

SYSTEM AND METHOD FOR MULTIPLE SITE BIPHASIC STIMULATION TO REVERT VENTRICULAR ARRHYTHMIAS

Field of the Invention

The present invention relates generally to an anti-reentry apparatus and method that can favorably influence the beating of ineffective hearts, especially hearts with pathological conditions that interfere with normal rhythmicity, electrical conduction, and/or contractility by causing ventricular fibrillation. The present invention particularly relates to an anti-reentry apparatus and method that provides simultaneous or progressive biphasic stimulation at multiple sites in one or both ventricles.

Background of the Invention

Heart disease and malfunction is a major killer of men and women in America. A variety of pathologies can affect the beating patterns of a heart, and thereby predispose it to developing ventricular fibrillation. Prior to the occurrence of such a severe and ineffective rhythm, conventional pacemakers can be used to treat, for example, such disorders as sino-atrial (SA) node block, A-V block, and multiple independent sites of contraction in the ventricles (also termed ectopic foci), which, in the extreme, can lead to life threatening ventricular fibrillation. Conventional pacemakers often will control and prevent the recurrence of ectopic foci by preprogrammed stimulation of (usually) the right ventricle via a single electrode. Some pacemakers also employ a second electrode that is dedicated to the left ventricle. In addition, conventional pacemakers utilize a range of circuit logic patterns to counter specific problems that are encountered in the more common pathologies.

However, conventional single ventricular electrode technologies, including the use of a separate single electrode to each ventricle, fail in cases in which ventricular fibrillation has

1 ensued (particularly when the fibrillation is due to multiple random reentry), and single site
2 stimulation does not entrain sufficiently large areas of surrounding tissue to produce the
3 concerted contraction that is necessary for optimal efficiency in pumping blood. In such
4 cases of ventricular fibrillation from multiple random reentry, the patient is put in grave
5 jeopardy for the basic reason that virtually all of the body functions depend on delivery of
6 blood to the tissues in order to supply oxygen and nutrients, and also to carry away metabolic
7 waste products. Failure to correct such a condition, where the rhythm is so far from optimal,
8 results in the patient being in substantial danger of dying in a very short period of time.
9 Though cardioversion/defibrillation may be employed, including that preprogrammed in the
10 control logic for automatic activation in some pacemaker-defibrillators, such protocols
11 typically require large doses of electrical energy to the patient. In addition to producing
12 extreme discomfort and sharp pain, these large doses of electrical energy often also produce
13 cardiac damage. The voltage for standard internal defibrillation/cardioversion is from 150 to
14 800 volts, corresponding to approximately 10-35 joules.

15 Several approaches to these problems have been disclosed. One approach is to
16 stimulate greater portions of ventricular myocardium by utilizing larger electrodes so that
17 greater portions of myocardium are simultaneously stimulated. For example, U.S. Patent No.
18 5,411,547 to Causey, III discloses the use of defibrillation electrode patches for more efficient
19 bipolar cardiac stimulation. In addition, the use of large, plate-like electrodes for
20 defibrillation and cardioversion is well known. However, the use of such larger electrodes
21 suffers from the problem of delivery of large doses of electrical energy that produce great
22 discomfort to the patient and the possibility of tissue damage.

23 Yet another approach is to use multiple individual electrodes appropriately placed
24 about the ventricles, as has been disclosed in the following U.S. patents: 5,649,966 to Noren.

1 et al., 5,391,185 to Kroll, 5,224,475 to Berg, et al., 5,181,511 to Nickolls, et al., and
2 5,111,811 to Smits. Though these patents disclose the use of multiple electrodes, they do not
3 disclose or suggest their use for gradually (yet quickly) entraining the various reentrant foci
4 that can exist in pathological ventricles by stimulating in a progressive pattern that mimics
5 the normal wave of depolarization that occurs in the heart.

6 Thus, a need exists for an anti-reentry apparatus and method that will require the use
7 of less electrical current/voltage than is typically used for defibrillation and cardioversion in
8 order to decrease the likelihood, or at least the severity, of tissue damage. A need also exists
9 for an anti-reentry apparatus and method that will simultaneously stimulate greater portions
10 of ventricular myocardium to increase the probability of ventricular conversion (particularly
11 in the presence of multiple random reentry), but with delivery of lower doses of electrical
12 energy per stimulation, which, consequently, will prolong the life of the apparatus's batteries
13 and decrease myocardial soft tissue damage. A need also exists for such an anti-reentry
14 apparatus and method that not only will produce the vitally needed improvement in cardiac
15 pumping efficiency, but additionally will simultaneously lower the probability of tissue
16 damage, and provide greater comfort for the patient. In addition, a need exists for an anti-
17 reentry apparatus and method that progressively stimulates the ventricles in a manner that
18 mimics the normal cardiac wave of depolarization, thereby providing rapid control and
19 reversion of cardiac rhythm to a normal beating pattern.

20 **Summary of the Invention**

21 In view of the foregoing limitations in the art, it therefore is an object of the present
22 invention to provide an apparatus and method that more efficiently and quickly entrains larger
23 areas of myocardium to promote ventricular conversion, particularly in patients suffering

1 from episodes of multiple random ventricular reentrant foci that produce, or may produce,
2 ventricular fibrillation.

3 It is another object of the present invention to provide an apparatus and method that,
4 while entraining larger areas of myocardium, does so with smaller doses of electrical energy
5 than typically are used in defibrillation and cardioversion.

6 It is yet another object of the present invention to provide an apparatus and method
7 that, while entraining larger areas of myocardium, does so by stimulating in a progressive
8 pattern that mimics the normal wave of depolarization of the heart.

9 It is a further object of the present invention to provide an apparatus and method that,
10 while entraining larger areas of myocardium, does so with less stress on the heart and greater
11 comfort to the patient.

12 It is yet another object of the present invention to provide an apparatus and method
13 that, while entraining larger areas of myocardium, does so with less damage to cardiac tissue.

14 It is yet a further object of the present invention to provide an apparatus and method
15 that, while entraining larger areas of myocardium, also promotes greater myocardial blood
16 pumping efficiency.

17 It is yet a further object of the present invention to provide an apparatus and method
18 that entrains larger areas of myocardium by using multiple electrodes that provide biphasic
19 stimulation.

20 Pacemakers, which utilize low energy stimulation pulses, constitute a separate and
21 distinct art from cardioverters/defibrillators, which utilize stimulation pulses of much larger
22 energy -- even when the electrodes are positioned directly on the heart. Thus, according to
23 conventional practice, more energy is required to entrain the entire heart
24 (cardioversion/defibrillation) than to exogenously employ the traditional pacemaker that

1 typically utilizes the natural cardiac conducting fibers and/or endogenous pacemaker(s) to
2 control the beating of a heart that is only slightly "out of synch" relative to the more
3 dangerous rhythmicity disorders that often result in extensive fibrillation.

4 An intermediate ground is demonstrated by the present invention. By using multiple
5 electrodes and applying biphasic stimulation, one or both ventricles may gradually (yet
6 quickly) be entrained to beat more normally in the face of multiple random reentry, even
7 though the stimulation energy level used is lower than that generally used for
8 cardioversion/defibrillation.

9 Thus, the present invention accomplishes the above objectives by utilizing multiple
10 electrodes that contact multiple ventricular areas 1) for *simultaneous* biphasic stimulation, or
11 2) for *progressive* biphasic stimulation, that is, the mimicking of the physiological patterns
12 of electrical current flows or waves of depolarization in the myocardium. The control circuit
13 logic can activate the multiple site, biphasic ventricular stimulation upon the occurrence of A-
14 V block in a patient known to be susceptible to multiple random ventricular reentrant foci, or
15 upon the direct or indirect sensing of ventricular fibrillation. For example, direct sensing of
16 ventricular fibrillation can be based on data from multiple ventricular sensing electrodes, and
17 indirect sensing can be based on any of various functional parameters, such as arterial blood
18 pressure, size and/or presence of an R wave, rate of the electrogram deflections, or the
19 probability density function (PDF) of the electrogram.

20 The present invention accomplishes the above objectives through the use of multiple
21 site, biphasic ventricular stimulation in one or both ventricles to 1) gradually (yet quickly)
22 entrain and interrupt substantially all of the multiple random reentrant circuits that are
23 present; or, failing that, 2) reduce the number of such reentrant circuits to a level at which
24 much smaller stimuli may be used than in conventional defibrillation/cardioversion to convert

1 the rhythms to more normal ones, and thereby produce coordinated and efficient cardiac
2 function.

3 The first and second phases of stimulation consist of an anodal pulse (first phase)
4 followed by a cathodal pulse (second phase). In a preferred embodiment, the first phase of
5 stimulation is an anodal pulse at maximum subthreshold amplitude and for a long duration in
6 order to precondition the myocardium for subsequent stimulation, and the second phase of
7 stimulation is a cathodal pulse with a short duration and a high amplitude. Additional
8 embodiments of the first phase include, but are not limited to, the use of ramped pulses, a
9 series of short duration square wave pulses, anodal pulses that are less than the maximum
10 subthreshold amplitude, and pulses whose magnitudes decay from an initial subthreshold
11 amplitude to a lower amplitude, where the shape of the decay can be linear or curvilinear. It
12 is to be understood that the use of the phrase "medium energy" stimulation or pulse refers to
13 electrical stimulation or electrical pulses in which the magnitude of the voltage of the
14 electrical stimulation/pulse is lower in magnitude than that used in typical
15 defibrillation/cardioversion.

16 **Brief Description of the Drawings**

17 Fig. 1-A depicts a heart with multiple ventricular electrodes that are introduced via the
18 vena cava.

19 Fig. 1-B depicts a heart with multiple ventricular electrodes that are connected to
20 external surfaces of the ventricles, and include a separate electrode set each for the right and
21 left ventricles.

22 Fig. 2 is a schematic representation of leading anodal biphasic stimulation.

23 Fig. 3 is a schematic representation of leading anodal stimulation of low level and
24 long duration, followed by cathodal stimulation.

1 Fig. 4 is a schematic representation of leading anodal stimulation of ramped low level
2 and long duration, followed by cathodal stimulation.

3 Fig. 5 is a schematic representation of leading anodal stimulation of low level and
4 short duration administered in a series, followed by cathodal stimulation.

5 **Description of the Preferred Embodiments**

6 The apparatus and method of the present invention may be understood with reference
7 to Figs. 1-A, 1-B, and 2 to 5.

8 Referring to Fig. 1-A, a diagram of the heart is shown connected to vena cava 103,
9 and having four chambers: right atrium (RA), left atrium (LA), right ventricle (RV), and left
10 ventricle (LV). Electrode lead 101 is shown feeding into the right ventricle via vena cava
11 103, the right atrium, and tricuspid valve 111. Individual electrodes 102, 104, 106, 108 and
12 110 are connected to electrode lead 101, and contact multiple internal sites of the right
13 ventricle. No set number, or absolute range as to the number, of individual electrodes is
14 required to practice this embodiment of the present invention. A typical range could
15 encompass 2 to 30 electrodes, though numbers greater than 30 are also contemplated. In
16 addition, there is no set placement of these electrodes. In a preferred embodiment, 6 or less
17 electrodes are used, 4 in the left ventricle and 2 in the right ventricle. It is noted that
18 stimulation of the atria through the practice of the present invention also is envisioned.

19 Referring to Fig. 1-B, a similar diagram of the heart is shown in which two sets of
20 multiple electrodes are depicted connected to external ventricular surfaces. Electrode lead
21 201, connected to individual electrodes 202, 204, 206, 208 and 210, is shown with the
22 individual electrodes connected to multiple points on the external surfaces of the right
23 ventricle. Electrode lead 301, connected to individual electrodes 302, 304, 306, 308 and 310,
24 is shown with the individual electrodes connected to multiple points on the external surfaces

1 of the left ventricle.

2 In alternative embodiments, the locations of the individual electrodes in Fig. 1-A
3 (102, 104, 106, 108 and 110), and in Fig. 1-B (202, 204, 206, 208 and 210; and 302, 304,
4 306, 308 and 310) may 1) follow a regular or relatively regular geometric pattern (e.g., an
5 orthogonal or other patterned grid) so as to cover well the ventricular surfaces in appropriate
6 locations; 2) be localized to a particular ventricular area that is known or suspected to be a
7 source of random reentry circuits; 3) be randomly placed about the selected ventricular
8 surfaces; and/or 4) be placed about the ventricular surfaces in a progressive pattern to
9 facilitate mimicking the normal physiological flow of the depolarization wave that leads to
10 the most efficient contraction of the particular ventricle(s).

11 The latter progressive stimulation embodiment, which mimics the normal
12 physiological flow of the normal ventricular depolarization wave, requires that areas closest
13 to (or at) the A-V node be the areas first stimulated during a given beat, and that areas farthest
14 from the A-V node

15 -- following the normal intrinsic conduction paths -- be the last areas to be stimulated.
16 Areas intermediate between these two extremes are appropriately stimulated on a scaled time
17 basis that, again, mimics the normal intrinsic conduction paths that facilitate the most
18 efficient cardiac contraction.

19 This progressive stimulation embodiment requires specific knowledge of the
20 placement of each electrode relative to each other electrode, as well as the placement relative
21 to the electrical conduction pathways in the heart. Thus, it is appropriate to contemplate
22 "classes" of electrodes, in which, for example, electrodes are identified or categorized
23 according to when they are fired. In a simplistic five tier system, e.g., the first tier electrodes
24 are designated as the first to be fired (i.e., the electrodes closest to the A-V node), followed

1 successively (and temporally progressively according to the normal conducting paths) by the
2 second, third, fourth, and fifth tier electrodes, where the fifth tier electrodes would be the last
3 to be fired, and whose locations on the ventricle(s) would correspond to the last areas to be
4 depolarized in the course of a normal ventricular contraction/beat. An even simpler (i.e., two,
5 three or four) tiered system may be used, or one more complex (i.e., one with greater than 5
6 tiers, or with any other basis of electrode placement, such as a honeycomb-like array in a
7 particular area with a known or suspected pathology as to rhythmicity, reentry, conduction,
8 contractility, etc.). Furthermore, multiple electrodes within a given tier may be numbered or
9 otherwise distinctly identified so that the practitioner may test and use electrodes with respect
10 to known locations in the heart, for example, to anticipate and/or bypass an area of electrical
11 blockage. This type of embodiment would require the use of multiple, small electrodes
12 pulsed in a physiologic sequential fashion. In application to atria, electrodes are
13 progressively placed from close to the SA node (first to be fired) to close to the AV node (last
14 to be fired), mimicking the normal intrinsic conduction paths.

15 Bypassing an area of electrical blockage is also anticipated by the present invention,
16 and can be effected by first identifying such areas, for example, by determining myocardial
17 resistance values between electrodes. Electrical pulses then are routed to those myocardial
18 areas with appropriately low resistances, following as closely as possible the lines of
19 conduction of the normal intrinsic conduction paths. Communication of, and control of,
20 measurements of resistance between electrodes, as well as developing a bypass protocol for a
21 particular patient, can be effected by an external computer. The external computer can
22 communicate with the pacemaker by any convenient method, for example, radiotelemetry,
23 direct coupling (as by connecting to an external wire from the pacemaker to the surface of the
24 skin of the patient), etc.

1 Figs. 2 through 5 depict a range of biphasic stimulation protocols. These protocols
2 have been disclosed in United States Patent Application No. 08/699,552 to Mower, which is
3 herein incorporated by reference in its entirety.

4 Fig. 2 depicts biphasic electrical stimulation in which a first stimulation phase
5 comprising anodal stimulus 202 is administered with amplitude 204 and duration 206. The
6 first stimulation phase is followed immediately by a second stimulation phase comprising
7 cathodal stimulus 208, which is of equal intensity and duration to those of anodal stimulus
8 202.

9 Fig. 3 depicts biphasic electrical stimulation wherein a first stimulation phase
10 comprising low level, long duration anodal stimulation 302 having amplitude 304 and
11 duration 306 is administered. This first stimulation phase is immediately followed by a
12 second stimulation phase comprising cathodal stimulation 308 of conventional intensity and
13 duration. In an alternative embodiment of the invention, anodal stimulation 302 is at
14 maximum subthreshold amplitude. In yet another alternative embodiment of the invention,
15 anodal stimulation 302 is less than three volts. In another alternative embodiment of the
16 invention, anodal stimulation 302 is a duration of approximately two to eight milliseconds.
17 In yet another alternative embodiment of the invention, cathodal stimulation 308 is of a short
18 duration. In another alternative embodiment of the invention, cathodal stimulation 308 is
19 approximately 0.3 to 1.5 milliseconds. In yet another alternative embodiment of the
20 invention, cathodal stimulation 308 is of a high amplitude. In another alternative
21 embodiment of the invention, cathodal stimulation 308 is in the approximate range of three to
22 twenty volts. In yet another alternative embodiment of the present invention, cathodal
23 stimulation 308 is of a duration less than 0.3 milliseconds and at a voltage greater than twenty

1 volts. In another alternative embodiment, anodal stimulation 302 is administered over 200
2 milliseconds post heart beat. In the manner disclosed by these embodiments, as well as those
3 alterations and modifications which may become obvious upon the reading of this
4 specification, a maximum membrane potential without activation is achieved in the first
5 phase of stimulation.

6 Fig. 4 depicts biphasic electrical stimulation wherein a first stimulation phase
7 comprising anodal stimulation 402 is administered over period 404 with rising intensity level
8 406. The ramp of rising intensity level 406 may be linear or non-linear, and the slope may
9 vary. This anodal stimulation is immediately followed by a second stimulation phase
10 comprising cathodal stimulation 408 of conventional intensity and duration. In an alternative
11 embodiment of the invention, anodal stimulation 402 rises to a maximum subthreshold
12 amplitude. In yet another alternative embodiment of the invention, anodal stimulation 402
13 rises to a maximum amplitude that is less than three volts. In another alternative embodiment
14 of the invention, anodal stimulation 402 is a duration of approximately two to eight
15 milliseconds. In yet another alternative embodiment of the invention, cathodal stimulation
16 408 is of a short duration. In another alternative embodiment of the invention, cathodal
17 stimulation 408 is approximately 0.3 to 1.5 milliseconds. In yet another alternative
18 embodiment of the invention, cathodal stimulation 408 is of a high amplitude. In another
19 alternative embodiment of the invention, cathodal stimulation 408 is in the approximate range
20 of three to twenty volts. In yet another alternative embodiment of the present invention,
21 cathodal stimulation 408 is of a duration less than 0.3 milliseconds and at a voltage greater
22 than twenty volts. In another alternative embodiment, anodal stimulation 402 is administered
23 over 200 milliseconds post heart beat. In the manner disclosed by these embodiments as well

1 as those alterations and modifications which may become obvious upon the reading of this
2 specification, a maximum membrane potential without activation is achieved in the first
3 phase of stimulation.

4 Fig. 5 depicts biphasic electrical stimulation wherein a first stimulation phase
5 comprising series 502 of anodal pulses is administered at amplitude 504. In one embodiment
6 rest period 506 is of equal duration to stimulation period 508 and is administered at baseline
7 amplitude. In an alternative embodiment, rest period 506 is of a differing duration than
8 stimulation period 508 and is administered at baseline amplitude. Rest period 506 occurs
9 after each stimulation period 508 with the exception that a second stimulation phase
10 comprising cathodal stimulation 510 of conventional intensity and duration immediately
11 follows the completion of series 502. In an alternative embodiment of the invention, the total
12 charge transferred through series 502 of anodal stimulation is at the maximum subthreshold
13 level. In yet another alternative embodiment of the invention, the first stimulation pulse of
14 series 502 is administered over 200 milliseconds post heart beat. In another alternative
15 embodiment of the invention, cathodal stimulation 510 is of a short duration. In yet another
16 alternative embodiment of the invention, cathodal stimulation 510 is approximately 0.3 to 1.5
17 milliseconds. In another alternative embodiment of the invention, cathodal stimulation 510 is
18 of a high amplitude. In yet another alternative embodiment of the invention, cathodal
19 stimulation 510 is in the approximate range of three to twenty volts. In another alternative
20 embodiment of the invention, cathodal stimulation 510 is of a duration less than 0.3
21 milliseconds and at a voltage greater than twenty volts. The individual pulses of the series of
22 pulses may be square waves, or they may be of any other shape, for example, pulses which
23 decay linearly or curvilinearly from an initial subthreshold amplitude, to a lower amplitude.

1 In the preferred biphasic stimulation protocol practiced by the present invention, the
2 magnitude of the anodal phase does not exceed the maximum subthreshold amplitude. The
3 anodal phase serves to precondition the stimulated myocardium, thereby lowering the
4 excitation threshold such that a cathodal stimulation of lesser intensity than normal will
5 produce depolarization leading to contraction.

6 The values of duration and amplitude will depend on factors such as the
7 placement/position of the particular electrode (including, e.g., whether the electrode is in
8 purely muscle tissue versus in specialized conducting or pacemaking tissue), whether
9 damaged/scarred tissue is in close vicinity to the electrode, depth of the electrode within the
10 tissue, local tissue resistance, presence or absence of any of a large range of local pathologies,
11 etc. Nonetheless, typical anodal phase durations often fall within the range from about two
12 milliseconds to about eight milliseconds, whereas typical cathodal durations often fall within
13 the range from about 0.3 millisecond to about 1.5 millisecond. Typical anodal phase
14 amplitudes (most commonly at the maximum subthreshold amplitude) often fall within the
15 range from about 0.5 volt to 3.5 volts, compared to typical cathodal phase amplitudes from
16 about 3 volts to about 20 volts.

17 The present invention also permits the physician to readily test ranges of stimulation
18 and other parameters (voltage, duration, shape of voltage versus time pulses, etc.) once the
19 anti-reentry system is in place in the patient. Thus, the ability to engage in trial and error
20 testing of pulsing parameters permits the physician not only to determine such a parameter as
21 maximum subthreshold amplitude, but also to optimize other stimulation parameters to fit a
22 given patient's condition, location of electrodes, etc. Furthermore, the physician may so
23 determine optimal parameters for each individual electrode in a set of multiple electrodes.

24 Such a system of testing could be related to defibrillation threshold testing, wherein

1 ventricular fibrillation is deliberately provoked and various levels of defibrillatory shocks are
2 given to determine the amount of energy needed. In the present application, testing is done
3 with the various patterns of pacing so as to find the one with the lowest requirement for
4 countershock energy.

5 Based on the examples provided herein, the skilled practitioner in the art will readily
6 appreciate that generalization of the teachings expands the scope of the present invention to
7 include stimulation time and voltage ranges to beyond those mentioned herein, as well as to
8 beyond the numbers of individual electrodes employed, and other parameters subject to
9 simple and quick experimentation in a specific situation not specifically addressed in the
10 verbiage presented on the practice of the present invention.

11 Having thus described the basic concept of the invention, it will be readily apparent to
12 those skilled in the art that the foregoing detailed disclosure is intended to be presented by
13 way of example only, and is not limiting. Various alterations, improvements and
14 modifications will occur and are intended to those skilled in the art, but are not expressly
15 stated herein. These modifications, alterations and improvements are intended to be
16 suggested hereby, and within the scope of the invention. Accordingly, the invention is
17 limited only by the following claims and equivalents thereto.

1 **Claims**

2 I claim:

3 1. A method for myocardial stimulation, comprising:

4 (a) at least two electrodes administering a first stimulation phase having a first
5 phase polarity, a first phase amplitude, a first phase shape, and a first phase duration for
6 preconditioning the myocardium to accept subsequent stimulation;7 (b) the at least two electrodes administering a second stimulation phase having a
8 second phase polarity, a second phase, a second phase shape, and a second phase duration;

9 (c) sensing for fibrillation; and

10 d) repeating steps a), b), and c) if fibrillation is sensed.

11 2. The method for myocardial stimulation according to claim 1, where the first
12 phase polarity is positive, and the second phase polarity is negative.13 3. The method for myocardial stimulation according to claim 2, where the first
14 phase amplitude is at a maximum subthreshold amplitude.15 4. The method for myocardial stimulation according to claim 3, where the
16 maximum subthreshold amplitude is about 0.5 volt to about 3.5 volts.17 5. The method for myocardial stimulation according to claim 1, where the first
18 phase shape is ramped from a baseline value to a second value.19 6. The method for myocardial stimulation according to claim 1, where the first
20 phase duration is about one millisecond to about nine milliseconds.21 7. The method for myocardial stimulation according to claim 1, where the second
22 phase amplitude is greater than the first phase amplitude.23 8. The method for myocardial stimulation according to claim 7, where the second
24 phase amplitude is about two volts to about twenty volts.

1 9. The method for myocardial stimulation according to claim 1, where the second
2 phase duration is about 0.2 millisecond to about 1.5 milliseconds.

3 10. The method for myocardial stimulation according to claim 1, where the first
4 stimulation phase further comprises a series of stimulating pulses of a predetermined
5 amplitude and duration, and a series of rest periods.

6 11. The method for myocardial stimulation according to claim 10, where applying
7 the first stimulation phase further comprises applying a rest period after at least one
8 stimulating pulse.

9 12. The method for myocardial stimulation according to claim 10, where the
10 predetermined duration is about 0.2 millisecond to about 1.5 milliseconds.

11 13. The method for myocardial stimulation according to claim 10, where the rest
12 period is about 0.2 millisecond to about 1.2 milliseconds.

13 14. The method for myocardial stimulation according to claim 1, where the first
14 phase shape is selected from the group consisting of square wave pulse, ramped pulse, and
15 series of short duration square wave pulses.

16 15. The method for myocardial stimulation according to claim 1, where at least
17 one of the electrodes is applied to an inner ventricular wall via the vena cava.

18 16. The method for myocardial stimulation according to claim 1, where at least
19 one of the electrodes is applied to an exterior ventricular wall.

20 17. The method for myocardial stimulation according to claim 5, where the second
21 value is not more than a maximum subthreshold amplitude.

22 18. The method for myocardial stimulation according to claim 1, where the
23 stimulation is applied to the electrodes progressively in a manner that mimics the normal flow
24 of electrical depolarization in a heart.



- 1 19. The method for myocardial stimulation according to claim 18, where:
2 the electrodes closest to the A-V node are the first to be fired;
3 the electrodes farthest from the A-V node, following the normal intrinsic conduction
4 paths, are the last to be fired; and
5 the electrodes intermediate between the electrodes closest to the A-V node and the
6 electrodes farthest from the A-V node are fired at an intermediate time that is
7 proportional to their intermediate position, following the normal intrinsic
8 conduction paths.
- 9 20. The method for myocardial stimulation according to claim 19, where the
10 electrodes are ordered by class according to their distance from the A-V node, following the
11 normal intrinsic conduction paths.
- 12 21. The method for myocardial stimulation according to claim 20, where the
13 number of classes is between two and about thirty.
- 14 22. The method for myocardial stimulation according to claim 1, where the second
15 phase amplitude is larger in absolute value than the first phase amplitude.
- 16 23. A method for reverting ventricular arrhythmias comprising:
17 at least two electrodes administering a first stimulation phase having a positive
18 polarity, a subthreshold amplitude, a duration of about one millisecond to about nine
19 milliseconds, and a shape, wherein the shape is selected from the group consisting of square
20 wave pulse, ramped pulse, and series of short duration square wave pulses; and
21 the at least two electrodes administering a second stimulation phase having a negative
22 polarity, an amplitude of about two volts to about twenty volts, and a duration of about 0.2
23 millisecond to about 1.5 milliseconds.

