

ISOLATED DISCOGENIC CELLS, METHODS OF USING, AND METHODS OF PREPARING SAME FROM MAMMALIAN TISSUE

CROSS REFERENCE TO RELATED APPLICATIONS

[1] This application claims benefit of priority pursuant to 35 U.S.C. § 119(e) of U.S.
5 provisional patent application no. 61/794,691 filed March 15, 2013, which is hereby incorporated by reference in its entirety.

FIELD OF INVENTION

[2] This disclosure relates to the isolation and methods of using intervertebral disc cells in the treatment of degenerative disc disease.

BACKGROUND OF THE INVENTION

[3] The mammalian spine serves two basic functions: (1) load bearing support for the upper body and (2) protection of the nerves comprising the spinal column. The spine is made up of interlocking vertebrae, separated by intervertebral discs. These discs act as shock absorbers and allow the spine to bend, compress, and twist. Spinal discs have two
15 basic parts: an outer fibrous structure (the annulus fibrosus), and a gel-like inner structure (the nucleus pulposus). A healthy nucleus pulposus in a young mammal is about 80% water. Over time, the nucleus pulposus loses its high water content and thus its ability to absorb shock. Additionally, intervertebral discs can be damaged through dehydration, disease, over-use, injury or trauma, resulting in rupture, bulging, herniation, etc. The
20 intervertebral disc is also susceptible to other diseases such as degenerative disc disease.

[4] In a healthy intervertebral disc, cells represent a small fraction of the total volume. Much of the disc volume is extracellular matrix (ECM; collagen and proteoglycans, which aid in retaining the large volume of water) produced by the cells, and much of the difference between the nucleus pulposus and annulus fibrosus is the
25 water content and the makeup of the ECM.

[5] Back pain resulting from degenerative disc disease is a major cause of morbidity, disability, and lost productivity. Back pain is frequently cited as limiting the activity of people under the age of 45 and a reason for physician visits, hospitalization, and surgical procedures. Chronic back conditions are reported by between 15%-45% of the population

each year, and in 70% to 85% of the population at some time in their lives. The financial impact in terms of health care dollars and lost work hours to society is high. More than one million spine surgery procedures are performed annually in United States.

Furthermore, the lumbar fusion segment of the spine surgery market is estimated at well
5 over \$1 billion in annual revenue.

[6] Despite continued improvements in both operative and non-operative treatment options for subjects suffering from back pain and spine disease, there is no solution to eliminate or consistently improve this condition. Current treatments for spine disease include steroid injections, physical therapy, discectomy and spinal fusions. Spinal
10 prostheses have been introduced by several companies. However, these prostheses differ greatly in their design, for example in the bearing surface, fixation to bone, number of articulations, material, constraint, mobility of rotation, and have seen little success in practice.

[7] Nucleus arthroplasty or nucleus replacement is also an option for treating
15 degenerative disc disease. In some cases, these devices consist of a hydrogel core center encased in a polyethylene sleeve that allows the device to shrink and swell during normal loading and unloading. This may partially help restore disc space height and aid in mimicking healthy human disc.

[8] Disc arthroplasty is not without complications. The most common complications
20 include adjacent level spinal disease, subsidence, and facet joint arthrosis. Furthermore, recent studies from clinical trials have demonstrated incidences of infection, vertebral body fracture, implant malposition, subsidence, mechanical failure, and paravertebral heterotopic ossification. More serious complications, including anterior dislocation of the implant, have been reported. Also, the issue of wear particles from the total disc
25 arthroplasty and the potential effects on the spinal cord are still not known.

[9] What is needed is a biologic treatment for intervertebral discs that can aid in repairing or replacing a subject's intervertebral disc.

SUMMARY OF THE INVENTION

[10] Disclosed herein are isolated discogenic cell populations, methods of using, and methods of preparing. The discogenic cell populations are used in the repair, regeneration, and replacement of damaged, injured, or diseased intervertebral discs. In some embodiments the discogenic cell population is derived from mammalian disc tissue and grown in vitro under anchorage independent conditions. In some embodiments the cells the culture comprises a media comprising one or more additives selected from the group consisting of EGF, bFGF, serum, fibroblast conditioned media, and a viscous non-reactive substance. In some cases, the cells are grown in a receptacle comprising a low adhesion coating. The disclosed discogenic cell populations may be used for autologous and/or non-autologous treatment of intervertebral disc in a subject in need thereof.

[11] The disclosed discogenic cells can be used to produce an artificial disc replacement in vitro or in vivo using non-resorbable material or resorbable material. The material may create an artificial annulus that serves to contain a discogenic cell population, which may or may not be combined with at least one of the following – scaffold material, matrix material, carrier material, growth factor(s), and/or other biologically active agents. The artificial outer annulus may incorporate attachment means so that it can be fixed to one or more vertebral bodies. For example, the artificial annulus may incorporate through-holes, cuffs, tabs, loops, or washers to allow for screw fixation to one or more vertebral bodies. The artificial disc can be surgically implanted in a subject to completely replace a spinal disc.

[12] Also disclosed are various methods for obtaining and preparing discogenic cells from autologous and non-autologous donors. A method of deriving a discogenic cell population is disclosed wherein the method comprises isolating one or more cells from tissue, passaging the one or more cells in an anchorage dependent culture media, and transferring the one or more cells to an anchorage independent culture media. Another method is disclosed using a discogenic cell to treat at least one disc in a subject in need thereof comprising, administering a therapeutic amount of a discogenic cell population to the subject, and thereby treating the subject. In various embodiments, the tissue is mammalian disc tissue, for example from a donated organ or spine. In some aspects, the

disclosed method includes wherein the cell population is passaged at least one time in the anchorage independent culture, and the population of cells produces extracellular matrix. In some aspects the population of cells produces one or more cell surface markers selected from the group comprising CD24, CD34, CD44, CD73, CD90, CD105, CD166, 5 Stro-1, HIF1, nestin, CK8, and HLA proteins, wherein the percentage of the cells in the population expressing a cell surface marker(s) is greater than 70% or less than 40%. In some aspects of the disclosed method the population of cells expresses one or more gene or gene products selected from the group comprising GAPDH, SDHA, HPRT1, B2M, Sox9, Aggrecan, Col1, Col2, nestin, CK8, Sox1, CD44, ALPI, PPARG, ADAMTS, 10 MMP, FMOD, IL.

[13] In another aspect, a population of Discogenic cells is described, wherein greater than 40 % of the cells produce cell surface markers CD44, CD73, CD90, HLA-A, B, or C, CD24, CD105, CD166, or combinations thereof, and less than about 20% of the cell population produces CD34, HLA-DR or –DQ, or STRO-1. In some embodiments, about 15 80-100 % of the population produces CD73, CD90, CD44, HLA ABC, or combinations thereof. In some embodiments, about 20-75 % of the population produces CD105, CD166, CD24, or combinations thereof.

[14] In accordance with one aspect, the invention provides methods for treating a subject having a disease of or damage to at least one intervertebral disc caused by 20 damage to the intervertebral disc induced by age, trauma, toxin exposure, drug exposure, radiation exposure, oxidation, immune-complex deposition, or transplant rejection.

[15] In accordance with another aspect, the invention provides kits for treating a subject having a disease of or damage to at least one intervertebral disc, comprising a pharmaceutically acceptable carrier, discogenic cells in an amount effective to treat the 25 disease or injury, spinal column tissue and wherein the cells are capable of expansion in culture and have the potential to differentiate. In some embodiments, the kit includes at least one agent.

BRIEF DESCRIPTION OF THE DRAWINGS

[16] The file of this patent contains at least one drawing/photograph executed in color. Copies of this patent with color drawing(s)/photograph(s) will be provided by the Office upon request and payment of the necessary fee.

5 [17] The subject matter regarded as the invention is particularly pointed out and distinctly claimed in the concluding portion of the specification. The invention, however, both as to organization and method of operation, together with objects, features, and advantages thereof, may best be understood by reference to the following detailed description when read with the accompanying drawings in which:

10 [18] FIG. 1 shows expression profile of various surface markers known to identify stem cells and chondrogenic cells. Eight cell types are explored: fibroblasts, chondrocytes, mesenchymal stem cells (MSCs), discogenic cells grown in monolayer (attachment dependent), and discogenic cells grown in suspension (attachment independent).

15 [19] FIG. 2 shows cell morphology of discogenic cells grown in suspension culture.

[20] FIG. 3 shows the chondrogenic potential of discogenic cells compared to MSCs.

[21] FIG. 4 shows the adipogenic and osteogenic potential of discogenic cells.

[22] FIG. 5 shows the viability of cells (green, or bright in B/W – alive, red or dark in B/W – dead) after being combined with a viscous 1% hyaluronic acid scaffold, and after
20 being extruded through a 27-gauge 1.5 inch surgical needle.

[23] FIG. 6 shows potential devices and formulations for the therapeutic use of discogenic cells to treat various forms of disc disease. Treatment can be by injection (top) or implantation (bottom). Cells and carrier/scaffold can be together, or combined immediately prior to use.

25 [24] FIG. 7 shows the in vivo efficacy to repair degenerated intervertebral discs (animal model) of discogenic cells formulated with a viscous scaffold carrier in rabbits.

[25] FIG. 8 is a schematic flow chart showing preparation and implantation of discogenic cells. Step a shows a diagram of an intervertebral disc. Step b shows fresh human intervertebral disc isolated from a spine. Step c shows nucleus pulposus cells after dissected and enzymatic digestion. Cells that adhere are expanded in the presence of EGF and FGF-2. Scale bar = 50 μ m. Step d is a micrograph of cells transitioned to a contact-inhibited culture environment containing methylcellulose, wherein clusters and spheres develop over about 2 weeks. Scale bar = 200 μ m. Step e is a syringe for injecting cells that have been washed free of methylcellulose-containing media and combined with uncross-linked hyaluronic acid scaffold. Step f shows the syringe being used to inject the mixture of cells and scaffold is into degenerated rabbit intervertebral discs, afterward, the safety and efficacy was assessed for about 1 month.

[26] FIGs. 9A-F shows discogenic cells assessed for Aggrecan and Collagen production. FIG. 9A is a phase image of hematoxylin and eosin staining. FIG. 9B is a phase image of alcian blue counterstained with nuclear fast red. Note presence of matrix around single cell (left). Scale bar = 10 μ m. FIG. 9C is a phase image of picrosirius red staining. Fig. 9D is a confocal image including actin (red) and cell nuclei (blue). Fig. 9E is a confocal image for aggrecan, collagen and actin (no nuclei) with additional magnification, wherein the black arrow indicates intracellular aggrecan, and the white arrow indicates extracellular aggrecan. Fig. 9F is a bar graph showing RT-PCR analysis of matrix molecules (aggrecan and collagen 2A) over time in culture, at day 14 harvest and after chondrogenic differentiation. Fold expression was calculated by normalized crossing threshold to housekeeping gene HRPT and baseline gene expression at day 0.

[27] FIGs. 10A-C depicts flow cytometry studies on discogenic cells. FIG. 10A is a forward and side scatter plot, showing gating applied to all subsequent analyses that includes 89% of the cell population. FIG. 10B is a bar graph showing expression levels (compared to isotype control) of discogenic cells from 5 distinct human donors. FIG. 10C shows representative histograms of surface marker expression.

[28] FIGs. 11A-E shows multipotency of discogenic cells. FIG. 11A shows osteogenic differentiation shown with alizarin red staining. FIG. 11B shows adipogenic differentiation shown with oil red O staining. FIG. 11C shows chondrogenic

differentiation after micromass formation (alcian blue and nuclear fast red). Scale bars = 100 μ m. FIG. 11D is a bar graph showing quantitative evaluation of soluble (media) and insoluble (micromass) GAG production after chondrogenic differentiation for articular chondrocytes (AC), adult fibroblasts (FB), bone marrow derived MSCs, and discogenic cells (DCs). FIG. 11E is a bar graph showing total GAG production normalized to DNA content for various cell types. Line indicates significant difference ($p < 0.01$, 1-way ANOVA with Bonferroni's post-hoc test).

[29] FIGs. 12A- show safety and efficacy assessments of treatment in a rabbit model of degenerative disc disease. FIG. 12A is a representative x-ray, which were taken every 2 weeks and used to calculate *disc height index* (DHI) based on 18 boney landmarks, as shown. FIG. 12B shows that rabbit body weight (therapy injected on day 14) for the duration of the study remained within normal ranges. FIG. 12C is a graph DHI over the 6 week treatment showing that treatment improved DHI compared to control conditions after 4 and 6 weeks, with low dose performing better than high dose. No improvement as noted in scaffold or injured control, whereas uninjured control disc height remained unchanged from week 0.

[30] FIGs. 13A-B show histological evaluation of treatment after 6 week Pilot Study. FIG. 13A is a cross-section of a healthy, injured and treated disc (Hematoxylin and eosin stain, scale bar = 2mm). FIG. 13B shows histology of various regions of the IVD after treatment, including bone marrow, annulus fibrosus (AF), cartilage endplate (CEP), and nucleus pulposus (NP); stained with hematoxylin and eosin (H&E) or alcian blue. (Scale bar = 100 μ m).

[31] FIGs. 14A-C show pilot safety and efficacy of treatment in pigs (12 weeks) FIG. 14A is a graph showing that doses of IDCT improved DHI compared to injured control ($p < 0.05$) in a sustained manner at 12 weeks; no improvement noted in scaffold or injured control, whereas uninjured control disc height remained unchanged from week 0. FIG. 14B is a fluoroscopic view of pig spine during treatment procedure (note collapsed disc space in injured discs). FIG. 14C is a histological evaluation of IDCT-treated discs, including nucleus pulposus (NP), cartilage endplate (CEP) and annulus fibrosus (AF) stained with Hematoxylin and eosin (H&E) and alcian blue.

DETAILED DESCRIPTION OF THE INVENTION

[32] The presently described discogenic cells are cells that are derived from disc tissue and may be used for the treatment and/or repair of intervertebral discs. In some cases, discogenic cells may be processed in vitro to provide for discogenic cells that are more
5 potent than other cells in repairing, replacing, or augmenting existing or damaged nucleus pulposus tissue. In various embodiments discogenic cells produce extracellular matrix. In some embodiments, discogenic cells produce proteoglycans. In other embodiments, discogenic cells produce collagens. In other embodiments, discogenic cells implanted adjacent native cells may aid in stimulating native cells through chemical, mechanical or
10 other forces. For example, discogenic cells may excrete growth factors, cytokines or other proteins.

[33] “Discogenic,” as used herein, refers to the ability to produce disc tissue in vivo. In some embodiments, discogenic cells are able to regenerate disc tissue that is diseased or damaged and/or has lost one or more properties of disc tissue in vivo. In some cases,
15 “discogenic” cells may produce disc tissue in vitro, for example where discogenic cells may be used to generate an artificial disc for implantation.

[34] “Maintained” as used herein where referring to cells grown in-vitro, is meant to encompass cells grown in culture for greater than 24 hours. In some cases, maintained cells are cells that have divided in cell culture.

20 [35] “Micromass” is formed by concentrating about 10,000 to 1,000,000 cells in a conical vessel that inhibits attachment, which results in cells forming a least one singular mass. The micromass may additionally contain extracellular matrix. It is based upon an assay to determine chondrogenic potential, and is also known as a pellet.

[36] “About” as used herein when referring to a measurable value such as an amount,
25 a temporal duration, and the like, is meant to encompass variations of less than about $\pm 20\%$. In some cases about may refer to variations of 10% or less, or $\pm 5\%$ or less. In some cases about may refer to variation of $\pm 1\%$ - $\pm 0.1\%$.

[37] “Derived” may be used to indicate that a cell has been obtained or isolated from its natural or previous biological state or situs and maintained, grown or expanded in

culture, or immortalized, or otherwise manipulated in vitro. For example, in some present embodiments, the disclosed discogenic population may be derived from disc tissue or cartilaginous tissue, and in some embodiments discspheres may be derived from a discogenic cell population.

5 [38] If a cell or molecule is “isolated” it has been removed from, or altered in relation to its natural state through human intervention.

[39] The term “express,” “expressed,” or “expression” refers to the biosynthesis of a gene product from a nucleic acid molecule or gene, for example, the biosynthesis of a polypeptide. A cell surface marker may be expressed on the surface of a cell if more or
10 less of that cell marker is present on the surface of the cell after some event, for example growth in vitro.

[40] “Damage” refers to any harm, injury, degeneration, or trauma to the intervertebral disc whether due to age, trauma, or disease.

[41] A “disease” is any deviation from, or impairment in the health, condition, or
15 functioning of a cell, tissue, organ, system, or organism on the whole, as measured by any means suitable in the art.

[42] “Treat,” treating” or “treatment” refer to any attenuation or amelioration of disease, damage, or condition, including any objective or subjective parameter such as abatement, remission, diminishing of symptoms or making the disease, damage, or
20 condition more tolerable to a subject, for example through pain reduction), slowing in the rate of degeneration or decline, making the final point of degeneration less debilitating, improving a subject's physical or mental well-being, or prolonging the length of survival. The treatment or amelioration of symptoms can be based on objective and/or subjective parameters; including the results of a physical examination, radiographic examination,
25 neurological examination, and/or psychiatric evaluations.

[43] “Effective amount” or “therapeutically effective amount” are used interchangeably herein, and refer to an amount of a compound, material, or composition, as described herein effective to achieve a particular biological result such as, but not limited to, biological results disclosed, described, or exemplified herein. Such results may

include, but are not limited to, the treatment of intervertebral disc disease or damage in a subject, as determined by any means suitable in the art.

[44] “Pharmaceutically acceptable” refers to those properties and/or substances which are acceptable to the subject from a pharmacological/toxicological point of view and to
5 the manufacturing pharmaceutical chemist from a physical/chemical point of view regarding composition, formulation, stability, subject acceptance and bioavailability.

[45] “Pharmaceutically acceptable carrier” refers to a medium that does not interfere with the effectiveness of the biological activity of the active ingredient(s) and is not toxic to the host to which it is administered. One embodiment of a pharmaceutically acceptable
10 carrier is hyaluronic acid.

[46] “Discosphere” is described in patent US 8,227,246 B2 and PCT application number PCT/US2012/025066, which are hereby incorporated by reference in their entireties.

Obtaining discogenic cells from disc tissue

15 [47] The presently described discogenic cells may be obtained from disc tissue. Disc tissue may include both nucleus pulposus tissue, transition zone tissue, and annulus fibrosis tissue. In some cases, discogenic cells may be obtained from cartilaginous endplate of the intervertebral disc. In other cases, discogenic cells may be obtained from other cartilaginous tissues in the body.

20 [48] In various embodiments the disc tissue may be obtained from living or deceased donors. The donor may be a mammal, for example a human. In some cases, the donor is a tissue donor and may be genetically unrelated to the recipient. The donor may be of any age, including neonatal, young, adult and senior.

[49] In various embodiments the disc tissue may be healthy disc tissue or may be
25 diseased or injured disc tissue. Injured or diseased disc tissue that may be used with the currently disclosed discogenic cells and methods include, for example, degenerated tissue, herniated tissue, tissue removed from painful discs, tissue removed from deceased donors.

[50] In various embodiments, the tissue is used directly to obtain cells. In other embodiments, the tissue is frozen prior to use, and used at a later date, for example by cryopreservation or vitrification. In other embodiments, the tissue is kept at 4°C in specialized media until the cells are extracted. Tissue can be maintained in media,
5 containing sugars, cryoprotectants, stabilizers, serum, etc.

Culturing discogenic cells

[51] Discogenic cells may be grown in mammalian cell culture. In most cases, the cell culture may allow anchorage to a substratum, or alternatively prevent anchorage to a substratum. In some cases, the cell culture may include a medium. Cell culture media
10 may be any media suitable for the growth of mammalian cells in culture, for example DMEM (Dulbecco's Modified Eagle's Medium), MEM (Modified Eagle's Medium), RPMI, RPMI 1640, etc. In some cases, the media may additionally include further additives such as a nutrient media, for example Ham's F12 (F12). In some case, the cell culture media may or may not include further additives.

[52] In various embodiments, serum may or may not be added to the culture media. Serum may refer to animal serum derived from a mammal, for example, cow, chicken, goat, equine, human, sheep, pig, rabbit, etc. In some cases, the serum may be derived from adult, newborn, or fetal animals, for example fetal bovine serum may be obtained from a cow or calf fetus. In some cases, a serum additive such as animal platelet lysates
20 (for example, human platelet lysates), serum-converted platelet lysate, animal serum albumin (bovine serum albumin), or conditioned media from another cell culture (for example neonatal foreskin fibroblast conditioned media) may be added with, or in the place of serum.

[53] In some cases, the serum or serum additive concentration in the culture media
25 may be greater than 0%, 1%, 2%, 3%, 4%, 5%, 6%, 7%, 8%, 9%, 10%, 11%, 12%, 13%, 14%, 15%, 16%, 17%, 18%, 19%, 20%, 25%, and 30%, and/or less than about 35%, 30%, 25%, 20 %, 19%, 18%, 17%, 16%, 15%, 14%, 13%, 12%, 11%, 10%, 9%, 8%, 7%, 6%, 5%, 4%, 3%, 2%, and 1% by volume. In some cases the serum concentration in the media may be 0%. In some embodiments the serum concentration is from 0-17%, 0-5%,
30 5-17% or 0-2.5%.

[54] In some cases, additional supplements may or may not be added to the culture media. In some cases, the supplement may be a hormone or growth factor. In some cases the hormone or growth factor may be Adrenomedullin (AM), Angiopoietin (Ang), Autocrine motility factor, Bone morphogenetic proteins (BMPs), Brain-derived neurotrophic factor (BDNF), Epidermal growth factor (EGF), Erythropoietin (EPO), basic Fibroblast growth factor (bFGF, FGF-2, or FGF- β), Glial cell line-derived neurotrophic factor (GDNF), Granulocyte colony-stimulating factor (G-CSF), Granulocyte macrophage colony-stimulating factor (GM-CSF), Growth differentiation factor-9 (GDF9), Hepatocyte growth factor (HGF), Hepatoma-derived growth factor (HDGF), Insulin-like growth factor (IGF), Migration-stimulating factor, Myostatin (GDF-8), Nerve growth factor (NGF) and other neurotrophins, Platelet-derived growth factor (PDGF), Thrombopoietin (TPO), Transforming growth factor alpha(TGF- α), Transforming growth factor beta (TGF- β), Tumor_necrosis_factor-alpha(TNF- α), Vascular endothelial growth factor (VEGF), placental growth factor (PlGF), IL-1, IL-2, IL-3, IL-4, IL-5, IL-6, IL-7, insulin, progesterone, putrescine, transferrin, sodium selenite. In many cases, the growth factors may be EGF and bFGF. In many cases the concentration of supplement in the cell culture media may be greater than about 1 ng/ml, 2 ng/ml, 3 ng/ml, 4 ng/ml, 5 ng/ml, 6 ng/ml, 7 ng/ml, ng/ml, 9 ng/ml, 10 ng/ml, 11 ng/ml, 12 ng/ml, 13 ng/ml, 14 ng/ml, 15 ng/ml, 20 ng/ml, 25 ng/ml, 30 ng/ml, 35 ng/ml, 40 ng/ml, 50 ng/ml, 100 ng/ml, 200 ng/ml, 300 ng/ml, 400 ng/ml, 500 ng/ml, 600 ng/ml, 700 ng/ml, 800 ng/ml, 900 ng/ml, 1 μ g/ml, 10 μ g/ml, 20 μ g/ml, 30 μ g/ml, 40 μ g/ml, 50 μ g/ml, 60 μ g/ml, 70 μ g/ml, 80 μ g/ml, 90 μ g/ml, 100 μ g/ml, 200 μ g/ml, 300 μ g/ml, 400 μ g/ml, 500 μ g/ml, 600 μ g/ml, 700 μ g/ml, 800 μ g/ml, 900 μ g/ml, and 1 mg/ml and/or less than about 1.1 mg/ml, 900 μ g/ml, 800 μ g/ml, 700 μ g/ml, 600 μ g/ml, 500 μ g/ml, 400 μ g/ml, 300 μ g/ml, 200 μ g/ml, 100 μ g/ml, 90 μ g/ml, 80 μ g/ml, 70 μ g/ml, 60 μ g/ml, 50 μ g/ml, 40 μ g/ml, 30 μ g/ml, 20 μ g/ml, 10 μ g/ml, 9 μ g/ml, 8 μ g/ml, 7 μ g/ml, 6 μ g/ml, 5 μ g/ml, 4 μ g/ml, 3 μ g/ml, 2 μ g/ml, 1 μ g/ml, 900 ng/ml, 800 ng/ml, 700 ng/ml, 600 ng/ml, 500 ng/ml, 400 ng/ml, 300 ng/ml, 200 ng/ml, 150 ng/ml, 100 ng/ml, 50 ng/ml, 40 ng/ml, 35 ng/ml, 30 ng/ml, 25 ng/ml, 20 ng/ml, 16 ng/ml, 15 ng/ml, 14 ng/ml, 13 ng/ml, 12 ng/ml, 11 ng/ml, 10 ng/ml, 9 ng/ml, 8 ng/ml, 7 ng/ml, 6 ng/ml, 5 ng/ml, 4 ng/ml, 3 ng/ml, 2 ng/ml, and 1 ng/ml. In some cases the concentration of supplement may be greater than

1 nM, 2 nM, 3 nM, 4 nM, 5 nM, 6 nM, 7 nM, 8 nM, 9 nM, 10 nM, 11 nM, 12 nM, 13 nM, 14 nM, 15 nM, 20 nM, 25 nM, 30 nM, 35 nM, 40 nM, 50 nM, 100 nM, 200 nM, 300 nM, 400 nM, 500 nM, 600 nM, 700 nM, 800 nM, 900 nM, 1 μ M, 10 μ M, 20 μ M, 30 μ M, 40 μ M, 50 μ M, 60 μ M, 70 μ M, 80 μ M, 90 μ M, 100 μ M, 200 μ M, 300 μ M, 400 μ M, 500 μ M, 600 μ M, 700 μ M, 800 μ M, 900 μ M, and 1 mM and/or less than about 1.1 mM, 1mM, 900 μ M, 800 μ M, 700 μ M, 600 μ M, 500 μ M, 400 μ M, 300 μ M, 200 μ M, 100 μ M, 90 μ M, 80 μ M, 70 μ M, 60 μ M, 50 μ M, 40 μ M, 30 μ M, 20 μ M, 10 μ M, 9 μ M, 8 μ M, 7 μ M, 6 μ M, 5 μ M, 4 μ M, 3 μ M, 2 μ M, 1 μ M, 900 nM, 800 nM, 700 nM, 600 nM, 500 nM, 400 nM, 300 nM, 200 nM, 150 nM, 100 nM, 50 nM, 40 nM, 35 nM, 30 nM, 25 nM, 20 nM, 16 nM, 15 nM, 14 nM, 13 nM, 12 nM, 11 nM, 10 nM, 9 nM, 8 nM, 7 nM, 6 nM, 5 nM, 4 nM, 3 nM, 2 nM, and 1 nM. In some cases the concentration is about 5-110 ng/ml, 5-15 ng/ml, or 90-110 ng/ml.

[55] In some case the cell culture media may or may not comprise a neuronal supplement. In some cases, the neuronal supplement may be a commercial neuronal supplement, for example B27, N2, or N10. In cases where a neuronal media supplement is added, the concentration of supplement in the cell culture media may be greater than about 0X, 1X, 2X, 3X, 4X, and 5X, and/or less than about 10X, 6X, 5X, 4X, 3X, 2X and 1X. Other commercial products include NeuroCult, ANS Neural Media supplement, Neurobasal supplement, B28, NS21, G5, N21, NS21, etc.

[56] In some cases, the cell culture media may also include other chemicals, molecules, supplements, or additives known to one of skill in the art of mammalian cell culture, for example, amino acids, peptides, salts, vitamins, antibiotics, antimycotics, antifungals, minerals, pH buffers, pH indicators, and sugars. In many cases the pH of the cell culture media may be greater than about 6.0, 6.1, 6.2, 6.3, 6.4, 6.5, 6.6, 6.7, 6.8, 6.9, 7.0, 7.1, 7.2, 7.3, 7.4, 7.5, 7.6, 7.7, 7.8, and 7.9 and/or less than about 8.0, 7.9, 7.8, 7.7, 7.6, 7.5, 7.4, 7.3, 7.2, 7.1, 7.0, 6.9, 6.8, 6.7, 6.5, 6.6, 6.4, 6.3, 6.2, 6.1, and 6.0. In various embodiments, the pH is from about 6.9-7.7, 7.0-7.4, or 7.3-7.7.

[57] Discogenic cells may be grown in a monolayer or in suspension. In some cases, cells may be grown in cell sterile receptacles, such as plates, dishes, flasks, roller flasks, and reactors for mammalian cell culture, which allow for the exchange of gases and

medium as needed. In various embodiments, cells may be grown while the receptacle may be stationary, or moving, for example by rotation or rolling of the receptacle. In some cases, the cell culture media may be agitated, for example by rotation, rocking, or rolling of the receptacle. The cell culture media may also be agitated through other
5 processes, for example physical movement of the cell culture media in a stationary receptacle, by, for example a stir rod, stir bar or other mechanical stirring mechanism within the cell culture media. In some cases the receptacle may include baffles for aiding in agitation of the media.

[58] In some cases, the receptacle may be treated, for example to aid or inhibit cell
10 attachment. Various culture methods may be used to grow cells under anchorage independent conditions. In general, cells that are able to grow in suspension may be grown in anchorage independent conditions. For example, cells that can grow and divide without attachment to a substratum may be anchorage independent. In some cases the receptacle may be coated to aid in attachment, for example with gelatin or collagen. In
15 some cases the receptacle may be coated to inhibit cells adhering or attaching to the receptacle surface, for example ultra-low attachment surface modifications. In some cases, receptacles may be commercially available, for example ultra-low attachment receptacles (Corning). Further, viscous non-reactive media additives, such as methylcellulose, poloxmer, or agar/agarose, may or may not be used to maintain a free-
20 floating suspension of cells.

[59] In some cases, where the cell culture media is supplemented to prevent or inhibit cell attachment, for example where a viscous, non-reactive substance is added, the concentration of the additive in the cell culture media may be greater than about 0.1%, 0.2%, 0.3%, 0.4%, 0.5%, 0.6%, 0.7%, 0.8%, 0.9%, 1.0%, 2%, 3%, 4%, 5%, 6%, 7%, 8%,
25 9%, 10%, and 15% and/or less than about 20%, 15%, 10%, 9%, 8%, 7%, 6%, 5%, 4%, 3%, 2%, 1.0%, 0.9%, 0.8%, 0.7%, 0.6%, 0.5%, 0.4%, 0.3%, 0.2%, and 0.1%. The final concentration of viscous reagent may depend on the reagent used, for example where methylcellulose is used the concentration may be between about 0.6-0.9%, 0.7-0.8%, or .75%. Where agarose is used, the concentration may be from about 1-5%, 2-4%, or 3%.

[60] In some cases, discogenic cells may be grown in atmospheres that have ambient levels of oxygen or higher or lower levels of oxygen. In many cases ambient levels of oxygen may be between 22 and 19 % oxygen. In some case the atmosphere that the cells are grown in has less than 22%, 21%, 20%, 19 %, 18 %, 17 %, 16 %, 15 %, 14 %, 13 %, 12 %, 11 %, 10 %, 9 %, 8 %, 7 %, 6 %, and 5 %, and/or greater than about 4 %, 5 %, 6 %, 7 %, 8 %, 9 %, 10 %, 11 %, 12 %, 13 %, 14 %, 15 %, 16 %, 17 %, 18 %, 19%, 20%, 21%, 22%. In some embodiments where hypoxia is desired the concentration of oxygen may be about 3-7%, 4-6%, 5% or 6%.

Surface Markers

[61] In some cases, discogenic cell populations may be characterized by the expression of cell-surface markers. In some cases the population of discogenic cells may or may not express one or more specific cell surface markers and/or cluster of differentiation proteins. In various embodiments, a population of discogenic cells may have a higher or lower percentage of gated cells with a specific marker than does a reference cells, for example a chondrocyte or a adipocyte. In other cases, a population of discogenic cells may have a percentage of gated cells with a cell surface marker. In some cases the percentage of gated cells having a specific cell surface marker is greater than 40%, 50%, 60%, 70%, 80%, and 90%, and/or less than about 100%, 90%, 80%, 70%, 60%, and 50%.

[62] Cell surface markers that may aid in characterizing a discogenic cell population may include, without limitation, CD24, CD34, CD44, CD73, CD90, CD105, CD166, Stro-1, HIF1, nestin, CK8, and HLA proteins (Human Leukocyte Antigen, e.g. HLA-A, -B, -C, HLA-DQ, and HLA-DR). In some cases, CD24 may be a glycoprotein expressed at the cell surface (e.g. lymphocytes, granulocytes, and neuroblasts), anchored via a glycosyl phosphatidylinositol (GPI) link to the cell surface. CD24 may also be referred to as Heat Stable Antigen (HSA). CD44 may refer to a cell-surface glycoprotein involved in cell-cell interactions, cell adhesion and migration. CD73 may also be referred to as 5'-ribonucleotide phosphohydrolase, and expressed on, for example B-cells, T-cells, endothelial cells, pericytes, follicular dendritic cells, fibroblasts, epithelial cells, cardiomyocytes, neurons, osteoblasts, trophoblasts and mesenchymal stem cells (MSCs). CD90 may refer to the glycoprotein Thy-1 Thymus cell antigen. CD105 may refer to

Endoglin, a glycoprotein component of the TGF-beta receptor complex. CD166 may refer to activated leukocyte cell adhesion molecule (ALCAM). Stro-1 may refer to a marker of immature mesenchymal stem cells. HIF-1 may refer to hypoxia inducible factor. Nestin may refer to a neural marker. CK8 may refer to a cytokeratin marker.

5 [63] In some cases, expression of cell surface proteins/markers on discogenic cells may be measured. In various embodiments, cell surface protein expression is measured by using fluorescent antibodies that recognize an epitope of the cell surface protein being measured. In some cases, the measurement is through the use of flow cytometry, including fluorescence-activated cell sorting (FACs), using standard techniques. As
10 measured by FACs, the expression is measured as a percentage of gated cells within a specified range, where gating is set using IgG controls. In some cases, expression is greater than about 70% for HLA-ABC, CD44, CD73 and CD90, and below 40% for CD24, CD105 and CD166.

[64] Additionally, in some cases surface markers may be used to sort, isolate or
15 concentrate a specific subpopulation or cells. For example, cell sorting using magnetic beads, fluorescent markers or other techniques may be used to select subpopulations within the population.

Gene Expression

[65] Additional to surface markers, in some cases genomic and genetic analysis is used
20 to identify the discogenic cells. Techniques include quantitative polymerase chains reaction, microarray analysis, western blot, etc. Expression of genes via measurements of DNA, mRNA, miRNA or protein, measured by fold-increase or decrease against house-keeping genes (GAPDH - Glyceraldehyde 3-phosphate dehydrogenase, SDHA - Succinate dehydrogenase complex, subunit A, HPRT1 - hypoxanthine
25 phosphoribosyltransferase, B2M - Beta-2 microglobulin, etc), may be used to identify discogenic cells, such as transcription factor Sox9, extra-cellular matrix component aggrecan, extra-cellular matrix components collagen 1 and 2, neural marker nestin, cytokeratin 8, transcription actor Sox1, CD44 (a receptor for hyaluronic acid), ALPI (alkaline phosphatase), PPARG (peroxisome proliferator-activated receptor gamma),

MMP (matrix metalloproteinase), ADAMTS (A disintegrin and metalloproteinase with thrombospondin motifs), FMOD (fibromodulin), interleukins, etc.

Cryopreservation

[66] In some cases, cells may be cryopreserved. Cells may be combined with
5 preformulated cryopreservation media, such as Cryostor, HyCryo, UltraCruz, Cyagen, etc. Or, cells may be combined with formulated cryopreservation media, which may or may not contain serum, albumin, dimethyl sulfoxide, trehalose, sucrose, other sugars, ethylene glycol, glycerol, propylene glycol, hyaluronic acid, collagen, matrigel, other natural extra-cellular matrix molecules etc. Cells may be frozen rapidly (vitrification) or
10 slowly (defined timecourse in various temperatures, or controlled-rate freezing machine). Cells may be frozen from 0.1-10 million cells/mL.

Isolating Discogenic Cell Populations

[67] Methods of deriving, obtaining, or isolating discogenic cells from disc tissue are described. In some cases the isolated discogenic cell population is derived from
15 autologous or non-autologous donors. An autologous donor may be where the discogenic cell population is derived from subject to be treated with the cells. A non-autologous donor, also known as an allogeneic donor, may be a different subject. Also disclosed are various methods for obtaining and preparing discogenic cells from living and or deceased donors.

20 [68] In most cases, the method of isolating discogenic cells from disc tissue comprises separating disc cells from extracellular matrix. In some cases, disc tissue may be broken up mechanically, chemically, and/or enzymatically. In some cases disc tissue may be chopped, sliced, or minced. In some cases, disc tissue is treated with an enzyme, for example collagenase. The treatment of disc tissue may aid in removal of extracellular
25 matrix. In some cases, the tissue is placed in tissue cultured-treated dishes with direct contact to the surface of the dish with media, and the cells migrate from the tissue and onto the plate. In other cases, the cells are separated from the tissue using a filter.

[69] Extracellular matrix may comprise collagen, proteoglycans, and other molecules. In some cases, collagen may refer to a group of naturally occurring proteins found in

animals, for example, mammals. Natural collagen may form of elongated fibrils comprising a triple helix. In most cases the three helices of collagen comprise two identical alpha 1 chains ($\alpha 1$) and one alpha 2 chain ($\alpha 2$). In most cases collagen is post transcriptionally modified by hydroxylation, cross-linking, glycosylation, cleavage, etc.

5 Collagen may be obtained from animals or animal cells. Collagen may be synthesized from a variety of cells include mammalian and non-mammalian cells such as bacteria using techniques well known to one of skill in the art. Proteoglycans may refer to proteins that are glycosylated. A proteoglycan may comprise one or more sulfated glycosaminoglycan (sGAG) chain(s) that may be attached at a Ser residue (generally in

10 the sequence -Ser-Gly-X-Gly-, where X can be any amino acid residue). Proteoglycan chains are, in general, long, linear, and negatively charged under physiological conditions. Glycosaminoglycans can be assayed as a method of determining proteoglycan production from a cell using, for example, a dimethylmethylene blue colorimetric assay or enzyme-linked immunosorbent assay. In many cases, both the

15 glycosaminoglycans in the micromass as well as the exchanged media (from media changes) are assayed. In some cases, discogenic cells grown in micromass culture in pro-chondrogenic media containing, for example TGF-B (transforming growth factor beta) or other growth factors known to produce a chondrogenic phenotype, may produce greater than about 1, 2, 3, 4, 5, 6, 7, 8, 9, 10 and less than about 20, 19, 18, 17, 16, 15, 14, 13, 12,

20 11 μg sGAG. In many cases, these results may be normalized to a quantification of cell number, DNA content or protein content, in order to determine proteoglycan production per cells. When normalized to cell number, the values may be greater than about .001, .005, .01, .02, .03, .04, .05, .06, and less than 1, .09, .08, .07 ng sGAG/cell. If the sGAG is normalized to its protein content, the values may be greater than 50, 60, 70, 80, 90,

25 100, 200, 300, 400 and less than about 1000, 900, 800, 700, 600, 500 ng sGAG/ μg protein. In many cases, discogenic cells produce more or less soluble proteoglycan than other cells, for example fibroblasts, mesenchymal stem cells, or disc cells that may or may not have been grown in attachment-independent conditions grown for similar amounts of time.

30 [70] Cells may be differentiated along adipogenic, osteogenic, and neurogenic lineages through techniques known to those skills in the art, such as the use of a StemPro multi-

potentiality kit by Life Technologies, wherein specialized media is used to differentiated adherent monolayer cultures of cells. After osteogenic differentiation, presence of mineralized bone is identified with Alizarin Red dye. After adipogenic differentiation, presence of fat is identified with Oil Red O dye. After neuronal differentiation, presence of neuronal morphologies is observed. Additionally, genetic markers can be tested to identify relevant phenotypic changes.

[71] Cells may be capable of self-renewal, which is defined as the ability to replicate without change to cell phenotype. This property can be identified in vitro via growth characterization over many passages, or in vivo through serial implantations and extractions.

[72] In some embodiments, discogenic cells may produce extra-cellular matrix molecules. In other embodiments, discogenic cells may produce proteins. In other embodiments, discogenic cells may produce growth factors. In other embodiments, discogenic cells may produce cytokines. In other embodiments, discogenic cells may produce hormones. In other embodiments, discogenic cells may produce sugars.

[73] After removal, reduction, or degradation of the extracellular matrix, cells from disc tissue may be transferred to either an attachment-dependent or attachment-independent culture system. In most cases, if disc tissue is submitted to an attachment dependent system, the receptacles may be treated with gelatin and/or collagen. In cases where the disc tissue cells are submitted to an attachment-independent culture system, the media may contain a viscous, non-reactive material to form a gel, such as methylcellulose.

[74] Once attached, cells are passaged (detached from a vessel, resuspended at a lower density and again attached on a vessel) up to about 10 times. In some cases, the cells never reach confluence in the vessel. In other cases, the cells reach confluence in the vessel. Cells are passaged using standard cell culture techniques. When sufficient cell quantities are obtained, but prior to proliferative 'drop-off' (meaning that the cells divide at a significantly slower rate), the cells are transitioned to a suspension culture containing a viscous, non-reactive medium for the desired period of time. When complete, the cells

are isolated, washed free of residual material, and further processed as needed for cryopreservation or direct therapeutic use.

[75] Growth of disc tissue derived cells on gelatin or collagen-coated receptacles may allow growth, expansion, and/or differentiation of the discogenic cells. Discogenic cells
5 can be added at 1,000-50,000 cells/cm². In these cases, cells may be grown in the presence of serum, EGF, and bFGF. In some cases, serum additives may added to the cell media, for example conditioned media from fibroblast cell cultures may also be added. In these cases, non-discogenic and less-discogenic cells may also grow, expand, and/or differentiate with the discogenic cells.

10 [76] Growth of disc tissue-derived cells on in anchorage independent conditions may aid in the growth, expansion, and/or selection of discogenic cells. Discogenic cells can be added at 10,000 cells/mL or up to 80,000 cells/mL. In some cases, neuronal supplements, bFGF, and EGF may also be added to the attachment-independent cell culture media. In some cases, serum may or may not be added to the media. In most cases, disc cells from
15 cultures of disc cells grown in attachment-dependent conditions produce less extracellular matrix than cells grown in attachment-independent conditions.

[77] In most cases, the cells may be washed free of cell culture media. In some cases, the cells are washed with PBS, additional media, cryoprotectant media, etc. In some cases, the vessels are chilled at 4C prior to washing to aid in dissolution of viscous
20 components. Sometimes, repeated centrifugation is used to remove unwanted components. Sometimes, the cells are attached to a new vessel to allow for the removal of unwanted components. Sometimes, the cells are attached to carriers to allow for removal of unwanted components. Sometimes, chemical agents are used to remove unwanted components.

25 [78] According to standard techniques, cells may or may not be detached from vessels or other cells using enzymes such as trypsin, recombinant trypsin, Accutase, HyQTase, TrypLE, etc. Cells may be combined with additional media and centrifuged at various speeds, including low speeds to allow for separation of different cell densities (single cells versus clusters), regular speeds to form a cell concentrate at the bottom, high speeds
30 to form a tight cell pellet.

[79] In many embodiments, a discogenic cell population is characterized by the ability to restore, regenerate, and/or grow disc tissue in vivo. For example, a discogenic cell population is able to restore damaged or diseased disc in a subject with a damaged or diseased disc. In many embodiments, the introduction of a discogenic cell population into a damaged disc of a subject will restore intervertebral disc height to about the pre-damage height.

Cell Morphology

[80] Discogenic cells may be mononucleated. Discogenic cells may be multi-nucleated. Discogenic cells may contain organelles, such as mitochondria, Golgi apparatus, and ribosomes. Discogenic cells may be demonstrated to be viable according to trypan blue, alamar blue, live/dead assay (Life Technologies), or other assay. Discogenic cells may be capable of proliferating. Discogenic cells may be capable of producing extra-cellular matrix.

[81] Discogenic cells grown under attachment-independent conditions may have a morphology that is different than discogenic cells grown as monolayers. For example, discogenic cells from attachment-independent cell culture may be isolated, rounded cells. Or, they may form cell clusters that are loosely associated with other cells and/or rounded cell associations. Or, they may form tight cell clusters known as Discospheres. Discogenic cell clusters may have at least one dimension that is greater than about 50 μm after sufficient growth. Discogenic cells may begin as isolated, rounded cells and over time, some cells may proliferate to form clusters or discospheres. This duration may be 1, 2, 3, 4, 5, 6, 7, 8, 10, 12, 14, 21, or 28 days. Discogenic cells may have associated extracellular matrix, such as proteoglycans and collagens.

Cell Population

[82] In one embodiment of a discogenic population, the cells may include without limitation one or more of the following cells: a chondrocyte, a fibroblast, a nucleus pulposus cell, an annulus fibrosus cell, a mesenchymal stem cell, a stem cell, a progenitor cell, a cartilage cell. In another embodiment of a discogenic population, the cells may include without limitation one or more of the following cells having been modified ex

vivo: a chondrocyte, a fibroblast, a nucleus pulposus cell, an annulus fibrosus cell, a mesenchymal stem cell, a stem cell, a progenitor cell, a cartilage cell. In another embodiment of a discogenic population, the population may include one or more of the following: isolated cells, or clusters of associated cells, or discospheres. In another
5 embodiment of a discogenic population, cells may be separated to form an isolated cell population. In another embodiment of a discogenic population, cells may be aggregated to form at least one micromass. In another embodiment, the discogenic cells may be more therapeutic prior to separating the cells.

Therapeutic use of Discogenic Cell Populations

10 [83] Discogenic cells may be delivered directly to a damaged tissue. Discogenic cells in an effective amount may or may not be combined with a biomaterial scaffold that is pharmaceutically acceptable to aid implantation. For example, a viscous natural material, containing hyaluronic acid, collagen or other extra-cellular matrix molecules, may be used. Or, a solid natural material may be used. In some cases, additives may be included
15 for stability. In some cases, additives may be included to aid in cryopreservation.

[84] It has been discovered in accordance with the present invention that damaged intervertebral discs can be repaired and/or the structure of the disc regenerated by administration of discogenic cells, thereby reversing or stabilizing degenerative disc disease and other disc injuries. It has also further been discovered that administration of
20 such cells to a subject with an injured disc partially restored pre-injury disc height. Accordingly, the invention features methods for isolating and growing discogenic cells, for use in treating subjects having a disease of or damage to at least one intervertebral disc. In general, the methods comprise administering to the subject a therapeutically effective amount of discogenic cells such that repair and/or regeneration of the afflicted
25 intervertebral disc(s) occur.

[85] In highly preferred aspects, the methods comprise administering cells obtained or isolated from spinal column or other cartilaginous tissue to a subject in need of treatment for at least one diseased or damaged intervertebral disc, wherein the cells are capable of self-renewal and/or expansion in culture. Cells isolated from spinal column and
30 cartilaginous tissue may be expanded or maintained in culture prior to administration.

[86] In the presently disclosed methods, the discogenic cells can be administered in conjunction with biologically active agent(s). The discogenic cells can be administered in sequence with, or co-administered with the agents. Lysates, soluble cell fractions, membrane-enriched cell fractions, proteins, growth factors, hormones, cell culture media (e.g., conditioned media), or extracellular matrix derived from spinal, disc, or cartilaginous tissue or discogenic cells can also be administered to subject as appropriate, including co-administered discogenic cells themselves, and additional cells or agents. The particular agent chosen can be delivered as part of a kit from the provider. Or, the particular agent can be at the discretion of the medical professional directing the treatment of the subject, and can vary according to the particular needs or condition of the subject. The agent chosen can be used for various purposes such as, but not limited to, facilitating the administration of the cells, improving the repair and/or regeneration of the intervertebral disc, improving the overall health of the subject, reducing pain, and/or enhancing survival of transplanted cells.

[87] The cells can be administered to the subject by injection. For example, the cells can be injected directly into one or more intervertebral discs of the subject. In many cases, the cells can be injected into the nucleus pulposus, transition zone, or annulus fibrosis of an intervertebral disc. The discogenic cells can be administered alone, or in combination with a biological or therapeutic, and/or scaffold or matrix agent(s).

[88] In some embodiments discogenic cell populations may be implanted into a subject. For example, discogenic cell populations may be surgically implanted into a damaged or diseased disc. In some embodiments, a discogenic cell population may be surgically implanted into an intervertebral disc space where all or part of a disc has been removed. In some embodiments, a discogenic cell population may be implanted into an intervertebral disc space as part of an artificial or replacement disc.

[89] The cells can also be administered as a scaffold- or matrix-cell complex. Scaffold and matrix compositions include but are not limited to proteins, hydrogels, synthetic polymers, and combinations thereof. Scaffold and matrix compositions may or may not be biodegradable. Such materials are known to one of skill in the art of therapeutic treatment, surgical repair, tissue engineering, and wound healing. Scaffold- and matrix-

cell compositions can be introduced into a subject's body in any way known in the art, including but not limited to implantation, injection, surgical attachment, transplantation with other tissue, and the like. In some embodiments, the matrices form in vivo, or even more preferably in situ, for example in situ polymerizable gels can be used in accordance
5 with the invention. Examples of such gels are known in the art.

[90] The discogenic cells can be mixed with scaffolds and matrices prior to implantation or seeded onto such compositions in vitro, which may allow the cells to proliferate and/or establish extracellular matrix. In some cases, the matrix may resemble mammalian intervertebral disc structure, and the matrix may replace an entire
10 intervertebral disc in a subject. In some cases, the matrices may include therapeutic agents.

[91] The discogenic cells can be used to produce an artificial disc replacement device in vitro or in vivo. In one example, an appropriate non-resorbable material, such as polyurethane, is used to create an artificial outer annulus. In another example, a
15 resorbable material, such as polyglycolic or polylactic acid, is used. This artificial annulus serves as a container for discogenic cells, which may or may not be combined with at least one of the following – scaffold material, matrix material, carrier material, growth factor(s), and/or other biologically active agents. In many embodiments, the artificial annulus structure may be porous and/or fibrous. The artificial outer annulus
20 may incorporate attachment means so that it can be fixed to one or more vertebral bodies. For example, the artificial annulus may incorporate through-holes, cuffs, tabs, loops, or washers to allow for screw fixation to one or more vertebral bodies. The artificial disc can be surgically implanted in a subject to completely replace a spinal disc.

[92] Artificial disc replacement devices may comprise the disclosed discogenic cells.
25 In various embodiments, discogenic cells may be inserted into an artificial annulus structure. The artificial annulus structure may be designed to provide a containment structure for the discogenic cells and may also further comprise attachment structures for securedly attaching the disc replacement device to one or more vertebral bodies.

[93] Discogenic cells are added to an artificial disc replacement device at or about the
30 time of insertion of the disc replacement device. In other embodiments, discogenic cells

are added to the disc replacement device well before insertion such that the cells can grow, divide, and provide a matrix or scaffold material. In many embodiments, scaffold material, matrix material, carrier material, growth factor(s), and/or other biologically active agents can be added to the disc replacement device before, after, or in combination
5 with addition of discogenic cells to the disc replacement device.

[94] Artificial disc replacement devices may comprise a resorbable or non-resorbable artificial annulus. In some embodiments, the artificial annulus may comprise matrix material, scaffolds, growth factors, or other biologically active agents to aid in supporting the growth of discogenic cells. In one embodiment, the artificial annulus supports the
10 growth and/or differentiation of annulus fibrosus cells, for example by providing a localized source of growth factors and/or cytokines that may promote discogenic cell differentiation into annulus fibrosus cells. In many embodiments, the artificial disc replacement device comprising the discogenic cells may have a cellular architecture similar to that of a non-artificial intervertebral discs, for example comprising an annulus
15 fibrosus, nucleus pulposus, and endplate tissue.

[95] In some cases, discogenic cells and scaffold may be cryopreserved in liquid nitrogen. Or, they may be stored at various sub-zero temperatures, such as -80C, -20C, or -1C. Additionally, the discogenic cells may be stored at 4°C or 37°C. The discogenic cells may be combined with a scaffold prior to storage. Or, the discogenic cells may or
20 may not be combined with a scaffold immediately prior to implantation.

[96] A successful treatment could thus comprise treatment of a subject with a disease, pathology, or trauma to the intervertebral disc with a therapeutic cell composition comprising the discogenic cells, in the presence or absence of therapeutic agents, matrices, or scaffolds. The cells are present in an effective amount to promote, for
25 example, direct proteoglycan production or stimulation of native cells to regenerate. The result could be regeneration, repair, or reconstruction of native tissue architecture, as shown in a rabbit model. This can be determined via medical imaging (x-ray, MRI) or reduction in pain. In the case of implantation into a human intervertebral disc, cell content can be 1,000-10,000,000, containing a discogenic cell population or a

subpopulation extracted from discogenic cells. The scaffold volume can also range, from 10 μ L to 1000 μ L, or 10 mg to 10 g, depending on the spatial needs of the subject.

[97] In some embodiments, one or more subpopulations of cells within discogenic cells may be implanted. This subpopulation may be isolated using magnetic beads for
5 sorting, fluorescent markers for sorting, density gradients, fluorescent genetic tagging for sorting, physical separation, filtration, etc. This subpopulation may be additionally therapeutic compared to the entire population.

[98] Upon treatment of a subject, discogenic cells may provide superior therapeutic effect compared to attachment-dependent discogenic cells or other cell populations, such
10 as nucleus pulposus cells, fibroblasts, chondrocytes, stem cells, progenitor cells, etc. This effect may or may not be improved through the use of scaffolds, carriers or other biomaterials.

[99] Also featured in accordance with the present invention are kits for practicing the disclosed methods. In one aspect, kits for treating a subject having a disease of or damage
15 to at least one intervertebral disc are provided. The kits may comprise a pharmaceutically acceptable carrier, discogenic cells in an amount effective to treat the disease or injury, and instructions for using the kit in a method for treating a subject having a disease of or damage to at least one intervertebral disc. The kits may further comprise at least one reagent and instructions for culturing the cells. The kits may further comprise at least one
20 biologically or therapeutically active agent. The kits may further comprise vials and syringes. The kits may further comprise discogram needles for direct entry into the intervertebral disc. The kits may further comprise a radiopaque agent to aid in imaging during the procedure.

EXPERIMENTAL DETAILS SECTION

25 **Materials and Methods**

Supplies/Reagents:

[100] In some cases, attachment dependent cell culture may be referred to as Expansion Conditions. Expansion Medium may be used for growth of cells under Expansion conditions. Expansion medium contained DMEM/F12 with 10% fetal bovine serum.

30% of this medium can be optionally pre-conditioned in the presence of neonatal foreskin fibroblasts for 3 days. This pre-conditioned portion was filtered prior to use. Prior to adding Expansion Medium to cells, bFGF and EGF are added to make the media 'complete' with final concentrations of 10 ng/mL bFGF and 10 ng/mL EGF, obtained
5 from stored 1000X stock solutions.

[101] In some cases, attachment independent cell culture may be referred to as Suspension Conditions. Suspension Medium may be used for growth of cells under Suspension conditions. Suspension medium contained 1% A4M Premium methylcellulose (Dow Chemical) in DMEM/F12 supplemented with 1X B27 (Life
10 Technologies), 5% fetal bovine serum, 10ng/ml EGF, and 10 ng/ml bFGF. Suspension feeding medium is the same, minus the methylcellulose.

[102] For expansion, Plates or Flasks coated with collagen, gelatin or another similar matrix protein were used. These vessels were purchased pre-made. Or, plates can be coated manually by dissolving 1 g powdered gelatin (Sigma) in 1 L ddH₂O (double
15 distilled water) or prepared from premade solutions (Sigma) by diluting the solution to 1% final concentration. If coating in the lab, the receptacles are then incubated at room temperature for 15 minutes or more. For suspension, ultra-low attachment vessels were used from Corning.

[103] In some cases, a viscous scaffold carrier was used to implant the cells in vivo.
20 Specifically, a pre-manufactured sterile gel composed of 1.7% hyaluronic acid (0.8-1.2 MDa) in PBS was diluted with human serum albumin and 30,000 discogenic cells to result in a 1% hyaluronic acid gel with 2.5% human serum albumin. This was loaded into sterile 50 ul glass Hamilton syringes. 27 gauge Precision Glide needles were fixed onto the Hamilton syringes via leur-lock for animal implantation.

25 Methods:

[104] Human adult nucleus pulposus tissue is obtained from consenting surgical donors using an IRB-approved protocol. In various embodiments, tissue may be obtained from a variety of sources and tissues, for example, living and deceased non-autologous donors, autologous donors, intervertebral tissue, or from other cartilaginous tissue. Non-nucleus

pulposus tissue, including annulus fibrosis and cartilaginous endplate, was manually resected and discarded. 2-7 grams of the resulting tissue was combined with 15 ml DMEM/F12 containing 300 units/ml of collagenase II in a T75 flask, and incubated overnight under standard tissue culture conditions (37 °C and 5% CO₂). Next, the
5 liberated cells were transferred into a 50ml tube, the cells spun down (4 min, 1200 rpm), supernatant removed and the cells resuspended in DMEM/F12 to a final concentration of about 10,000 cells/ml.

[105] Alternatively, cells were obtained from cryopreserved vial (stored in liquid nitrogen) by thawing in 37C water bath and immediately transferring to a 15 mL tube
10 containing 10 mL of expansion medium. The cell mixture was then centrifuged for 4 minutes at 1200 rpm. The supernatant was then aspirated and the cells resuspended in expansion medium to count cells and confirm viability.

[106] For expansion, cells were plated at 10,000 cells/mL in complete expansion medium, and the medium replaced every 2-3 days. Prior to confluence, cells are washed
15 with PBS, incubated with 0.25% trypsin for 7 minutes, and then removed from the plate with expansion medium and transferred into a 50ml conical tube. The tube was spun as previously described, and the supernatant containing trypsin was removed. The cells were then resuspended in expansion medium to allow for cell counting and viability check. At this point, cells could be cryopreserved by combining 0.5-3 million cells with 90%
20 FBS/10% DMSO in a cryovial. To freeze the contents, the cryovial was stored overnight in -80 and then moved to vapor phase of liquid nitrogen for long-term storage.

[107] For suspension culture, the cells were combined with suspension medium at 10,000 cells/ml (15 ml total) and added to ultra-low attachment 100 mm dishes (Corning). Every 2 days, 300 µl of suspension feeding medium was added to the dish.
25 Images were captured at 4X, 10X and 20X using a phase-contrast light microscope, as in Figure 2. After 2 weeks, the cells were harvested by incubating the plates at 4°C for 20 minutes, followed by diluting the gel-like medium with 15 mL PBS. The contents of the plate were then transferred to a 15mL tube, the volume increased to 50mL with additional PBS, and spun at 1200 rpm for 4 minutes, and the supernatant removed. The cells were
30 then washed twice more in this manner, to ensure removal of methylcellulose. Finally,

the discogenic cells were resuspended in expansion medium to allow for cell count and viability check.

[108] To determine surface marker expression, the discogenic cells are combined with 0.25% trypsin for 7 minutes to create a single-cell suspension. Additional cell populations tested included: pre-discogenic cells from expansion culture, adult human mesenchymal stem cells (CET Company), human neonatal foreskin fibroblasts (AATC), and human adult articular chondrocytes (ScienCell). The cells were analyzed using standard techniques with a Partec Cyflow flow cytometer. Briefly, 70,000 cells and antibodies (one type per tube) were diluted in 200ul PBS containing 0.5% bovine serum albumin and incubated for 30 minutes at 4C in dark conditions. The following antibodies were used: CD73-PE (BD), CD90-PE-Cy5 (BD), CD105-PE (Miltenyi), CD166-PE (BD), HLA-ABC (BD) and control. The work was performed in duplicate, with one representative data set shown.

[109] To determine proteoglycan production in a pro-chondrogenic environment, a standard chondrogenic pellet assay was utilized. Briefly, 200,000 cells (discogenic cells or mesenchymal stem cells) were added to individual conical wells in a 96-well plate with 200ul pro-chondrogenic media (StemPro media, Life Technologies) and spun briefly in a centrifuge. Every 3 days, the media was removed, collected and replaced, taking care not to aspirate the central micromass that formed. After 2 weeks (see Figure 3), the cell pellets were harvested, dried and digested overnight in 250ul of papain (Sigma) in a buffer at 60C. The digestate, along with the collected media, was assay for sGAG content, a component of proteoglycan, using a standard DMMB assay. Specifically, to make the DMMB working solution, 1g of sodium formate was dissolved in 490mL of deionized water and 1mL of formic acid was added. In another tube dissolve, 8 mg of dimethylmethylene blue (DMMB) powder (Sigma) was combined with 2.5 mL of ethanol, and the two solutions were combined. Then, water was added to bring the volume up to 500mL. To make the chondroitin-6-sulfate standards, 40mg of chondroitin-6-sulfate (C6S) was mixed with 40mL of water to make 1mg/mL of standard solution (stock). Then, the stock was diluted to make 0, 1.0, 2.0, 3.0, 4.0, 5.0, 6.0, 7.0, 8.0, 9.0, 10.0 µg/mL. To assay for proteoglycan content, 100ul of standards or sample and 100 ul

of DMMB working solution was combined in a 96-well clear plate and read within 5 minutes at 525 nm. The standards were used to determine a concentration of the sample, and then the concentration was normalized to original volume to determine mass.

Additionally, the digestate was assayed for protein content using the Pierce protein assay per manufacturer instructions, or DNA content using the Quanti-IT PicoGreen Assay (Life Technologies) per manufacturer instructions..

[110] To determine Osteogenic and Adipogenic potential (Figure 4), discogenic cells were grown using the StemPro Osteogenic and Adipogenic Kits (Life Technologies) according to the manufacturers instructions.

[111] To assess in vivo efficacy, 30,000 discogenic cells were combined with 25ul of 1% hyaluronic acid (0.8-1.2M Da) in PBS containing 2.5% human serum albumin. The cell and viscous scaffold were loaded into 50ul glass Hamilton syringes. Using a previously internally validated model of degenerative disc disease, lumbar discs of 3 New Zealand rabbits were accessed surgically and punctured with a needle to induce degeneration (n=4 discs/animal, approved by local IACUC). After two weeks, either cell therapy or a cell-free scaffold control was injected into the injured discs. Additionally, injured and uninjured control lumbar discs were maintained in each animal. Every 2 week for 6 weeks, disc height was measured using plain radiographs and normalized to week 0 values, resulting in a Disc Height Index (DHI). After 6 weeks, the rabbits were euthanized; the discs were harvested and prepared for histology. Sections were stained with H&E or Alcian blue, and blindly scored for abnormality, assigning a score of 0 to 2 for AF/NP border, AF organization, NP extra-cellular matrix, and NP cellularity (AF – annulus fibrosus; NP – nucleus pulposus) and summing the 4 results (0=normal, 8=abnormal).

EXAMPLES

[112] The following examples describe in detail the preparation and properties of embodiments of the microenvironments of the present disclosure. It will be apparent to those skilled in the art that many modifications, both to configurations, materials, and methods, may be practiced without departing from the scope of the disclosure.

Example 1 – Surface Marker Expression of various cell types via FACs analysis.

[113] Cell surface marker expression was analyzed for a variety of cells including discogenic cells. As shown in Figure 1, Human cell types tested include adult mesenchymal stem cells, neonatal foreskin fibroblasts (Fibro.), articular chondrocytes (Chondro.), anchorage-dependent discogenic cells (AD-DC), anchorage-independent discogenic cells (AI-DC). Figure 1A shows the percent of cells expressing given marker based on standard IgG gating. DC cells show lowered expression of CD105 and CD166 compared to other cell types. Other surface markers are being explored to identify unique a unique phenotypic fingerprint for DCs. Figure 1B presents representative flow cytometry analysis of cell markers in MSCs, pre-discogenic cells and discogenic cells for CD105 and CD166 showed unique pattern of discogenic cells.

Example 2 – Morphology of discogenic cells in suspension culture

[114] The morphology of discogenic cells was investigated. Figure 2A is a micrograph showing discogenic cells consist of tight spheroids known as discospheres (*), loose aggregates of cells (**), and single cells (***). Figure 2B shows that discosphere and aggregate diameters vary, and are typically larger than 50um.

Example 3 – Chondrogenic potential of mesenchymal stem cells (MSCs) and discogenic cells (DCs).

[115] Figure 3 shows analysis of chondrogenic potential of mesenchymal stem cells and discogenic cells. Figure 3A shows morphology of cells after 2 weeks of growth in pro-chondrogenic environment, as shown here, the DCs produce larger micromasses than the MSCs. Figure 3B are graphs comparing MSC and DC production of protein and GAGs. As shown here, DC produced more sGAG (a component of proteoglycan), collected from both the media and digested micromass, compared to MSCs, despite comparable protein content in each micromass.

Example 4 – Adipogenic and Osteogenic potential of discogenic cell population.

[116] Figure 4A shows that adipogenesis was confirmed via Oil Red O staining counterstained with hematoxylin after differentiation in adipogenic media using manufacturer's instructions (Life Technologies). Figure 4B shows that osteogenesis was

confirmed via Alizarin Red staining after differentiation in osteogenic media using manufacturer's instructions (LifeTechnologies). (Scale = 50 μ m)

Example 5 – Discogenic cell viability at various stages prior to therapeutic use, via live/dead assay (green indicates live, red indicates dead).

- 5 [117] Figure 5A shows that viability is confirmed after 24 hours in viscous hyaluronic acid scaffold, where the majority of cells are alive (bright cells; some dead cells are indicated with arrows). Figure 5B shows that viability (bright cells) is again confirmed after being extruded through 27 gauge surgical needle used for surgical implantation into rabbits.

10 **Example 6 – Therapeutic use of discogenic cells with a viscous scaffold carrier.**

- [118] Figure 6 is a schematic diagram showing embodiments and steps of using discogenic cells to treat DDD. Figure 6(A) I depicts the use of discogenic cells by injection. Cells and scaffold may be mixed and pre-filled into a syringe or vial. Cells and scaffold may be separate and mixed immediately prior to injection. Product can be delivered from a syringe or false-bottom vial. Product is injected directly into a
15 degenerated disc. (B) IFigure 6B depicts discogenic cell implantation. Cells and scaffold may be combined, or delivered separately. Prior to implantation, the material is cut or modified to fill the intended implantation area. The material is then implanted into the defect. The cells may be added after implantation (not shown). For the in vivo animal
20 study presented, cells and scaffold were mixed and shipped at 4°C in a vial, and the contents loaded immediately prior to injection into a syringe.

Example 7 – Efficacy of discogenic cells in a viscous scaffold carrier in a rabbit model of degenerative disc disease.

- [119] Figure 7A is a graph showing that discogenic cell therapy within a scaffold carrier
25 restored disc height index (DHI) compared to control groups at 4 and 6 weeks. All control groups were consistent between week 2, 4 and 6. N=3, * indicates $p < 0.001$ compared to both Scaffold Control and Injured Control using 2-way ANOVA and Tukey's Post-hoc test. Figure 7B is a graph showing the histological score. Blind scoring of discs at week 6 (0-8, 0= normal). The cell therapy showed significant improvement

compared to injured control and scaffold control. ** indicates $p < 0.05$ compared to scaffold control, * indicates $p < 0.05$ compared to injured control, using t-test.

Example 8 – production and assessment of discogenic cells

Materials and Methods

5 Production of Discogenic cells

[120] The described procedure is depicted in the flow diagram in Figure 8. First, discarded adult human nucleus pulposus tissue from discectomy procedures was procured with IRB from consenting donors (Baptist Hospital, TN). Fibrous, annulus material and other tissue contaminants were removed via dissection. The remaining material was then
10 washed three times with 2X antibiotic-antimycotic (ABAM, HyClone by Thermo Scientific) in PBS, and digested overnight in 300 units recombinant type-2 collagenase (Life Technologies) in DMEM/F12 (Life Technologies) with 1X ABAM. The isolated cells were plated onto gelatin-coated flasks in expansion medium (DMEM/F12 with 10% FBS by Hyclone, 10 ng/mL EGF and 10 ng/mL FGF-2 by Peprotech), and over time a
15 subpopulation of cells attached to the plates, composed of stem/progenitor cells. These cells were expanded for up to 4 passages.

[121] Next, the cells were combined with suspension medium (10,000 cells/cm² in DMEM/F12 with 5% FBS, 10 ng/mL EGF and 10 ng/mL FGF-2) in the presence of 1% methylcellulose (A4M Premium, Dow Chemical), adding 1.5 mL of media per cm² in
20 ultra-low attachment vessels (Corning; Corning, NY). After 2 weeks, the cells were harvested from methylcellulose for further use with 3 washes in phosphate buffered saline (PBS, Corning CellGro; Manassas VA. Discogenic cells were produced from 5 distinct human don A suspension of discogenic cells was fixed in 10% formalin (Sigma-Aldrich; St. Louis, MO) for 15 minutes.

25 Histology of Discogenic Cells

[122] Next, the cells were washed with PBS 3 times and cells resuspended in 37°C agarose (approximately 1×10^7 cells per 0.5mL in 1% BioReagent low gelling temperature agarose by Sigma). When the agarose solidified, the pellet was kept in PBS until frozen in OCT medium (Sakura Tissue-Tek; Torrance, CA) in a Leica Cryostat and

sectioned 6 μ m thick onto charged slides. Samples were then stained with either hematoxylin and eosin, alcian blue counterstained with nuclear fast red, or picrosirius red, according to standard protocols. Histology shown is from one representative donor.

Confocal Microscopy

- 5 [123] A suspension of discogenic cells was fixed for 15 minutes in 10% formalin and washed 3 times in PBST (PBS + .1% Triton 100X by Sigma; St. Louis MO). Subsequently, primary antibodies in PBSTA (PBST + 0.5% human albumin by Baxter Healthcare; Westlake Village CA) were diluted 1:100 for anti-human aggrecan (Santa Cruz Biotechnology; Dallas TX) and 1:20 for anti-collagen II (Developmental Studies
- 10 Hybridoma Bank; Iowa City, Iowa) with fixed cells and incubated for 2 hours at room temp followed by 3 washes in PBST. The cells were then conjugated with Alexa-Fluor 488 for aggrecan and 633 for Collagen II in (both in PBSTA) for 1 hour at room temp and washed again 3 times with PBST. Finally, cells were then counterstained with DAPI and Phalloidin (Molecular Probes by Life Technologies) for cell nuclei and actin,
- 15 respectively, and imaged using an Olympus FV1000 confocal microscope. Histology shown is from one representative donor.

Flow Cytometry

- [124] Cell surface antigen expression of discogenic cells were analyzed by flow cytometry using the following fluorescence-conjugated mouse anti-human monoclonal
- 20 antibodies: CD105-phycoerythrin (PE, Miltenyi Biotec, Inc., Auburn, CA, USA); Stro-1-Alexa Fluor 647 (BioLegend, San Diego, CA, USA); CD166-PE, CD73-APC, and CD90-FITC, CD44-FITC, CD-24-PerCP-Cy5.5, CD34-PE, HLA-DRDP-FITC, HLA-ABC-FITC (all from BD Biosciences, San Jose, CA, USA). Appropriate isotype controls were run in parallel. The cells were incubated for 30 minutes at 4°C in PBS containing
- 25 50% mouse serum and subsequently washed and re-suspended in PBS containing 1.0% bovine serum albumen. DAPI, dilactate (Life Technologies, Carlsbad, CA, USA) was used to assess viable cell content. A minimum of 20,000 events were collected on a FACSCanto flow cytometer (BD Biosciences, San Jose, CA, USA) using FlowJo Software for data acquisition and analysis.

Multipotentiality

[125] Osteogenesis and adipogenesis was induced using kits supplied by Life Technologies (Grand Island, NY). Briefly, discogenic cells were dissociated for 15 minutes using TrypLE (Gibco by Life Technologies) to form a single-cell suspension and plated onto tissue-culture treated dishes at 20,000 cells/cm². Dishes of cells for osteogenic and adipogenic differentiation were maintained in DMEM with 10% FBS for 3 days, and then fed with the appropriate supplied differentiation media for 3 weeks. After differentiation, the monolayers were stained for calcification with Alizarin red or lipid content with Oil Red O as instructed. Phase images were captured at various magnifications. The study was performed on 4 distinct donors.

[126] Chondrogenesis was induced as described in [Johnstone 1998]. Briefly, 250,000 cells were added to individual conical wells in a 96-well plate with 200 µl chondrogenic media (Life Technologies) and spun briefly. Every 3 days, the media was completely removed, collected and replaced, taking care not to aspirate the central micromass that formed. After 2 weeks, the cell micromasses were harvested, dried and digested overnight in 250ul of papain (Sigma-Aldrich) at 60°C. The digestate, along with the collected media, was assay for GAG content as in [Farnesdale]. Additionally, the digestate was assayed for DNA content using the Quant-IT PicoGreen Assay (Invitrogen), and the results normalized to assess GAG/DNA.

[127] For comparison, the chondrogenic potential of discogenic cells was assessed with other, well-known adult human cell lines. Articular chondrocytes (Sciencell; Carlsbad, CA), bone-marrow derived mesenchymal stem cells (CET by Thermo Scientific; Waltham, MA), and dermal fibroblasts (ATCC; Manassas VA) were purchased and expanded according to provided instructions.

Results

Histology, Confocal Microscopy and Gene Expression

[128] The discogenic cells were assayed for aggrecan and collagen production. After 2 weeks in non-plastic adherent culture, individual NP-derived stem/progenitor cells proliferated into clusters of various sizes embedded in extracellular matrix (ECM). The

matrix was composed of proteoglycan and various collagens (Figure 9A-C), the main components of nucleus pulposus tissue. Confocal imaging was used to identify aggrecan and collagen 2 in the ECM, with variability in matrix content noted across clusters (Figure 9D). Further, high-magnification imaging revealed both intra- and extra-cellular
5 aggrecan (Figure 9E), suggesting active transport of ECM molecules out of the cell at the time of imaging.

[129] As depicted in Figure 9F, the expression of aggrecan and collagen 2 was assessed over time in culture (day 3 and 7), at cell harvest (day 14) and after chondrogenic differentiation compared to the housekeeping gene HPRT. During the course of the
10 culture period, expression of the extracellular matrix molecules increased significantly, with approximately 20-fold more expression of aggrecan and 70-fold more expression collagen 2 compared to the plastic-adherent cells. Gene expression was further upregulated upon chondrogenic differentiation.

Flow Cytometry

[130] Discogenic cells were dissociated to form a single-cell suspension, and analyzed for various surface markers using flow cytometry. The population was homogenous in terms of size and internal structure, as seen in the forward/side scatter plot shown in Figure 10A). A broad range of surface markers were tested and compared to mesenchymal stem cells (as a control). The expression of these markers was generally
20 uniform across 5 distinct human donors ($p > 0.05$) with >80% expression of CD44, CD73, CD90, HLA-ABC, and <10% expression of CD34, HLA-DR/DQ and STRO-1 compared to isotype controls (Figure 10B and 10C). CD24, CD105 and CD166 were expressed at approximately 40%, with slightly more variability.

Multipotency

[131] The discogenic cells were tested for their ability to form bone, fat and cartilage according to standard differentiation protocols. Robust adipogenesis and osteogenesis was observed by staining monolayers of differentiated cells (Figure 11A-B). Discogenic cells formed large, hard micromasses of cartilage upon differentiation that stained positively for proteoglycan (Figure 11C).

[132] After 3 weeks in culture, the media and micromasses were assayed quantitatively for proteoglycan content. Discogenic cells (DCs) were compared to articular chondrocytes (AC), fibroblasts (FB) and mesenchymal stem cells (MSCs). As depicted in Figure 11E, while the insoluble (micromass) quantity of proteoglycan did not change significantly by cell type, the amount of soluble proteoglycan measured varied considerably. When these two sources of matrix were combined and normalized to DNA content (Figure 11D), discogenic cells were shown to produce more proteoglycan/DNA than fibroblasts ($p < 0.01$), comparable levels to MSCs and articular chondrocytes.

Example 9 - In vivo Pilot Study in Rabbits

[133] Female New Zealand White rabbits (3-4 kg) were used for these studies, under approval by private IACUC. Three rabbits were fasted overnight prior to surgery. For the first surgery to induce degeneration, animals were anesthetized intravenously and the surgical site prepared for aseptic surgery. An 8-10 cm longitudinal incision was made on the left lateral abdomen between the iliac crest and the last rib. The lumbar vertebral discs were accessed via retroperitoneal approach using blunt dissection. An 18 gauge needle was then inserted at least 5mm into the lumbar vertebral discs of interest to create disk injury. Discs L2-L3, L3-L4, L4-L5, L5-L6 were injured using this technique. L5-L6 was left undisturbed. Muscle and skin were then closed in two or three layers using sutures, and the animals monitored during recovery. Prior to this study, 6 rabbits were injured and assessed for 8 weeks to confirm the creation of a stable and appropriate defect (data not shown).

[134] After 2 weeks, the rabbits were again prepared for surgery and anesthetized, and the discs accessed as previously described. A 27-gauge needle was used to inject 25ul of either the cell therapy containing 30,000 cells (L5-L6) or scaffold alone (L4-L5). For high dose therapy, 300,000 cells were injected. One disc was not modified to serve as injured control (L2-L3). The injection was held in place for 5 seconds, and when the needle was removed no material was observed to leak out.

[135] For 6 weeks, animals were monitored for any adverse events or health concerns. Body weight was measured every week. Additionally, x-rays of the lumbar spine were performed every 2 weeks by anesthetizing the animal briefly (Figure 12A). The distances

between boney landmarks (three measurements of the disc space, 3 measurements of the left-adjacent vertebral bone) were measured on plain radiographs using a micrometer by a single individual and normalized against week 0 distances to assess Disc Height Index (DHI) percent.

- 5 [136] After 6 weeks, the rabbits were euthanized; the discs were harvested and prepared for histology using paraffin. Sections (4µm) were stained with a mixture of hematoxylin and eosin, or Alcian blue and eosin.

Statistical Analysis

- [137] All statistical analyses were performed using StatPlus software (AnalystSoft; 10 Vancouver, Canada) utilizing Tukey's post-hoc tests. The p-values for significance were indicated in each study. A one-way analysis of variance (ANOVA) was used to compare the surface marker expression and proteoglycan production of discogenic cells in vitro. A two-way ANOVA was used to analyze the disc height data over time, and a one-way ANOVA was utilized for comparing histological scores at week 6. Graphs show mean 15 with error bars representing standard deviation of the data set.

Results

- [138] As described above, surgical puncture was used to induce 'degeneration' in the intervertebral discs of 3 New Zealand White rabbits (n=3 condition). At the time of injury, nucleus pulposus material was noted to extrude from the needle track after the 20 needle was removed. After surgery, animals did not show any unusual signs of distress due to the injury. After 2 weeks, an injection of low dose, high dose, or scaffold control was administered to the discs, with pressure held for 5 seconds before removing the needle. After injection, some material was noted to extrude out of the disc after the needle was removed.

- 25 [139] Over the 6 week course of the study, no safety issues were noted. No major changes in body weight were observed after initial injury (Figure 12B; 3.4 grams pre-injury to 3.3 grams at day 7, on average) or after cell therapy injection (Figure 12B; 3.2 grams on average). No health or behavioral issues were reported by the animal care staff regarding any of the animals after receiving injections of the human cells into their discs.

[140] Notable changes in disc height were measured over the course of 6 weeks by x-ray (a representative x-ray is at Figure 12A, which also shows measurement methods).

Two weeks after injury, all discs decreased to 59% of the original height on average. At week 4 and 6, the discs treated with low dose and high dose showed significant

5 improvements in disc height compared to both scaffold and injured controls, from 70% and 64% DHI to 94% and 71% DHI, respectively ($p < 0.001$) (Figure 12C). Injection of the scaffold control was slightly better than no injection at all, represented by the injured control (64 % DHI and 53% DHI, respectively; $p < 0.01$).

[141] Histologically, H&E images revealed normalization of disc architecture with cell
10 therapy treatment. The height of the nucleus pulposus, from end-plate to end-plate, decreased with injury and increased with cell therapy (Figure 13A). As shown in Figure 13B, no immune reaction or abnormal tissue formation was noted in bone marrow, annulus fibrosus (AF), cartilage endplates (CEP) or nucleus pulposus (NP) after treatment. The nucleus pulposus remained dense with proteoglycan, as shown by alcian
15 blue staining (Figure 13B).

Example 10 - In vivo Pilot Study in Pigs

[142] Female Gottingen minipigs (10-15 kg) were used for this research, under approval by private IACUC. Two pigs were fasted overnight prior to surgery. The injury surgery was performed as described above in Example 9, with addition of fluoroscopic imaging
20 to confirm proper placement of needles, injuring discs L2-L3, L3-L4 and L3-L4. L5-L6 was left undisturbed. Muscle and skin were then closed in two or three layers using sutures, and the animals monitored during recovery. Prior to this study, 6 minipigs were injured and assessed for 12 weeks to confirm the creation of a stable and appropriate defect (data not shown).

25 [143] After 2 weeks, the pigs were again prepared for surgery and anesthetized, and the discs accessed as previously described. A 27-gauge needle was used to injected 150ul of either the cell therapy containing 100,000 cells (High Dose; animal 1: L3-L4, L4-L5), 500,000 cells (Low Dose; animal 2: L3-L4, L4-L5) or scaffold alone (L2-L3). One disc was not modified to serve as injured control (L2-L3). The injection was held in place for
30 5 seconds, and when the needle was removed no material was observed to leak out.

[144] For an additional 10 weeks, animal were monitored for any adverse events or health concerns. Body weight was measured every week. Additionally, x-rays of the lumbar spine were performed at 4, 8 and 12 weeks by anesthetizing the animal briefly, and DHI determined as described above. After 12 weeks, the pigs were euthanized; the discs were harvested and prepared for histology using paraffin as previously described.

Results

[145] Injury of pig discs resulted in 20-30% reduction in disc height. Upon treatment, both the low and high dose treatment resulted in immediate improvement in disc height that was sustained to week 12 and was better than the injured control ($p < 0.05$). The scaffold and injured control did not improve over time (Figure 14A). Fluoroscopic imaging was used to aid in correct needle placement, and shows the differences in disc height along an injured spine (Figure 14B). Similar to the rabbit pilot study described above, no immune reaction or abnormal tissue formation was noted in nucleus pulposus, cartilage endplate, annulus fibrosus or bone marrow (not shown), with the nucleus pulposus staining for proteoglycan similarly to untreated tissue (Figure 14C).

[146] It is noted that there are alternative ways of implementing the embodiments disclosed herein. While a number of exemplary aspects and embodiments have been discussed above, those of skill in the art will recognize certain modifications, permutations, additions and sub combinations thereof. Accordingly, the present embodiments are to be considered as illustrative and not restrictive. Furthermore, the claims are not to be limited to the details given herein, and are entitled their full scope and equivalents thereof.

WHAT IS CLAIMED IS:

1. A discogenic cell population comprising:
at least one cell derived from mammalian disc tissue and grown in vitro in anchorage independent culture.
2. A heterogeneous cell population comprising:
one or more cells derived from a cartilaginous tissue, wherein at least one cell has been maintained in anchorage independent culture.
3. The cell population of one of claims 1 or 2, where the culture comprises a media comprising one or more additives selected from the group consisting of EGF, bFGF, serum, fibroblast conditioned media, and viscous non-reactive substance.
4. The cell population of claim 3, wherein the cell population is passaged in a culture receptacle comprising a low adhesion coating.
5. The cell population of one of claims 1-4, wherein the cell population is passaged at least one time in the anchorage independent culture.
6. The cell population of one of claims 1-5, wherein the population produces extracellular matrix.
7. The cell population of one of claims 1-6, wherein the population produces one or more cell surface markers selected from the group comprising CD24, CD34, CD44, CD73, CD90, CD105, CD166, Stro-1, HIF1, FIT-1, nestin, CK8, and HLA proteins.
8. The cell population of claim 7, wherein the percentage of the cells producing the cell surface marker(s) is greater than 70% or less than 40%.
9. The cell population of one of claims 1-6, wherein the population expresses one or more gene or gene products selected from the group comprising GAPDH, SDHA, HPRT1, B2M, Sox9, Aggrecan, Col1, Col2, nestin, CK8, Sox1, CD44, ALPI, and PPARG.
10. The cell population of one of claims 1-9, wherein the cell population is obtained from intervertebral disc tissue.
11. A method of deriving a discogenic cell population comprising:
isolating one or more cells from tissue;

- passaging the one or more cells in an anchorage dependent culture media;
transferring the one or more cells to an anchorage independent culture media.
12. A method of using a discogenic cell to treat at least one disc in a subject in need thereof comprising:
administering a therapeutic amount of a discogenic cell population to the subject, thereby
treating the subject.
 13. A method of treating a subject having at least one diseased, damaged, or missing intervertebral disc, comprising:
administering to the subject a discogenic cell population, in an amount effective to treat the disease or damage.
 14. A method of treating an indication selected from the group consisting of degenerative disc disease, herniated disc, injured disc, comprising:
administering a therapeutic amount of a discogenic cell population, thereby treating the indication.
 15. The method of claim 11, wherein the tissue is mammalian disc tissue.
 16. The method of claim 15, wherein the tissue is donated organ tissue.
 17. The method of one of claims 11-14, wherein the cell population is passaged at least one time in the anchorage independent culture.
 18. The method of one of claims 11-15, wherein the population of cells produces extracellular matrix.
 19. The method of one of claims 11-16, wherein the population of cells produces one or more cell surface markers selected from the group comprising CD24, CD34, CD44, CD73, CD90, CD105, CD166, Stro-1, HIF1, nestin, CK8, and HLA proteins.
 20. The method of one of claims 11-17, wherein the percentage of the cells in the population producing a cell surface marker(s) is greater than 70% or less than 40%.
 21. The method of one of claims 11-16, wherein the population expresses one or more gene or gene products selected from the group comprising GAPDH, SDHA,

HPRT1, B2M, Sox9, Aggrecan, Col1, Col2, nestin, CK8, Sox1, CD44, ALPI, PPARG, ADAMTS, MMP, FMOD, IL.

22. A device for treating a diseased or injured intervertebral disc comprising:
a discogenic cell population derived from cartilaginous tissue, wherein at least one cell has been grown in anchorage independent culture; and
a scaffold, matrix, or implantable structure.
23. The device of claim 22, wherein the device further comprises a biological active agent.
24. An artificial disc replacement device comprising:
an artificial outer annulus
a discogenic cell population.
25. The artificial disc replacement device of claim 24, wherein the outer annulus is comprised of a non-resorbable material.
26. The artificial disc replacement device of claim 25, wherein the non-resorbable material is polyurethane.
27. The artificial disc replacement device of claim 24, wherein the outer annulus is comprised of a resorbable material.
28. The artificial disc replacement device of claim 27, wherein the resorbable material is polyglycolic acid or polylactic acid, or a combination thereof.
29. The artificial disc replacement device of claim 24, wherein the discogenic population of cells further comprises one or more of the following a scaffold material, a matrix material, a carrier material, a growth factor(s), and other biologically active agents.
30. The artificial disc replacement device of claim 24, further comprising an attachment means for fixedly securing the device to one or more vertebral bodies.
31. The artificial disc replacement device of claim 30, further comprising through-holes, cuffs, tabs, loops, or washers to allow for screw fixation to one or more vertebral bodies.
32. A method of replacing a spinal disc comprising:
producing the artificial disc of claim 24 in vitro; and

surgically implanting the artificial disc into a subject, thereby replacing a spinal disc.