects, the signal conditioning circuitry and/or processor may be configured to determine a neural signal relationship during changes in the thermal state of the tissues in the vicinity thereof, or at a remote stimulatory site on the body.

[00420] In aspects, the processor may be coupled to the electromagnetic field generator, and/or the thermal stimulatory component, wherein the patch may be programmed so as to perform a stress test upon the subject therewith.

[00421] In aspects, the processor may be configured to export a neurological state and/or neuro-thermal relationship based signal recordings related to one or more of microelectrode, optical, and/or bioimpedance recordings captured during the stress test.

[00422] In aspects, the gel layer may be precooled (e.g. such as refrigerated, frozen, etc.) so as to provide a cooling effect to an adjacent tissue site upon engagement with the skin. The stress test may include monitoring traffic or a surrogate thereof at one or more sites on the subject and optionally monitoring local tissue temperature while the gel warms towards body temperature after coupling thereto.

[00423] In aspects, the gel layer may include an ionic conductivity or polarity such that it may be subsequently heated with the thermal stimulatory / thermo regulatory component (e.g. by current delivery there through, RF heating, dielectric heating, electromagnetic heating, etc.).

[00424] According to aspects there is provided, a system to assess an ANS state and/or ANS – stress response of a subject including one or more patches, modules,

and/or devices, each in accordance with the present disclosure, and a processor, the processor configured to assess signals and/or metrics obtained from one or more of the patches, the eye assessment device, or the like to determine the ANS state and/or ANS – stress response.

[00425] In aspects, the processor may be configured to drive one or more thermal, electromagnetic, electrical, and/or tactile stimulatory components in one or more of the patches/devices to coordinate a stress test on the subject.

[00426] In aspects, the system may be arranged to measure one or more of, but not limited to, respiration (breathing rate, breathing volume, lung stress or load, or the like), blood pressure, blood oxygen level, heart rate variability, heat flux, galvanic skin response, core body temperature, skin temperature, sympathetic or parasympathetic response, combinations thereof, or the like in order to assess the function of the ANS or changes therein during an assessment, before, during, and/or after a stress test, before, during, and/or after a procedure, etc.

[00427] In aspects, the system may be configured to capture a plurality of signals, signals from multiple sites on the body, signals from multiple skin types, etc. in order to establish relationships between aspects of a neural system and the stress test, between aspects of the neural system of the subject in general, etc.

[00428] Discussion related specifically to the Figures follows, the discussion above may be applied where ever applicable to a particular Figure reference.

[00429] Figs. 1a-d show aspects of modular physiologic monitoring systems in accordance with the present disclosure. Fig. 1a shows a subject 1 with a series of patches and/or patch/module pairs 5 – 137 each in accordance with the present disclosure, a host device 145 in accordance with the present disclosure, a feedback/user device 147 in accordance with the present disclosure displaying some data 148 based upon signals obtained from the subject 1, an one or more feedback devices 135, 140, in accordance with the present disclosure configured to convey to the subject one or more aspects of the signals or information gleaned therefrom. The host device 145, the user device 147 the patches and/or patch module pairs 5 – 137, and/or the feedback devices 135, 140 may be configured for wireless communication 146, 149 during a monitoring session.

[00430] In aspects, a patch/module pair may be adapted for placement almost anywhere on the body of a subject 1. As shown in Fig. 1a, some sites may include attachment to the cranium or forehead 131, the temple, the ear or behind the ear 50, the neck, the front, side, or back of the neck 137, a shoulder 105, a chest region with minimal muscle mass 100, integrated into a piece of ornamental jewelry 55 (may be a host, a hub, a feedback device, etc.), arrangement on the torso 110a-c, arrangement on the abdomen 80 for monitoring movement or breathing, below the rib cage 90 for monitoring respiration (generally on the right side of the body to substantially reduce EKG influences on the measurements), on a muscle such as a bicep 85, on a wrist 135 or in combination with a wearable computing device 60 on the wrist (e.g. a smart watch, a fitness band, etc.), on a buttocks 25, on a thigh 75, on a calf muscle 70, on a knee 35 particularly for proprioception based studies and impact studies, on a shin 30 primarily for impact studies, on an ankle 65, over an Achilles tendon 20, on the front or top of the foot 15, on a heel 5, or around the bottom of a foot or toes 10. Other sites for placement of such devices are envisioned. Selection of the monitoring sites is generally determined based upon the intended application of the patch/module pairs described herein.

[00431] Additional placement sites on the abdomen, perineal region 142a – c, genitals, urogenital triangle, anal triangle, sacral region, inner thigh 143, or the like may be advantageous in the assessment of autonomic neural function of a subject. Such placements regions may be advantageous for assessment of PNS activity, somatosensory function, assessment of SNS functionality, etc.

[00432] Placement sites on the wrist 144a, hand 144b or the like may advantageous for interacting with a subject, such as via performing a stress test, performing a thermal stress test, performing a tactile stress test, monitoring outflow, afferent traffic, efferent traffic, etc.

[00433] Placement sites on the nipples, areola, lips, labia, clitoris, penis, the anal sphincter, levator ani muscle, over the ischiocavernous muscle, deep transverse perineal muscle, labium minus, labium majus, one or more nerves near the surface thereof, posterior scrotal nerves, perineal membrane, perineal nerves, superficial transverse perineal nerves, dorsal nerves, inferior rectal nerves, etc. Such placement may be advantageous for assessment of autonomic neural ablation procedures,

autonomic neural modulation procedures, assessment of the PNS of a subject, assessment of sexual dysfunction of a subject, etc.

[00434] Placement sites on the face 141, over ocular muscles, near the eye, over a facial muscle (e.g. a nasalis, temporalis, zygonaticus minor/major, orbicularis oculi, occipitofrontalis), near a nasal canal, over a facial bone (e.g. frontal process, zygomatic bone/surface, zygomaticofacial foreman, malar bone, nasal bone, frontal bone, maxilla, temporal bone, occipital bone, etc.), may be advantageous to assess ocular function, salivary function, sinus function, interaction with the lips, interaction with one or more nerves of the PNS (e.g. interacting with the vagus nerve within, on, and/or near the ear of the subject), etc.

[00435] In aspects, a system in accordance with the present disclosure may be configured to monitor one or more physiologic parameters of the subject 1 before, during, and/or after one or more of, a stress test, consumption of a medication, exercise, a rehabilitation session, a massage, driving, a movie, an amusement park ride, sleep, intercourse, a surgical, interventional, or non-invasive procedure, a neural remodeling procedure, a denervation procedure, a sympathectomy, a neural ablation, a peripheral nerve ablation, a radio-surgical procedure, an interventional procedure, a cardiac repair, administration of an analgesic, a combination thereof, or the like. In aspects, a system in accordance with the present disclosure may be configured to monitor one or more aspects of an autonomic neural response to a procedure, confirm completion of the procedure, select candidates for a procedure, follow up on a subject after having received procedure, assess the durability of a procedure, or the like (e.g. such as wherein the procedure is a renal denervation procedure, a carotid body denervation procedure, a hepatic artery denervation procedure, a LUTs treatment, a bladder denervation procedure, a urethral treatment, a prostate ablation, a prostate nerve denervation procedure, a cancer treatment, a pain block, a neural block, a bronchial denervation procedure, a carotid sinus neuromodulation procedure, implantation of a neuromodulation device, tuning of a neuromodulation device, etc.).

[00436] Fig. 1b shows a series of patch/module pairs 150a-e each in accordance with the present disclosure placed upon a subject 2 as part of a monitoring session in accordance with the present disclosure, in this case an EKG monitoring session. An image 152 of the subject 2 has been taken and may be analyzed in accordance with

the present disclosure to calculate one or more standard lead configurations from the arrangement of patch/modules 150a-e shown.

[00437] Fig. 1c shows aspects of communication between subjects 155, 160 and non-subject users 156, 161 partaking in a monitoring session in accordance with the present disclosure. In a first aspect, the subject 155 is wearing a series of patches and modules each in accordance with the present disclosure configured to communicate with one or more of a host device 158, a display 157b, a HUD, a pair of virtual reality goggles, a Google Glasses™ based feedback device 157a (i.e. potentially via a smartphone hub), and/or a wristwatch 157c to communicate one or more feedback signals in accordance with the present disclosure to the user 156.

[00438] In aspects, the subject 160 may wear a series of patches and modules each in accordance with the present disclosure configured to communicate with one or more of a host device 163, a display 162b, a virtual reality headset, a HUD, a Google Glasses™ based feedback device 162a (i.e. via a smartphone hub), a wristwatch 162c, and/or one or more patches and/or modules configured upon the body of the user 161 to communicate one or more feedback signals in accordance with the present disclosure to the user 161 or to convey one or more sensations to the body of the user 161 (i.e. via the attached patches). In aspects, the ocular feedback device 162a, may be used to perform a visual and/or audible stress test on the subject, one or more aspects of the feedback device 162a, or an associated patch configured to monitor the response of one or more aspects of the ANS to the stress test.

[00439] In aspects, the communication between the subjects 155, 160 and the users 156, 161 may be bidirectional (i.e. the subject 155, 160 may also receive information corresponding to physiologic and/or physical information obtained from the user 156, 161).

[00440] Fig. 1d shows a time series example of an identification process for a patch/module pair in accordance with the present disclosure. In the non-limiting example shown, the modules are equipped with radios, each radio capable of sending a signal with a predetermined signal strength (left vertical axis) and equipped with a barometer or altimeter calibrated to give a relative height signal (right vertical axis). In aspects, a user, a program on a feedback device, on a host device, on a user device, etc. may prompt a subject or user to cover a particular patch on the body of the subject

(e.g. communicated to the subject or user via an instruction pamphlet, via a screen prompt, etc.). When the prompted module is covered (period between timestamps 180, 185) the signal strength of the module is changed by a substantially identifiable amount 175. Thus a simple procedure may be used to identify a module in the BAN, correlate a module on a particular site on the subject with a corresponding ID received by the host device, etc. Other aspects and variants of a localization and/or orientation procedure are discussed throughout this disclosure.

[00441] Figs. 2a – d show aspects of a patch and a corresponding module in accordance with the present disclosure. Fig. 2a shows aspects of a patch 210 in accordance with the present disclosure coupled to a module 215 in accordance with the present disclosure.

[00442] The patch 210 may include a substrate formed from a flexible, stretchable material. In aspects, the patch 210 may be stretchy, elastically deformable, or the like. In aspects, once interfaced with a subject, the patch 210 may continue to function (i.e. provide working interfaces between an interconnect provided thereupon and one or more sensors/electrodes provided thereupon) at a stretch of greater than 25%, greater than 50%, greater than 80%, etc.

[00443] Fig. 2b shows aspects of a patch 225 and a corresponding module 235 each in accordance with the present disclosure. The patch includes a substrate 245, an adhesive layer 250, multiple patch interconnects 230a, b, and multiple electrodes 255a, b, each in accordance with the present disclosure. The patch interconnects 230a,b and the electrodes 255a,b are electrically coupled together via one or more conducting traces located on the substrate 245, embedded into the substrate 245, etc.

[00444] The module 235 includes multiple module interconnects 240a,b in accordance with the present disclosure configured, dimensioned, and arranged so as to mate with the corresponding patch interconnects 230a,b. The interconnects 230a,b 240a,b may include snap elements, magnetic elements, etc. as discussed throughout the present disclosure.

[00445] Fig. 2b also shows a module 215 coupled to a corresponding patch 210, the module and patch containing a series of mating interconnects 219a-g, the patch including one or more sensors and/or electrodes 218a-d (e.g. here shown as circular electrode elements spaced along the length of the patch 210). Also shown are two

axes, a lengthwise axis 217 oriented along a direction of expected stretch for the patch 210 during use, and a substantially perpendicular axis 216 oriented along a directly substantially perpendicular to the direction of expected stretch for the patch 210. In aspects, the mating interconnects 219a-g may be oriented along the patch, such that they are aligned with the perpendicular axis 216 when the module 215 is mated with the patch 210 during use. Such a configuration may be advantageous for limiting stress and/or movement around one or more of the interconnects 219a-g during a typical usage case.

[00446] Fig. 2c shows aspects of a module 260 in accordance with the present disclosure. The module 260 includes a housing 265, a portion of which is provided by a printed circuit board 280 in accordance with the present disclosure. The housing 265 includes a dome shaped component upon which a three dimensional antenna 287 is formed, embedded, printed upon, etc. In aspects, the antenna 287 may be formed in the shape of a helical structure, a multipolar structure, a curved dipole structure, a monopolar structure etc. In aspects, the antenna 287 may include one or more interconnects which may be patterned so as to mate with corresponding connectors on the printed circuit board 280 during an assembly procedure. In aspects, the antenna 287 may be formed from a free standing electrically conducting structure and attached to the printed circuit board 280 during assembly.

[00447] In aspects, the antenna 287 may be formed as a curved dipolar antenna. The trace of the antenna 287 arranged so as to extend from contacts on the circuit board 280 onto the housing 265 and outward to form an airplane wing-like shape. Such a configuration may be advantageous for simplifying interface between the antenna and an amplifier, for shaping the RF waves emitted from the module 260 when adjacent to a subject, as well as for decoupling the RF circuitry from a ground plane of the circuit board 280, so as to minimize DC charge accumulation during RF transmissions from the module 260.

[00448] The module 260 in accordance with the present disclosure may include one or more components 270 (e.g. microcircuits, sensors, transducers, etc. optionally stacked / embedded into PCBs, etc.), interconnects 285, and a power source 275 each in accordance with the present disclosure.

[00449] The module 260 may be hermetically sealed, etc. so as to isolate components 270 therein from the surroundings. The module 260 may include gasket 285 in accordance with the present disclosure to minimize transfer of fluids into the region of contact between the module 260, the interconnects 285, and a corresponding patch in accordance with the present disclosure.

[00450] Fig. 2d shows aspects of orientation markings 294, 296, 298 on a module 292 and a patch 290 in accordance with the present disclosure. Such markings 294, 296, 298 may be advantageous to assist a user with alignment and attachment of a module 292 to a patch 290 as well as may be useful for a vision algorithm to extract orientation, functional (e.g. by color or pattern of the markings 294, 296, 298), and/or identification information from an image of the patch 290 and module 292 as placed onto a subject as part of a monitoring session.

[00451] Figs. 3a – f show aspects of patches in accordance with the present disclosure.

[00452] Fig. 3a shows a patch 301 coupled to a module 305 each in accordance with the present disclosure. The patch 301 includes a plurality of electrodes 303a-e for interfacing with a subject. The electrodes 303a-e are arranged in a somewhat linear fashion along the patch 301, for assessing electrocardiographic signals at sites in the vicinity of the heart of a subject, etc. In aspects, one or more of the electrodes 330a-e may include a microelectrode in accordance with the present disclosure, the microelectrode configured to capture one or more single-unit, or multi-unit neural signals from within the skin of the subject after placement thereupon.

[00453] Fig. 3b shows a patch 306 coupled to a module 310 each in accordance with the present disclosure. The patch 306 includes a bipolar electrode arrangement 307a,b for interfacing with a subject. Such an arrangement may be advantageous for monitoring heart-rate, a signal channel EKG, EMG, respiration rate, acoustic monitoring coupled with EMG of one or more neck or facial muscles, EEG monitoring coupled with kinematic sensors behind the ear, a stimulation device, etc. of a subject as part of a monitoring session. In aspects, an EEG monitoring component may be coupled with a stimulatory component (e.g. a tactile stimulatory component, a vibratory component, a thermal regulating component, a chemical stimulating / blocking component, an electrical stimulation component, an radio

frequency stimulatory component, an electromagnetic field generator, etc.), to provide a stress test and/or monitor the effects of a stress test on a subject in accordance with the present disclosure.

[00454] Fig. 3c shows a patch 311 coupled to a module 315 each in accordance with the present disclosure. The patch 311 includes multiple electrodes 313a-f for interfacing with a subject. The electrodes 313a-f may be arranged so as to interface with one or more muscle groups along an appendage of a subject, as part of an EMG study in accordance with the present disclosure. The patch 311 may be formed so as to maintain an electrical interface between the electrodes and the module 315 while undergoing stretch during the monitoring session (e.g. in aspects where the length between the electrodes may change substantially during the monitoring session, etc.). In aspects, such a configuration may be advantageous for spanning the length of a muscle, crossing a joint, wrapping around the curvature of a limb, a foot, etc. of a subject as part of a monitoring session.

[00455] Fig. 3d shows a patch 316 coupled to a module 320 each in accordance with the present disclosure. The patch 316 includes three electrodes 318a-c for interfacing with a subject. The electrodes 318a-c may be arranged so as to provide multi-site capture of electrophysiological activity on the subject. Such an arrangement may be advantageous for generating a field vector in the vicinity of the patch 316. In aspects, the module 320 may include an algorithm configured to analyze incoming data from the electrodes 318a-c (i.e. perhaps with a signal electrode in the set selected as a reference etc.) and to calculate field vector with orientation and magnitude in relationship to the orientation of the electrodes on the patch 316. In aspects, such a configuration may be advantageous for calculating a vector electrocardiogram on a subject, calculating muscle action potential vectors, muscle signal orientation, calculating electric field propagation directions along the surface of the subject, or the like.

[00456] Fig. 3e shows a patch 322 in accordance with the present disclosure. The patch 322 includes a window 324 through which a sensor (e.g. an optical sensor, etc.) included in an associated module (not explicitly shown) may interface with the subject during a monitoring procedure. The patch 322 may include further electrodes, sensors, etc. (not explicitly shown) to enable additional monitoring functionality from the same module. Such a configuration may be advantageous for monitoring local

blood flow optically, perform colorimetric assessment of underlying tissues, provide seamless entry of radiation into the tissues, etc. during a monitoring session in accordance with the present disclosure.

[00457] Fig. 3f shows a patch 328 in accordance with the present disclosure. The patch 328 includes multiple sensors 333a,b, coupled with a plurality of conducting members (e.g. electrical traces, printed conductors, spun nanotube fibers, stretchable conducting traces, etc.) 336a-c, and a plurality of interconnects 331a-c each in accordance with the present disclose. The sensors 333a,b may be arranged, configured, and dimensioned so as to interface with and/or monitor an underlying tissue structure, property, and/or tissue or fluid analyte (such as tissue composition, fat content, water content, blood oxygen saturation levels, etc.), of a subject during a monitoring procedure.

[00458] Figs. 4a-f show top, side, and isometric views of a mated patch 401, 405 and module 403, 407 each in accordance with the present disclosure. Such a configuration may be advantageous for providing robust, comfortable monitoring of a subject in accordance with the present disclosure.

[00459] Fig. 5 shows a perspective view of aspects of a mated patch 503 and module 505 forming a patch/module pair 501 each in accordance with the present disclosure. The patch/module pair 501 is shown beside a coin 16 (a US quarter) for perspective.

[00460] Fig. 6 shows a schematic of aspects of a module 601 in accordance with the present disclosure. The module 601 includes one or more of interconnects, sensors, optical source(s), optical detector(s), a radio, an antenna, a sensor communication circuit, a signal conditioning circuit, a processor, a memory device, a controller, a power supply, power management circuit, and/or energy harvesting circuit, and one or more peripherals each in accordance with the present disclosure. The module 601 is shown in wireless communication 615, 625, 620 with an additional module 605 (e.g. perhaps situated in the same monitoring system, on the same subject, etc.), and a host device 610. Further aspects of the module 601 are discussed throughout this disclosure.

[00461] Fig. 7 shows a schematic of a patch/module pair 701 attached to a subject 3 in accordance with the present disclosure. The patch/module pair 701 includes a

module 705 coupled to a corresponding patch 710, the patch includes multiple electrodes 720a-c arranged for interfacing with the skin of the subject 3. The patch/module pair 701 is shown in wireless communication 725 with a host device (not explicitly shown). Such a configuration may highlight how a stretchy patch 710 may be able to maintain monitoring of the subject during periods of movement, changes in shape or stretching along the surface of the skin of the subject 3, etc. Such a configuration may be advantageous to provide a soft comfortable monitor, with a low cost disposable patch 710 and a miniature reusable module 705.

[00462] Figs. 8a-e show aspects of non-limiting examples of patch electrode layouts in accordance with the present disclosure.

[00463] Fig. 8a shows a patch 801 coupled to a module 803 each in accordance with the present disclosure. The patch 801 includes a plurality of electrodes 805a-b for interfacing with a subject. The electrodes 805a-b are arranged in a very tight bipolar arrangement suitable for obtaining a bipolar electrical reading from the surface of a subject with a very small profile. In aspects, one or more of the electrodes 805a,b may include an electrode feature in accordance with the present disclosure for enhancing the electrical coupling between the module 803 and the underlying tissues of a subject. In aspects, pressure applied to the top of an attached module 803 may be suitable for engaging an electrode feature with the underlying tissue of the subject. Such an arrangement may be advantageous for providing an ultra-miniature heart-rate monitor, a pediatric heart-rate monitor, an EMG sensor for placement near a sexual organ, an electrophysiological monitor behind an ear, on a neck, etc.

[00464] Fig. 8b shows a patch 807 coupled to a module 809 each in accordance with the present disclosure. The patch 807 includes a bipolar electrode arrangement 811a,b for interfacing with a subject. Such an arrangement may be advantageous for monitoring heart-rate, a signal channel EKG, respiration rate, etc. of a subject as part of a monitoring session. A plurality of such patches 807 may be applied to a subject to simultaneously extract a higher level or spatially distributed electrical field over the body of the subject.

[00465] Fig. 8c shows a patch 813 coupled to a module 815 each in accordance with the present disclosure. The patch 813 includes three electrodes 817a-c for

interfacing with a subject. The electrodes 817a-c may be arranged so as to allow for multi-site capture of electrophysiological activity on the subject. Such an arrangement may be advantageous for generating a field vector in the vicinity of the patch 813.

[00466] Fig. 8d shows a patch 819 coupled to a module 821 each in accordance with the present disclosure. The patch 819 includes a quadripolar electrode arrangement 823a-d for interfacing with a subject. The quadripolar electrodes 823a-d may be arranged so as to allow for multi-site capture of electrophysiological activity on the subject. Such an arrangement may be advantageous for generating a field vector in the vicinity of the patch 819, for mapping electric field propagation across the surface of the subject, etc.

[00467] Fig. 8e shows a patch 825 coupled to a module 827 each in accordance with the present disclosure. The patch 825 includes a plurality of electrodes 829a-b for interfacing with a subject. The electrodes 829a-b are shown in a bipolar arrangement connected to stretchable conducting elements 831a,b. In aspects, such a configuration may be advantageous to freely flex and stretch 833 along with the nearby tissues of the subject during a monitoring session. The stretchable conducting elements 831a,b may be arranged so as to repeat ably change impedance during stretch. Such a configuration may be advantageous for assessing movement under the patch (e.g. due to muscle movement, breathing, etc.) in conjunction with one or more physiologic signals (e.g. such as electrophysiological signals, stretch related artifact, etc.) in accordance with the present disclosure. Such a configuration may be suitable for physiotherapy monitoring sessions (e.g. combined proprioceptive monitoring in conjunction with EMG, assessing breathing in conjunction with EKG, gait assessment, a running gait correction system, etc.).

[00468] In aspects, such a configuration may be advantageous for a training and/or physiotherapy system. Dual implementation of movement (e.g. across joints, cross-wise to a joint, etc.) in conjunction with EMG related information (e.g. exertion, muscle fatigue, etc.) may be advantageous to detect risk of injury, map progression of therapy, etc.

[00469] Fig. 9 shows aspects of an impact sensing patch 900 and a feedback component 925 in accordance with the present disclosure. The impact sensing patch

900, here shown integrated into a knee brace on a subject 4, may include one or more piezoresistive materials (i.e. materials that change electrical properties or charge storage thereupon in relation to strains placed thereupon), a capacitive stretch sensor in accordance with the present disclosure, a pressure sensitive nano-composite structure, or the like. Upon impact 905 of the patch 900 with an object 5, a coupled module 910 may send one or more signals 915 to a feedback device 925, a host device, etc. in accordance with the present disclosure. In aspects, the feedback device 925 may accept the signal 935 and produce a feedback signal (e.g. an audio signal, a vibration signal, a tactile signal, a visual signal, etc.) for delivery to a user in accordance with the present disclosure. In this non-limiting example, the feedback device 925 produces an audible feedback signal in the ear 6 of the subject via a loudspeaker 930. Such a system may be advantageous for monitoring impacts on a subject with neuropathy (e.g. lack of sensation in an extremity, for assistance with gait analysis, for providing feedback during exercise, etc.) so as to provide the subject with a transferred sensation of touch in a region of their body that still has sensation (e.g. via a tactile feedback component, audible cue, visual cue, etc.), or for formation of a feedback loop to a touch related event.

[00470] Figs. 10a – e show aspects of a patch in accordance with the present disclosure. Fig. 10a shows a patch including a substrate 1005 (or adhesive layer) formed from a conducting, self-adhering material (i.e. so as to provide combined electrical and mechanical coupling to an adjacent body), a conducting trace 1010 configured to electrically couple a region of the substrate 1005 to a corresponding interconnect (not explicitly shown), and including a dielectric layer 1015 configured to isolate one or more regions of the substrate 1005 from one or more regions of the conducting trace 1010. In aspects, the substrate 1005 may initiate charge transfer 1025 with an adjacent surface during use (e.g. so as to electrically couple thereto, to form a suitable sensory interface thereto, to provide electroporation thereto, to provide current flow thereto, to monitor aspects thereof, etc.). The substrate 1005 is shown with a predetermined thickness 1020 in accordance with the present disclosure. In aspects, to maintain a breathable, flexible interface, the thickness 1020 may be less than 200um, less than 100um, less than 50um, less than 25um, less than 12um, less than 6um, etc. In aspects, the thickness 1020 may be sufficiently thick so as to retain a tear strength sufficient to allow for removal of the patch from the subject without

tearing. The tear strength may be greater than 0.5N/mm, greater than 1N/mm, greater than 2N/mm, etc. (i.e. as measured in accordance with ASTM standard ASTM-D-624 DIE tear strength).

[00471] Fig. 10b shows aspects of a patch in accordance with the present disclosure including an adhesive layer 1035 coupled with a substrate 1030 each in accordance with the present disclosure. The patch includes a microstud 1040 interconnect coupled through the thickness of the substrate 1030. In the non-limiting example shown the microstud 1040 also includes an electrode 1050 and corresponding electrode features 1055 for interfacing with the body of a subject. The microstud 1040 may also provide a top side interconnect for interfacing with a module, etc.

[00472] In aspects, the electrode 1050 may be suitable for providing charge transfer 1055 with an adjacent surface during use (e.g. so as to electrically couple thereto, to provide electroporation thereto, to provide current flow thereto, to monitor aspects thereof, etc.). The patch is shown with a predetermined thickness 1045 in accordance with the present disclosure. In aspects, to maintain a breathable, flexible interface, the thickness 1045 may be less than 200um, less than 100um, less than 50um, less than 25um, less than 12um, less than 6um, etc.

[00473] Fig. 10c shows aspects of a patch in accordance with the present disclosure including an adhesive layer 1060 coupled with a substrate 1065 each in accordance with the present disclosure. The patch includes a rivet style stud 1070 coupled through the thickness of the substrate 1065. In the non-limiting example shown the rivet style stud 1070 also includes an electrode and a cap 1075 to hold it tightly to the substrate 1065 in use. The patch includes a conducting layer 1080 (in this case shown coating the stud 1070 but may alternatively be applied between the cap 1075 and the substrate 1065) for communicating with an interconnect (not explicitly shown). The patient side of the stud 1070 is shown coated with a thin electroconducting gel 1085 in accordance with the present disclosure. In aspects, the rivet style stud 1070 may include a button top, a snap interconnect, etc. for directly interfacing with a corresponding module or the like.

[00474] In aspects, the gel 1085 may be suitable for providing charge transfer 1095 with an adjacent surface during use (e.g. so as to electrically couple thereto, to

provide electroporation thereto, to provide current flow thereto, to monitor aspects thereof, etc.). The patch is shown with a predetermined thickness 1090 in accordance with the present disclosure. In aspects, to maintain a breathable, flexible interface, the thickness 1090 may be less than 200um, less than 100um, less than 50um, less than 25um, less than 12um, less than 6um, etc.

[00475] Fig. 10d shows aspects of a patch in accordance with the present disclosure in a substantially fully printed form. An exaggerated through thickness schematic is shown of a region of a patch attached to a liner 1012, the liner being removed 1018 from the patch. The patch includes one or more electrically insulating layers 1014a,b,1022a,b, which may be formed into substantially thin films with a patterning process (e.g. such as printing, physical vapor deposition, sputter deposition, jet based printing, screen based printing, flexographic printing, etc.). The electrically insulating layers 1014a,b,1022a,b may be formed from a soft biocompatible material (e.g. a polyurethane, a copolyester, a silicone, a thermoplastic elastomer, etc.). The patch includes one or more electrically conducting layers 1024, 1026, etc. The electrically conducting layers 1024, 1026 may include a flexible, stretchable material, a ferromagnetic conducting material, a conductive gel 1016, a bioadhesive gel, or the like. The patch may include a patient side ionic and/or thin electroconducting gel adhesive 1028 in accordance with the present disclosure. The ionic and/or electroconducting gel adhesive 1028 is shown covering the entire patient side face of the patch. In this configuration, the ionic and/or electroconducting gel adhesive 1028 may provide a substantially strong ionic or electrical connection between one or more of the conducting materials 1024, 1026 and a surface of the subject when interfaced therewith, which having a conductivity that is essentially low enough so as to not influence the lateral conductivity along the patch (such capability may also be achieved using an gel 1028 with anisotropic properties). In aspects, the overall thickness 1016 of the patch may be less than 50um, less than 25um, less than 12um, less than 8um, less than 6um, less than 4um, etc. In aspects, the structure shown may be advantageous to generate an essentially connector-less patch with embedded and encapsulated traces, electrodes, sensors, or the like. In aspects, one or more of the materials 1014a,b, 1022a,b, 1024,1026,1028 may be formulated as an ink, the ink printed upon the liner 1012 or each successive layer to from the resulting patch

structure. Such an approach may be advantageous for forming the patches in a roll to roll process.

[00476] Fig. 10e shows aspects of a patch in accordance with the present disclosure in a substantially fully printed form. An exaggerated through thickness schematic is shown of a region of a patch attached to a liner 1032, the liner being removed 1038 from the patch. The patch includes one or more electrically insulating layers 1034a,b,1042a,b, which may be formed into substantially thin films with a patterning process (e.g. such as printing, physical vapor deposition, sputter deposition, jet based printing, screen based printing, flexographic printing, etc.). The electrically insulating layers 1034a,b,1042a,b may be formed from a soft biocompatible material (e.g. a polyurethane, a copolyester, a silicone, a thermoplastic elastomer, etc.). The patch may include one or more electrically conducting layers 1052, 1046, etc. The electrically conducting layers 1052, 1046 may include a flexible, stretchable material, a ferromagnetic conducting material, a conductive gel, a bioadhesive gel, or the like. The patch may include a patient side bioadhesive 1048a,b in accordance with the present disclosure. In aspects, the overall thickness 1036 of the patch may be less than 50um, less than 25um, less than 12um, less than 8um, less than 6um, less than 4um, etc. In aspects, the structure shown may be advantageous to generate an essentially connector-less patch with embedded and encapsulated traces, electrodes, sensors, or the like. In aspects, one or more of the materials 1034a,b, 1042a,b, 1046, 1048a,b may be formulated as an ink, the ink printed upon the liner 1032 or each successive layer to from the resulting patch structure. Such an approach may be advantageous for forming the patches in a roll to roll process.

[00477] Figs. 11a-b show aspects of patches and modules in accordance with the present disclosure.

[00478] Fig. 11a shows aspects of a patch 1105 and a module 1115 in accordance with the present disclosure. The patch 1105 includes an adhesive layer 1107, a substrate 1109, and electrodes 1110a,b. In the non-limiting example shown, the electrodes are configured to directly interface with a corresponding module 1115. The module 1115 is shown with a plurality of corresponding interconnects 1120a,b each including caps to isolate one or more regions of the module 1115 from the elements. The module 1115 may be configured for wireless communication 1130 in accordance with the present disclosure.

[00479] Fig. 11b shows aspects of a patch 1140 and a corresponding module 1160 each in accordance with the present disclosure. The patch 1140 may include a substrate 1145 coupled to a patient facing adhesive layer 1155, and a module interfacing adhesive 1150 (i.e. patterned over a region thereof so as to interface with a module 1160 placed thereupon). The module 1160 includes an optical source 1165 for emitting energy towards 1172 a subject, and an optical sensor 1170 for receiving energy from 1175 a subject. The module 1160 may be configured and dimensioned for placement onto the corresponding patch 1140. One or more layers of the patch 1140 may be transparent to the radiation, so as to facilitate interaction of the module 1160 with an adjacent subject. The module 1160 may be configured for wireless communication 1180 in accordance with the present disclosure.

[00480] Figs. 12a – c show aspects of patches and modules in accordance with the present disclosure.

[00481] Fig. 12a shows aspects of a patch 1201 in accordance with the present disclosure including a substrate 1203 and an adhesive layer 1205 coupled thereto (in this case, optionally an ionically conducting adhesive). The patch 1201 includes an optional sensor 1207 embedded into the substrate 1203 and a patch interconnect 1211a,b for coupling to a module 1215 and provided in electrical communication with the sensor 1207 (e.g. via electrical interconnects coupled to the substrate 1203, optionally printed along the face of the substrate 1203 facing the adhesive layer 1205). The sensor 1207 is configured for delivery of energy 1209a into and sensing of energy 1209b received from a subject during use (e.g. such as an optical sensor, a spectrometer, etc.). Fig. 12a shows a module 1215 in accordance with the present disclosure configured and dimensioned to mate with the patch 1201. The module 1215 includes a module interconnect 1217a,b arranged to mesh with the corresponding patch interconnect 1211a,b. In the non-limiting example shown, the patch interconnect 1211a,b may include a ferromagnetic material and the module interconnect 1217a,b may include magnetic materials configured such than a substantially strong magnetic field 1219a,b is formed there between when placed in close proximity to each other (i.e. during assembly). The interconnects 1211a,b, 1217a,b may be configured to form an electrically conducting interface between the patch 1201 and the module 1215 upon assembly, alternatively an inductive interface, or a capacitive interface in accordance with the present disclosure.

[00482] Also shown, the patch 1201 includes a plurality of electrodes 1208a,b each in accordance with the present disclosure. The electrodes 1208a,b are coupled to the patch interconnect 1211a,b with one or more stretchy electrically conducting traces each in accordance with the present disclosure (e.g. in this case provided by a stretchable electrically conducting ink, printed onto a surface 1212 of the substrate 1203 so as to be sandwiched between the substrate 1203 and the adhesive 1205). The electrodes 1208a,b may include one or more electrically conducting and/or ionically conducting substances each in accordance with the present disclosure for interfacing with an adjacent subject during use.

[00483] Fig. 12b shows aspects of a patch 1227 coupled to a module 1225 in accordance with the present disclosure. The patch 1227 includes a substrate 1230 and adhesive 1235 for interfacing with a subject, each of which are substantially transparent to energy transferred to and/or from 1245 the module 1225 into the subject during use. Such energy transfer 1245 may be completed by a transducer 1240 and may be monitored by the same or an equivalent sensitive transducer 1240.

[00484] Fig. 12c shows aspects of a patch 1251 and a corresponding module 1265 each in accordance with the present disclosure. The patch 1251 includes a substrate 1253 and electrodes 1255a-c for interfacing with a subject. The patch 1251 includes multiple microstuds 1271a-c in accordance with the present disclosure. The studs 1271a-c may be electrically coupled with the electrodes 1255a-c via traces included in the substrate 1253 (e.g. patterned on, coated on, printed onto, embedded into the substrate 1253, etc.). The studs 1271a-c may coated with an insulating layer 1260, in this case configured so as to both insulate the studs 1271a-c from the subject but also to add local rigidity to the region of the patch 1251 in the vicinity of the interconnect (i.e. the collection of microstuds 1271a-c).

[00485] The module 1265 may include a plurality of corresponding connectors 1275a-c for interfacing with the microstuds 1271a-c. The connectors 1275a-c may be integrated into a printed circuit board 1270, which duals as a portion of the housing of the module 1265. In aspects, the connectors 1275a-c may include solderable components for securement to the PCB 1270. In aspects, the connectors 1275a-c may be substantially formed from through hole interconnects in the PCB 1270 (i.e. so as to simplify the assembly process thereof). In aspects, the PCB 1270 may include an

undercoating, gasket, etc. in order to substantially isolate the interconnects 1271a-c, 1275a-c from the surroundings during use.

[00486] Fig. 13 shows a schematic of a patch/module pair 1300 configured to apply energy 1325 (in this case tactile stimulus, vibrational energy, stroking, poking, circular movement, etc.) to the surface 7 a subject 1301 in accordance with the present disclosure. The patch/module pair 1300 includes an adhesive layer 1310 (i.e. potentially formed from a patch coupled to the module) for securement to the subject 7. The module 1315 includes a transducer 1305 configured to for generate vibrational energy 1325 for transfer 1330 into the subject 1301. The transducer 1305 may be controlled and/or powered by an electronics unit 1320 included in the module 1315. In the non-limiting example shown, the transducer 1305 may be piezoelectric material (e.g. polymer, ceramic, etc.). Additionally, alternatively, or in combination the transducer 1305 may be a motor with an unbalanced shaft, a stroking actuator, etc.

[00487] Fig. 14 shows a patch/module pair 1400 for applying thermal energy 1430 to a subject 1402 in accordance with the present disclosure. The patch/module pair 1400 includes an adhesive layer 1410 for attachment to a skin surface 1402 of the subject 1401. The module 1415 includes one or more heater bands 1405 or RF heating circuits, and thermocouples 1406 coupled to an electronics unit 1420 including a power source, a microcircuit, etc. via one or more electronic interconnects 1408. Alternatively, additionally, or in combination the module 1415 may include one or more thermoelectric units, a Peltier device, an RF heating circuit, an ultrasound source, a component for providing a Seebeck effect, Peltier effect, and/or Thomson effect in the adjacent tissues upon engagement. Such a configuration may be advantageous for generating a transfer function between thermal stress and afferent traffic, efferent outflow, etc. associated with one or more neural structures of the subject. The thermoelectric unit may be configured to heat, cool, or substantially maintain the temperature of an adjacent tissue 1401 during use. The patch 1400 may include one or more sensors, microelectrodes, etc. each in accordance with the present disclosure, configured to interface with the tissues 1401 while the thermoelectric unit, heater bands 1405, etc. heat or cool the tissues 1401.

[00488] Fig. 15 shows a vibrating patch/module pair 1500 configured to apply a tactile input 1525 to the surface 1502 of a subject 1501 in accordance with the present disclosure. The patch/module pair 1500 includes an adhesive layer 1510 (i.e.

potentially formed from a patch coupled to the module) for securement to the subject 1501. The module 1515 includes a transducer 1505 configured to for generate the torsional energy 1525 for transfer 1530, 1535, 1540 into the subject 1501. The transducer 1505 may be controlled and/or powered by an electronics unit 1520 included in the module 1515. In the non-limiting example shown, the transducer 1505 may be an electric motor with an eccentricity on the output shaft thereof. The transfer 1530, 1535, 1540 of energy into the surface 1502 of the subject 1501 may induce a range of sensations, from poking, to rubbing, dependent upon the amplitude, frequency, duration, duty cycle of the transducer 1505 as well as the physical configuration of the patch/module pair 1500 and the choice of adhesive layer 1510, if such a layer is used in the embodiment in question.

[00489] Figs. 16a – c show non-limiting arrangements of patches on a subject 2 for performing and generating a multi-lead equivalent EKG in accordance with the present disclosure. Fig. 16a shows an arrangement of three patches 1610a-c in accordance with the present disclosure arranged on the torso of a subject 2 so as to cover equilateral vectors passing out from the heart of the subject 2.

[00490] Fig. 16b shows an arrangement of four patches 1620a-d in accordance with the present disclosure arranged on the torso of a subject so as to cover vectors with trajectories leading away from the heart (patches 1620a,c,d) and to capture one or more signals near to the heart 1620c of the subject 2.

[00491] Fig. 16c shows an arrangement of three patches 1630a-c in accordance with the present disclosure arrange on the torso of the subject forming roughly a linear arrangement spanning from the sternum of the subject 2 across the left torso thereof.

[00492] The patches 1610a-c, 1620a-d, 1630a-c may include bipolar, tripolar, quadripolar, and/or multipolar electrode arrangements in accordance with the present disclosure. In aspects, the patches 1610a-c, 1620a-d, 1630a-c may include optical sensors positioned so as to determine local blood flow dynamics below each patch 1610a-c, 1620a-d, 1630a-c and may include one or more physical sensor (e.g. accelerometer, gyroscope, etc.) for purposes of addressing movement artifacts in the captured signals, etc.

[00493] Figs. 17a-c show aspects of electrode features and methods for engaging such features with skin in accordance with the present disclosure.

[00494] Fig. 17a shows aspects of a patch 1701 in accordance with the present disclosure including a conductor 1710 and an electrode with one or more electrode features 1715 in accordance with the present disclosure thereupon. The electrode features 1715 may be forced into engagement with an adjacent tissue surface via a bias force 1716 (i.e. as may be applied by a thumb over top thereof). In aspects, one or more of the electrode features 1715 may be electrically isolated from the others, so as to form a microelectrode in accordance with the present disclosure.

[00495] Fig. 17b shows a 3 part sequence for engaging a series of electrode features 1721 with a skin surface 13 in accordance with the present disclosure. Initially (in frame 1), the electrode features 1721 are included on a conducting surface, in this case a bistable structure 1719 (i.e. a snap dome in this example). The bistable structure 1719 and the electrode features 1721 are electrically and mechanically connected into a substrate 1717 in accordance with the present disclosure, thus forming part of a patch in accordance with the present disclosure.

[00496] After placement onto the skin 13 of a subject, referring to frame 2, a bias force 1723 may be applied to the bistable structure 1719 causing the electrode features 1721 to engage with the skin 13 (e.g. to embed into the skin 13, penetrating 1725 one or more of the electrode features 1721 into the stratum corneum thereof).

[00497] Referring to frame 3, upon release of the bias force 1723, depending on the nature of the bistable structure 1719 the electrode features 1721 may disengage from the skin 13 leaving behind a series of micropunctures 1727, thus lowering the impedance of the electrical connection between the patch and the subject (i.e. via the skin 13). In aspects, the bistable structure 1719 may be configured to remain in the deformed position (i.e. plastically deform so as to engage one or more of the electrode features 1721 with the skin 13 long term during the monitoring process).

[00498] In aspects, the patch may include a hydrogel (not explicitly shown) located in and around the electrode features 1721 to maintain conductivity after retraction of the electrode features 1721 from the skin 13. The hydrogel may be intrinsically and/or ionically conducting, and may be patterned underneath the electrode, printed onto the electrode features 1721, etc.

[00499] Fig. 17c shows a 3 part sequence for engaging a series of electrode features 1747 with a skin surface 14 in accordance with the present disclosure.

Initially (in frame 1), the electrode features 1747 are included on a conducting surface, in this case a conducting trace 1745 coupled with a substrate 1741 in accordance with the present disclosure, thus forming part of a patch 1740 in accordance with the present disclosure. The patch further includes a gel adhesive 1743 (such as an intrinsically or ionically conducting gel adhesive, hydrogel adhesive, etc. in accordance with the present disclosure). In aspects, the gel adhesive 1743 may be patterned, formed, or foamed so as to provide compressible aspects (i.e. so as to allow for volumetric changes during compression thereof). In aspects, the gel adhesive 1743 may be patterned onto the substrate 1741 and conducting trace 1745 so as to embed the electrode features 1747 within the gel adhesive 1743 (i.e. so as to just cover the electrode features 1747 such that upon placement of a skin surface 14, the electrode features 1747 do not engage with the skin surface 14).

[00500] After placement onto the skin 14 of a subject, referring to frame 2, a bias force 1749 may be applied to patch 1740 in the general vicinity of the electrode features 1747 causing the electrode features 1747 to engage with the skin 14 (e.g. to embed into the skin 14, penetrating 1753 into the stratum corneum thereof). During application of the bias force 1749, the gel adhesive 1743 may deform 1751 allowing for the electrode features 1747 to penetrate into the skin 14.

[00501] Referring to frame 3, upon release of the bias force 1749, the gel adhesive 1743 may (may be slowly) return 1757 to a somewhat undeformed state and thus the electrode features 1747 embedded therein may disengage from the skin 14 leaving behind a series of micropunctures 1759, thus lowering the impedance of the electrical connection between the patch and the subject (i.e. via the skin 14). In aspects, the gel adhesive 1743 may be configured to maintain a relatively humid and biofriendly environment in the vicinity of the micropunctures 1759, thus maintaining the micropunctures for a prolonged monitoring period without irritating the skin 14.

[00502] One or more of the electrode features 1715, 1721, 1747 may be configured as a microelectrode in accordance with the present disclosure.

[00503] Fig. 18 illustrates an isolation patch 1801 in accordance with the present disclosure. The isolation patch 1801 may be configured to allow a user to electrically interface a handheld monitoring device 1805 (e.g. a handheld EKG monitor, an Alivecor™ device, etc.) with a plurality of electrodes 1807a,b with a subject 15

without bringing the device 1805 into physical contact therewith. In aspects, the isolation patch 1801 may include a substrate in accordance with the present disclosure and include one or more patterned regions 1803 thereupon through which a conduction adhesive layer patterned on the substrate of patch 1801 may be accessed from the surface of the substrate 1801 facing away from the subject 15 upon placement. To monitor the patient, the user may position the handheld monitoring device 1805 against the isolation patch 1801 such that electrodes 1807a,b touch the patch 1801 on the regions 1803.

[00504] In aspects, the adhesive layer may be formed from a conducting hydrogel adhesive, from a z-axis adhesive, etc. in accordance with the present disclosure.

[00505] Fig. 19 shows aspects of a modular physiologic monitoring kit in accordance with the present disclosure. The kit includes one or more modules 1905, patches 1901 (or equivalently sets of patches provided for implementation in specific monitoring functions), a recharging bay 1915 and optionally one or more accessories (e.g. such as a substantially pain free patch remover 1935, skin prep pads, etc.) each in accordance with the present disclosure. In aspects, the recharging bay 1915 may also provide functionality of a host device 1915. As a host device 1915, the recharging bay 1915 may include an antenna 1925 for communicating 1927 with one or more modules 1905 during a monitoring session.

[00506] To interface with the modules 1905, the recharging bay 1915 may include one or more docking receptacles 1907 each including a connector 1911 for interfacing with a module 1905 docked therein.

[00507] In aspects, the recharging bay 1915 may include one or more diagnostic indicators 1920 and a power supply 1930 in accordance with the present disclosure. In aspects, the recharging bay 1915 may be integrated into a monitoring display (e.g. a bedside monitor for placement within an ICU, OR, hospital, hospice, or homecare setting), etc.

[00508] In aspects, the recharging bay 1915 may include an alarm clock, for placement beside the bed of a subject (e.g. for easy swap out of modules 1905 during a prolonged monitoring session, for ease of use during sleep studies, etc.).

[00509] Figs. 20a – l illustrate aspects of a multi-sensor patch including a plurality of microelectrodes in accordance with the present disclosure. The non-limiting

aspects of the patches are implemented so as to facilitate, enhance, and/or simplify the monitoring of neural activity from a tissue region (e.g. skin, mucous tissue, etc.), upon which the patch has been placed. Such implementations may be advantageous for monitoring one or more neural activities in the adjacent tissue (e.g. skin SNA, SNS, PNS, somatosensory function, etc.). Non-limiting examples of sub-dermal and dermal microelectrodes are illustrated throughout Figs. 20a – j. Such microelectrodes may be advantageous to interact locally with subcutaneous structures in an adjacent volume of tissue. The microelectrodes may establish local monitoring of signals between different regions or individual sensors in an array (so as to remove artifacts and to tease out neural signals from those electrodes which happen to be closest to the SNA traffic, irrespective of placement).

[00510] Fig. 20a illustrates a patch 2001 in accordance with the present disclosure, the patch 2001 includes a substrate 2003, a plurality of microelectrodes 2005, coupled with the substrate, arranged on the substrate 2003 so as to interface with an adjacent tissue surface, the microelectrodes 2005 coupled to a connector/microcircuit 2009 by a plurality of electrical traces 2007 (the electrical traces 2007 mechanically coupled to the substrate 2003). The connector/microcircuit 2009 may be configured as a connector, the connector arranged such that a microcircuit in a corresponding module (not explicitly shown), may be configured so as to interface with the microelectrodes through the connector when coupled thereto. The connector/microcircuit 2009 may be configured as a microcircuit 2009, the microcircuit 2009 configured to accept signals from one or more of the microelectrodes 2005, and to generate one or more signals or metrics therefrom, convey one or more of the signals to a corresponding processor, etc.

[00511] The microelectrodes 2005 may be patterned over the substrate 2003 and coupled to the microcircuit 2009 (within the patch 2001 or within an associated module), the pattern formed so as to generate an activity map, to increase the probability of interacting with a specific neural structure in the tissue, to allow for readings to be made from two substantially different types of tissue (glabrous skin, mucous membranes, hairy skin, skin with proportionally high concentrations of sympathetic nerves, high concentrations of parasympathetic nerves, somatosensory nerves, etc.).

[00512] Fig. 20b illustrates a side view of the patch 2001 illustrating the isolation of each microelectrode 2005 from one another, also illustrating a non-limiting example of vias 2011 to interconnect the microelectrodes 2005 with the interconnects 2007. The patch 2001 is shown with a connector/microcircuit 2009 coupled to the interconnects 2007, the connector/microcircuit 2009 optionally potted within a housing 2013 so as to protect the circuit 2009 from the surroundings, to protect the interconnects 2007, etc.

[00513] In aspects, the overall thickness of the patch 2001 may be substantially thin, breathable, stretchable, etc. so as to minimize environmental impact on the underlying tissues during use (so as to minimize hydration of the tissues, limit potential tissue maceration, etc.).

[00514] As shown in Fig. 20a,b the microelectrodes 2005 may be configured as gel electrodes, as ionically and/or electrically conducting adhesives, etc. The microelectrodes 2005 may be generally electrically isolated from each other while isolated from an adjacent tissue surface. The microelectrodes 2005 may be individually connected to signal conditioning circuitry or collectively coupled together (e.g. two or more microelectrodes 2005 electrically coupled together so as to form a larger overall electrode surface). Such coupling may be completed physically (e.g. by layout thereof, through coupling with a connector network, etc.), or virtually (e.g. via algorithmic combination post signal conditioning, via a signal fusion algorithm, through coupling to a neural network, etc.).

[00515] Fig. 20c illustrates a schematic of aspects of a patch 2015 in accordance with the present disclosure, the patch 2015 including a plurality of microelectrodes 2019, each microelectrode 2019 coupled via an interconnect 2023 to a connector/microcircuit 2025 in accordance with the present disclosure. The microelectrodes 2019, interconnects 2023, and connector/microcircuit 2025 coupled to a substrate 2017 of the patch 2015. The patch 2015 may include a macroelectrode 2021, the macroelectrode 2021 having an area substantially larger than the average area of one of the microelectrodes 2019, the macroelectrode 2021 being electrically coupled to the connector/microcircuit 2025. The macroelectrode 2021 may be used as a reference electrode, as part of bioimpedance circuit, as a ground electrode, as a guard electrode, as a stretch determining electrode, etc.

[00516] Fig. 20d illustrates aspects of the patch 2015 from a side view so as to show a non-limiting relationship and pattern of the microelectrodes 2019 and the macroelectrode 2021 onto the substrate 2017. The electrodes 2019, 2021 are coupled to the connector/microcircuit 2025, optionally potted in a protective material 2026 to prevent corrosion, water ingress, environmental exposure, etc.

[00517] Fig. 20e illustrates aspects of a side view of a patch in accordance with the present disclosure, the patch including a plurality of microneedle-based microelectrodes 2030, the microelectrodes 2030 including a shank with an exposed region 2028, the exposed region 2028 arranged so as to interface with a target tissue region, a neural structure, a site at a particular depth under the skin, etc. and an isolated region 2029 arranged along the microelectrodes 2030 so as to electrically and/or ionically isolate that region of the microelectrode 2030 from the surrounding tissues, patch, other electrodes 2030, etc. One or more of the microelectrodes 2030 may be electrically coupled to an associated connector/microcircuit 2033 via one or more interconnects 2031. The connector/microcircuit 2033 may optionally be potted in a protective material 2035 in accordance with the present disclosure.

[00518] In aspects, the microneedle-based microelectrodes 2030 may be implemented with a length 2037 (optionally different between electrodes 2030 in the array), the length being defined so as to access a particular neural structure within an adjacent tissues surface. One or more of the electrodes 2030 may include an isolating layer 2029 arranged along a first length 2041 thereof, the remaining length arranged as an exposed region 2028 (e.g. a region for interfacing electrically with an adjacent tissue site, optionally treated, coated, etc.). In aspects, the microneedle-based microelectrodes 2030 may be implemented with a penetrating length 2039 into an adjacent tissue volume, the penetrating length 2039 arranged so as to place the exposed region 2028 thereof near to a neural structure (e.g. a follicular unit, a follicular bulb, arrector pili follicular muscle, a sweat gland, a somatosensory receptor, a sebaceous gland, a Merkel cell, a Ruffini cell, a Meissner corpuscle, a longitudinal lanceolate ending, a Pacinian corpuscle, a free nerve ending, combinations thereof, or the like). In aspects, the length 2037 of one or more microelectrodes 2030 may be in the range of 20 – 3000um, 30 – 1000um, 40 – 500um, 50 – 200um, etc. so as to penetrate the skin and interact with a neural structure therein. In aspects, the length of one or more of the microelectrodes 2030

may be tailored so as to align with a target neural structure within a particular type of tissue under study.

[00519] Fig. 20f illustrates aspects of a patch with a plurality of needle-like microelectrodes 2030 coupled with a substrate 2043, and one or more interconnects 2049 each in accordance with the present disclosure. The microelectrodes 2030 include an exposed region 2044 and an insulated region 2045 so as to interact with one or more tissue sites with an adjacent volume of tissue during use. The patch includes a controller/microcircuit 2051 coupled to the interconnects 2049, and optionally a protective coating 2053. The patch may include a macroelectrode 2047, coupled to the substrate 2043, the macroelectrode 2047 arranged among, or around the micoelectrodes 2030 so as to provide a reference electrode, a guard electrode, to minimize stretch-based artifacts, movement based artifacts, etc. from corrupting signals taken by one or more of the microelectrodes 2030.

[00520] Fig. 20g shows aspects of a patch with a plurality of microelectrodes 2057 each in accordance with the present disclosure, each microelectrode 2057 coupled to the substrate 2055. The patch also including a ring-like macroelectrode 2059 surrounding one or more regions of the patch that are patterned with microelectrodes 2057. The macroelectrode 2059 may provide a reference electrode, a guard electrode, a stretch insensitive electrode configuration, etc. for use in the monitoring of electrophysiological function from a skin surface of a subject.

[00521] Fig. 20h shows aspects of a patch with a plurality of needle-like microelectrodes 2068 each in accordance with the present disclosure, each microelectrode 2068 coupled to the substrate 2065. The patch also including an inter-dispersed macroelectrode 2069 intimately surrounding the microelectrodes 2068 (e.g. a region of microelectrodes 2068, a single microelectrode 2068, a group of microelectrodes 2068, clusters of microelectrodes 2068, all microelectrodes 2068, etc.), etc. In aspects, the macroelectrode 2069 may be patterned such that small gaps 2067 are formed between the macroelectrode 2069 and the microelectrodes 2068. Additionally, alternatively, or in combination the macroelectrode 2069 may be closely coupled to the isolating region of one or more microelectrodes 2068, the exposed region of the microelectrodes 2068 being substantially removed from the region occupied by the macroelectrode 2069. The macroelectrode 2069 may provide a reference electrode, a guard electrode, a stretch insensitive electrode configuration,

etc. for use in the monitoring of electrophysiological function from a skin surface of a subject.

[00522] Fig. 20i illustrates aspects of a patch, demonstrating a plurality of micoroelectrodes 2057 coupled to a substrate 2055 and a ring-like macroelectrode 2059 coupled with the substrate 2055, a collection of vias 2063 connecting one or more of the microelectrodes 2057, the macroelectrode 2059, an additional sensor in accordance with the present disclosure, or the like to an interconnect 2061, a shield or guard (not explicitly shown), etc. Such a configuration may be advantageous for isolating one or more of the microelectrodes 2057 from surrounding electromagnetic interference, for substantially forming a Faraday cage around one or more of the microelectrodes 2057, etc.

[00523] Fig. 20j shows aspects of a patch 2070 in accordance with the present disclosure, the patch 2070 including a substrate 2071 for providing interconnection of one or more components thereof, and a plurality of macroelectrodes 2075, 2079, each surrounding a group of microelectrodes 2073, 2077 respectively. The macroelectrodes 2075, 2079 and microelectrodes 2073, 2077 may be coupled to a switch network, a connector, a microcircuit, etc. each in accordance with the present disclosure. The macroelectrodes 2075, 2079 may be configured to measure a macro-electrophysiological signal (e.g. an EKG, an EMG, an EEG, an ERG, etc.), act as a reference electrode, as a guard electrode, to provide a ground electrode, etc. while the microelectrodes 2073, 2077 may be configured to monitor one or more neural activity signals from a nearby neural structure.

[00524] In aspects, such a configuration may be advantageous for simultaneously monitoring multiple tissue types located near to each other on a subject. Such tissue sites may include monitoring a first region of tissue served primarily by a first neural branch, and a second region of tissue served primarily by a second neural branch, the branches having different SNS, PNS, somatosensory innervation or receptor concentration, etc. The patch 2070 may be placed such that the macroelectrdoes 2075, 2079 are placed against the different zones respectively. The signals may be harvested from the electrodes and compared region for region to determine the differential response to a stress test thereupon, to locate region boundaries and to subsequently simulate only a first region and not a second region, etc.

[00525] Fig. 20k illustrates a patch 2080 in accordance with the present disclosure, the patch 2080 including a substrate 2081 and a variety of components coupled thereto (either directly or via inclusion in an associated module in accordance with the present disclosure). The patch 2080 includes a region 2083 coupled to the substrate 2081 configured to monitor local neural traffic (e.g. skin SNA, PNA, somatosensory response, etc.), from an adjacent tissue surface, a perfusion sensor 2085 in accordance with the present disclosure, a temperature sensor 2087, and/or a heat transfer sensor 2089 each in accordance with the present disclosure. The perfusion sensor 2085 may be configured to monitor changes in optical absorption, reflection, etc. caused by local constriction or dilation of microvasculature in an adjacent skin surface. The temperature sensor 2087 may be configured to analyze changes in the temperature of a volume of tissue under the patch 2080. The heat transfer sensor 2089 may include a plurality of temperature sensors, a plurality of sensors amid varying heat transfer pathways, an airflow sensor, a ambient environment heat transfer estimator (e.g. a thermal mass transfer estimating sensor), or the like.

[00526] Fig. 20l illustrates a patch 2090 in accordance with the present disclosure including substrate 2091 upon which one or more components is arranged, coupled to, etc. The patch 2090 includes a region 2093 configured to measure neural traffic, skin neural traffic, etc. in accordance with the present disclosure, a stimulator 2095 coupled to the substrate 2091 (e.g. either directly or via an associated module), and an optional feedback sensor 2097 coupled to the substrate 2091 (e.g. either directly or via an associated module). The stimulator 2095 may be configured to deliver a stimulus to an adjacent tissue volume upon interaction therewith, such stimulation may include electrical stimulation, thermal stimulation, heating, cooling, tactile stimulation, vibration, stroking, scratching, penetrating, chemical irritation of tissues, chemical block of neural traffic, tissue function, or the like. The feedback sensor 2097 may incorporate one or more sensors such as temperature sensors, vibration sensors, accelerometers, current sensors, or the like, so as to provide feedback to a circuit, processor, etc. coupled thereto. The feedback sensor 2097 configured to convey a feedback signal to the circuit, processor, etc. during use. The feedback signal used by an associated processor, etc. to more precisely control the stimulus, confirm delivery of the stimulus, adjust the amplitude of the stimulus, etc.

[00527] In aspects, a patch, module, or patch/module pair in accordance with the present disclosure may include one or more sensors, physiologic sensors, electrodes, electrophysiological sensors, environmental sensors, kinematic sensors, kinetic sensors, proprioceptive sensors, analyte sensors, perfusion sensors, galvanic skin response sensors, bioimpendance sensing, combinations thereof, or the like as outlined throughout the present disclosure.

[00528] In aspects, one or more microelectrodes and/or exposed regions of one or more microneedle based microelectrodes each in accordance with the present disclosure may be formed with an active surface area of less than $4mm^2$, less than $1mm^2$, less than $100{,}000um^2$, less than $10{,}000um^2$, less than $1{,}000um^2$, less than $100um^2$, or the like. Such sizing may be selected so as to focus the range that the microelectrode may interact with a nearby tissue volume, provide a means for multi-unit and/or single unit study of neural structures in the skin, etc.

[00529] In aspects, one or more macroelectrodes in accordance with the present disclosure may be formed with an active surface area of greater $5mm^2$, greater than $50mm^2$, greater than $100mm^2$, greater than $500mm^2$, or the like. Such sizing may be advantageous for capturing artifacts, movement artifacts, monitoring macroscopic electrophysiologic fields, local electrical potentials, bioimpedance of adjacent tissues, providing reference electrode function, providing a guard electrode function, providing a drive electrode function, combinations thereof, or the like.

[00530] Fig. 21 shows aspects of a circuit schematic to interface with a plurality of microelectrodes in accordance with the present disclosure. The circuit schematic is arranged such that signals monitored locally by one or more electrodes in the array may be aggregated, accentuated, extracted from the group behavior, combined with grouped reference signals, compared against each other, compared against algorithmic or transfer function generated metrics, signals, etc. The circuit schematic illustrates electrical coupling to one or more microelectrode banks 2101a,b, and/or one or more macroelectrodes 2103a,b. Each electrode 2101a,b, 2103a,b may be coupled to a subject during a monitoring session. The microelectrodes in a bank 2101a,b, macroelectrodes 2103a,b, etc. may be positioned against a subject so as to cross different regions of neural innervation, to interact with different neural structures (e.g. based upon penetration depth into the tissues, based upon location on the body, a neural plexus near to the surface of the tissues, etc.). In aspects, one or more

electrodes 2101a,b, 2103a,b may be nearer or farther from a target neural structure to capture the activity thereof during a monitoring session. The microcircuit may be configured to generate one or more output signals 2121 in order to interface with one or more stimulatory transducers, heating elements, thermoelectric units, electrodes, coils, etc. so as to perform a stress test on a region of the body. The microcircuit may be configured to couple with one or more sensors 2117 in accordance with the present disclosure, the sensors 2117 configured to monitor one or more physiologic signals, environmental signals, a stimulatory feedback signal, etc. as part of a monitoring process performed on a subject.

[00531] The microcircuit may include one or more processing banks 2109a,b the banks configured to interface with a corresponding microelectrode bank 2101a,b and/or one or more macroelectrodes 2103a,b each in accordance with the present disclosure. The processing bank 2109a,b including a preamplification / switch bank (PASB) 2105a,b and a signal conditioning / analog to digital conversion circuit (SCADC) 2107a,b. The PASB 2105a,b may include a switch bank configured to interconnect one or more electrodes 2101a,b, 2103a,b to each other or to a corresponding preamplifier bank and/or a preamplifier bank configured to extract the low level physiologic signals from the overall signals acquired by the electrodes 2101a,b, 2103a,b during use. The switch bank may be coupled to a processor via a control line, and may be adaptively configured to select individual signals from microelectrodes, electrically tie multiple microelectrode signals together, create a low noise analog interconnection between a microelectrode, a macroelectrode, and one or more inputs to a preamplifier, a second stage amplifier, etc. In aspects, the switch bank and/or preamplifier bank may include one or more conditioning circuits, configured so as to substantially reduce signal content from an extracted signal outside of a target bandwidth, to extract stretch related, movement related, artifacts, to compare a signal against a reference, etc.

[00532] In aspects, the preamplifier bank may be configured to interface a plurality of microelectrode inputs against a single reference input (such as may be obtained from a macroelectrode), etc. The switch bank may be configured so as to select the microelectrode inputs that are coupled to the preamplifier bank. In aspects, the preamplifier bank may include an input for each microclectrode in a microelectrode

bank 2101a,b such that the individual signals may be analyzed, compared, selected, etc. during a monitoring process.

[00533] The microcircuit and/or processing banks 2109a,b may include a SCADC circuit 2107a,b coupled to the PASB 2105a,b, the SCADC 2107a,b configured so as to convert one or more signals to a digital form, metrics, or the like, include one or more signal conditioning circuits, hardware descriptive forms, and/or programs to perform one or more analog signal conditioning functions, one or more digital signal conditioning functions, etc., the SCADC 2107a,b and/or PASB 2105a,b coupled to a processor 2111 so as to communicate the associated signals thereto. The processor 2111 may include or be coupled to one or more peripherals, a radio, a battery, etc. in order to further communicate 2113, wirelessly communicate 2115, a signal, metric, etc. related to the monitoring, to a display, processor, an operator, the subject, a feedback system, a report generator, etc.

[00534] The processor 2111 may be coupled to one or more sensor conditioning circuits, control circuits, and/or ADC circuits 2119, so as to couple with one or more sensors 2117 in the patch, module, or patch/module pair, or the like. The processor 2111 may be configured to condition and/or compare one or more sensory signals with one or more activities extracted from the electrodes 2101a,b, 2103a,b to generate a metric, remove an artifact, to generate a report, a diagnostic signal, etc.

[00535] The processor 2111 may be coupled with a controller 2132, the controller including one or more circuits to interface with one or more stimulators 2121, energy delivery units, sensor energy delivery aspects, etc. in accordance with the present disclosure. The processor 2111 may be programmed with a control algorithm to control the output 2121 based upon one or more of the inputs 2117, 2101a,b, 2103a,b, etc.

[00536] In aspects, the SCADC 2107a,b and/or PASB 2105a,b may be arranged so as to highlight and/or extract one or more key signal components from the overall signals obtained from the electrodes 2101a,b, 2103a,b. In aspects, such information as individual action potential properties, cluster dynamics of successive action potentials, relationships between the state of a stimulatory input and a train of related action potentials, etc. may be extracted for further use, for generation of a metric, etc.

[00537] In aspects, the microcircuit may be arranged to separate one or more of the microelectrode 2101a,b signals from the group, or to generate a metric from one or more signals, to compare the signals against an algorithmic function of the overall signal population, compare signals across multiple tissue sites, across multiple tissue types, across alternative types of tissue coupled with the same patch/module pair, combinations thereof, or the like.

[00538] Figs. 22a – c show aspects of multi-site monitoring, stimulation, stress application, and/or treatments applied to a subject each in accordance with the present disclosure. Fig. 22a illustrates a subject 25 adorned with a plurality of patches 2201, 2203, 2205, 2207, 2209, 2211, 2213 each patch configured to interface with the subject, measure one or more physiologic parameter from the subject, apply one or more stimulatory input to a subject, or the like. The subject has a temporally applied patch 2201, arranged near to the eye of the subject. A temporally applied patch 2201 may be configured to monitor one or more ocular inputs, facial muscle tone, ocular muscle tone, neural traffic associated with the eye, the retina, the pupil, a neural ganglion (e.g. such as may be related to the onset of cluster migraine headaches, etc.) via inclusion of one or more sensors each in accordance with the present disclosure. In aspects, the temporally applied patch 2201 may include one or more energy or stimulus delivery elements, a thermal regulating unit, an electrical stimulator, a light source, a tactile stimulator, etc. in order to stress the subject near to the ocular circuits. A neck applied patch 2203 has been applied to the subject 25. The neck applied patch 2203 may be configured so as to monitor one or more muscular activities, thyroid and/or parathyroid activities, neural traffic along the carotid artery, activity around the carotid sinus, near the carotid body, muscular tone along the larynx, trachea, swallowing activity, choking, occlusion, etc. In aspects, the neck applied patch 2203 may include one or more energy or stimulus delivery elements, a vibratory stimulating element, a tactile stimulating element, an electrical stimulator, a thermal regulating unit, etc. in order to stimulate one or more neural structures in the neck of the subject 25. Such stimulation may be advantageous to interact and/or stimulate one or more neural structures, nerves, and/or receptors such as near to or within a carotid sinus, a carotid body, a vagus nerve plexus, a baroreceptor, a chemoreceptor, a cutaneously innervated region of tissue, or the like located in the neck of the subject 25.

[00539] A groin applied patch 2205 has been applied to the subject 25. The groin applied patch 2205 may be configured to monitor one or more of an autonomic nerve activity, a peroneal nerve activity, a pudendal nerve activity, a lumbar sympathetic nerve activity, a dorsal nerve activity, a splanchnic nerve activity, a hypogastric plexus activity, a femoral nerve activity, a popliteal nerve activity, a scrotal nerve activity, activity in a cutaneously innervated volume of tissue, SNS activity, PNS activity, somatosensory activity, local tissue perfusion, local sweating, local EMG, local smooth muscle EMG, etc. In aspects, the groin applied patch 2205 may include one or more energy or stimulus delivery elements, a vibratory stimulating element, a tactile stimulating element, an electrical stimulator, a thermal regulating unit, etc. in order to stimulate one or more neural structures in the groin of the subject 25.

[00540] A thigh applied patch 2207 has been applied to the subject 25. The thigh applied patch 2207 may be positioned so as to record cutaneous innervation related to a obturator plexus, an anterior femoral plexus, a lateral femoral plexus, a branch thereof, of the like. In aspects, the thigh applied patch 2207 may include one or more energy or stimulus delivery elements, a vibratory stimulating element, a tactile stimulating element, an electrical stimulator, a thermal regulating unit, etc. in order to stimulate one or more neural structures in the neck of the subject 25. In aspects, the thigh applied patch 2207 may be arranged such that physiologic signals associated with substantially different neural plexuses may be simultaneously recorded on the subject 25, the differential response measured between the different plexuses may be used to characterize the state, the stress-state response, the health of the ANS of the subject, to assess a local neural block (e.g. to one of the two plexuses, etc.), assess a state or progression of peripheral neuropathy, etc.

[00541] In aspects, one or more patches 2215 (not explicitly shown) in accordance with the present disclosure, may be applied to the ankle, lower limb, foot, or hand of the subject 25.

[00542] A torso applied patch 2209 is shown applied to the subject 25. The torso applied patch 2209 may be configured to monitor one or more physiologic parameters, EKG, heart rate, heart rate variability, cutaneous nerve activity, nipple, areola, near a sebaceous gland, traffic associated with a branch or receptor coupled to the thoraco-dorsal nerve, the thoracic nerve, branches from the second, third, fourth, fifth, and/or sixth intercostal nerves, tissue within the superficial fascia, the subdermal

tissue of the areola, the intercostal brachial nerve, neural structures coupled thereto, innervation near the axilla, nerve traffic near the axilla, the axillary nerve, ulnar nerve, intercostalis nerve, or the like.

[00543] In aspects, one or more patches in accordance with the present disclosure may be used to plan a plastic surgical procedure. In one non-limiting example, the innervation to the breast of a subject varies widely from person to person. Assessment of somatosensory innervation of breast tissues with one or more patches in accordance with the present disclosure may be used to develop a personalized nerve map, to determine which nerves are critical to preserving sensory function of the nipple, areola, etc. of a subject, or the like. Based upon the neural activity mapped around the breast, the surgical approach may be planned so as to avoid key nerve plexuses associated with the sensory function to be preserved. Such a process may be advantageous for performing nerve sparing plastic surgeries, restoration of sensation to a tissue volume in a region of a subject, nerve sparing tumor excision surgeries, etc.

[00544] An abdominally applied patch 2211 has been attached to the subject 25. The abdominally applied patch 2211 may be configured to monitor respiration, posture, movement, to generate a feedback signal associated with the respiration to help guide the subject in breathing (e.g. to help control the breathing rate, breathing depth, diaphragmatic breathing of the subject, etc.), skin neural activity, autonomic neural activity, etc. In aspects, the abdominally applied patch 2211 may include one or more energy or stimulus delivery elements, a vibratory stimulating element, a tactile stimulating element, an electrical stimulator, a thermal regulating unit, etc. in order to stimulate one or more neural structures in the abdominal region of the subject 25.

[00545] A hand applied patch 2213 has been attached to the subject 25. The hand applied patch 2213 may include one or more sensors each in accordance with the present disclosure, arranged so as to interface with one or more regions of the hand (e.g. palm, wrist, fingers, median nerve branches, radial nerve branches, ulnar nerve branches, and the like). In aspects, the hand applied patch 2213 may be integrated into a glove, a wrist band, etc. so as to be worn by the subject 25. In aspects, the hand applied patch 2213 may include one or more energy or stimulus delivery elements, a vibratory stimulating element, a tactile stimulating element, an electrical stimulator, a thermal regulating unit, etc. in order to stimulate one or more neural structures in the

hand or wrist of the subject 25. In aspects, the patch 2213 or patch/module pair may include a thermoelectric thermoregulating device, the thermoelectric device configured so as to heat, cool, and/or maintain a temperature of one or more skin surfaces on the hand or wrist. In aspects, the thermoelectric device includes a Peltier element, a power supply, and a controller, the thermoelectric device configured so as to cool the tissues of the hand, warm the tissues of the hand, etc.

[00546] In aspects, a stimulus applied to one or more regions of the body of the subject 25 and the resulting physiologic changes thereof may be monitored by one or more of the patches 2201, 2203, 2205, 2207, 2209, 2211, 2213, etc. Such a multi-site monitoring and/or stimulating configuration may be advantageous to assess the functional relationship between a stress input at one site on the body, to an afferent response to the stress at the site, to an efferent response at one or more sites on the body (i.e. sites innervated to varying degrees by one or more autonomic and/or somatosensory branches), etc.

[00547] In aspects, a stimulus may be applied to the subject via a neck applied patch 2203, the stimulus of sufficient amplitude so as to elicit a response from a baroreceptor in the carotid sinus of the subject 25. The patches 2201, 2203, 2205, 2207, 2209, 2211, 2213 configured so as to monitor local responses to the stimulus, signals generated from one or more of the patches 2201, 2203, 2205, 2207, 2209, 2211, 2213 to be communicated to a processor in accordance with the present disclosure. The processor may be programmed with machine readable code so as to accept the signals, condition the signals, analyze the signals, generate one or more metrics therefrom, compare the metrics against a patient history, a patient population, a database of disease state responses, etc. so as to perform an assessment on the subject 25. In aspects, such a procedure may be advantageous to assess the cardiac baroreflex sensitivity (BRS) of the subject (e.g. such a procedure may be a predictor of the response of a subject to a renal denervation procedure <RDN>, etc.).

[00548] Fig. 22b shows a multi-site system for assessing the response of a subject 27 to a stress test, assessment of the ANS of the subject 27, response of the subject 27 to an interventional procedure, state of completion of an ANS affecting interventional procedure, etc. The subject 27 is adorned with a plurality of patches in accordance with the present disclosure 2201, 2203, 2205, 2207, 2209, 2211, 2213, 2215, etc. A delivery tool 2221 may be subcutaneously, endovascularly, percutaneously,

transcutanesouly, etc. interfaced with the subject 27 so as to perform a procedure, deliver a substance, perform a stress test, etc. upon the subject 27, the patches 2201, 2203, 2205, 2207, 2209, 2211, 2213, 2215, configured to monitor the response to the procedure, delivery, stress test, etc. on the subject 27 and, in aspects, to stimulate and/or apply one or more additional stress tests to the subject 27. The delivery tool 2221 may be used to deliver 2223 a substance in accordance with the present disclosure, energy (e.g. as part of a neural blocking, neurostimulation, denervation, etc.), or the like as part of a stress test, procedure, etc. Although illustrated on the wrist of the subject 27, the delivery tool 2221 may interface with substantially any artery, vein, or vessel in the subject 27.

[00549] Fig. 22c shows a multi-site system for assessing the response of a subject 29 to a stress test, assessment of the ANS of the subject 29, response of the subject 29 to an procedure, state of completion of an ANS affecting procedure, etc. The subject 29 is adorned with a plurality of patches in accordance with the present disclosure 2201, 2203, 2205, 2207, 2209, 2215, etc. The subject 29 is positioned near to one or more energy delivery transducers 2225a,b (e.g. HIFU transducers, MR guided HIFU transducers, radiosurgical transducers, proton therapy, x-ray therapy, etc.). As shown in Fig. 22c, the subject 29 is interfaced with a pair of HIFU delivery transducers 2225a,b (could be a single tranducer, a transducer array, multiple transducer arrays, etc.), and a focused delivery of energy 2227a,b towards a target site 2229 in the body of the subject 29. The patches 2201, 2203, 2205, 2207, 2209, 2215 may be configured to monitor one or more aspects of the energy delivery 2227a,b (e.g. such as time of flight assistance for the HIFU delivery transducers 2225a,b, assessment of reflections, assessment of energy delivery levels near critical tissue sites, etc.), and/or the response of the subject 29 to the procedure, assess the completion of the procedure 29 via a method in accordance with the present disclosure, etc. In aspects, the energy delivery 2227a,b may be part of an ablation procedure, a tumor treatment, administration of a neural block, a sympathectomy, a peripheral nerve treatment, a treatment for inflammation at a site in the subject, a neuromodulation procedure, combinations thereof, or the like. In aspects, such a system and/or method may be advantageous to confirm completion of, follow up on, partial completion of, a patient response to, etc. a denervation procedure, a renal denervation procedure, ablation of a renal nerve, ablation of renal artery, an accessory renal artery, or the like.

[00550] According to aspects there is provided a method for treating one or more neural structures in the vicinity of an artery, a renal artery, an accessory renal artery, or the like, including monitoring autonomic neural activity and/or a closely tied surrogate thereof, at one or more sites on or in the body, applying a test bolus of energy (e.g. a substantially low dosage of ultrasound energy, radiation, thermal energy, etc. so as to affect but not substantially damage tissues), in the vicinity of a suspected treatment site (e.g. a site where a target vessel, neural structure, etc. is suspected but not entirely confirmed due to a lack of adequate imaging in the vicinity of the target vessel, lack of distinguishing features of the suspected target area, lack of confirmation of the destination for nerves traveling through the target area, etc.), and assessing the response to the test bolus to determine if the suspected treatment site includes the target neural structures (e.g. autonomic nerves, vessels innervated with such nerves, one or more ganglia, etc.), and if so, performing a substantially durable treatment at the now confirmed treatment site, if not testing another suspected treatment site, or finishing the procedure. In aspects, one or more steps included in a method in accordance with the present disclosure may be applied so as to test various aspects of the treatment, the subject response to the treatment, predict outcome of the treatment, select patients for suitability of performing a treatment, etc.

[00551] Fig. 23 illustrates aspects of a range of different types of skin, innervation, vasculature, and hair follicles thereof and approaches for interaction therewith in accordance with the present disclosure. In aspects, a system, device, and/or method in accordance with the present disclosure may be adapted to interface with a mucosal tissue 31, a glabrous skin site 33, a hairy skin site 35, a neural structure within the skin, near to an arteriole, a venule, an artery, a vein, a hair follicle 37, a Merkel cell 39, a vessel 41, 45, a Pacinian corpuscle 43, near a sweat gland, near a sebaceous gland, an arrector pili follicular muscle, a free nerve ending in the epidermis, dermis, hypodermis, the stratum corneum, stratum lucidum, stratum glanulosum, stratum spinosum, stratum basale, or the like. A device, patch, and/or patch/module pair in accordance with the present disclosure may be configured for placement 2305, 2310 onto the surface of the tissue, one or more microelectrodes in the patch configured so as to penetrate into the skin surface, to a epidermal depth 2320, a dermal depth 2325, a hypodermal depth 2330, etc. so as to interface with a neural structure therein. In aspects, one or more microelectrodes may be sized and shaped so as to penetrate 2315

down a follicle 37 root, so as to interface with a sweat gland, a sebaceous gland, a muscle, a vessel, etc. Depending on the intended application, one or more of the microelectrodes can be sized so as to penetrate into the tissues of the skin to a depth 2320, 2325, 2330 suitable for interfacing with the intended neural structure.

[00552] In aspects, a placed patch 2305, 2310 may include a plurality of microelectrodes (e.g. non-penetrating, penetrating, etc.), the sizing, spacing and patterning of the microelectrodes designed so as to substantially optimally interface with the target neural structures in the tissues.

[00553] Figs. 24a – c illustrate perineal region, cutaneous innervation of the perineal skin, and associated patches in accordance with the present disclosure for interaction therewith. Fig. 24a illustrates a low lumbar region of a subject 49, the subject 49 positioned so as to expose a region 2401, 2403 of the pelvis suitable for monitoring neural traffic near the skin thereof. The foot and lower leg 51 of the subject 49 is elevated and placed against the opposing knee so as to form a space of access along the anterior superior iliac spine and the pubic tubercle along or near to the inguinal ligament. In the vicinity of such a region 2401, 2403, 2409, the femoral nerves, obturator nerves, femoral cutaneous nerves, etc. travel very near to the surface of the skin, providing cutaneous or subcutaneous opportunities to monitor traffic associated therewith (e.g. and cutaneous innervation emerging therefrom), etc. Some non-limiting examples of patch placement locations are shown at sites 2405, 2411 around the inguinal ligament, the perineal region 2407, the inner thigh 2413, the ankle 2415, the foot 2419, 2421, the calf 2417, or the like. Such placement may be advantageously arranged on the subject 49 so as to pick-up more or less innervated regions of tissue, SNS heavy innervated regions, PNS innervated regions, regions dense in sweat glands, regions couple to particular neural plexuses, etc.

[00554] Fig. 24b illustrates some loosely defined regions associated with cutaneous innervation of the pelvis and perineal region of a female subject 53. The non-limiting examples of highlighted regions include a region innervated by the lateral cutaneous branch of the iliohypogastric nerve plexus 2425, the lateral femoral cutaneous nerve plexus 2427a,b, the cutaneous branch of the obturator nerve, 2429a,b, the posterior femoral cutaneous nerve 2431, the inferior rectal/perineal/pudendal nerves 2433, the internal labial nerves 2437a,b, the posterior labial nerves 2435, the femoral branch of the genitofemoral nerve 2439a,b, dorsal/perineal nerves 2441, etc. One or more

patches in accordance with the present disclosure may be interfaced with the regions so as to monitor afferent/efferent traffic, sympathetic and/or parasympathetic outflow, somatosensory function, or the like associated with each region of the pelvis of the subject 53. The patches may be placed generally within regions, across regions, etc. In aspects, the patch monitoring may be used to identify one or more neural branches coupled to a region of cutaneous innervation of a particular subject. In aspects, a patch monitoring procedure may be used to assess neural function in the pelvis, assess autonomic function of the subject 53, response to a stress test, response to a procedure, a stimulus, a medical procedure, a surgical procedure, etc.

[00555] Fig. 24c illustrates regional innervation of the perineal region of a female subject 55 with reference to the vagina 56, and the anus 57 thereof. The perineal region includes branches of the perineal neural plexus 2451, pudendal neural plexus 2453, the posterior cutaneous nerve 2455, the genitofemoral nerve 2457, and the ilioingual nerve 2459. Also illustrated are non-limiting examples of patch placements 2461, 2463, 2465, 2467, placed around the perineal region of the subject 55 so as to interface with one or more branches of the above neural plexuses.

[00556] Figs. 25a – b illustrate cranial and ear cutaneous innervation, and associated patches in accordance with the present disclosure for interaction therewith. Approximate regions of cutaneous innervation are shown on the face and neck, illustrating regions of skin on the face and neck that are innervated by alternative branches of parent neural plexuses. Some non-limiting regions highlighted include regions of skin innervated by one or more branches of a branch of the frontal, orbital and the trigeminal nerves 2511; branches of the zygomatic, infraorbital, maxillary and the trigeminal nerves 2513; occipital nerve branches 2515; ushno-occipital branches, chin, and mandibular trigeminal nerves 2517; occipital nerve branches 2519; auricular nerve branches 2521; subcutaneous branches of the dorsal nerve of scapula 2523; transverse nerves of the neck 2525; and supraclavicular nerves 2527a,b. Also illustrated are non-limiting regions for placement of a device, patch, or the like each in accordance with the present disclosure. A first placement zone 2531 is shown, generally positioned on, within, near to, and/or behind the ear of the subject, a patch placed in this zone 2531 may be advantageously positioned to interact with vagus nerve receptors of the subject, both in terms of sensing vagus nerve and cutaneous vagus nerve outflow, but also in order to apply a stimulus in accordance with the

present disclosure to the vagus nerve, so as to influence traffic, perform a stress test, treat a condition, etc.

[00557] A second placement zone 2533 is shown along the neck near the carotid artery bifurcation of the subject. A patch placed in this zone 2533 may be advantageously positioned in order to interact with the carotid body, neural receptors along the internal carotid artery, external carotid artery, coupled to the carotid body, a nerve running along the carotid artery (e.g. common, internal, external, etc.), a parasympathetic receptor, a glomus cell, a baroreceptor, a sympathetic nerve, a nerve coupled to the cervical ganglion, a carotid sinus nerve, a vagus nerve, a visceral sensory or parasympathetic motor nerve, or the like. Such placement may be substantially optimally oriented so as to interact with, sense traffic associated with, and/or stimulate/stress one or more such neural structures in the neck of the subject.

[00558] A third placement zone 2535 is shown near to the eye of the subject. The third placement zone 2535 may be substantially optimal for monitoring one or more ocular neural or muscular activity, and/or applying a stimulus/stress test to one or more such tissues. Such placement may be advantageous for monitoring one or more of ocular neural tone, facial muscular tone, electroretinography, nasalis muscular tone, temporalis tone, zygonaticus tone, orbicularis tone, occipitofrontalis tone, etc. Such tone may be assessed and change as the relationship between the overall SNS and PNS of a subject change, during the stress state, during a procedure, after completion of a procedure, etc.

[00559] A fourth placement zone 2537 is shown near to the mouth, and optionally over the lips of the subject. Such placement may be substantially optimal for monitoring one or more aspect of PNS innervation, facial muscle activity, facial muscle tone, assessing movement of the tongue, interfacing with a salivary gland, monitoring outflow from a salivary gland, etc.

[00560] Fig. 25b illustrates some non-limiting regions of cutaneous innervation of the ear of a subject. The non-limiting regions include regions innervated by one or more branches of the vagus nerve 2551; great auricular nerve 2553a,b; lesser occipital nerve 2555; and auriculotemporal nerve 2557a,b. Placement and interaction of one or more portions of a device, stimulator, patch, patch/module pair, or the like in accordance with the present disclosure may be interfaced with one or more such

regions 2551, 2553a,b, 2555, 2557a,b in order to interact therewith and assess, stimulate, and/or interact with one or more aspects of the autonomic nervous system, and/or the somatosensory nervous system of the subject.

[00561] Figs. 26a – b illustrate lower lumbar extremity and plantar cutaneous innervation, and associated patches placement regions in accordance with the present disclosure for interaction therewith. Some approximately defined regions include tissue innervated by one or more branches of the lateral femoral cutaneous nerves 2601a,b, lumbionguinal branch of the genitofemoral nerve 2603, ilioinguinal nerve 2605, anterior femoral cutaneous nerves 2607a,b, obturator nerve 2609, peroneal nerve 2611, saphenous nerve 2613a,b, superficial peroneal nerve 2615, sural lateral dorsal cutaneous nerve 2617, deep peroneal nerve 2619, tibial medial calcaneal nerve 2621, lateral calcaneal/dorsal/sural cutaneous nerves 2623, peroneal nerve 2625, common peroneal lateral sural cutaneous nerves 2627, posterior femoral cutaneous nerve 2629, middle cluneal nerve 2631, superior cluneal nerve 2633, iliohypogastric nerve 2635, and the thoracic nerve 2637. Such regions of innervation may be interfaced with, monitored, and/or studied in accordance with the present disclosure. Some non-limiting patch locations are shown on the subject so as to interface with nerves from a particular region, between regions, etc. Non-limiting patch locations are shown including placement of one or more perineal patches 2641, for monitoring one or more nerves in that region in accordance with the present disclosure. Another thigh located patch 2643 is shown attached to the inner thigh of the subject. The thigh located patch 2643 may be advantageous for assessing neural traffic associated with the obturator nerve, etc. Another foot located patch 2645 is shown attached to the upper foot of the subject near to one or more branches of the deep peroneal nerve 2619. A gluteal located patch 2647 is shown attached to near to the buttock of the subject. The gluteal located patch 2647 may be advantageous for assessing neural traffic along the perineal region of the subject, near the perianal tissues, near the pudendal nerve, etc. A peritoneal located patch 2649 is shown attached near the back of the knee of the subject. The peritoneal patch 2649 may be advantageous for monitoring and/or stimulating activity in the vicinity of the peritoneal nerve, the tibial nerve, branches thereof, cutaneous innervation coupled thereto, etc. Such locations may be selected so as to assist with assessing a form of neuropathy, assessing neural traffic to a particular region of tissue, assisting with a nerve blocking procedure, etc.

[00562] Fig. 26b illustrates approximate regions relating to plantar cutaneous innervation, for which one or more patches in accordance with the present disclosure may interact with as part of a procedure, stress test, assessment, etc. Some approximately defined regions of cutaneous innervation associated with one or more branches of parent nerve plexuses include a region coupled to the lateral plantar nerve 2651; the medial plantar nerve 2653; the saphenous nerve 2655; the medial calcaneal nerve branches 2657; and the sural nerve 2659. Such locations may be selected to interact with one or more devices, patches, or patch/module pairs in accordance with the present disclosure so as to assist with assessing a form of neuropathy, assessing neural traffic to a particular region of tissue, assisting with a nerve blocking procedure, assessing the ANS of the subject, monitoring a surgical procedure, monitoring and/or administering a stress test, etc.

[00563] Figs. 27a – b illustrate arm and hand cutaneous innervation, and associated patches in accordance with the present disclosure for interaction therewith. Fig. 27a illustrates cutaneous innervation of the arms of a subject along with non-limiting examples of regions for placement of devices, patches, patch/module pairs in accordance with the present disclosure. Some approximately defined regions of cutaneous innervation associated with one or more branches of parent nerve plexuses including a region coupled to branches of the supra-clavicular nerve 2701a,b, axillary cutaneous nerves 2703a,b, intercostobrachial nerves 2705a,b, medial brachial cutaneous nerves 2707a,b, medial antebrachial cutaneous nerves 2709a,b, radial dorsal antebrachial cutaneous nerves 2711a,b, lateral antebrachial cutaneous nerves 2713a,b, radial superficial nerve 2715a,b, palmar/palmar digital nerves 2717a,b, and the dorsal palmar/palmar digital nerves 2719a,b. An axillary located patch 2721a,b is shown attached near the axilla of the subject. The axillary located patch 2721a,b may be advantageous for monitoring and/or stimulating activity in the vicinity of the axillary nerve, ulnar nerve, sweat glands, perfusion, changes in sweating, local changes in hydration, etc. A palm located patch 2723 is shown attached to the palm of the subject. The palm located patch 2723 may be advantageously located for monitoring and/or stimulating activity in the vicinity of the ulnar nerve, palmar nerves, radial nerves, branches thereof, somatosensory receptors, sweat gland activity, sweating, changes in hydration, onset of sweating, changes in skin conductivity associated therewith, etc. A back-hand located patch 2725 is shown attached to the

backside of the hand of a subject. In aspects, a back-hand located patch 2725 may be used differentially with a palm located patch 2723 in order to assess specific PNS activity, sweat gland activity, changes in perfusion, or the like between the two regions of cutaneous innervation on the hand. Such locations may be selected so as to assist with assessing a form of neuropathy, assessing neural traffic to a particular region of tissue, assisting with a nerve blocking procedure, assessing the ANS of the subject, monitoring a surgical procedure, monitoring and/or administering a stress test, etc.

[00564] Fig. 27b illustrates cutaneous innervation of the hands of a subject along with non-limiting examples of regions for placement of devices, patches, patch/module pairs in accordance with the present disclosure. Some approximately defined regions of cutaneous innervation associated with one or more branches of parent nerve plexuses including a region coupled to branches of the medial antebrachial cutaneous nerves 2709b, radial dorsal antebrachial cutaneous nerves 2711b, lateral antebrachial cutaneous nerves 2713a,b, radial superficial nerve and dorsal digital branches 2715a,b, palmar/palmar digital nerves 2717a,b, and the dorsal palmar/palmar digital nerves 2719a,b, the palmar branch 2731, the median nerve 27, with the palmar nerve branches coupling to the ulnar nerve. Placement of devices, patches, and/or patch/module pairs may be advantageous for interfacing with branches of the radial nerve, ulnar nerve, median nerve, etc. A first hand located patch 2735a,b is shown extending from the palm side of the hand around the 5th metacarpal bone onto the bank of the hand. A second hand located patch 2737a,b is shown extending from the palm side of the hand around the 2nd metacarpal bone on to the bank of the hand. The first and second hand located patches 2735a,b, 2737a,b may be advantageous for establishing differential activities of the palm and back of the hand during interaction therewith. The second hand located patch 2737a,b, may also be advantageous for interacting with the autonomous area of the radial nerve 2745.

[00565] A wrist mounted patch 2739 is shown attached to the wrist of the subject. The wrist mounted patch 2739 may be advantageous for interfacing with one or more activities associated with the median and/or ulnar nerves or branches thereof.

[00566] Also shown are regions for interfacing with the median nerve 2741 and the ulnar nerve 2743 on the digits of the hand. A substantially small patch or device may

be placed around one or more digits of the subject so as to interface with such regions. Such locations may be selected so as to assist with assessing a form of neuropathy, assessing neural traffic to a particular region of tissue, assisting with a nerve blocking procedure, assessing the ANS of the subject, monitoring a surgical procedure, monitoring and/or administering a stress test, etc.

[00567] Figs. 28a – b illustrate torso cutaneous innervation regional guidelines for consideration when interfacing with the body via a system, device, patch, patch/module pair, and/or method in accordance with the present disclosure for interaction therewith. Fig. 28a shows some approximately defined regions of cutaneous innervation associated with one or more branches of parent nerve plexuses including a region coupled to branches of the ophthalmic nerve 2801, maxillary nerve 2803, mandibular nerve 2805, transverse cervical nerve 2807, superclavicular nerves 2809, axillary nerve 2811, anterior cutaneous branches of the intercostal nerves 2813, lateral cutaneous branches of the intercostal nerves 2815, inferior lateral branchial cutaneous nerves 2817, medial brachial cutaneous nerves 2819, iliohypogastric nerve and medial antebrachial cutaneous branches thereof 2821, musculocutaneous nerves and lateral antebrachial cutaneous branches thereof 2823, superficial branch of the radial nerve 2825, median nerve and palmar branches thereof 2827, ulnar nerve, superficial and dorsal branches thereof 2829, lateral and anterior cutaneous branches of the iliohypogastric nerve 2831, femoral branches of the genitofemoral nerve 2833, anterior scrotal branches of the ilioinguinal nerve 2835, dorsal nerves of the penis 2837, femoral nerve branches 2839, and lateral femoral cutaneous nerves 2841. The back side image shows some approximately defined regions of cutaneous innervation associated with one or more branches of parent nerve plexuses including a region coupled to branches of the greater occipital nerve 2802, lesser occipital nerve 2804, greater auricular nerve 2806, dorsal branches of the rami of the thoracic nerve 2808, medial branches of the rami of the thoracic nerve 2810, intercostal nerves 2812, supraclavicular nerves 2814, axillary nerves and superior lateral brachial cutaneous nerves 2816, inferior lateral branchial cutaneous nerves 2820, medial brachial cutaneous nerves and intercostobrachial nerves 2822, posterior cutaneous branches of rami from the radial nerve 2824, iliohypogastric nerve and medial antebrachial cutaneous branches thereof 2826, posterior antebrachial cutaneous nerve 2828, musculocutaneous nerves and lateral antebrachial cutaneous branches thereof 2830,

superficial branch of the radial nerve 2832, median nerve and palmar branches thereof 2834, ulnar nerve, superficial and dorsal branches thereof 2836, lateral and anterior cutaneous branches of the iliohypogastric nerve 2838, lateral branches of dorsal rami of thoracic and lumbar nerves 2840, superior cluneal nerves 2842, middle cluneal nerve 2844, inferior cluneal nerve 2846, lateral femoral cutaneous nerve 2848, and the illioinguinal nerve 2850. Such locations may be selected so as to assist with assessing a form of neuropathy, assessing neural traffic to a particular region of tissue, assisting with a nerve blocking procedure, assessing the ANS of the subject, monitoring a surgical procedure, monitoring and/or administering a stress test, etc.

[00568] Fig. 28b illustrates torso cutaneous innervation regional guidelines for consideration when interfacing with the body via a system, device, patch, patch/module pair, and/or method in accordance with the present disclosure for interaction therewith. Fig. 28a shows some approximately defined regions of cutaneous innervation associated with one or more branches of parent nerves coupled to the indicated spinal nerves, C2 2851a,b, C3 2853a,b, C4 2855a-c, C5 2857a-c, C6 2859a-c, C7 2861a,b, C8 2863a-c, T1 2864, T2 2865a,b, T3 2867a,b, T4 2869a,b, T5 2871a,b, T6 2873a,b, T7 2875a,b, T8 2877a,b, T9 2879a,b, T10 2881a,b, T11 2883a,b, T12 2885a,b, L1 2887a,b, L2 2889a,b, L3 2892, S1 2893, S2 2894, S3 2891a,b, S4 2896, and S5/Col1 2897. Such locations may be selected so as to assist with assessing a form of neuropathy, assessing neural traffic to a particular region of tissue, assisting with a nerve blocking procedure, assessing the ANS of the subject, monitoring a surgical procedure, monitoring and/or administering a stress test, etc.

[00569] Figs 29 a – d, illustrate temporal readings of metrics associated with stress testing and procedures in accordance with the present disclosure. Fig. 29a illustrates a time – signal graph of a metric derived from a neural activity signal 2900 in accordance with the present disclosure as generated by one or more systems, devices, patches, patch/module pairs, or the like. The signal 2900 may be derived from a skin SNA recording (e.g. derived from an absolute value thereof, from a pulse-per-second calculation, a filtered version thereof, from a pulse-per-minute calculation of a predetermined signal type, from a count of afferent signals per unit time, a count of efferent signals per unit time, a count of somatosensory nerve action potentials per unit time, SNS, PNS, a signal relating to vasodilation, vasoconstriction, local blood perfusion, sweat, hydration, etc.). The graph shows a first procedure 2901 in

accordance with the present disclosure being applied to one or more sites on or within the body of the subject (e.g. a stress test, a nerve block, an ablation procedure, a neuromodulation procedure, etc.), for a first period of time 2903, the signal 2900 demonstrating an initial increase in activity and then an overall decrease in activity over a time period following the first procedure 2901. After a delay 2904, a second procedure 2905 is applied to one or more sites on or within the body of the subject. Following this second procedure 2905 the signal 2900 does not substantially change, thus indicating that the first procedure 2901 affected the ANS in a manner that was substantially durable over the timeframe considered.

[00570] Fig. 29b illustrates a time – signal graph of a metric derived from a neural activity signal 2906 in accordance with the present disclosure as generated by one or more systems, devices, patches, patch/module pairs, or the like. The signal 2906 may be derived from a skin SNA recording (e.g. derived from an absolute value thereof, from a pulse-per-second calculation, a filtered version thereof, from a pulse-per-minute calculation of a predetermined signal type, from a count of afferent signals per unit time, a count of efferent signals per unit time, a count of somatosensory nerve action potentials per unit time, SNS, PNS, a signal relating to vasodilation, vasoconstriction, local blood perfusion, sweat, hydration, etc.). The graph shows a first procedure 2907 in accordance with the present disclosure being applied to one or more sites on or within the body of the subject (e.g. a stress test, a nerve block, an ablation procedure, a neuromodulation procedure, etc.), for a first period of time 2909, the signal 2901 demonstrating an initial increase in activity and then an overall decrease in activity over a time period following the first procedure 2907. After a delay 2910, a second procedure 2911 is applied to one or more sites on or within the body of the subject. Following this second procedure 2911 the signal 2906 substantially changes again, demonstrating that the first procedure 2907 did not significantly affect the ANS of the subject in a manner that was substantially durable over the timeframe considered. A third procedure of higher dosage, longer duration, etc. may be attempted to form a durable procedure if that is the desired affect for the given example.

[00571] Fig. 29c illustrates a time – signal graph of a metric derived from a neural activity signal 2912 in accordance with the present disclosure as generated by one or more systems, devices, patches, patch/module pairs, or the like. The signal 2912 may

be derived from a skin SNA recording (e.g. derived from an absolute value thereof, from a pulse-per-second calculation, a filtered version thereof, from a pulse-per-minute calculation of a predetermined signal type, from a count of afferent signals per unit time, a count of efferent signals per unit time, a count of somatosensory nerve action potentials per unit time, SNS, PNS, a signal relating to vasodilation, vasoconstriction, local blood perfusion, sweat, hydration, etc.). The graph shows a first stress test 2913 in accordance with the present disclosure being applied to one or more sites on or within the body of the subject (e.g. a stress test, delivery of a medication, local administration of a chemical specie, consumption of a drug, questioning the subject, stimulating one or more sites on the subject's body, presenting a visual and/or auditory input to the subject, etc.), for a first period of time 2915, the signal 2912 demonstrating a small change in signal level 2917 over a monitoring period following application of the first stress test 2913. After completion of the monitoring period, one or more additional stress tests may be performed on the subject so as to gauge other metrics, ANS relationships, to generate dose response relationships, etc. In this non-limiting example, the signal 2912 did not change substantially during the monitoring period. Such small changes may indicate that the subject is not a suitable candidate for a procedure, the subject's ANS or the aspect monitored thereof is not substantially influenced by the stress test 2913, etc.

[00572] Fig. 29d illustrates a time – signal graph of a metric derived from a neural activity signal 2918 in accordance with the present disclosure as generated by one or more systems, devices, patches, patch/module pairs, or the like. The signal 2918 may be derived from a skin SNA recording (e.g. derived from an absolute value thereof, from a pulse-per-second calculation, a filtered version thereof, from a pulse-per-minute calculation of a predetermined signal type, from a count of afferent signals per unit time, a count of efferent signals per unit time, a count of somatosensory nerve action potentials per unit time, SNS, PNS, a signal relating to vasodilation, vasoconstriction, local blood perfusion, sweat, hydration, etc.). The graph shows a first stress test 2919 in accordance with the present disclosure being applied to one or more sites on or within the body of the subject (e.g. a stress test, delivery of a medication, local administration of a chemical specie, consumption of a drug, questioning the subject, stimulating one or more sites on the subject's body, presenting a visual and/or auditory input to the subject, etc.), for a first period of time

2921, the signal 2918 demonstrating a strong change in response to the stress test 2919. The signal 2918 begins to change after a short delay, the change continues towards a maximum value within a first timeframe 2927, the maximum change in value being registered 2923, and then recovering back to substantially the resting value over a second timeframe 2925 (herein shown the time between half the maximum change on the up-wing and the down-swing of the signal change). The timeframes 2927, 2925, the maximum change 2923, comparison with other stress tests, other response metrics, or the like, may be advantageous in determining if the subject is a suitable candidate for a procedure, a medical procedure, if the subject has a particular disease state, the extent of the disease state, that subject's ANS or the aspect monitored thereof is substantially influenced by the stress test 2919, etc.

[00573] Fig. 30 illustrates spatial readings of skin neural activity with a mult-sensor patch in accordance with the present disclosure. An array of monitoring sites 3010 is shown in a spatially representative grid (the actual electrode pattern on the patch, between patches, etc. need not be grid like, and may take on substantially any pattern). The individual monitoring sites 3010 may be referenced against a common reference signal, against an algorithmically derived reference, against each other, or the like. Individually obtained signals 3005 are shown at each site 3010, some signals 3005 clearly picking up a physiologic signal near the monitoring site 3010 while other signals 3006 are essentially representative of a baseline reading, an artifact, etc. Clusters 3020, 3025, 3030 of activity are shown at sites within the array. The clusters 3020, 3025, 3030 may be mined, combined, isolated, etc. in accordance with the present disclosure. In aspects, a microcircuit coupled to the array may be adjusted such that signals coming from the clusters 3020, 3025, 3030 are emphasized, sent onward to a processor, processed further, coupled together, combined, compared against on or more other signals, etc.

[00574] Such a configuration, microcircuit, and/or method may include establishing an initial connectivity map between the microelectrodes such that each microelectrode signal can be monitored independently from the others, determining which microelectrodes are picking up the desired activity, and reorganizing the connectivity so as to: collectively couple microelectrodes that are monitoring similar activity, reinforcing a reference signal for those microelectrodes that are recording the

desired activity, providing a guard function for the microelectrodes that are recording the desired activity, etc.

[00575] Figs. 31a – c illustrate aspects of methods for monitoring, stressing, and/or treating one or more regions of a subject each in accordance with the present disclosure. Fig. 31a illustrates aspects of a method for modulating or assessing neural traffic in accordance with the present disclosure. The method includes interfacing one or more systems, devices, patches, and/or patch/module pairs in accordance with the present disclosure to a subject, optionally accessing one or more target sites within a body, applying a stress test in accordance to the present disclosure to the subject, treating one or more tissues in accordance with the present disclosure, administering a medication to the subject, treating the optional target site, stimulating, sensing, or ablating one or more nerves in the subject, augmenting neural activity, treating the afferent nerves and/or receptors, and optionally evaluating one or more physiologic responses, nerve activity, specific nerve activity, a combination thereof, or the like, pre/post stress test, procedure, treatment, etc. to determine if the traffic has been modulated. In aspects, the evaluation maybe performed by comparing a physiologic and/or nerve activity metric before and after treatment (e.g. a change in integrated activity level, a change in phasic response such a shift from a biphasic polarity to a monophasic polarity, a change in action potential firing rate, a change in the spectral content of the firing, etc. associated with the local neural tissues), a differential response between metrics, combinations thereof, or the like. In aspects, the method may include performing a neural block, a temporary neural block, varying a pressure applied to one or more nerves in the subject, stimulating the nerves, and/or receptors, and monitoring afferent nerve activity during such changes in block, stimulus, applied pressure (i.e. monitoring activity during a variable pressure compression block), etc.

[00576] Additionally, alternatively, or in combination with the monitoring of physiologic response and/or electrophysiological activity, the method may include monitoring one or more additional physiologic parameters in accordance with the present disclosure and assessing changes in the parameters before, during, or for a period of time following application of a procedure to the target tissues. In aspects, the additional physiologic parameter may be monitored from a catheter, a pressure sensing catheter, an analyte sensing catheter, etc.

[00577] One or more of the steps may be completed with a guidewire or surgical tool in accordance with the present disclosure. One or more steps may be completed with a radiosurgical system, a HIFU system, a proton therapy device, an ablation catheter, an ablation system, a chemical delivery catheter, combinations thereof, and the like.

[00578] Fig. 31b illustrates a method for assessing the neural structures in the vicinity of a target organ. The method includes interfacing one or more systems, devices, patches, and/or patch/module pairs in accordance with the present disclosure to a subject, optionally accessing one or more target sites within a body, accessing/monitoring (such as communicating with, recording activity from, etc.) one or more neural structures in the body, the nerves associated with the target organ, a related ANS circuit, a skin sympathetic nerve, a skin parasympathetic nerve, a somatosensory nerve, a physiologic parameter, one or more sites related to the disease state to be treated, etc. The method may include monitoring an initial activity level, signal character, periodic element to a signal, afferent or efferent traffic proportion of the neural traffic, etc. The method may include monitoring such activity or metrics associated there with during a stress test in accordance with the present disclosure as applied to the organ, or subject as a whole, a vessel, a skin surface, a tissue volume, etc. The method may include generating and/or analyzing a metric associated with the change in the monitored activity and determining a suitability of the subject for performing a surgical procedure, determining a proportion of nerve types amongst the captured responses, determining if the nerves require treatment, determining the influence of the stressor on the locally measured electrophysiological activity, or the like.

[00579] The method may include modulating a functionality of, neural activity from, afferent activity from, or the like of the target organ of a subject, the method may include selectively stimulating and/or stressing one or more regions of the target organ and monitoring the physiologic response at one or more sites nearby and/or systemically to the stimulus/stress. In aspects, the stimulus/stress response maybe used to identify regions of the target organ that are suitable for neuromodulation to treat a particular condition. In aspects, the method may include selectively treating one or more sites within or in the vicinity of the target organ. In aspects, the method may include monitoring activity and/or local physiologic response to the treatment at

one or more of the sites to determine the extent of the procedure, to evaluate when the procedure has been completed, to decide whether or not to continue with the procedure, etc. The method may include ablating a portion of the organ, or a neurological structure coupled thereto, in accordance with the present disclosure. In aspects, the method may include using surgical system, an interventional device, a guidewire, a catheter, an ablation catheter, and/or surgical device in accordance with the present disclosure to perform one or more of the above steps.

[00580] Fig. 31c shows aspects of a method for treating one or more neural structures in at a site within a subject. The method including accessing the target site (e.g. with a catheter, a guidewire, via a focused energy delivery system, with a chemical substance, etc.), optionally monitoring activity in one or more regions around the target site, treating the nerves, and assessing based on a change in the activity if the treatment was successful. In aspects, the assessment may be determined based on a change in activity level (e.g. pulses per unit of time, drop out of pulses associated with a particular nerve type, changes in traffic associated with a neural circuit biorhythm, a change in subject pain levels, etc.), a shift in the polarity of the signals (i.e. a transition from a biphasic signal related to multi-directional traffic near the vessel, to a monophasic signal related to changes more representative of a uni-directional traffic near the vessel), a drop off in periodic behavior in the captured signals, or the like. In aspects, the, assessment may be determined by combining and/or comparing activity measured at multiple sites on or within the subject, associated vessels, or the like. Such comparison may include assessing a change in coherence between two signals collected from different nearby sites, from a change in one signal wrt the other signal collected from nearby sites, a change in a representative transfer function representative of a correlation between the traffic at one site and the other site, etc.

[00581] If, after the first test, procedure, treatment dose, etc. a response was recognized, a subsequent test, procedure, treatment dose, may be performed to confirm completion of the first test, procedure, treatment dose, etc. If the response occurs again, if a substantial change is monitored after the subsequent test, procedure, treatment dose, etc. further tests, procedures, treatments, etc. may be required to complete the intended task (e.g. neural block, substantially durable neural block, neural remodeling, neuromodulation, stimulating neural block, or the like). If the

response was not observed after the subsequent test, procedure, treatment dose, etc. then the second test, etc. substantially served as a confirmation of adequate dose, etc. applied to the target site(s) of the subject.

[00582] The assessment may include determining if a change in one or more homeostatic functions of the organ have changed in a desired direction, if the response of the neural traffic to a stress test has changed as desired by the therapy, assessing if the subject feels the same, increased, or decreased pain compared with an assessment made before the procedure. If the treatment has been finished, complete the procedure, pull out any system component in the subject, etc. otherwise, monitor activity, continue with treatment, and/or move to a new treatment site in the vicinity of the target site.

[00583] In aspects, an ablation may be performed so as to minimize damage to surrounding tissues. The ablation may include delivering energy to the local tissues in an amount just sufficient to induce irreversible damage to one or more adjacent nerves, but not in an amount sufficient to irreversibly damage other surrounding tissues.

[00584] In aspects, a method in accordance with the present disclosure may be used to assess the durability of a previously applied treatment to a subject. In aspects, a system, device, patch, patch/module pair, and/or method in accordance with the present disclosure may be used for non-invasive sensing of neurological tone or closely coupled surrogates thereof as pertaining to diagnostics, patient selection, procedural feedback, and follow-up assessment of autonomic neural ablation procedures.

[00585] It will be appreciated that additional advantages and modifications will readily occur to those skilled in the art. Therefore, the disclosures presented herein and broader aspects thereof are not limited to the specific details and representative embodiments shown and described herein. Accordingly, many modifications, equivalents, and improvements may be included without departing from the spirit or scope of the general inventive concept as defined by the appended claims and their equivalents.

WHAT IS CLAIMED IS:

1. A patch interface for monitoring neural activity from the skin of a subject, comprising:

a substrate with a surface;

an adhesive coupled to the substrate formulated for attachment to the skin of a subject;

an interconnect embedded into or coupled to the substrate for attachment of the patch to a microcircuit; and

a plurality microelectrodes attached to or embedded into the surface of the substrate, the microelectrodes arranged, configured, and1 dimensioned to interface with the subject when the adhesive is attached thereto, the microelectrodes electrically coupled with the interconnect and/or the microcircuit.

2. The patch interface in accordance with claim 1, wherein the adhesive is patterned onto the substrate so as to form one or more exposed regions of the substrate, one or more of the microelectrodes arranged within the exposed regions.

3. The patch interface in accordance with claim 1 or 2, wherein the adhesive comprises an electrically or ionically conducting component.

4. The patch interface in accordance with claim 3, wherein the adhesive provides a macroelectrode function for interfacing with the subject when attached thereto, the macroelectrode electrically coupled to the interconnect and/or the microcircuit.

5. The patch interface in accordance with claim 4, wherein the macroelectrode provides a guard, a reference, and/or a counter electrode function for one or more of the microelectrodes.

6. The patch interface in accordance with any preceding claim, wherein one or more of the microelectrodes is configured to penetrate into the skin when the adhesive is attached to the subject.

7. The patch interface in accordance with any preceding claim, wherein one or more of the microelectrodes has a needle-like shape, the microelectrode comprising an insulated region and an exposed region.

8. The patch interface in accordance with claim 7, wherein the microelectrode has a length such that the exposed region is positioned within an epidermal layer, a dermal layer, and/or a hypodermal layer of the skin upon attachment of the patch to the surface thereof.

9. The patch interface in accordance with one of claims 7 or 8, wherein one or more of the microelectrodes is shaped so as to penetrate down a root of a follicle, to reach a sweat gland, a sebaceous gland, a muscle, a vessel, and/or an arrector pili follicular muscle upon attachment of the patch interface to the skin.

10. The patch interface in accordance with any preceding claim, wherein one or more of the microelectrodes comprises an electrode feature arranged so as to improve the electrical connection between the microelectrode or an exposed region thereof and the skin upon placement on a subject.

11. The patch interface in accordance with claim 10, wherein the improved electrical connection is achieved after pressure is applied to the microelectrode.

12. The patch interface in accordance with any preceding claim, wherein the microcircuit comprises a switch bank and an amplifier with two or more inputs, the switch bank adaptably configurable to couple one or more of the microelectrodes with an input of the amplifier, and one or more alternative microelectrodes and/or the macroelectrode to another input of the amplifier.

13. The patch interface in accordance with claim 12, wherein the switch bank is adaptively configurable to connect one or more clusters of microelectrodes to a single input of the amplifier.

14. The patch interface in accordance with any preceding claim, wherein one or more of the microelectrodes are configured and shaped so as to penetrate less than 3mm, less than 2mm into the skin, less than 1mm, less than 0.5mm, or less than 0.2mm into the skin during engagement therewith.

15. The patch interface in accordance with any preceding claim, wherein one or more of the microelectrodes and/or exposed regions thereof has an active surface area of less than $4mm^2$, less than $1mm^2$, less than $100,000um^2$, less than $10,000um^2$, less than $1,000um^2$, or less than $100um^2$.

16. The patch interface in accordance with any one of claims 4 – 15, wherein one or more macroelectrodes has an active surface area of greater $5mm^2$, greater than $50mm^2$, greater than $100mm^2$, or greater than $500mm^2$.

17. The patch interface in accordance with claim 16, wherein the macroelectrode is shaped and patterned on the substrate so as to reduce movement artifacts, monitor a macroscopic electrophysiologic field, provide a reference electrode function, provide a guard electrode function, provide a drive electrode function, or a combination thereof during use.

18. The patch interface in accordance with any preceding claim, comprising a sensor coupled with the substrate, and electrically coupled with the microcircuit, the sensor configured to monitor one or more physiologic parameters of the subject when coupled thereto.

19. The patch interface in accordance with claim 18, wherein the sensor comprises an electrophysiologic sensor, a temperature sensor, a thermal gradient sensor, a barometer, an altimeter, an accelerometer, a gyroscope, a humidity sensor, a magnetometer, an inclinometer, an oximeter, a colorimetric monitor, a sweat analyte sensor, a perfusion sensor, a galvanic skin response sensor, a hydration sensor, an interfacial pressure sensor, a flow sensor, a stretch sensor, a microphone, or a combination thereof.

20. The patch interface in accordance with any preceding claim, comprising a stimulating device selected from an electrical stimulator, a thermoregulating device, a heating coil, a thermoelectric device, a Peltier device, a tactile stimulating component, a vibratory stimulating element, or a combination thereof arranged along the substrate so as to interface with the skin of the subject when the adhesive is attached thereto.

21. The patch interface in accordance with claim 20, comprising a feedback sensor coupled with the stimulating device, configured and arranged so as to monitor the interaction of the stimulating device with the skin during activation thereof.

22. The patch interface in accordance with any preceding claim, comprising one or more stretchable electrically conducting traces attached to the substrate, arranged so as to couple one or more of the sensors, microelectrodes, and/or macroelectrodes with the interconnect or microcircuit.

23. The patch interface in accordance with any preceding claim, wherein the patch interface is sufficiently physically frail such that it cannot retain a predetermined shape in a free standing state.

24. The patch interface in accordance with claim 23, comprising a temporary stiffening member attached to the substrate, the temporary stiffening member configured to provide retention of the shape of the patch interface prior to attachment to the subject, the stiffening member being removable from the substrate after attachment to the subject.

25. The patch interface in accordance with claim 24, wherein after removal of the stiffening member, the retention of the shape of the patch interface is provided by the skin of the subject.

26. The patch interface in accordance with any one of claims 23 – 25, wherein removal of the patch interface from the skin of the subject results in a permanent loss in the shape of the patch interface without tearing of the patch interface.

27. The patch interface in accordance with any one of claims 23 – 26, wherein the interconnect is sufficiently frail such that removal of the patch interface from the skin of the subject results in a permanent loss of shape of the interconnect.

28. The patch interface in accordance with any preceding claim, wherein the adhesive has a peel tack to mammalian skin of greater than 0.02N/mm, greater than 0.1N/mm, greater than 0.25N/mm, greater than 0.50N/mm, or greater than 0.75N/mm.

29. The patch interface in accordance with any preceding claim, wherein the patch has a tear strength of greater than 0.5N/mm, greater than 1N/mm, greater than 2N/mm, or greater than 8N/mm.

30. The patch interface in accordance with any one of claims 23 – 29, wherein the ratio between the tear strength of the patch and the peel tack of the adhesive to mammalian skin is greater than 8:1, greater than 4:1, or greater than 2:1.

31. The patch interface in accordance with any preceding claim wherein the substrate is formed from a soft pseudo-elastic material and the patch interface is configured to maintain operation when stretched to more than 25%, more than 50%, or more than 80%.

32. The patch interface in accordance with any preceding claim, wherein the patch interface is configured with a moisture vapor transmission rate of between 200g/m^2/24hrs and 20,000g/m^2/24hrs, between 500g/m^2/24hrs and 12,000g/m^2/24hrs, or between 1,000g/m^2/24hrs and 8,000g/m^2/24hrs.

33. The patch interface in accordance with any preceding claim, wherein the adhesive includes an endothermic-reactive specie, an exothermic-reactive specie, a local analgesic, a neuro-blocker, a neurotoxin, a neural antagonist, or a neural agonist.

34. A module for monitoring one or more physiologic signals, physical signals, skin based neural activity, and/or electrophysiological signals from a subject, comprising:

a housing;

a circuit board comprising one or more microcircuits, the circuit board coupled to the housing; and

a module interconnect coupled to one or more of the microcircuits configured for placement and coupling of the module onto a patch interface in accordance with any one of claims 1 – 33.

35. The module in accordance with claim 34, wherein the module interconnect is embedded into the circuit board.

36. The module in accordance with claim 34 or 35, wherein the circuit board constitutes at least a portion of the housing.

37. The module in accordance with any one of claims 34 – 36, comprising a three dimensional antenna coupled to one or more of the microcircuits, the microcircuits comprising a transceiver or transmitter coupled to the antenna.

38. The module in accordance with claim 37, wherein the antenna is printed on an interior wall of or embedded into the housing, the circuit board providing a ground plane for the antenna.

39. The module in accordance with claim 37 or 38, wherein the housing is shaped like a dome and the antenna is patterned into a spiraling helix centered within the dome.

40. The module in accordance with any one of claims 34 – 39 comprising a sensor coupled with one or more of the microcircuits, the sensor configured to interface with the subject upon attachment of the module to the patch interface.

41. The module in accordance with claim 40 wherein the module comprises a sensor and/or microelectronics configured to interface with a sensor included on the patch interface, one or more of the sensors comprising an electrophysiologic sensor, a temperature sensor, a thermal gradient sensor, a barometer, an altimeter, an accelerometer, a gyroscope, a humidity sensor, a magnetometer, an inclinometer, an oximeter, a colorimetric monitor, a perfusion sensor, a sweat analyte sensor, a galvanic skin response sensor, an interfacial pressure sensor, a flow sensor, a stretch sensor, or a microphone.

42. The module in accordance with any one of claims 34 – 41, comprising a stimulating device selected from an electrical stimulator, a thermoregulating device, a heating coil, a thermoelectric device, a Peltier device, a tactile stimulating component, a vibratory stimulating element, or a combination thereof arranged along the substrate so as to interface with the skin of the subject when coupled to the patch interface.

43. The module in accordance with any one of claims 34 – 42, wherein the module is hermetically sealed.

44. The module in accordance with any one of claims 34 – 43, wherein the module comprises a gasket coupled to the circuit board, the gasket shaped so as to isolate the region formed by the module interconnect and the patch from a surrounding environment, when the module is coupled with the patch.

45. A device for monitoring one or more physiologic, physical, and/or electrophysiologic signals from a subject comprising:

a patch interface in accordance with any one of claims 1 – 33; and

a module in accordance with any one of claims 34 – 44, the module interconnect comprised within the module, sized and dimensioned to interface with the interconnect comprised within the patch interface, wherein to form an operable interconnection between the patch interface and the module, the patch interface must first be coupled to the subject.

46. The device in accordance with claim 45, wherein the module interconnect comprises an electrically conducting magnetic element, and the patch interface comprises one or more ferromagnetic regions coupled to the substrate, the magnetic elements arranged so as to physically and electrically couple the module to the patch interface when the magnetic elements are aligned with the ferromagnetic regions.

47. The device in accordance with claim 46, wherein the ferromagnetic regions are formed from stretchable pseudo elastic material and are printed onto the substrate.

48. The device in accordance with any one of claims 45 – 47, wherein the module and/or the patch interface comprise one or more fiducial markings to visually assist with the alignment of the module to the patch during coupling thereof.

49. A system for monitoring one or more physiologic signals, physical signals, and electrophysiological signals from a subject comprising:

a patch interface in accordance with any one of claims 1 - 33 configured for attachment to the subject; and

a module in accordance with any one of claims 34 - 44 configured and dimensioned to mate with the patch, and to interface with the subject there through, the module configured to convey and/or store one or more physiologic signals, neural activity, and/or physical signals, a signal derived therefrom, and/or a metric derived therefrom obtained via the interface with the subject.

50. The system in accordance with claim 49, comprising a host device coupled in wireless communication or physical communication with the module, configured to receive one or more of the signals and/or metrics therefrom.

51. The system in accordance with claim 50, wherein the host device comprises features for recharging and/or performing diagnostic tests on one or more of the modules.

52. The system in accordance with any one of claims 49 – 51, wherein the system comprises a plurality of modules, the modules being hot swappable with the patch interface, so as to maintain a nearly continuous operation thereof.

53. The system in accordance with any one of claims 49 – 52, wherein the system comprises a plurality of modules and associated patch interfaces for placement onto a single subject, the host device, and/or one or more of the modules configured to coordinate synchronous monitoring of the signals amongst the modules on the subject.

54. The system in accordance with any one of claims 49 – 53, wherein the host device is integrated into a bedside alarm clock, housed in an accessory, within a purse, a backpack, a wallet, is or is included in a mobile computing device, a smartphone, a tablet computer, a pager, a laptop, a local router, a data recorder, a network hub, a server, a secondary mobile computing device, a repeater, or a combination thereof.

55. Use of a device in accordance with any one of claims 45 – 48, or a system in accordance with any one of claims 49 – 54, to confirm completion of, follow up on, confirm partial completion of, monitor a patient response to, or patient selection in connection with, a denervation procedure, a renal denervation procedure, ablation of a renal nerve, ablation of renal artery, or ablation of an accessory renal artery.

56. A method for assessing an autonomic nervous system of a subject, comprising:

monitoring neural activity at one or more sites on the skin of the subject to generate one or more signals;

performing a stress test on the subject; and

analyzing the signals obtained before the stress test, during the stress test, and/or after the stress test to generate a metric, a diagnostic, a report, and/or additional signals therefrom relating to the autonomic nervous system of the subject.

57. The method in accordance with claim 56, wherein the stress test includes administration of a chemical, a drug, medicament, a hormone, an enzyme, a diuretic, a solution, electrolytes, a peptide, steroid, saline, a hypotonic solution, a hypertonic solution, or a combination thereof to the subject.

58. The method in accordance with claim 57 wherein the administration is topical, systemic, intravenous, intra-arterial, intra-parenchymal, sub-dermal delivery, transdermal delivery, rectal, via vaginal suppositories, via urethral suppositories, via nasal suppositories, via rectal suppositories, inhaled, or a combination thereof.

59. The method in accordance with any one of claims 56 – 58, wherein the stress test includes delivery of energy, stimulation of, electrical stimulation of, presenting an audio field to, application of thermal stress to, presenting a light field to, presenting an image to, asking a question to, or playing music for, the subject.

60. The method in accordance with any one of claims 56 – 59, wherein the stress test includes providing a tactile input to one or more sites on the subject.

61. The method in accordance with any one of claims 56 – 60, wherein the stress test includes stimulating one or more of a carotid sinus, a carotid body, a baroreceptor, a vagus nerve receptor, or an erogenous zone of the subject.

62. The method in accordance with any one of claims 56 – 61, wherein the stress test includes applying an electromagnetic field to, injecting a current into, applying pressure to, applying stroking to, or applying a change in barometric pressure surrounding the subject.

63. The method in accordance with any one of claims 56 – 62, wherein the stress test includes having the subject sleep, cry, laugh, lie down, jump, walk, run, change posture, exercise, perform a breath holding exercise, or climb stairs.

64. The method in accordance with any one of claims 56 – 63, wherein one or more of the sites is coupled to the perineal region, the perianal region, the pubic

region, the inner thigh region, the posterior knee region, the neck, the ear, the ocular region, the breast, the axilla, the elbow, the wrist, the palm, the foot, the lips, and/or an erogenous zone of the subject.

65. The method in accordance with any one of claims 56 – 64, wherein one or more of the steps is performed at least in part by a device in accordance with one of claims 45 – 48, a system in accordance with any one of claims 49 – 54, a module in accordance with any one of claims 34 – 44, and/or a patch interface in accordance with any one of claims 1 - 33.

66. A system for performing a neuromodulation and/or an ablation procedure on a target site within a subject comprising:

a treatment system for delivering energy or a chemical to the target site;

one or more monitoring devices configured to interface with the skin of the subject at one or more monitoring sites on the subject, each of the monitoring devices configured to measure one or more electrophysiological signals, neural traffic signals, and physiologic parameters from the monitoring sites so as to produce an activity signal; and

a processor comprised in or coupled to one or more of the monitoring devices, the processor configured to receive the activity signal(s), and/or one or more signals generated therefrom, the processor comprising an algorithm, the algorithm configured to analyze the activity signal(s) to determine the effect of the treatment system on the target site.

67. The system for performing a neuromodulation and/or an ablation procedure on a target site within a subject in accordance with claim 66, wherein the algorithm is configured to compare activity signal(s) generated by a plurality of monitoring devices against each other to determine the effect of the treatment system on the target site.

68. The system for performing a neuromodulation and/or an ablation procedure on a target site within a subject in accordance with claim 66 or 67, wherein the treatment system includes one or more of an ablation system, a neuromodulation device, a neuromodulation implant, an ablation catheter, a focused energy delivery device, a radio frequency ablation system or catheter, a microwave ablation system or catheter, an ultrasound energy delivery system, a high intensity focused ultrasound [HIFU] delivery system or catheter, a cryoablation system or

catheter, a chemical ablation system or catheter, a radiosurgical system, an optical ablation system, an infrared ablation system, a laser ablation system, or an MR guided HIFU system.

69. The system for performing a neuromodulation and/or an ablation procedure on a target site within a subject in accordance with claim 66 - 68, wherein the algorithm is configured to indicate when only a partial neuromodulation and/or ablation procedure has been performed on a target site, and/or when a complete procedure has been performed on the target site.

70. The system for performing a neuromodulation and/or an ablation procedure on a target site within a subject in accordance with claim 66 – 69, wherein one or more of the monitoring devices includes a chemical agent selected from an endothermic reactive specie, an exothermic reactive specie, a local analgesic, a neuro-blocker, a neurotoxin, or a neural agonist for delivery to the associated monitoring site when coupled thereto, the algorithm configured to compensate for the presence of the chemical agent in the analysis of the activity signal(s).

71. The system for performing a neuromodulation and/or an ablation procedure on a target site within a subject in accordance with claim 66 – 70, comprising a stimulating device selected from an electrical stimulator, a thermoregulating device, a heating coil, a thermoelectric device, a Peltier device, a tactile stimulating component, a vibratory stimulating element, or a combination thereof, the stimulating device configured to stimulate the subject at one or more stimulation sites, the algorithm configured to compensate for the stimulation in the analysis.

72. The system for performing a neuromodulation and/or an ablation procedure on a target site within a subject in accordance with claim 66 – 71, wherein one or more of the monitoring devices and/or one or more of the stimulating devices are embedded in the treatment system.

73. The system for performing a neuromodulation and/or an ablation procedure on a target site within a subject in accordance with claim 66 – 72, wherein the treatment system is configured to deliver a stimulating energy or chemical agent to the target site, and/or one or more stimulatory sites within the subject.