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(54) Title: CONDITIONED MEDIUM AND EXTRACELLULAR MATRIX COMPOSITIONS AND USES THEREOF

(57) Abstract: The present invention is directed to a methods for the stimulation of hair, lash and/or nail growth by the administration of extracellular matrix (ECM) and/or conditioned culture media (CCM) compositions in combination with active agents such as minoxidil and bimatoprost. The combination of hypoxic CCM and an active agent acts to promote hair, lash and/or nail growth and/or promote hair follicle development and/or activate or stimulate on an area of the skin when administered to the region of skin or tissue in need of growth or repair in a subject.



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CONDITIONED MEDIUM AND EXTRACELLULAR MATRIX COMPOSITIONS AND USES THEREOF

CROSS-REFERENCE TO RELATED APPLICATIONS

[001] This application claims benefit of priority under 35 U.S.C. §119(e) of U.S. Serial No. 62/713,984, filed August 2, 2018, the entire contents of which is incorporated herein by reference in its entirety.

BACKGROUND OF THE INVENTION

FIELD OF THE INVENTION

[002] The present invention relates generally to the production and use of growth factors and/or conditioned culture medium compositions and more specifically to compositions for hair, lash and/or nail growth.

BACKGROUND INFORMATION

[003] The extracellular matrix (ECM) is a complex structural entity surrounding and supporting cells that are found *in vivo* within mammalian tissues. The ECM is often referred to as the connective tissue. The ECM is primarily composed of three major classes of biomolecules including structural proteins such as collagens and elastins, specialized proteins such as fibrillins, fibronectins, and laminins, and proteoglycans. Conditioned culture medium (CCM) contains biologically active components obtained from previously cultured cells or tissues that have released into the media substances affecting certain cell function. It has been found that ECM and CCM compositions derived *in vitro* from cells grown under hypoxic or normoxic conditions have therapeutic properties beneficial for treating certain conditions.

[004] Growth of ECM compositions *in vitro* and their use in a variety of therapeutic and medical applications have been described in the art. One therapeutic application of such ECM compositions includes treatment and repair of soft tissue and skin defects such as wrinkles and scars.

[005] The repair or augmentation of soft tissue defects caused by defects, such as, acne, surgical scarring or aging has proven to be very difficult. A number of materials have been used to correct soft tissue defects with varying degrees of success, however, no material has been completely safe and effective. For example, silicon causes a variety of physiological and clinical problems including long term side effects, such as nodules, recurring cellulitis and skin ulcers.

[006] Collagen compositions have also been used as an injectable material for soft tissue augmentation. Collagen is the main protein of connective tissue and the most abundant protein in mammals, making up about 25% of the total protein content. There are currently 28 types of collagen described in literature (see, *e.g.*, Tables 1 and 2 *infra*, for a detailed listing). However, over 90% of the collagen in the body are Collagens I, II, III, and IV.

[007] Different collagen materials have been used for treatment of soft tissue defects, such as reconstituted injectable bovine collagen, crosslinked collagen, or other xenogeneic collagens. However, several problems exist with such collagens. A common problem includes the complexity and high cost of producing the implant materials to remove potentially immunogenic substances to avoid allergic reactions in the subject. Additionally, treatments using such collagens have not proven long lasting.

[008] Other materials have also been described that may be used for soft tissue repair or augmentation, such as, biocompatible ceramic particles in aqueous gels (U.S. Pat. No. 5,204,382), thermoplastic and/or thermosetting materials (U.S. Pat. No. 5,278,202), and lactic acid based polymer blends (U.S. Pat. No. 4,235,312). Additionally, use of naturally secreted ECM compositions have also been described (U.S. Pat. No. 6,284,284). However, such materials have all proven to have limitations.

[009] *In vitro* cultured ECM compositions can additionally be used to repair and/or regenerate damaged cells or tissue, such as chondral or osteochondral cells. Osteochondral tissue is any tissue that relates to or contains bone or cartilage. The compositions of the present invention are useful for treatment of osteochondral defects, such as degenerative connective tissue diseases, such as rheumatoid and/or osteoarthritis as well as defects in patients who have cartilage defects due to trauma.

[0010] *In vitro* cultured ECM compositions are also useful in tissue culture systems for generation of engineered tissue implants. The field of tissue engineering involves the use of cell culture technology to generate new biological tissues or repair damaged tissues. Fueled in part, by the stem cell revolution, tissue engineering technology offers the promise of tissue regeneration and replacement following trauma or treatment of degenerative diseases. It can also be used in the context of cosmetic procedures.

[0011] Culture medium compositions typically include essential amino acids, salts, vitamins, minerals, trace metals, sugars, lipids and nucleosides. Cell culture medium attempts to supply the components necessary to meet the nutritional needs required to grow cells in a controlled, artificial and *in vitro* environment. Nutrient formulations, pH, and osmolarity vary

in accordance with parameters such as cell type, cell density, and the culture system employed. Many cell culture medium formulations are documented in the literature and a number of media are commercially available. Once the culture medium is incubated with cells, it is known to those skilled in the art as "spent" or "conditioned medium". Conditioned medium contains many of the original components of the medium, as well as a variety of cellular metabolites and secreted proteins, including, for example, biologically active growth factors, inflammatory mediators and other extracellular proteins. Cell lines grown as a monolayer or on beads, as opposed to cells grown in three-dimensions, lack the cell-cell and cell-matrix interactions characteristic of whole tissue *in vivo*. Consequently, such cells secrete a variety of cellular metabolites although they do not necessarily secrete these metabolites and secreted proteins at levels that approach physiological levels. Conventional conditioned cell culture medium, medium cultured by cell-lines grown as a monolayer or on beads, is usually discarded or occasionally used in culture manipulations such as reducing cell densities.

[0012] The majority of vertebrate cell cultures *in vitro* are grown as monolayers on an artificial substrate bathed in culture medium. The nature of the substrate on which the monolayers grow may be solid, such as plastic, or semisolid gels, such as collagen or agar. Disposable plastics have become the preferred substrate used in modern-day tissue or cell culture.

[0013] A few researchers have explored the use of natural substrates related to basement membrane components. Basement membranes comprise a mixture of glycoproteins and proteoglycans that surround most cells *in vivo*. For example, Reid and Rojkund, 1979, In, *Methods in Enzymology*, Vol. 57, Cell Culture, Jakoby & Pasten, eds., New York, Acad. Press, pp. 263 278; Vlodavsky et al., 1980, *Cell* 19:607 617; Yang et al., 1979, *Proc. Natl. Acad. Sci. USA* 76:3401 have used collagen for culturing hepatocytes, epithelial cells and endothelial tissue. Growth of cells on floating collagen (Michalopoulos and Pitot, 1975, *Fed. Proc.* 34:826) and cellulose nitrate membranes (Savage and Bonney, 1978, *Exp. Cell Res.* 114:307 315) have been used in attempts to promote terminal differentiation. However, prolonged cellular regeneration and the culture of such tissues in such systems has not heretofore been achieved.

[0014] Cultures of mouse embryo fibroblasts have been used to enhance growth of cells, particularly at low densities. This effect is thought to be due partly to supplementation of the medium but may also be due to conditioning of the substrate by cell products. In these

systems, feeder layers of fibroblasts are grown as confluent monolayers which make the surface suitable for attachment of other cells. For example, the growth of glioma on confluent feeder layers of normal fetal intestine has been reported (Lindsay, 1979, *Nature* 228:80).

[0015] While the growth of cells in two dimensions is a convenient method for preparing, observing and studying cells in culture, allowing a high rate of cell proliferation, it lacks characteristic of whole tissue *in vivo*. In order to study such functional and morphological interactions, a few investigators have explored the use of three-dimensional substrates such as collagen gel (Douglas et al., 1980, *In Vitro* 16:306 312; Yang et al., 1979, *Proc. Natl. Acad. Sci.* 76:3401; Yang et al., 1980, *Proc. Natl. Acad. Sci.* 77:2088 2092; Yang et al., 1981, *Cancer Res.* 41:1021 1027); cellulose sponge, alone (Leighton et al., 1951, *J. Natl. Cancer Inst.* 12:545 561) or collagen coated (Leighton et al., 1968, *Cancer Res.* 28:286 296); a gelatin sponge, Gelfoam (Sorour et al., 1975, *J. Neurosurg.* 43:742 749).

[0016] In general, these three-dimensional substrates are inoculated with the cells to be cultured. Many of the cell types have been reported to penetrate the matrix and establish a "tissue-like" histology. For example, three-dimensional collagen gels have been utilized to culture breast epithelium (Yang et al., 1981, *Cancer Res.* 41:1021 1027) and sympathetic neurons (Ebendal, 1976, *Exp. Cell Res.* 98:159 169). Additionally, various attempts have been made to regenerate tissue-like architecture from dispersed monolayer cultures. (Kruse and Miedema, 1965, *J. Cell Biol.* 27:273) reported that perfused monolayers could grow to more than ten cells deep and organoid structures can develop in multilayered cultures if kept supplied with appropriate medium (see also Schneider et al., 1963, *Exp. Cell. Res.* 30:449 459; Bell et al., 1979, *Proc. Natl. Acad. Sci. USA* 76:1274 1279; Green, 1978, *Science* 200:1385 1388). It has been reported that human epidermal keratinocytes may form dermatoglyphs (friction ridges if kept for several weeks without transfer; Folkman and Haudenschild (1980, *Nature* 288:551 556) reported the formation of capillary tubules in cultures of vascular endothelial cells cultured in the presence of endothelial growth factor and medium conditioned by tumor cells; and Sirica et al. (1979, *Proc. Natl. Acad. Sci. USA* 76:283 287; 1980, *Cancer Res.* 40:3259 3267) maintained hepatocytes in primary culture for about 10 13 days on nylon meshes coated with a thin layer of collagen.

[0017] Synthetic matrices composed of biodegradable, biocompatible copolymers of polyesters and amino acids have also been designed as scaffolding for cell growth (U.S. Pat. Nos. 5,654,381; 5,709,854). Non-biodegradable scaffolds are likewise capable of supporting cell growth. Three-dimensional cell culture systems have also been designed which are

composed of a stromal matrix which supports the growth of cells from any desired tissue into an adult tissue (Naughton et al., U.S. Pat. Nos. 4,721,096 and 5,032,508). Another approach involves slowly polymerizing hydrogels containing large numbers of the desired cell type which harden into a matrix once administered to a patient (U.S. Pat. No. 5,709,854). Extracellular matrix preparations have been designed which are composed of stromal cells which provide a three dimensional cell culture system for a desired cell type which may be injected into the patient for precise placement of the biomaterial (Naughton et al., WO 96/39101).

[0018] The secretion of extracellular proteins into conditioned cell media such as growth factors, cytokines, and stress proteins opens new possibilities in the preparation of products for use in a large variety of areas including tissue repair, e.g., in the treatment of wounds and other tissue defects such as cosmetic defects as well as human and animal feed supplements. For example, growth factors are known to play an important role in the wound healing process. In general, it is thought desirable in the treatment of wounds to enhance the supply of growth factors by direct addition of these factors.

[0019] Cellular cytokines and growth factors are involved in a number of critical cellular processes including cell proliferation, adhesion, morphologic appearance, differentiation, migration, inflammatory responses, angiogenesis, and cell death. Studies have demonstrated that hypoxic stress and injury to cells induce responses including increased levels of mRNA and proteins corresponding to growth factors such as PDGF (platelet-derived growth factor), VEGF (vascular endothelial growth factor), FGF (fibroblast growth factor), and IGF (insulin-like growth factor) (Gonzalez-Rubio, M. et al., 1996, *Kidney Int.* 50(1):164 73; Abramovitch, R. et al., 1997, *Int J. Exp. Pathol.* 78(2):57 70; Stein, I. et al., 1995, *Mol Cell Biol.* 15(10):5363 8; Yang, W. et al., 1997, *FEBS Lett.* 403(2):139 42; West, N. R. et al., 1995, *J. Neurosci. Res.* 40(5):647 59).

[0020] Growth factors, such as transforming growth factor- β , also known in the art as TGF- β , are induced by certain stress proteins during wound healing. Two known stress proteins are GRP78 and HSP90. These proteins stabilize cellular structures and render the cells resistant to adverse conditions. The TGF- β family of dimeric proteins includes TGF- β .1, TGF- β .2, and TGF- β .3 and regulates the growth and differentiation of many cell types. Furthermore, this family of proteins exhibits a range of biological effects, stimulating the growth of some cell types (Noda et al., 1989, *Endocrinology* 124:2991 2995) and inhibiting the growth of other cell types (Goey et al., 1989, *J. Immunol.* 143:877 880;

Pietenpol et al., 1990, Proc. Natl. Acad. Sci. USA 87:3758 3762). TGF- β has also been shown to increase the expression of extracellular matrix proteins including collagen and fibronectin (Ignatz et al., 1986, J. Biol. Chem. 261:4337 4345) and to accelerate the healing of wounds (Mustoe et al., 1987, Science 237:1333 1335).

[0021] Another such growth factor is PDGF. PDGF was originally found to be a potent mitogen for mesenchymal-derived cells (Ross R. et al., 1974, Proc. Natl. Acad. Sci. USA 71(4):1207 1210; Kohler N. et al., 1974, Exp. Cell Res. 87:297 301). Further studies have shown that PDGF increases the rate of cellularity and granulation in tissue formation. Wounds treated with PDGF have the appearance of an early stage inflammatory response including an increase in neutrophils and macrophage cell types at the wound site. These wounds also show enhanced fibroblast function (Pierce, G. F. et al., 1988, J. Exp. Med. 167:974 987). Both PDGF and TGF- β have been shown to increase collagen formation, DNA content, and protein levels in animal studies (Grotendorst, G. R. et al., 1985, J. Clin. Invest. 76:2323 2329; Sporn, M. B. et al., 1983, Science (Wash D.C.) 219:1329). PDGF has been shown to be effective in the treatment of human wounds. In human wounds, PDGF-AA expression is increased within pressure ulcers undergoing healing. The increase of PDGF-AA corresponds to an increase in activated fibroblasts, extracellular matrix deposition, and active vascularization of the wound. Furthermore, such an increase in PDGF-AA is not seen in chronic non-healing wounds (Principles of Tissue Engineering, R. Lanza et al. (eds.), pp. 133 141 (R.G. Landes Co. TX 1997). A number of other growth factors having the ability to induce angiogenesis and wound healing include VEGF, KGF and basic FGF.

[0022] Androgenic alopecia (AGA) is characterized by hereditary thinning of the hair induced by androgens in genetically susceptible men and women. This condition is also known as male pattern hair loss or common baldness in men and as female pattern hair loss in women. Drug therapies specifically approved for treating AGA are limited to minoxidil and finasteride as major category products. Several other drugs are also used off label and a plethora of treatments with unsubstantiated hair growth claims can be obtained, however, looking at the number of treatment options currently available to patients with AGA, though the clinical data supporting their use is often very limited.

[0023] Minoxidil (i.e., 2,4-diamino-6-piperidinylpyrimidine-3 -oxide) is the active ingredient of the brand Rogaine® (in USA and Canada) and Regaine® (in Europe and Asia Pacific) as a treatment and prevention for androgenic alopecia (male and female pattern baldness) available as 5% minoxidil solution designed for men and 2% solutions designed for

women. The preparation of minoxidil is described in U.S. Pat. No. 3,461,461. Methods and topical preparations for using the compound to grow hair and to treat male and female pattern baldness are described and claimed in U.S. Pat. Nos. 4,139,619 and 4,596,812.

[0024] Pharmaceutical compositions for topical application may take a variety of forms including, for example, solutions, gels, suspensions, and the like. Generally speaking, improved absorption may be achieved when the topical compositions are in the form of a solution or gel, i.e., where the active ingredient, for example, minoxidil, is dissolved in the carrier solution, in contrast to topical compositions which are in the form of suspensions, i.e., where the active ingredient is merely suspended in the composition.

[0025] Bimatoprost (i.e. Latisse®) is a prostaglandin analog used topically (as eye drops) to control the progression of glaucoma and in the management of ocular hypertension. Bimatoprost is a structural analog of prostaglandin F_{2α} (PGF_{2α}). Like other PGF_{2α} analogs such as travoprost, latanoprost and tafluprost, it increases the outflow of aqueous fluid from the eye and lowers intraocular pressure. However, in contrast to these it does not act on the prostaglandin F receptor, nor on any other known prostaglandin receptor.

[0026] Eyelash growth cycle consists of three phases: anagen, catagen, and telogen. During the growth phase, i.e. anagen, only about 40% of lashes are actively growing. This is followed by a 2-3 week transitional phase, i.e. catagen, during which the lashes stop growing and the follicle shrinks. The final phase, i.e. telogen. Is a resting phase which lasts over 100 days before the lash falls out. This cycle indicates that at any one time only 40% of lashes will be growing and that there is a long period of time between growth phases. Cosmetically, it is desired to increase the growth rate of lashes for a more aesthetically pleasing appearance. The cosmetics industry has responded to this desire by marketing products designed to increase the growth of eyelashes. As one example, Bimatoprost has been shown to increase the growth of eye lashes when applied in a gel suspension at the base of the upper eyelid lashes.

[0027] Brittle nails are a common problem seen by dermatologists. Brittle nails include nails that are splitting; brittle; soft or thin; peeling at nail tips; easily broke, cracked or chipped and nails that are difficult to grow longer. As of now, as long as there is no underlying medical condition, brittle nails are treated by reducing exposure to water and irritants and moisturizing the nails and hands more frequently. Recently however, Bimatoprost has been shown to have adnexal activity and was observed to increase nail growth.

SUMMARY OF THE INVENTION

[0028] The present invention is based in part on the seminal discovery that cells cultured on surfaces (*e.g.*, in monolayers or layers on one-dimensional surfaces; two-dimensional or three-dimensional surfaces) produce ECM compositions and CCM compositions. The ECM and CCM compositions produced by culturing cells under normal, or normoxic, or under hypoxic conditions containing one or more embryonic proteins have a variety of beneficial applications, including stimulating the growth of hair, nail or lash growth.

[0029] In one embodiment, the present invention provides for the topical application of a composition containing growth factors and optionally vitamins, *e.g.*, Vitamin D, niotin, biotin, Vitamin A, Vitamin C, B Vitamins including Vitamin B12 or other supplements, or CCM, optionally along with penetration enhancers to the cuticle to stimulate stronger nail growth and structure, particularly for people with brittle nails.

[0030] In an additional embodiment, the present invention provides for the administration of a combination of a growth factor composition or CCM and minoxidil as a topical hair growth product. The growth factors and soluble matrix proteins in the growth factor composition or CCM enhance the hair growth and contribute to scalp health. The growth factors or CCM can be used alone or as a hair maintenance product after a hair transplant or other hair growth treatments (PRP, HSC, etc). Additionally, the combination of growth factors or CCM and minoxidil formulation can be utilized to enhance facial hair and eyebrows.

[0031] In a further embodiment, the present invention provides for the administration of a combination of growth factors or CCM and bimatoprost for the stimulation of eye lash growth.

[0032] In one embodiment, the present invention provides a method of making ECM or CCM compositions containing one or more proteins. The method includes culturing cells under normoxic or hypoxic conditions on a surface (*e.g.*, one-, two-dimensional or three-dimensional) in a suitable growth medium to produce a soluble and non-soluble fraction. In various aspects, the compositions include the soluble or non-soluble fraction separately, as well as combinations of the soluble and insoluble fraction. In various aspects, the compositions produced include upregulation of gene expression and production of laminins, collagens and Wnt factors. In other aspects the compositions produced include downregulation of gene expression of laminins, collagens and Wnt factors. In other aspects, the compositions are species specific and include cells and/or biological material from a

single animal species. While *in vitro* cultured ECM compositions are useful in the treatment of humans, such compositions may be applied to other species of animals. Accordingly, such compositions are well suited for veterinary applications.

[0033] In another embodiment, the compositions of the present invention can be used to provide a method of treating damaged tissue. The method includes contacting the damaged tissue with a composition generated by culturing cells under culture conditions on a one-dimensional, two-dimensional or three-dimensional surface containing one or more proteins under conditions that allow for treatment of the damaged tissue or treating keratinocytes, e.g., hair growth, lashes, nails.

[0034] In another embodiment, the present invention provides a method for stimulating or promoting hair growth. The method includes contacting a cell with the compositions described herein. In an exemplary aspect, the cell is a keratinocyte or hair follicle cell. In various aspects the cell may be contacted *in vivo* or *ex vivo*.

DETAILED DESCRIPTION OF THE INVENTION

[0035] The present invention relates to a method for making and using growth factor compositions, including but not limited to CCM. In particular the compositions are generated by culturing cells (e.g., fibroblasts) under culture conditions on a surface (e.g., one-dimensional, two-dimensional or three-dimensional) in a suitable growth medium. Culturing methods produce both ECM and CCM fractions may be used separately or in combination for a variety of applications.

[0036] The division, differentiation, and function of stem cells and multipotent progenitors are influenced by complex signals in the microenvironment, including oxygen availability. Regions of severe oxygen deprivation (hypoxia) arise in tumors for example due to rapid cell division and aberrant blood vessel formation. The hypoxia-inducible factors (HIFs) mediate transcriptional responses to localized hypoxia in normal tissues and in cancers and can promote tumor progression by altering cellular metabolism and stimulating angiogenesis. Recently, HIFs have been shown to activate specific signaling pathways such as Notch and the expression of transcription factors such as Oct4 that control stem cell self-renewal and multipotency. As many cancers are thought to develop from a small number of transformed, self-renewing, and multipotent "cancer stem cells," these results suggest new roles for HIFs in tumor progression. The data shown in the present examples indicate that the cells cultured under hypoxic conditions express genes typically associated with pluripotent cells, such as Oct4, NANOG, Sox2, KLF4 and cMyc, for example.

[0037] The compositions of the present invention have a variety of applications including, but not limited to, promoting repair and/or regeneration of damaged cells or tissues, use in patches and implants to promote tissue regeneration (*e.g.*, hernial repair, pelvic floor repair, rotator cuff repair, and wound repair), use in tissue culture systems for culturing cells, such as stem cells, use in surface coatings used in association with implantable devices (*e.g.*, pacemakers, stents, stent grafts, vascular prostheses, heart valves, shunts, drug delivery ports or catheters, hernial and pelvic floor repair patches), promoting soft tissue repair, augmentation, and/or improvement of a skin surface, such as wrinkles, post-traumatic skin applications (*e.g.*, post-laser), hair growth, use as a biological anti-adhesion agent, as a biological vehicle for cell delivery or maintenance at a site of delivery, stimulating hair, nail or lash growth and/or promoting hair follicle development and/or activation or stimulation on an area of the skin of a subject comprising contacting the hair (scalp), nail or lash or adjacent areas thereof.

[0038] The invention is based in part, on the discovery that cells cultured on beads or three-dimensional surfaces under conditions that stimulate the early embryonic environment (hypoxia and reduced gravitational forces) prior to angiogenesis produces ECM compositions with fetal properties, including generation of embryonic proteins. Growth of cells under hypoxic conditions demonstrate a unique ECM with fetal properties and growth factor expression and a unique CCM. Unlike the culturing of ECM under traditional culture conditions, over 5000 genes are differentially expressed in ECM cultured under hypoxic conditions. This results in a cultured ECM that has different properties and a different biological composition. For example, an ECM produced under hypoxic conditions is similar to fetal mesenchymal tissue in that it is relatively rich in collagens type III, IV, and V, and glycoproteins such as fibronectin, SPARC, thrombospondin, and hyaluronic acid.

[0039] Hypoxia also enhances expression of factors which regulate wound healing and organogenesis, such as VEGF, FGF-7, and TGF- β , as well as multiple Wnt factors including Wnts 2b, 4, 7a, 10a, and 11. Cultured embryonic human ECM also stimulates an increase of metabolic activity in human fibroblasts *in vitro*, as measured by increased enzymatic activity. Additionally, there is an increase in cell number in response to the cultured embryonic ECM.

[0040] Before the present compositions and methods are described, it is to be understood that this invention is not limited to particular compositions, methods, and experimental conditions described, as such compositions, methods, and conditions may vary. It is also to be understood that the terminology used herein is for purposes of describing particular

embodiments only, and is not intended to be limiting, since the scope of the present invention will be limited only in the appended claims.

[0041] As used in this specification and the appended claims, the singular forms “a”, “an”, and “the” include plural references unless the context clearly dictates otherwise. Thus, for example, references to “the method” includes one or more methods, and/or steps of the type described herein which will become apparent to those persons skilled in the art upon reading this disclosure and so forth.

[0042] In various embodiments, the present invention involves methods for making ECM compositions that include one or more embryonic proteins and applications thereof. In particular the compositions are generated by culturing cells under normoxic or hypoxic conditions on a one-dimensional, two-dimensional or three-dimensional surface in a suitable growth medium. The compositions are derived by growing cells on a three-dimensional framework resulting in a multi-layer cell culture system. Cells grown on a three-dimensional framework support, in accordance with the present invention, grow in multiple layers, forming a cellular matrix. Growth of the cultured cells under hypoxic conditions results in differential gene expression as the result of hypoxic culturing conditions versus traditional culture in the ECM and the conditioned medium.

[0043] ECM is a composition of proteins and biopolymers that substantially comprise tissue that is produced by cultivation of cells. Stromal cells, such as fibroblasts, are an anchorage dependent cell type requiring growth while attached to materials and surfaces suitable for cell culture. The ECM materials produced by the cultured cells are deposited in a three-dimensional arrangement providing spaces for the formation of tissue-like structures.

[0044] The cultivation materials providing three-dimensional architectures are referred to as scaffolds. Spaces for deposition of ECM are in the form of openings within, for example woven mesh or interstitial spaces created in a compacted configuration of spherical beads, called microcarriers.

[0045] The methods described herein provide both a non-soluble ECM composition and a soluble CCM composition. The non-soluble composition includes those secreted ECM proteins and biological components that are deposited on the support or scaffold. The soluble composition includes culture media or conditioned media in which cells have been cultured and into which the cells have secreted active agent(s) and includes those proteins and biological components not deposited on the scaffold. Both compositions may be collected,

and optionally further processed, and used individually or in combination in a variety of applications as described herein.

[0046] The three-dimensional support or scaffold used to culture stromal cells may be of any material and/or shape that: (a) allows cells to attach to it (or can be modified to allow cells to attach to it); and (b) allows cells to grow in more than one layer (*i.e.*, form a three dimensional tissue). In other embodiments, a substantially two-dimensional sheet or membrane or beads may be used to culture cells that are sufficiently three dimensional in form.

[0047] The biocompatible material is formed into a three-dimensional structure or scaffold, where the structure has interstitial spaces for attachment and growth of cells into a three dimensional tissue. The openings and/or interstitial spaces of the framework in some embodiments are of an appropriate size to allow the cells to stretch across the openings or spaces. Maintaining actively growing cells stretched across the framework appears to enhance production of the repertoire of growth factors responsible for the activities described herein. If the openings are too small, the cells may rapidly achieve confluence but be unable to easily exit from the mesh. These trapped cells may exhibit contact inhibition and cease production of the appropriate factors necessary to support proliferation and maintain long term cultures. If the openings are too large, the cells may be unable to stretch across the opening, which may lead to a decrease in stromal cell production of the appropriate factors necessary to support proliferation and maintain long term cultures. Typically, the interstitial spaces are at least about 100 μm , at least 140 μm , at least about 150 μm , at least about 180 μm , at least about 200 μm , or at least about 220 μm . When using a mesh type of matrix, as exemplified herein, we have found that openings ranging from about 100 μm to about 220 μm will work satisfactorily. However, depending upon the three-dimensional structure and intricacy of the framework, other sizes are permissible. Any shape or structure that allows the cells to stretch and continue to replicate and grow for lengthy time periods may function to elaborate the cellular factors in accordance with the methods herein.

[0048] In some aspects, the three dimensional framework is formed from polymers or threads that are braided, woven, knitted or otherwise arranged to form a framework, such as a mesh or fabric. The materials may also be formed by casting of the material or fabrication into a foam, matrix, or sponge-like scaffold. In other aspects, the three dimensional framework is in the form of matted fibers made by pressing polymers or other fibers together to generate a material with interstitial spaces. The three dimensional framework may take

any form or geometry for the growth of cells in culture. Thus, other forms of the framework, as further described below, may suffice for generating the appropriate conditioned medium.

[0049] A number of different materials may be used to form the scaffold or framework. These materials include non-polymeric and polymeric materials. Polymers, when used, may be any type of polymer, such as homopolymers, random polymers, copolymers, block polymers, coblock polymers (*e.g.*, di, tri, etc.), linear or branched polymers, and crosslinked or non-crosslinked polymers. Non-limiting examples of materials for use as scaffolds or frameworks include, among others, glass fibers, polyethylenes, polypropylenes, polyamides (*e.g.*, nylon), polyesters (*e.g.*, dacron), polystyrenes, polyacrylates, polyvinyl compounds (*e.g.*, polyvinylchloride; PVC), polycarbonates, polytetrafluoroethylenes (PTFE; TEFLON), thermanox (TPX), nitrocellulose, polysaccharides (*e.g.*, celluloses, chitosan, agarose), polypeptides (*e.g.*, silk, gelatin, collagen), polyglycolic acid (PGA), and dextran.

[0050] In some aspects, the framework or beads may be made of materials that degrade over time under the conditions of use. Biodegradable also refers to absorbability or degradation of a compound or composition when administered *in vivo* or under *in vitro* conditions. Biodegradation may occur through the action of biological agents, either directly or indirectly. Non-limiting examples of biodegradable materials include, among others, polylactide, polyglycolide, poly(trimethylene carbonate), poly(lactide-co-glycolide) (*i.e.*, PLGA), polyethylene terephthalate (PET), polycaprolactone, catgut suture material, collagen (*e.g.*, equine collagen foam), polylactic acid, or hyaluronic acid. For example, these materials may be woven into a three-dimensional framework such as a collagen sponge or collagen gel.

[0051] In other aspects, where the cultures are to be maintained for long periods of time, cryopreserved, and/or where additional structural integrity is desired, the three dimensional framework may be comprised of a nonbiodegradable material. As used herein, a nonbiodegradable material refers to a material that does not degrade or decompose significantly under the conditions in the culture medium. Exemplary nondegradable materials include, as non-limiting examples, nylon, dacron, polystyrene, polyacrylates, polyvinyls, polytetrafluoroethylenes (PTFE), expanded PTFE (ePTFE), and cellulose. An exemplary nondegrading three dimensional framework comprises a nylon mesh, available under the tradename Nitex®, a nylon filtration mesh having an average pore size of 140 µm and an average nylon fiber diameter of 90 µm (#3-210/36, Tetko, Inc., N.Y.).

[0052] In other aspects, the beads, scaffold or framework is a combination of biodegradable and non-biodegradable materials. The non-biodegradable material provides

stability to the three dimensional scaffold during culturing while the biodegradable material allows formation of interstitial spaces sufficient for generating cell networks that produce the cellular factors sufficient for therapeutic applications. The biodegradable material may be coated onto the non-biodegradable material or woven, braided or formed into a mesh. Various combinations of biodegradable and non-biodegradable materials may be used. An exemplary combination is poly(ethylene terephthalate) (PET) fabrics coated with a thin biodegradable polymer film, poly[D-L-lactic-co-glycolic acid], in order to obtain a polar structure.

[0053] In various aspects, the scaffold or framework material may be pre-treated prior to inoculation with cells to enhance cell attachment. For example, prior to inoculation with cells, nylon screens in some embodiments are treated with 0.1 M acetic acid, and incubated in polylysine, fetal bovine serum, and/or collagen to coat the nylon. Polystyrene could be similarly treated using sulfuric acid. In other embodiments, the growth of cells in the presence of the three-dimensional support framework may be further enhanced by adding to the framework or coating it with proteins (*e.g.*, collagens, elastin fibers, reticular fibers), glycoproteins, glycosaminoglycans (*e.g.*, heparan sulfate, chondroitin-4-sulfate, chondroitin-6-sulfate, dermatan sulfate, keratan sulfate, etc.), fibronectins, and/or glycopolymer (poly[N-vinylbenzyl-D-lactamide], PVLA) in order to improve cell attachment. Treatment of the scaffold or framework is useful where the material is a poor substrate for the attachment of cells.

[0054] In one aspect, mesh is used for production of ECM. The mesh is a woven nylon 6 material in a plain weave form with approximately 100 μm openings and approximately 125 μm thick. In culture, fibroblast cells attach to the nylon through charged protein interactions and grow into the voids of the mesh while producing and depositing ECM proteins. Mesh openings that are excessively large or small may not be effective but could differ from those above without substantially altering the ability to produce or deposit ECM. In another aspect, other woven materials are used for ECM production, such as polyolefin's, in weave configurations giving adequate geometry for cell growth and ECM deposition.

[0055] For example, nylon mesh is prepared for cultivation in any of the steps of the invention by cutting to the desired size, washing with 0.1 – 0.5M acetic acid followed by rinsing with high purity water and then steam sterilized. For use as a three-dimensional scaffold for ECM production the mesh is sized into squares approximately 10 cm x 10 cm. However, the mesh could be any size appropriate to the intended application and may be used

in any of the methods of the present invention, including cultivation methods for inoculation, cell growth and ECM production and preparation of the final form.

[0056] In other aspects, the scaffold for generating the cultured tissues is composed of microcarriers, which are beads or particles. The beads may be microscopic or macroscopic and may further be dimensioned so as to permit penetration into tissues or compacted to form a particular geometry. In some tissue penetrating embodiments, the framework for the cell cultures comprises particles that, in combination with the cells, form a three dimensional tissue. The cells attach to the particles and to each other to form a three dimensional tissue. The complex of the particles and cells is of sufficient size to be administered into tissues or organs, such as by injection or catheter. Beads or microcarriers are typically considered a two-dimensional system or scaffold.

[0057] As used herein, a “microcarriers” refers to a particle having size of nanometers to micrometers, where the particles may be any shape or geometry, being irregular, non-spherical, spherical, or ellipsoid.

[0058] The size of the microcarriers suitable for the purposes herein can be of any size suitable for the particular application. In some embodiments, the size of microcarriers suitable for the three dimensional tissues may be those administrable by injection. In some embodiments, the microcarriers have a particle size range of at least about 1 μm , at least about 10 μm , at least about 25 μm , at least about 50 μm , at least about 100 μm , at least about 200 μm , at least about 300 μm , at least about 400 μm , at least about 500 μm , at least about 600 μm , at least about 700 μm , at least about 800 μm , at least about 900 μm , at least about 1000 μm .

[0059] In some aspects in which the microcarriers are made of biodegradable materials. In some aspects, microcarriers comprising two or more layers of different biodegradable polymers may be used. In some embodiments, at least an outer first layer has biodegradable properties for forming the three dimensional tissues in culture, while at least a biodegradable inner second layer, with properties different from the first layer, is made to erode when administered into a tissue or organ.

[0060] In some aspects, the microcarriers are porous microcarriers. Porous microcarriers refer to microcarriers having interstices through which molecules may diffuse in or out from the microparticle. In other embodiments, the microcarriers are non-porous microcarriers. A nonporous microparticle refers to a microparticle in which molecules of a select size do not diffuse in or out of the microparticle.

[0061] Microcarriers for use in the compositions are biocompatible and have low or no toxicity to cells. Suitable microcarriers may be chosen depending on the tissue to be treated, type of damage to be treated, the length of treatment desired, longevity of the cell culture *in vivo*, and time required to form the three dimensional tissues. The microcarriers may comprise various polymers, natural or synthetic, charged (*i.e.*, anionic or cationic) or uncharged, biodegradable, or nonbiodegradable. The polymers may be homopolymers, random copolymers, block copolymers, graft copolymers, and branched polymers.

[0062] In some aspects, the microcarriers comprise non-biodegradable microcarriers. Non-biodegradable microcapsules and microcarriers include, but not limited to, those made of polysulfones, poly(acrylonitrile-co-vinyl chloride), ethylene-vinyl acetate, hydroxyethylmethacrylate-methyl-methacrylate copolymers. These are useful to provide tissue bulking properties or in embodiments where the microcarriers are eliminated by the body.

[0063] In some aspects, the microcarriers comprise degradable scaffolds. These include microcarriers made from naturally occurring polymers, non-limiting example of which include, among others, fibrin, casein, serum albumin, collagen, gelatin, lecithin, chitosan, alginate or poly-amino acids such as poly-lysine. In other aspects, the degradable microcarriers are made of synthetic polymers, non-limiting examples of which include, among others, polylactide (PLA), polyglycolide (PGA), poly(lactide-co-glycolide) (PLGA), poly(caprolactone), polydioxanone trimethylene carbonate, polyhydroxyalkonates (*e.g.*, poly(hydroxybutyrate), poly(ethyl glutamate), poly(DTH iminocarbony(bisphenol A iminocarbonate), poly(ortho ester), and polycyanoacrylates.

[0064] In some aspects, the microcarriers comprise hydrogels, which are typically hydrophilic polymer networks filled with water. Hydrogels have the advantage of selective trigger of polymer swelling. Depending on the composition of the polymer network, swelling of the microparticle may be triggered by a variety of stimuli, including pH, ionic strength, thermal, electrical, ultrasound, and enzyme activities. Non-limiting examples of polymers useful in hydrogel compositions include, among others, those formed from polymers of poly(lactide-co-glycolide); poly(N-isopropylacrylamide); poly(methacrylic acid-g-polyethylene glycol); polyacrylic acid and poly(oxypropylene-co-oxyethylene) glycol; and natural compounds such as chondroitin sulfate, chitosan, gelatin, fibrinogen, or mixtures of synthetic and natural polymers, for example chitosan-poly (ethylene oxide). The polymers

may be crosslinked reversibly or irreversibly to form gels adaptable for forming three dimensional tissues.

[0065] In exemplary aspects, the microcarriers or beads for use in the present invention are composed wholly or composed partly of dextran.

[0066] In accordance with the present invention the culturing method is applicable to proliferation of different types of cells, including stromal cells, such as fibroblasts, and particularly primary human neonatal foreskin fibroblasts. In various aspects, the cells inoculated onto the scaffold or framework can be stromal cells comprising fibroblasts, with or without other cells, as further described below. In some embodiments, the cells are stromal cells that are typically derived from connective tissue, including, but not limited to: (1) bone; (2) loose connective tissue, including collagen and elastin; (3) the fibrous connective tissue that forms ligaments and tendons, (4) cartilage; (5) the ECM of blood; (6) adipose tissue, which comprises adipocytes; and (7) fibroblasts.

[0067] Stromal cells can be derived from various tissues or organs, such as skin, heart, blood vessels, bone marrow, skeletal muscle, liver, pancreas, brain, foreskin, which can be obtained by biopsy (where appropriate) or upon autopsy. In one aspect, fetal fibroblasts can be obtained in high quantity from foreskin, such as neonatal foreskins.

[0068] In some aspects, the cells comprise fibroblasts, which can be from a fetal, neonatal, adult origin, or a combination thereof. In some aspects, the stromal cells comprise fetal fibroblasts, which can support the growth of a variety of different cells and/or tissues. As used herein, a fetal fibroblast refers to fibroblasts derived from fetal sources. As used herein, neonatal fibroblast refers to fibroblasts derived from newborn sources. Under appropriate conditions, fibroblasts can give rise to other cells, such as bone cells, fat cells, and smooth muscle cells and other cells of mesodermal origin. In some embodiments, the fibroblasts comprise dermal fibroblasts, which are fibroblasts derived from skin. Normal human dermal fibroblasts can be isolated from neonatal foreskin. These cells are typically cryopreserved at the end of the primary culture.

[0069] In other aspects, the three-dimensional tissue can be made using stem or progenitor cells, either alone, or in combination with any of the cell types discussed herein. Stem and progenitor cells include, by way of example and not limitation, embryonic stem cells, hematopoietic stem cells, neuronal stem cells, epidermal stem cells, and mesenchymal stem cells.

[0070] In some embodiments, a “specific” three-dimensional tissue can be prepared by inoculating the three-dimensional scaffold with cells derived from a particular organ, *i.e.*, skin, heart, and/or from a particular individual who is later to receive the cells and/or tissues grown in culture in accordance with the methods described herein.

[0071] For certain uses *in vivo* it is preferable to obtain the stromal cells from the patient's own tissues. The growth of cells in the presence of the three-dimensional stromal support framework can be further enhanced by adding to the framework, or coating the framework support with proteins, *e.g.*, collagens, laminins, elastic fibers, reticular fibers, glycoproteins; glycosaminoglycans, *e.g.*, heparin sulfate, chondroitin-4-sulfate, chondroitin-6-sulfate, dermatan sulfate, keratan sulfate, etc.; a cellular matrix, and/or other materials.

[0072] Thus, since the two-dimensional or three-dimensional culture systems described herein are suitable for growth of diverse cell types and tissues, and depending upon the tissue to be cultured and the collagen types desired, the appropriate stromal cells may be selected to inoculate the framework.

[0073] While the methods and applications of the present invention are suitable for use with different cell types, such as tissue specific cells or different types of stromal cells as discussed herein, derivation of the cells for use with the present invention may also be species specific. Accordingly, ECM compositions may be generated that are species specific. For example, the cells for use in the present invention may include human cells. For example, the cells may be human fibroblasts. Likewise, the cells are from another species of animal, such as equine (horse), canine (dog) or feline (cat) cells. Additionally, cells from one species or strain of species may be used to generate ECM compositions for use in other species or related strains (*e.g.*, allogeneic, syngeneic and xenogeneic). It is also to be appreciated that cells derived from various species may be combined to generate multi-species ECM compositions.

[0074] Accordingly, the methods and compositions of the present invention are suitable in applications involving non-human animals. As used herein, “veterinary” refers to the medical science concerned or connected with the medical or surgical treatment of animals, especially domestic animals. Common veterinary animals may include mammals, amphibians, avians, reptiles and fishes. For example, typical mammals may include dogs, cats, horses, rabbits, primates, rodents, and farm animals, such as cows, horses, goats, sheep, and pigs.

[0075] As discussed above, additional cells may be present in the culture with the stromal cells. These additional cells may have a number of beneficial effects, including, among others, supporting long term growth in culture, enhancing synthesis of growth factors, and promoting attachment of cells to the scaffold. Additional cell types include as non-limiting examples, smooth muscle cells, cardiac muscle cells, endothelial cells, skeletal muscle cells, endothelial cells, pericytes, macrophages, monocytes, and adipocytes. Such cells may be inoculated onto the framework along with fibroblasts, or in some aspects, in the absence of fibroblasts. These stromal cells may be derived from appropriate tissues or organs, including, by way of example and not limitation, skin, heart, blood vessels, bone marrow, skeletal muscle, liver, pancreas, and brain. In other aspects, one or more other cell types, excluding fibroblasts, are inoculated onto the scaffold. In still other aspects, the scaffolds are inoculated only with fibroblast cells.

[0076] Fibroblasts may be readily isolated by disaggregating an appropriate organ or tissue which is to serve as the source of the fibroblasts. For example, the tissue or organ can be disaggregated mechanically and/or treated with digestive enzymes and/or chelating agents that weaken the connections between neighboring cells making it possible to disperse the tissue into a suspension of individual cells without appreciable cell breakage. Enzymatic dissociation can be accomplished by mincing the tissue and treating the minced tissue with any of a number of digestive enzymes either alone or in combination. These include but are not limited to trypsin, chymotrypsin, collagenase, elastase, hyaluronidase, DNase, pronase, and/or dispase, etc. Mechanical disruption can also be accomplished by a number of methods including, but not limited to the use of grinders, blenders, sieves, homogenizers, pressure cells, or insonators to name but a few. In one aspect, excised foreskin tissue is treated using digestive enzymes, typically collagenase and/or trypsinase to disassociate the cells from encapsulating structures.

[0077] The isolation of fibroblasts, for example, can be carried out as follows: fresh tissue samples are thoroughly washed and minced in Hanks' balanced salt solution (HBSS) in order to remove serum. The minced tissue is incubated from 1-12 hours in a freshly prepared solution of a dissociating enzyme such as trypsin. After such incubation, the dissociated cells are suspended, pelleted by centrifugation and plated onto culture dishes. All fibroblasts will attach before other cells, therefore, appropriate stromal cells can be selectively isolated and grown. The isolated fibroblasts can then be grown to confluency, lifted from the confluent culture and inoculated onto the three-dimensional framework, see Naughton et al., 1987, J.

Med. 18(3&4):219-250. Inoculation of the three-dimensional framework with a high concentration of stromal cells, *e.g.*, approximately 10^6 to 5×10^7 cells/ml, will result in the establishment of the three-dimensional stromal support in shorter periods of time.

[0078] Once the tissue has been reduced to a suspension of individual cells, the suspension can be fractionated into subpopulations from which the fibroblasts and/or other stromal cells and/or elements can be obtained. This also may be accomplished using standard techniques for cell separation including, but not limited to, cloning and selection of specific cell types, selective destruction of unwanted cells (negative selection), separation based upon differential cell agglutinability in the mixed population, freeze-thaw procedures, differential adherence properties of the cells in the mixed population, filtration, conventional and zonal centrifugation, centrifugal elutriation (counter-streaming centrifugation), unit gravity separation, countercurrent distribution, electrophoresis and fluorescence-activated cell sorting. For a review of clonal selection and cell separation techniques, see Freshney, *Culture of Animal Cells: A Manual of Basic Techniques*, 2d Ed., A. R. Liss, Inc., New York, 1987, Ch. 11 and 12, pp. 137-168.

[0079] In one aspect, isolated fibroblast cells can be grown to produce cell banks. Cell banks are created to allow for initiating various quantities and timing of cultivation batches and to allow preemptive testing of cells for contaminants and specific cellular characteristics. Fibroblasts from the cell banks are subsequently grown to increase cell number to appropriate levels for seeding scaffolds. Operations involving environmental exposure of cells and cell contacting materials are performed by aseptic practices to reduce the potential for contamination of foreign materials or undesirable microbes.

[0080] In another aspect of the invention, after isolation, cells can be grown through several passages to a quantity suitable for building master cell banks. The cell banks can then be harvested and filled into appropriate vessels and preserved in cryogenic conditions. Cells in frozen vials from master cell banks can be thawed and grown through additional passages (usually two or more). The cells can then be used to prepare cryogenically preserved working cell banks.

[0081] A cell expansion step uses vials of cells at the working cell bank stage to further increase cell numbers for inoculating three-dimensional scaffolds or supports, such as mesh or microcarriers. Each passage is a series of sub-culture steps that include inoculating cell growth surfaces, incubation, feeding the cells and harvesting.

[0082] Cultivation for cell banks and cell expansion can be conducted by inoculating culture vessels, such as culture flasks, roller bottles or microcarriers. Stromal cells, such as fibroblasts, attach to the intended growth surfaces and grow in the presence of culture media. Culture vessels, such as culture flasks, roller bottles and microcarriers are specifically configured for cell culture and are commonly made from various plastic materials qualified for intended applications. Microcarriers typically are microscopic or macroscopic beads and are typically made of various plastic materials. However, they can be made from other materials such as glasses or solid/semi-solid biologically based materials such as collagens or other materials such as Dextran, a modified sugar complex as discussed above.

[0083] During cultivation, expended media is periodically replaced with fresh media during the course of cell growth to maintain adequate availability of nutrients and removal of inhibitory products of cultivation. Culture flasks and roller bottles provide a surface for the cells to grow onto and are typically used for cultivation of anchorage dependent cells.

[0084] In one aspect, incubation is performed in a chamber heated at 37°C. Cultivation topologies requiring communication of media and the chamber environment use a 5% CO₂ v/v with air in the chamber gas space to aid in regulation of pH. Alternately, vessels equipped to maintain cultivation temperature and pH can be used for both cell expansion and ECM production operations. Temperatures below 35°C or above 38°C and CO₂ concentrations below 3% or above 12% may not be appropriate.

[0085] Harvesting cells from attachment surfaces can be conducted by removal of growth media and rinsing the cells with a buffered salt solution to reduce enzyme competing protein, application of disassociating enzymes then neutralization of the enzymes after cell detachment. Harvested cell suspension is collected and harvest fluids are separated by centrifugation. Cell suspensions from sub-culture harvests can be sampled to assess the quantity of cells recovered and other cellular attributes and are subsequently combined with fresh media and applied as inoculums. The number of passages used for preparing cell banks and scaffold inoculum is critical with regard to achieving acceptable ECM characteristics.

[0086] After an appropriate three-dimensional scaffold is prepared, it is inoculated by seeding with the prepared stromal cells. Inoculation of the scaffold may be done in a variety of ways, such as sedimentation. Mesh prepared for culture of ECM under aerobic conditions are prepared in the same manner as for hypoxic grown mesh with the exception that an anaerobic chamber is not used to create hypoxic conditions.

[0087] For example, for both mesh prepared for culture of ECM under both aerobic and hypoxic conditions, prepared and sterilized mesh is placed in sterile 150 mm diameter x 15 mm deep petri dishes and stacked to a thickness of approximately 10 pieces. Stacks of mesh are then inoculated by sedimentation. Cells are added to fresh media to obtain the appropriate concentration of cells for inoculum. Inoculum is added to the stack of mesh where cells settle onto the nylon fibers and attach while in incubated conditions. After an adequate time, individually seeded mesh sheets can be aseptically separated from the stack and placed individually into separate 150 mm x 15 mm petri dishes containing approximately 50 ml of growth media.

[0088] Incubation of the inoculated culture is performed under hypoxic conditions, which is discovered to produce an ECM and CCM with unique properties as compared to ECM and CCM generated under normal culture conditions. As used herein, hypoxic conditions are characterized by a lower oxygen concentration as compared to the oxygen concentration of ambient air (approximately 15%-20% oxygen). In one aspect, hypoxic conditions are characterized by an oxygen concentration less than about 10%. In another aspect hypoxic conditions are characterized by an oxygen concentration of about 1% to 10%, 1% to 9%, 1% to 8%, 1% to 7%, 1% to 6%, 1% to 5%, 1% to 4%, 1% to 3%, or 1% to 2%. In a certain aspect, the system maintains about 1-3% oxygen within the culture vessel. Hypoxic conditions can be created and maintained by using a culture apparatus that allows one to control ambient gas concentrations, for example, an anaerobic chamber.

[0089] Incubation of cell cultures is typically performed in normal atmosphere with 15-20% oxygen and 5% CO₂ for expansion and seeding, at which point low oxygen cultures are split to an airtight chamber that is flooded with 95% nitrogen/ 5% CO₂ so that a hypoxic environment is created within the culture medium.

[0090] For example, petri dishes with mesh cultured for producing ECM under hypoxic conditions are initially grown in incubation at 37°C and 95% air/5% CO₂ for 2-3 weeks. Following the period of near atmospheric cultivation, the petri dishes of mesh are incubated in a chamber designed for anaerobic cultivation that is purged with a gas mixture of approximately 95% nitrogen and 5% CO₂. Expended growth media is replaced with fresh media at atmospheric oxygen level through the culture period and after media is exchanged the mesh filled petri dishes are placed in the anaerobic chamber, the chamber is purged with 95% nitrogen/5% CO₂ then incubated at 37°C. Cultured mesh are harvested when they reach the desired size or contain the desired biological components.

[0091] During the incubation period, the stromal cells will grow linearly along and envelop the three-dimensional framework before beginning to grow into the openings of the framework. The growing cells produce a myriad of growth factors, regulatory factors and proteins, some of which are secreted in the surrounding media, and others that are deposited on the support to make up the ECM more fully discussed below. Growth and regulatory factors can be added to the culture, but are not necessary. Culture of the stromal cells produces both non-soluble and soluble fractions. The cells are grown to an appropriate degree to allow for adequate deposition of ECM proteins.

[0092] During culturing of the three-dimensional tissues, proliferating cells may be released from the framework and stick to the walls of the culture vessel where they may continue to proliferate and form a confluent monolayer. To minimize this occurrence, which may affect the growth of cells, released cells may be removed during feeding or by transferring the three-dimensional cell culture to a new culture vessel. Removal of the confluent monolayer or transfer of the cultured tissue to fresh media in a new vessel maintains or restores proliferative activity of the three-dimensional cultures. In some aspects, removal or transfers may be done in a culture vessel which has a monolayer of cultured cells exceeding 25% confluency. Alternatively, the culture in some embodiments is agitated to prevent the released cells from sticking; in others, fresh media is infused continuously through the system. In some aspects, two or more cell types can be cultured together either at the same time or one first followed by the second (*e.g.*, fibroblasts and smooth muscle cells or endothelial cells).

[0093] After inoculation of the three dimensional scaffolds, the cell culture is incubated in an appropriate nutrient medium and incubation conditions that supports growth of cells into the three dimensional tissues. Many commercially available media such as Dulbecco's Modified Eagles Medium (DMEM), RPMI 1640, Fisher's, Iscove's, and McCoy's, may be suitable for supporting the growth of the cell cultures. The medium may be supplemented with additional salts, carbon sources, amino acids, serum and serum components, vitamins, minerals, reducing agents, buffering agents, lipids, nucleosides, antibiotics, attachment factors, and growth factors. Formulations for different types of culture media are described in various reference works available to the skilled artisan (*e.g.*, *Methods for Preparation of Media, Supplements and Substrates for Serum Free Animal Cell Cultures*, Alan R. Liss, New York (1984); *Tissue Culture: Laboratory Procedures*, John Wiley & Sons, Chichester,

England (1996); Culture of Animal Cells, A Manual of Basic Techniques, 4 th Ed., Wiley-Liss (2000)).

[0094] The growth or culture media used in any of the culturing steps of the present invention, whether under aerobic or hypoxic conditions, may include serum, or be serum free. In one aspect, the media is Dulbecco's Modified Eagle Medium with 4.5 g/L glucose, alanyl-L-glutamine, Eq 2mM, and nominally supplemented with 10% fetal bovine serum. In another aspect, the media is a serum free media and is Dulbecco's Modified Eagle Medium with 4.5 g/L glucose base medium with Glutamax®, supplemented with 0.5% serum albumin, 2 µg/ml heparin, 1 µg/ml recombinant basic FGF, 1 µg/ml soybean trypsin inhibitor, 1X ITS supplement (insulin-transferrin-selenium, Sigma Cat. No. I3146), 1:1000 diluted fatty acid supplement (Sigma Cat. No. 7050), and 1:1000 diluted cholesterol. Additionally, the same media can be used for both hypoxic and aerobic cultivation. In one aspect, the growth media is changed from serum based media to serum free media after seeding and the first week of growth.

[0095] Incubation conditions will be under appropriate conditions of pH, temperature, and gas (*e.g.*, O₂, CO₂, etc) to maintain an hypoxic growth condition. In some embodiments, the three-dimensional cell culture can be suspended in the medium during the incubation period in order to maximize proliferative activity and generate factors that facilitate the desired biological activities of the fractions. In addition, the culture may be "fed" periodically to remove the spent media, depopulate released cells, and add new nutrient source. During the incubation period, the cultured cells grow linearly along and envelop the filaments of the three-dimensional scaffold before beginning to grow into the openings of the scaffold.

[0096] During incubation under hypoxic conditions, as compared to incubation under normal atmospheric oxygen concentrations of about 15-20%, thousands of genes are differentially expressed. Several genes have been found to be upregulated or downregulated in such compositions, most notably certain laminin species, collagen species and Wnt factors. In various aspects, the three dimensional ECM may be defined by the characteristic fingerprint or suite of cellular products produced by the cells due to growth in hypoxic condition as compared with growth under normal conditions. In the ECM compositions specifically exemplified herein, the three-dimensional tissues and surrounding media are characterized by expression and/or secretion of various factors.

[0097] The three dimensional tissues and compositions described herein have ECM that is present on the scaffold or framework. In some aspects, the ECM includes various laminin

and collagen types due to growth under hypoxic conditions and selection of cells grown on the support. The proportions of ECM proteins deposited can be manipulated or enhanced by selecting fibroblasts which elaborate the appropriate collagen type as well as growing the cells under hypoxic conditions in which expression of specific laminin and collagen species are upregulated or downregulated.

[0098] Selection of fibroblasts can be accomplished in some aspects using monoclonal antibodies of an appropriate isotype or subclass that define particular collagen types. In other aspects, solid substrates, such as magnetic beads, may be used to select or eliminate cells that have bound antibody. Combination of these antibodies can be used to select (positively or negatively) the fibroblasts which express the desired collagen type. Alternatively, the stroma used to inoculate the framework can be a mixture of cells which synthesize the desired collagen types. The distribution and origins of the exemplary type of collagen are shown in **Table 1**.

Table 1. Distributions and Origins of Various Types of Collagen

Collagen Type	Principle Tissue Distribution	Cells of Origin
I	Loose and dense ordinary connective tissue; collagen fibers	Fibroblasts and reticular cells; smooth muscle cells
	Fibrocartilage	
	Bone	Osteoblasts
	Dentin	Odontoblasts
II	Hyaline and elastic cartilage	Chondrocytes
	Vitreous body of eye	Retinal cells
III	Loose connective tissue; reticular fibers	Fibroblasts and reticular cells
	Papillary layer of dermis	Smooth muscle cells; endothelial cells
	Blood vessels	
IV	Basement membranes	Epithelial and endothelial cells
	Lens capsule of eye	Lens fibers
V	Fetal membranes; placenta	Fibroblasts

	Basement membranes	
	Bone	
	Smooth muscle	Smooth muscle cells
IV	Basement membranes	Epithelial and endothelial cells
	Lens capsule of the eye	Lens fiber
V	Fetal membranes; placenta	Fibroblasts
	Basement membranes	
	Bone	
	Smooth muscle	Smooth muscle cells
VI	Connective tissue	Fibroblasts
VII	Epithelial basement membranes	Fibroblasts
	anchoring fibrils	keratinocytes
VIII	Cornea	Corneal fibroblasts
IX	Cartilage	
X	Hypertrophic cartilage	
XI	Cartilage	
XII	Papillary dermis	Fibroblasts
XIV (undulin)	Reticular dermis	Fibroblasts
XVII	P170 bullous pemphigoid antigen	Keratinocytes

[0099] Additional types of collagen that may be present in ECM compositions are shown in Table 2.

Table 2. Types of Collagen and Corresponding Gene(s)

Collagen Type	Gene(s)
I	<i>COL1A1, COL1A2</i>
II	<i>COL2A1</i>
III	<i>COL3A1</i>
IV	<i>COL4A1, COL4A2, COL4A3, COL4A4, COL4A5, COL4A6</i>
V	<i>COL5A1, COL5A2, COL5A3</i>
VI	<i>COL6A1, COL6A2, COL6A3</i>
VII	<i>COL7A1</i>
VIII	<i>COL8A1, COL8A2</i>
IX	<i>COL9A1, COL9A2, COL9A3</i>
X	<i>COL10A1</i>
XI	<i>COL11A1, COL11A2</i>
XII	<i>COL12A1</i>
XIII	<i>COL13A1</i>
XIV	<i>COL14A1</i>
XV	<i>COL15A1</i>
XVI	<i>COL16A1</i>
XVII	<i>COL17A1</i>
XVIII	<i>COL18A1</i>
XIX	<i>COL19A1</i>
XX	<i>COL20A1</i>
XXI	<i>COL21A1</i>
XXII	<i>COL22A1</i>
XXIII	<i>COL23A1</i>
XXIV	<i>COL24A1</i>

Collagen Type	Gene(s)
XXV	<i>COL25A1</i>
XXVI	<i>EMID2</i>
XXVII	<i>COL27A1</i>
XXVIII	<i>COL28A1</i>

[00100] As discussed above the ECM compositions described herein include various collagens. As shown in Table 3 of Example 1, expression of several species of collagen are found to be upregulated in hypoxic cultured ECM compositions. Accordingly, in one aspect of the present invention, the ECM composition including one or more embryonic proteins, includes upregulation of collagen species as compared with that produced in oxygen conditions of about 15-20% oxygen. In another aspect, the upregulated collagen species are type V alpha 1; IX alpha 1; IX alpha 2; VI alpha 2; VIII alpha 1; IV, alpha 5; VII alpha 1; XVIII alpha 1; and XII alpha 1.

[00101] In addition to various collagens, the ECM composition described herein include various laminins. Laminins are a family of glycoprotein heterotrimers composed of an alpha, beta, and gamma chain subunit joined together through a coiled-coil domain. To date, 5 alpha, 4 beta, and 3 gamma laminin chains have been identified that can combine to form 15 different isoforms. Within this structure are identifiable domains that possess binding activity towards other laminin and basal lamina molecules, and membrane-bound receptors. Domains VI, IVb, and IVa form globular structures, and domains V, IIIb, and IIIa (which contain cysteine-rich EGF-like elements) form rod-like structures. Domains I and II of the three chains participate in the formation of a triple-stranded coiled-coil structure (the long arm).

[00102] Laminin chains possess shared and unique functions and are expressed with specific temporal (developmental) and spatial (tissue-site specific) patterns. The laminin alpha-chains are considered to be the functionally important portion of the heterotrimers, as they exhibit tissue-specific distribution patterns and contain the major cell interaction sites. Vascular endothelium is known to express two laminin isoforms, with varied expression depending on the developmental stage, vessel type, and the activation state of the endothelium.

[00103] Accordingly, in one aspect of the present invention, the ECM composition including one or more embryonic proteins, includes upregulation or downregulation of various laminin species as compared with that produced in oxygen conditions of about 15-20% oxygen.

[00104] Laminin 8, is composed of alpha-4, beta-1, and gamma-1 laminin chains. The laminin alpha-4 chain is widely distributed both in adults and during development. In adults it can be identified in the basement membrane surrounding cardiac, skeletal, and smooth muscle fibers, and in lung alveolar septa. It is also known to exist in the endothelial basement membrane both in capillaries and larger vessels, and in the perineurial basement membrane of peripheral nerves, as well as in intersinusoidal spaces, large arteries, and smaller arterioles of bone marrow. Laminin 8 is a major laminin isoform in the vascular endothelium that is expressed and adhered to by platelets and is synthesized in 3T3-L1 adipocytes, with its level of synthesis shown to increase upon adipose conversion of the cells. Laminin 8 is thought to be the laminin isoform generally expressed in mesenchymal cell lineages to induce microvessels in connective tissues. Laminin 8 has also been identified in mouse bone marrow primary cell cultures, arteriolar walls, and intersinusoidal spaces where it is the major laminin isoform in the developing bone marrow. Due to its localization in adult bone marrow adjacent to hematopoietic cells, laminin isoforms containing the alpha-4 chain are likely to have biologically relevant interactions with developing hematopoietic cells.

[00105] Accordingly, in another aspect of the present invention the ECM includes upregulation of laminin species, such as laminin 8. In another aspect, laminins produced by the three dimensional tissues of the present invention, includes at least laminin 8, which defines a characteristic or signature of the laminin proteins present in the composition.

[00106] The ECM compositions described herein can include various Wnt factors. Wnt family factors are signaling molecules having roles in a myriad of cellular pathways and cell-cell interaction processes. Wnt signaling has been implicated in tumorigenesis, early mesodermal patterning of the embryo, morphogenesis of the brain and kidneys, regulation of mammary gland proliferation, and Alzheimer's disease. As shown in Table 4 of Example 1, expression of several species of Wnt proteins are found to be upregulated in hypoxic cultured ECM compositions. Accordingly, in one aspect of the present invention, the ECM composition including one or more embryonic proteins, includes upregulation of Wnt species as compared with that produced in oxygen conditions of about 15-20% oxygen. In another

aspect, the upregulated Wnt species are wnt 7a and wnt 11. In another aspect, Wnt factors produced by the three dimensional tissues of the present invention, include at least wnt7a, and wnt11, which defines a characteristic or signature of the Wnt proteins present in the composition.

[00107] The culturing methods described herein, including culture under hypoxic conditions, have also been shown to upregulate expression of various growth factors. Accordingly, the ECM compositions described herein can include various growth factors, such as a vascular endothelial growth factor (VEGF). As used herein, a VEGF is intended to include all known VEGF family members. VEGFs are a sub-family of growth factors, more specifically of platelet-derived growth factor family of cystine-knot growth factors. VEGFs have a well known role in both vasculogenesis and angiogenesis. Several VEGFs are known, including VEGF-A, which was formerly known as VEGF before the discovery of other VEGF species. Other VEGF species include placenta growth factor (PlGF), VEGF-B, VEGF-C and VEGF-D. Additionally, several isoforms of human VEGF are well known.

[00108] In accordance with the increased production of Wnt proteins as well as growth factors by culturing under hypoxic conditions as described herein, the present invention further provides a method of producing a Wnt protein and a vascular endothelial growth factor (VEGF). The method can include culturing cells under hypoxic conditions as described herein, on a three-dimensional surface in a suitable growth medium, to produce the Wnt protein and the VEGF. In an exemplary aspect, the Wnt species are wnt 7a and wnt 11 and the VEGF is VEGF-A. The proteins may be further processed or harvested as described further herein or by methods known in the art.

[00109] As discussed throughout, the ECM compositions of the present invention includes both soluble and non-soluble fractions or any portion thereof. It is to be understood that the compositions of the present invention may include either or both fractions, as well as any combination thereof. Additionally, individual components may be isolated from the fractions to be used individually or in combination with other isolates or known compositions. Such compositions can be produced under normoxic or hypoxic conditions when CCM or ECM is desired for the composition.

[00110] Accordingly, in various aspects, the compositions produced using the methods of the present invention may be used directly or processed in various ways, the methods of which may be applicable to both the ECM and CCM compositions. The CCM, including the cell-free supernatant and media, may be subject to lyophilization for preserving and/or

concentrating the factors. Various biocompatible preservatives, cryoprotectives, and stabilizer agents may be used to preserve activity where required. Examples of biocompatible agents include, among others, glycerol, dimethyl sulfoxide, and trehalose. The lyophilizate may also have one or more excipients such as buffers, bulking agents, and tonicity modifiers. The freeze-dried media may be reconstituted by addition of a suitable solution or pharmaceutical diluent, as further described below.

[00111] In other aspects, the CCM is dialyzed. Dialysis is one of the most commonly used techniques to separate sample components based on selective diffusion across a porous membrane. The pore size determines molecular-weight cutoff (MWCO) of the membrane that is characterized by the molecular-weight at which 90% of the solute is retained by the membrane. In certain aspects membranes with any pore size is contemplated depending on the desired cutoff. Typical cutoffs are 5,000 Daltons, 10,000 Daltons, 30,000 Daltons, and 100,000 Daltons, however all sizes are contemplated.

[00112] In some aspects, the CCM may be processed by precipitating the active components (*e.g.*, growth factors) in the media. Precipitation may use various procedures, such as salting out with ammonium sulfate or use of hydrophilic polymers, for example polyethylene glycol.

[00113] In other aspects, the CCM is subject to filtration using various selective filters. Processing the CCM by filtering is useful in concentrating the factors present in the fraction and also removing small molecules and solutes used in the soluble fraction. Filters with selectivity for specified molecular weights include <5000 Daltons, <10,000 Daltons, and <15,000 Daltons. Other filters may be used and the processed media assayed for therapeutic activity as described herein. Exemplary filters and concentrator system include those based on, among others, hollow fiber filters, filter disks, and filter probes (see, *e.g.*, Amicon Stirred Ultrafiltration Cells).

[00114] In still other aspects, the CCM is subject to chromatography to remove salts, impurities, or fractionate various components of the medium. Various chromatographic techniques may be employed, such as molecular sieving, ion exchange, reverse phase, and affinity chromatographic techniques. For processing conditioned medium without significant loss of bioactivity, mild chromatographic media may be used. Non-limiting examples include, among others, dextran, agarose, polyacrylamide based separation media (*e.g.*, available under various tradenames, such as Sephadex, Sepharose, and Sephacryl).

[00115] In still other aspects, the CCM is formulated as liposomes. The growth factors may be introduced or encapsulated into the lumen of liposomes for delivery and for extending life time of the active factors. As known in the art, liposomes can be categorized into various types: multilamellar (MLV), stable plurilamellar (SPLV), small unilamellar (SUV) or large unilamellar (LUV) vesicles. Liposomes can be prepared from various lipid compounds, which may be synthetic or naturally occurring, including phosphatidyl ethers and esters, such as phosphatidylserine, phosphatidylcholine, phosphatidyl ethanolamine, phosphatidylinositol, dimyristoylphosphatidylcholine; steroids such as cholesterol; cerebroside; sphingomyelin; glycerolipids; and other lipids (see, *e.g.*, U.S. Patent No. 5,833, 948).

[00116] The ECM and/or CCM may be used directly without additional additives, or prepared as pharmaceutical compositions with various pharmaceutically acceptable excipients, vehicles or carriers. A “pharmaceutical composition” refers to a form of the soluble and/or non-soluble fractions and at least one pharmaceutically acceptable vehicle, carrier, or excipient. For intradermal, subcutaneous or intramuscular administration, the compositions may be prepared in sterile suspension, solutions or emulsions of the ECM compositions in aqueous or oily vehicles. The compositions may also contain formulating agents, such as suspending, stabilizing or dispersing agents. Formulations for injection may be presented in unit dosage form, ampules in multidose containers, with or without preservatives. Alternatively, the compositions may be presented in powder form for reconstitution with a suitable vehicle including, by way of example and not limitation, sterile pyrogen free water, saline, buffer, or dextrose solution.

[00117] In other aspects, the three dimensional tissues are cryopreserved preparations, which are thawed prior to use. Pharmaceutically acceptable cryopreservatives include, among others, glycerol, saccharides, polyols, methylcellulose, and dimethyl sulfoxide. Saccharide agents include monosaccharides, disaccharides, and other oligosaccharides with glass transition temperature of the maximally freeze-concentrated solution (T_g) that is at least -60 , -50 , -40 , -30 , -20 , -10 , or 0° C. An exemplary saccharide for use in cryopreservation is trehalose.

[00118] In some aspects, the three dimensional tissues are treated to kill the cells prior to use. In some aspects, the ECM deposited on the scaffolds may be collected and processed for administration (see U.S. Pat. Nos. 5,830,708 and 6,280,284, incorporated herein by reference).

[00119] In other embodiments, the three dimensional tissue may be concentrated and washed with a pharmaceutically acceptable medium for administration. Various techniques for concentrating the compositions are available in the art, such as centrifugation or filtering. Examples include, dextran sedimentation and differential centrifugation. Formulation of the three dimensional tissues may also involve adjusting the ionic strength of the suspension to isotonicity (i.e., about 0.1 to 0.2) and to physiological pH (i.e., pH 6.8 to 7.5). The formulation may also contain lubricants or other excipients to aid in administration or stability of the cell suspension. These include, among others, saccharides (e.g., maltose) and organic polymers, such as polyethylene glycol and hyaluronic acid. Additional details for preparation of various formulations are described in U.S. Patent Publication No. 2002/0038152, incorporated herein by reference.

[00120] As discussed above, the ECM and/or CCM compositions of the present invention may be processed in a number of ways depending on the anticipated application and appropriate delivery or administration of the ECM and/or CCM and/or composition. For example, the compositions may be delivered as three-dimensional scaffolds or implants, or the compositions may be formulated for injection as described above. The terms “administration” or “administering” are defined to include an act of providing a compound or pharmaceutical composition of the invention to a subject in need of treatment, including topical administration. The phrases “parenteral administration” and “administered parenterally” as used herein means modes of administration other than enteral and topical administration, usually by injection, and includes, without limitation, intravenous, intramuscular, intraarterial, intrathecal, intracapsular, intraorbital, intracardiac, intradermal, intraperitoneal, transtracheal, subcutaneous, subcuticular, intraarticular, subcapsular, subarachnoid, intraspinal and intrasternal injection and infusion. The phrases “systemic administration,” “administered systemically,” “peripheral administration” and “administered peripherally” as used herein mean the administration of a compound, drug or other material other than directly into the central nervous system, such that it enters the subject’s system and, thus, is subject to metabolism and other like processes, for example, subcutaneous administration.

[00121] Pharmaceutical compositions for topical application may take a variety of forms including, for example, solutions, gels, suspensions, and the like. Generally speaking, improved absorption may be achieved when the topical compositions are in the form of a solution or gel, i.e., where the active ingredient, for example, minoxidil, is dissolved in the

carrier solution, in contrast to topical compositions which are in the form of suspensions, i.e., where the active ingredient is merely suspended in the composition.

[00122] Penetration enhancers (also called sorption promoters or accelerants) which penetrate into skin to reversibly decrease the barrier resistance and improve transdermal drug delivery. Numerous compounds have been evaluated for penetration enhancing activity, including sulfoxides (such as dimethylsulphoxide, DMSO), Azones (e.g. laurocapram), pyrrolidones (for example 2-pyrrolidone, 2P), alcohols and alkanols (ethanol, or decanol), glycols (for example propylene glycol, PG, a common excipient in topically applied dosage forms), surfactants (also common in dosage forms) and terpenes.

[00123] The term “subject” as used herein refers to any individual or patient to which the subject methods are performed. Generally the subject is human, although as will be appreciated by those in the art, the subject may be an animal. Thus other animals, including mammals such as rodents (including mice, rats, hamsters and guinea pigs), cats, dogs, rabbits, farm animals including cows, horses, goats, sheep, pigs, etc., and primates (including monkeys, chimpanzees, orangutans and gorillas) are included within the definition of subject.

[00124] The ECM and/or CCM compositions of the present invention have a variety of applications including, but not limited to, promoting repair and/or regeneration of damaged cells or tissues, use in patches to promote tissue regeneration, use in tissue culture systems for culturing cells, such as stem cells, use in surface coatings used in association with implantable devices (e.g., pacemakers, stents, stent grafts, vascular prostheses, heart valves, shunts, drug delivery ports or catheters), promoting soft tissue repair, augmentation, and/or improvement of a skin surface, such as wrinkles, use as a biological anti-adhesion agent, as a biological vehicle for cell delivery or maintenance at a site of delivery, stimulating hair, nail or lash growth and/or promoting hair follicle development and/or activation or stimulation on an area of the skin of a subject comprising contacting the hair (scalp), nail or lash or adjacent areas thereof.

[00125] Additionally, the ECM and/or CCM compositions derived from culturing cells as described in any method herein, may be used in any other application or method of the present invention. For example, the ECM and/or CCM compositions generated by culturing cells using the tissue culture system of the present invention may be used, for example, in the repair and/or regeneration of cells, use in patches to promote tissue regeneration, use in tissue culture systems for culturing cells, such as stem cells, use in surface coatings used in association with implantable devices (e.g., pacemakers, stents, stent grafts, vascular

prostheses, heart valves, shunts, drug delivery ports or catheters), promoting soft tissue repair, augmentation, and/or improvement of a skin surface, such as wrinkles, use as a biological anti-adhesion agent or as a biological vehicle for cell delivery or maintenance at a site of delivery.

[00126] In various embodiments, the present invention includes methods for repair and/or regeneration of cells or tissue and promoting soft tissue repair. One embodiment includes a method of repair and/or regeneration of cells by contacting cells to be repaired or regenerated with the ECM and/or CCM compositions of the present invention. The method may be used for repair and/or regeneration of a variety of cells as discussed herein, including osteochondral cells.

[00127] In one aspect, the method contemplates repair of osteochondral defects. As used herein, “osteochondral cells” refers to cells which belong to either the chondrogenic or osteogenic lineage or which can undergo differentiation into either the chondrogenic or osteogenic lineage, depending on the environmental signals. This potential can be tested in vitro or in vivo by known techniques. Thus, in one aspect, the ECM compositions of the present invention are used to repair and/or regenerate, chondrogenic cells, for example, cells which are capable of producing cartilage or cells which themselves differentiate into cells producing cartilage, including chondrocytes and cells which themselves differentiate into chondrocytes (e.g., chondrocyte precursor cells). Thus, in another aspect, the compositions of the present invention are useful in repair and/or regeneration of connective tissue. As used herein, “connective tissue” refers to any of a number of structural tissues in the body of a mammal including but not limited to bone, cartilage, ligament, tendon, meniscus, dermis, hyperdermis, muscle, fatty tissue, joint capsule.

[00128] The ECM and/or CCM compositions of the present invention may be used for treating osteochondral defects of a diarthroidal joint, such as knee, an ankle, an elbow, a hip, a wrist, a knuckle of either a finger or toe, or a temporomandibular joint. Such osteochondral defects can be the result of traumatic injury (e.g., a sports injury or excessive wear) or a disease such as osteoarthritis. A particular aspect relates to the use of the matrix of the present invention in the treatment or prevention of superficial lesions of osteoarthritic cartilage. Additionally the present invention is of use in the treatment or prevention of osteochondral defects which result from ageing or from giving birth. Osteochondral defects in the context of the present invention should also be understood to comprise those conditions where repair of cartilage and/or bone is required in the context of surgery such as, but not

limited to, cosmetic surgery (e.g., nose, ear). Thus such defects can occur anywhere in the body where cartilage or bone formation is disrupted or where cartilage or bone are damaged or non-existent due to a genetic defect.

[00129] As discussed above, growth factors or other biological agents which induce or stimulate growth of particular cells may be included in the ECM compositions of the present invention. The type of growth factors will be dependent on the cell-type and application for which the composition is intended. For example, in the case of osteochondral cells, additional bioactive agents may be present such as cellular growth factors (e.g., TGF- β), substances that stimulate chondrogenesis (e.g., BMPs that stimulate cartilage formation such as BMP-2, BMP-12 and BMP-13), factors that stimulate migration of stromal cells to the scaffold, factors that stimulate matrix deposition, anti-inflammatories (e.g., non-steroidal anti-inflammatories), immunosuppressants (e.g., cyclosporins). Other proteins may also be included, such as other growth factors such as platelet derived growth factors (PDGF), insulin-like growth factors (IGF), fibroblast growth factors (FGF), epidermal growth factor (EGF), human endothelial cell growth factor (ECGF), granulocyte macrophage colony stimulating factor (GM-CSF), vascular endothelial growth factor (VEGF), cartilage derived morphogenetic protein (CDMP), other bone morphogenetic proteins such as OP-1, OP-2, BMP3, BMP4, BMP9, BMP11, BMP14, DPP, Vg-1, 60A, and Vgr-1, collagens, elastic fibers, reticular fibers, glycoproteins or glycosaminoglycans, such as heparin sulfate, chondroitin-4-sulfate, chondroitin-6-sulfate, dermatan sulfate, keratin sulfate, etc. For example, growth factors such as TGF- β , with ascorbate, have been found to trigger chondrocyte differentiation and cartilage formation by chondrocytes. In addition, hyaluronic acid is a good substrate for the attachment of chondrocytes and other stromal cells and can be incorporated as part of the scaffold or coated onto the scaffold.

[00130] Additionally, other factors which influence the growth and/or activity of particular cells may also be used. For example, in the case of chondrocytes, a factor such as a chondroitinase which stimulates cartilage production by chondrocytes can be added to the matrix in order to maintain chondrocytes in a hypertrophic state as described in U.S. Patent Application No. 2002/0122790 incorporated herein by reference. In another aspect, the methods of the present invention include the presence of polysulphated alginates or other polysulphated polysaccharides such as polysulphated cyclodextrin and/or polysulphated inulin, or other components capable of stimulating production of ECM of connective tissue

cells as described in International Patent Publication No. WO 2005/054446 incorporated herein by reference.

[00131] The cell or tissue to be repaired and/or regenerated may be contacted in vivo or in vitro by any of the methods described herein. For example, the ECM compositions may be injected or implanted (e.g., via ECM tissue, a patch or coated device of the present invention) into the subject. In another aspect, the tissue or cells to be repaired and/or regenerated may be harvested from the subject and cultured in vitro and subsequently implanted or administered to the subject using known surgical techniques.

[00132] As discussed above, the ECM compositions of the present invention may be processed in a variety of ways. Accordingly, in one embodiment, the present invention includes a tissue culture system. In various aspects, the culture system is composed of the ECM compositions described herein. The ECM compositions of the present invention may be incorporated into the tissue culture system in a variety of ways. For example, compositions may be incorporated as coatings, by impregnating three-dimensional scaffold materials as described herein, or as additives to media for culturing cells. Accordingly, in one aspect, the culture system can include three-dimensional support materials impregnated with any of the ECM compositions described herein, such as growth factors or embryonic proteins.

[00133] The ECM compositions described herein may serve as a support or three-dimensional support for the growth of various cell types. Any cell type capable of cell culture is contemplated. In one aspect, the culture system can be used to support the growth of stem cells. In another aspect, the stem cells are embryonic stem cells, mesenchymal stem cells or neuronal stem cells.

[00134] The tissue culture system may be used for generating additional ECM compositions, such as implantable tissue. Accordingly, culturing of cells using the tissue culture system of the present invention may be performed in vivo or in vitro. For example, the tissue culture system of the present invention may be used to generate ECM compositions for injection or implantation into a subject. The ECM compositions generated by the tissue culture system may be processed and used in any method described herein.

[00135] The ECM compositions of the present invention may be used as a biological vehicle for cell delivery. As described herein, the ECM compositions may include both soluble and/or non-soluble fractions. As such, in another embodiment of the present invention, a biological vehicle for cell delivery or maintenance at a site of delivery including

the ECM compositions of the present invention, is described. The ECM compositions of the present invention, including cells and three-dimensional tissue compositions, may be used to promote and/or support growth of cells in vivo. The vehicle can be used in any appropriate application, for example to support injections of cells, such as stem cells, into damaged heart muscle or for tendon and ligament repair as described above.

[00136] Appropriate cell compositions (e.g., isolated ECM cells of the present invention and/or additional biological agents) can be administered before, after or during the ECM compositions are implanted or administered. For example, the cells can be seeded into the site of administration, defect, and/or implantation before the culture system or biological delivery vehicle is implanted into the subject. Alternatively, the appropriate cell compositions can be administered after (e.g., by injection into the site). The cells act therein to induce tissue regeneration and/or cell repair. The cells can be seeded by any means that allows administration of the cells to the defect site, for example, by injection. Injection of the cells can be by any means that maintains the viability of the cells, such as, by syringe or arthroscope.

[00137] ECM compositions have been described for promoting angiogenesis in organs and tissues by administering such compositions to promote endothelialization and vascularization in the heart and related tissues. Accordingly, in yet another embodiment, the present invention includes a surface coating used in association with implantation of a device in a subject including the ECM compositions described herein. The coating may be applied to any device used in implantation or penetration of a subject, such as a pacemaker, a stent, a stent graft, a vascular prosthesis, a heart valve, a shunt, a drug delivery port or a catheter. In certain aspects, the coating can be used for modifying wound healing, modifying inflammation, modifying a fibrous capsule formation, modifying tissue ingrowth, or modifying cell ingrowth. In another embodiment, the present invention includes a for treatment of damaged tissue, such as heart, intestinal, infarcted or ischemic tissue. Presented below are examples discussing generation of ECM compositions contemplated for such applications. The preparation and use of ECM compositions grown under normal oxygen conditions is described in U.S. Patent Application No. 2004/0219134 incorporated herein by reference.

[00138] In another embodiment, the present invention includes various implantable devices and tissue regeneration patches including the ECM compositions described herein which allow for benefits, such as tissue ingrowth. As discussed herein, the ECM compositions may

serve as coatings on medical devices, such as patches or other implantable devices. In various aspects, such devices are useful for wound repair, hernia repair, pelvic floor repair (e.g., pelvic organ prolapse), rotator cuff repair and the like. In related aspects, coatings are useful for orthopedic implants, cardiovascular implants, urinary slings and pacemaker slings.

[00139] For example, the basic manifestation of a hernia is a protrusion of the abdominal contents into a defect within the fascia. Surgical approaches toward hernia repair is focused on reducing the hernial contents into the peritoneal cavity and producing a firm closure of the fascial defect either by using prosthetic, allogeneic or autogenous materials. A number of techniques have been used to produce this closure, however, drawbacks to current products and procedures include hernia recurrence, where the closure weakens again, allowing the abdominal contents back into the defect. In herniorrhaphy, a corrective tissue regeneration patch, such as a bioresorbable or synthetic mesh coated with ECM compositions could be used.

[00140] A variety of techniques are known in the art for applying biological coatings to medical device surfaces that may be utilized with the present invention. For example, ECM compositions may be coated using photoactive crosslinkers allowing for permanent covalent bonding to device surfaces upon activation of the crosslinker by applying ultraviolet radiation. An exemplary crosslinker is TriLite™ crosslinker, which has been shown to be non-cytotoxic, non-irritating to biological tissue and non-sensitizing. ECM materials may be unseparated or separated into individual components, such as human collagens, hyaluronic acid (HA), fibronectin, and the like before coating or incorporation into various implantable devices. Further, additional growth factors and such may be incorporated to allow for beneficial implantation characteristics, such as improved cell infiltration.

[00141] In various related embodiments, the present invention provides methods and devices applicable in cosmetic/cosmeceutical applications, such as, but not limited to anti-aging, anti-wrinkle, skin fillers, moisturizers, pigmentation augmentation, skin firming, and the like. Accordingly, in one embodiment the present invention includes a method for improvement of a skin surface in a subject including administering to the subject at the site of a wrinkle, the ECM compositions described herein. In a related embodiment, the present invention includes a method for soft tissue repair or augmentation in a subject including administering to the subject at the site of a wrinkle, the ECM compositions described herein. In various cosmetic applications, the compositions may be formulated as appropriate, such as injectable and topical formulations. As discussed further in the Examples included herein,

ECM compositions formulated as topicals have been shown to be effective in various skin aesthetics applications, such as anti-wrinkle, anti-aging applications as well as an adjunct to ablative laser surgery. Several beneficial characteristics of ECM containing topicals have been shown. Such benefits include 1) facilitating re-epithelization following resurfacing; 2) reduction of non-ablative and ablative fractional laser resurfacing symptoms (e.g., erythema, edema, crusting, and sensorial discomfort); 3) generating smooth, even textured skin; 4) generating skin moisturization; 5) reducing appearance of fine lines/wrinkles; 6) increasing skin firmness and suppleness; 7) reducing skin dyspigmentation and 8) reducing red, blotchy skin.

[00142] In one embodiment, the present invention provides for compositions comprising a conditioned culture media (CCM) in combination with an active agent to promote hair, lash and/or nail growth in a subject. In various aspects, the CCM is produced by the methods described herein. In one aspect, the active agent is a hair growth promoting agent. In certain aspects the CCM is a hypoxic CCM. In another aspect, the composition comprises the CCM and minoxidil. In certain aspects, the minoxidil is present at a concentration from about 0.5% to about 5% by weight and is in the form of a pharmaceutically acceptable derivative (i.e. pharmaceutically acceptable salts, solvates, hydrates, isomers, esters, tautomers, anhydrides, enantiomers, complexes, polymorphs or prodrugs). In an additional aspect, the composition comprises CCM and bimatoprost. In a further aspect, the bimatoprost is present at a concentration from about 0.01% to about 5% by weight and the bimatoprost is provided as a pharmaceutically acceptable salt. In certain aspects, the compositions also have penetration enhancers and/or a pharmaceutically acceptable excipient. In various aspects, the composition is adapted for topical application to mammalian skin as a foam, wherein said foam comprises bimatoprost and/or minoxidil, and at least one surfactant, wherein the surfactant optionally includes a foam stabilizer; an aqueous-alcohol solvent, and wherein said aqueous-alcohol solvent comprises water and an alcohol

[00143] In an additional embodiment, the present invention provides methods of stimulating hair, nail or lash growth and/or promoting hair follicle development and/or activation or stimulation on an area of the skin of a subject comprising contacting the hair (scalp), nail or lash or adjacent areas thereof with a composition of any of claims 6-24 under conditions that allow for hair, nail or lash growth and/or promoting hair follicle development and/or activation or stimulation on an area of the skin in the subject. In certain aspects, the scalp, dermis or hair follicle is contacted with the composition.

[00144] In a further embodiment, the present invention provides a topical pharmaceutical composition comprising minoxidil and/or bimatoprost, a CCM and at least one or more pharmaceutically acceptable excipients. In one aspect, the minoxidil is in the form of a pharmaceutically acceptable derivative (i.e. pharmaceutically acceptable salts, solvates, hydrates, isomers, esters, tautomers, anhydrates, enantiomers, complexes, polymorphs or prodrugs). In certain aspects, the minoxidil is present at a concentration from about 0.5% to about 5% by weight. In specific aspects, the minoxidil is present at a concentration of about 1%, 2% or 5% by weight. In various aspects, the bimatoprost is present at a concentration from about 0.01% to about 5% by weight. In a particular aspect, the bimatoprost is present at a concentration of about 0.1%, 1%, 3% or 5% by weight.

[00145] The compositions of the present invention may be prepared as known in the art, however employing the innovative culture methods described herein (e.g., culture under hypoxic conditions). The preparation and use of ECM compositions created under normal oxygen culture conditions for the repair and/or regeneration of cells, improvement of skin surfaces, and soft tissue repair are described in U.S. Patent No. 5,830,708, U.S. Patent No. 6,284,284, U.S. Patent Application No. 2002/0019339 and U.S. Patent Application No. 2002/0038152 incorporated herein by reference.

[00146] In another embodiment, the present invention includes a biological anti-adhesion agent including the ECM compositions described herein. The agent can be used in such applications as anti-adhesion patches used after the creation of intestinal or blood vessel anastomoses.

[00147] The compositions or active components used herein, will generally be used in an amount effective to treat or prevent the particular disease being treated. The compositions may be administered therapeutically to achieve therapeutic benefit or prophylactically to achieve prophylactic benefit. By therapeutic benefit is meant eradication or amelioration of the underlying condition or disorder being treated. Therapeutic benefit also includes halting or slowing the progression of the disease, regardless of whether improvement is realized.

[00148] The amount of the composition administered will depend upon a variety of factors, including, for example, the type of composition, the particular indication being treated, the mode of administration, whether the desired benefit is prophylactic or therapeutic, the severity of the indication being treated and the age and weight of the patient, and effectiveness of the dosage form. Determination of an effective dosage is well within the capabilities of those skilled in the art.

[00149] Initial dosages may be estimated initially from in vitro assays. Initial dosages can also be estimated from in vivo data, such as animal models. Animals models useful for testing the efficacy of compositions for enhancing hair growth include, among others, rodents, primates, and other mammals. The skilled artisans can determine dosages suitable for human administration by extrapolation from the in vitro and animal data.

[00150] Dosage amounts will depend upon, among other factors, the activity of the conditioned media, the mode of administration, the condition being treated, and various factors discussed above. Dosage amount and interval may be adjusted individually to provide levels sufficient to maintain the therapeutic or prophylactic effect.

[00151] Presented below are examples discussing generation of ECM compositions contemplated for the discussed applications. The following examples are provided to further illustrate the embodiments of the present invention, but are not intended to limit the scope of the invention. While they are typical of those that might be used, other procedures, methodologies, or techniques known to those skilled in the art may alternatively be used.

EXAMPLES

EXAMPLE 1

DIFFERENTIAL GENE EXPRESSION IN ECM COMPOSITIONS GROWN UNDER HYPOXIC CONDITIONS

[00152] Primary human neonatal foreskin fibroblasts were cultured as standard monolayers in tissue culture flasks and compared to three-dimensional fibroblast cultures, within a naturally deposited, fetal-like ECM. The cultures were grown as disclosed herein. To assess differential expression of genes, samples of total RNA were completed using Agilent Whole Human Genome Oligo Microarrays® for global gene expression (including less than 40,000 genes) following the manufacturer's protocol.

[00153] Upon comparison, fibroblasts were found to regulate collagen and ECM gene expression in three-dimensional cultures within a hypoxic cultured naturally secreted ECM. Upregulation and downregulation of expression of various collagen and ECM genes are evident in Table 3.

Table 3. Differential Collagen and ECM Expression in Hypoxic Three-Dimensional Fibroblast Cultures

GENE	FOLD INCREASE	FOLD DECREASE
COL4A1	17.2	
COL20A1	6.88	
COL19A1	5.22	
COL9A1	4.81	
COL10A1	4.45	
COL6A3	3.48	
COL9A2	2.48	
COL14A1	2	
SPARC	2.74	
COL1A2		3.45
COL13A1		4
COL18A1		4.76
COL1A2		7.14

[00154] Upon comparison, fibroblasts were found to regulate gene expression of Wnt pathway genes in three-dimensional cultures within a hypoxic cultured naturally secreted ECM. Upregulation and downregulation of expression of various Wnt pathway genes are evident in Table 4.

Table 4. Differential Wnt Expression in Hypoxic Three-Dimensional Fibroblast Cultures

GENE	FOLD INCREASE	FOLD DECREASE
WNT 4	5.94	
WNT 7a	5.43	
WNT 7b	4.05	
WNT 2b	3.95	
WNT 10a	3.86	
WNT 8b	3.48	
WNT 6	3.36	
WNT 3a	3.19	
WNT 9b	3.06	
WNT 9a	3.02	
WNT 11	2.89	
WNT 5a		8.33
WNT 2		7.14
WNT 5b		5.26
LRP6	3.43	
LRP3	2.27	
LRP11		10
LRP12		7.69
DKK1		50
DKK3		5.88
FSZD5	4.48	
FRZ9	3.85	
FRZB	3.36	
FRZD1	2.94	
SFRP2	2.95	
FRZD1	2.92	

FRZD3	2.84	
AXIN2	4.4	
KREMEN2	4.24	
KREMEN1	3.45	
b-CATENIN		4.76
GSK3b		11.1
GSK3a		6.67
bFGF		50

[00155] Upon comparison, fibroblasts were found to regulate gene expression of bone morphogenetic protein (BMP) pathway genes in three-dimensional cultures within a hypoxic cultured naturally secreted ECM. Upregulation and downregulation of expression of various BMP pathway genes are evident in Table 5.

Table 5. Differential BMP Expression in Hypoxic Three-Dimensional Fibroblast Cultures

GENE	FOLD INCREASE	FOLD DECREASE
BMP7/OP1	4.88	
BMP2	4.19	
BMP5	3.49	
BMP3	3.44	
BMPrecIb	3.37	
BMP8b	3.36	
BMP 8a	3.15	
BMP 10	2.86	
BMP1	2.12	
BMPrecIa		2.5
Osteocalcin		2.5
Osteopontin		6.25
BMPrecII		6.25

[00156] Upon comparison, fibroblasts were found to regulate expression of additional genes in three-dimensional cultures within a the hypoxic cultured naturally secreted ECM. Upregulation and downregulation of expression of additional genes are evident in Table 6. Results indicate that hypoxic culture conditions result in a 14.78-fold increase in mRNA expression for hypoxia-inducible factor (HIF 1A) and a 4.9 decrease in its respective inhibitor. This suggests that the hypoxic cultured conditioned medium is experiencing a low oxygen tension environment (hypoxia) because the messenger RNA for HIF 1A that codes for the translation of the protein is up-regulated and its inhibitor is down-regulated. Further, VEGFB (4.33-fold increase), KGF (11.51-fold increase), and IL-8 (5.81-fold increase) levels were also up-regulated under hypoxic culture conditions.

Table 6. Additional Gene Expression Changes Resulting from Low Oxygen Culture of Fibroblast ECM *In Vivo*.

GENES	FOLD INCREASE		FOLD DECREASE
	Collagens		
COL5A1	6.21		
COL9A2	3.96		
COL6A2	3.78		
COL6A2	3.21		
COL11A1	3.07		
COL8A1	2.78		
COL4A5	2.45		
COL7A1	2.45		
COL18A1	2.41		
COL12A1	2.04		
COL1A2			0.5
COL14A1			0.45
COL4A1			0.45
COL5A2			0.23
COL6A1			0.16

	Matrix Metalloproteinases (MMPs)		
MMP23B	2.75		
MMP27			0.24
MMP28			0.17
MMP10			0.16
MMP1			0.16
MMP7			0.1
MMP14			0.08
MMP3			0.06
MMP12			0.05
	Other ECM		
HAPLN3	8.11		
ACAN L12234	6.48		
AGC1	3.32		
LAMA3	2.92		
LAMA1	2.14		
LAMA5	2.14		
	Additional Genes		
HIF 1A	14.18		
HIF 1AN			4.9
VEGFB	4.33		
VEGFC	3.84		
KGF	11.51		
IL-8	5.81		

Table 7. Stem Cell Related Gene Expression

GENES	FOLD INCREASE	FOLD DECREASE
Oct4	5.1	
Sox2	8.2	
NANOG	4.9	
KLF4	21.0	
cMyc	7.1	

EXAMPLE 2

**PRODUCTION OF HYPOXIC ECM
USING PRIMARY HUMAN NEONATAL FORESKIN FIBROBLASTS**

[00157] Two examples are provided for hypoxic culture of ECM using primary human neonatal foreskin fibroblasts.

[00158] Primary human neonatal foreskin fibroblasts were expanded in tissue culture flasks in the presence of 10% fetal bovine serum, 90% High Glucose DMEM with 2 mM L-glutamine (10%FBS/DMEM). Cells were subcultured using 0.05% trypsin/EDTA solution until the 3rd passage at which time they were seeded to either Cytodex-1 dextran beads at 0.04 mgs dry beads/ml of medium (5×10^6 cells/10mgs beads in a 125 ml spinner flask filled with 100-120 mls), or to nylon mesh (25×10^6 cells/6 x 100cm² nylon). All cultures were kept in normal atmosphere and 5% CO₂ for expansion and seeding, at which point low oxygen cultures were split to an airtight chamber which was flooded with 95% nitrogen/ 5% CO₂ so that a hypoxic environment could be created within the culture medium. This system is maintains about 1-5% oxygen within the culture vessel. Cells were mixed well into the minimum volume needed to cover nylon or beads for seeding, and were subsequently mixed once after 30 minutes, then allowed to sit overnight in a humidified 37°C incubator. Cultures were fed 10%FBS/DMEM for 2-4 weeks with a 50-70% media exchange, every 2-3 days while cells proliferated and then began depositing ECM. Cultures were fed for another 4-6 weeks using 10% bovine calf serum with iron supplement, and 20 ug/ml ascorbic acid in place of FBS. Spinner flasks were mixed at 15-25 rpm initially and for about 2-4 weeks, at which time they were increased to 45 rpm and maintained at this rate thereafter. Bead cultures formed large amorphous structures containing ECM of as much as 0.5 to 1.0 cm in

width and diameter after 4 weeks, and these cultures were therefore hypoxic due to gas diffusion and high metabolic requirements.

[00159] In an additional example, primary human neonatal foreskin fibroblasts were expanded in monolayer flasks, and then cultured on nylon mesh scaffolds to support development of an ECM *in vitro*. Fibroblasts were expanded in DMEM with high glucose, 2 mM L-glutamine, and 10% (v/v) fetal bovine serum. Cultures were also supplemented with 20 µg/ml ascorbic acid. After 3 weeks in ambient oxygen (approximately 16%-20% oxygen) duplicate ECM-containing cultures were switched to hypoxic culture conditions (1%-5% oxygen) in a sealed chamber flushed extensively with 95% nitrogen/5% carbon dioxide (Cat.#MC-101, Billups-Rothenberg, Inc., Del Mar, CA). To ensure depletion of atmospheric oxygen from the culture medium, 2-3 hours later the atmosphere was replaced to ensure that the medium contained approximately 1-3% oxygen. Both sets of ECM-containing cultures were grown with twice weekly feedings for another 4 weeks, and then cultures were prepared for RNA isolation. Total cellular RNA was isolated using a commercially available kit according to the manufacturers instructions (Cat.# Z3100, Promega, Inc.). Purified RNA samples were stored at -80°C, prior to processing for microarray analysis of gene expression using Agilent Whole Human Genome Oligo Microarrays®.

[00160] In analyzing the results, there were approximately 5,500 differentially expressed transcripts detected from probes prepared from ambient oxygen in comparison to probes from low oxygen cultures, using Agilent Whole Human Genome Oligo Microarrays®. Of these, about half (2,500) were greater than 2.0 fold increased by low oxygen, and about half (2,500) were decreased greater than 2.0 fold in low oxygen. This indicates that low oxygen led to significant changes in gene expression *in vitro*. Of particular interest, transcripts for ECM proteins, particularly a number of collagen genes were up-regulated, while a number of genes for matrix-degrading enzymes were down-regulated.

EXAMPLE 3

TISSUE-ENGINEERED HUMAN EMBRYONIC EXTRACELLULAR MATRIX FOR THERAPEUTIC APPLICATIONS

[00161] The embryonic ECM creates an environment conducive to rapid cell proliferation and healing without the formation of scars or adhesions. It was hypothesized that the growth of human neonatal fibroblasts in 3 dimensions under conditions that simulate the early embryonic environment prior to angiogenesis (hypoxia and reduced gravitational forces) would generate an ECM with fetal properties. Gene chip array analysis showed the

differential expression of over 5000 genes under the hypoxic versus traditional tissue culture conditions. The ECM produced was similar to fetal mesenchymal tissue in that it is relatively rich in collagens type III, IV, and V, and glycoproteins such as fibronectin, SPARC, thrombospondin, and hyaluronic acid. Since the ECM also plays an important regulatory role in binding and presenting growth factors in putative niches which support regenerative stem cell populations with key growth factors, we evaluated the effects of hypoxia on growth factor expression during the development of the fetal-like ECM in culture. Hypoxia can also enhance expression of factors which regulate wound healing and organogenesis, such as VEGF, FGF-7, and TGF- β , as well as multiple wnts including wnts 2b,4,7a,10a, and 11. The embryonic human ECM also stimulated an increase of metabolic activity in human fibroblasts *in vitro*, as measured by increased enzymatic activity using the MTT assay. Additionally, we detected an increase in cell number in response to human ECM. This human ECM can be used as a biological surface coating, and tissue filler treatment in various therapeutic applications where new tissue growth and healing without scarring or adhesions.

EXAMPLE 4

PRODUCTION OF NATURALLY-SOLUBLE WNT ACTIVITY FOR REGENERATIVE MEDICINE APPLICATIONS

[00162] Stem or progenitor cells that can regenerate adult tissues, such as skin or blood, recapitulate embryonic development to some extent to accomplish this regeneration. A growing number of studies have shown that key regulators of stem cell pluripotency and lineage-specific differentiation active during embryogenesis are re-expressed in the adult under certain circumstances. The WNT family of secreted morphogenetic growth and development factors is among the growth factors which can potentially provide valuable research tools and eventually therapeutic treatments in the clinic. However, Wnt's have proven refractory to standard recombinant expression and purification techniques to date on a commercial scale, and there are no reports of large-scale WNT protein production to enable clinical development of WNT-based products. Techniques have been developed for growing fetal-like ECM in culture using neonatal human dermal fibroblasts on various scaffolds in culture to generate three-dimensional tissue-equivalents. In this process, it was discovered that these cultures can provide a commercial-scale source of bioactive WNT's contained in the serum-free conditioned medium used for ECM production. Here we present data on this WNT product candidate.

[00163] Gene expression analysis of the cells demonstrated that at least 3 WNT genes were expressed (wnt 5a, wnt 7a, and wnt 11), and a small number of genes related to wnt signaling were expressed as well; however, their function is not completely understood. The gene expression data was extended to an in vitro bioassay for wnt-signaling (nuclear translocation of β -catenin in primary human epidermal keratinocytes) and wnt activity on blood stem cells was evaluated. Both assays demonstrated activity consistent with canonical wnt activity. Furthermore, conditioned media from these cultures showed wnt activity when injected into the skin of mice, inducing hair follicle stem cells to enter anagen, thus causing hair growth. This indicates that the stabilized WNT activity within the defined and serum-free condition medium did not require purification. This product can be used for hair follicle regeneration and as a valuable research tool for the culture of various human stem cells.

EXAMPLE 5

HYPOXIC FIBROBLASTS DEMONSTRATE UNIQUE ECM PRODUCTION AND GROWTH FACTOR EXPRESSION

[00164] Human neonatal dermal fibroblasts produce an ECM when cultured in vitro, which closely mimics the dermis and which can replace the damaged dermis in regenerative medicine applications such as wound healing. Since the process of wound healing also recapitulates embryonic development, by simulating the embryonic environment we hypothesize that the ECM produced will provide an enhanced ECM for tissue regeneration applications. Therefore, human neonatal fibroblast-derived ECM were grown under hypoxic conditions in culture, to simulate the hypoxia which exists in the early embryo prior to angiogenesis. The goal was to generate an ECM with fetal properties using hypoxic conditions during tissue development in culture.

[00165] The ECM produced in these hypoxic cultures was similar to fetal mesenchymal tissue in that it is relatively rich in collagens type III and V, and glycoproteins such as fibronectin, SPARC, thrombospondin, and hyaluronic acid. Since the ECM also plays an important regulatory role in binding and presenting growth factors in putative niches which support regenerative stem cell populations with key growth factors, we evaluated the effects of hypoxia on growth factor expression during the development of the fetal-like ECM in culture. It was shown that hypoxia can also enhance expression of factors which regulate wound healing and organogenesis, such as VEGF, FGF-7, and TGF- β .

[00166] The human ECM also stimulated an increase of metabolic activity in human fibroblasts in vitro, as measured by increased enzymatic activity using the MTT assay.

Additionally, an increase in cell number in response to human ECM was detected. These results support the use of this human ECM as a coating/scaffold in embryonic cell cultures and as a biological surface coating/filler in various therapeutic applications or medical devices.

EXAMPLE 6

HUMAN EXTRACELLULAR MATRIX (HECM) COATED BIOMEDICAL MATERIALS

[00167] ECM has been reported to create an environment conducive to rapid cell proliferation and healing without the formation of scars or adhesions. Using methods described herein, unique, embryonic like, human ECM (hECM) was generated by culturing neonatal fibroblasts in low oxygen and specific gravity. Results included: angiogenesis when hECM is placed on the chorioallantoic membrane, and reduced inflammatory cell migration when hECM is coated on nylon mesh and implanted on the flank position in the subcutaneous region of SCID mice. Based upon these results, it was hypothesized that coating our hECM on polypropylene mesh would illicit reduced inflammatory cell migration and fibrous encapsulation at the material-biological interface in the subcutaneous region of SCID mice.

[00168] ECM compositions generated using human derived materials (hECM) were coated onto propylene mesh using a photoactive crosslinker. hECM was coated on 6mm biopsy punched polypropylene by UV covalent bonding mechanism (Innovative Surface Technologies (ISurTec) TriLite™ Crosslinker). Coated and uncoated hECM 6mm biopsy punches of polypropylene were sterilized utilizing E-Beam™ (BeamOne LLC E-BEAM™) or ethylene oxide (ETO) (described by ETO Flagstaff Medical Center). Next, each 6mm polypropylene disc was split into two symmetrical semi circular inserts. Finally, polypropylene implants were placed bilaterally utilizing aseptic technique on the flank position in the subcutaneous region. Samples were explanted at the two and five week endpoint for histology.

[00169] Anti-fibronectin immunofluorescent stains of hECM-coated polypropylene mesh showed that the ECM materials bound to and formed a uniform coating on the fibers of the mesh as compared to uncoated mesh. HECM coated mesh is suitable for implantable patches for medical applications, such as hernia repair and pelvic floor repair. The ECM materials were shown to coat the individual fibers of the mesh as shown through immunofluorescent staining with fibronectin antibodies which allows for improved cellular ingrowth.

[00170] hECM was implanted onto the chick chorioallantoic membrane (CAM) and stimulated a microvascular response as evidenced by new microvasculature growth. Additionally, hECM-coated nylon mesh subcutaneously implanted into mice for four weeks demonstrated improved biocompatibility versus uncoated nylon mesh. Specifically, fewer inflammatory cells and a thinner fibrous capsule were observed with the hECM-coated nylon fibers.

[00171] Biocompatibility evaluations were performed at two weeks and five weeks after implantation of hECM coated polypropylene mesh using hematoxylin and eosin stained samples. For FBGC analysis samples were blind-coded, evaluated using morphometry, separated into groups, statistically evaluated, then decoded. The number of foreign-body giant cells (FBGCs) were examined. A reduction in FBGCs for hECM coated polypropylene mesh as compared with non-coated mesh is evident. At the two week time point the mean FBGC count per sample was determined to be statistically higher (ANOVA bonferroni post-hoc analysis $p < 0.05$) for uncoated polypropylene (9.20 ± 2.03) versus hECM coated polypropylene (4.53 ± 0.89). At the five week time point the mean FBGC count per sample was determined to be higher, although not statistically significant, for uncoated polypropylene (10.95 ± 2.15) versus hECM coated polypropylene (8.17 ± 1.41).

[00172] The results indicated hECM coated polypropylene may reduce fibrous capsules. Fibrous encapsulation was evaluated at the two and five week time point using Trichrome stained samples. For capsule analysis samples were blind-coded, evaluated using morphometry, separated into groups, statistically evaluated, then decoded. At the two week time point the mean fibrous capsule thickness was not determined to be statistically higher (ANOVA bonferroni post-hoc analysis $p < 0.05$) for hECM coated polypropylene ($23.70 \pm 2.70 \mu\text{M}$), versus uncoated polypropylene ($19.70 \pm 3.00 \mu\text{M}$). At the five week time point, the mean fibrous capsule thickness was determined to be ($10.40 \pm 1.10 \mu\text{M}$) for hECM coated polypropylene, versus uncoated polypropylene ($12.30 \pm 1.20 \mu\text{M}$). Again, the differences in the hECM coated, versus uncoated polypropylene, were not determined to be statistically significant. Although, an important observation was found when evaluating the average percentage difference in fibrous encapsulation from the two to five week time points. The average percentage decrease in fibrous encapsulation from the two week to five week time points was 37.6% for hECM uncoated polypropylene, versus 56.1% for hECM coated polypropylene.

[00173] A mechanism of FBGC formation is the result of macrophage fusion in an immune response to implantable biomaterials such as polypropylene. These large multinucleated cells provide an effective means to quantitatively assess the inflammatory response to implantable biomaterials. A significant reduction in FBGC count per sample with human ECM coated versus uncoated polypropylene was observed at the two week time point. This data suggests that the human ECM surface coating may serve as an application for a variety of implantable devices.

[00174] Historically, the effectiveness and longevity of implantable devices have been challenged by specific immune responses including FBGC and fibrous capsule formation. Specifically FBGCs can excrete degradative agents such as superoxides and free radicals, as well as other degradative agents challenge device effectiveness, and longevity. These negative effects are especially significant since FBGCs are known to remain localized immediately around the implant for the duration of the presence of the implant. Fibrous capsule formation, which arises as a firm vascular collagen encapsulation around an implant, is designed to isolate foreign implantables from the host or host tissue. This response not only may cause discomfort for the patient in certain cases, but may shorten length of device viability and even diminish device effectiveness. Thus a coating that reduces FBGC and fibrous encapsulation is a highly desirable outcome for the longevity and function of implantable devices.

[00175] Findings that coating hECM on polypropylene will reduce FBGC presence, and potential reduction in fibrous encapsulation at the material-biological interface supports the need for further experimentation. Future evaluation may include time points of a longer duration to observe the changes in thickness of fibrous encapsulation with hECM coated and uncoated biomaterials. Additionally, evaluation with a continuum of biomaterials including dacron, nylon, stainless steel, and titanium in various in vivo environments, may elucidate further desirable hECM coated biomaterial outcomes.

EXAMPLE 7**USE OF EXTRACELLULAR MATRIX COMPOSITIONS FOR STIMULATION OF HAIR GROWTH**

[00176] This example illustrates the stimulation of hair growth by administration of ECM compositions.

[00177] Human hair follicle cells and cells taken from hair follicles were obtained to determine the ability of the ECM compositions described herein to stimulate and maintain hair forming ability. Hair follicle cells were obtained from Alderans Research International. Cells were cultivated in the presence of ECM. Analysis of the cells at four weeks and eight weeks of culture showed structures that resembled hair follicles as well as structures that resembled hair shafts. After two months of continuous culture, the cells remained alive and growing.

[00178] Cells cultured for four weeks in the presence of ECM composition were transplanted into mice. Four weeks after transplantation the cultured human hair follicles formed many large follicles as compared to control cells which showed only normal numbers of small resting hair follicles as observed using microscopic image analysis.

[00179] Based on these findings an in vivo hair growth trial was performed using human subject to determine de novo follicular hair regeneration. The study enrolled 24 men from the ages of 18 to 45 years having male pattern hair loss (MPHL). All members of the study group were without prior invasive or minimally invasive topical scalp surgery or topical treatment with MinoxidilTM or FinasterideTM. A Palomar Starluz 550p laser was used (1540-non-ablative and 2940 ablative). Study duration was designed upwards to 12 months for follow-up, baseline, and 5 months (following single sc injection). Following injection a three day wash-out period was observed and subjects only used CetaphilTM shampoo throughout study. A combination of lasing and microdermabrasion was performed prior to injection.

[00180] Subjects were administered vehicle admixed with hECM determined to have wnt protein activity and to include wnt 7a transdermally along with control vehicle and saline. End points of the study included a 7 point clinical grading system (3 blinded hair transplant surgeons), clinical macrophotography (follicle counts), 2mm punch biopsies and subject self assessment questionnaires.

[00181] Individual follicular units were analyzed for subjects. The follicles were counted at 12 weeks and compared to the baseline observed for the same individual at the start of the

study. Increases in follicular units were observed in subjects administered hECM without perturbation. For example, in one subject, treatment increased the number of apparent hair follicles from a baseline of 217 to 265 at 12 weeks. Total hair counts for the subject showed an increase from 307 to 360, an approximately 20% increase overall. The following increases in follicular counts were observed in other subjects as follows: subject 009 (baseline hair count 179, 12 weeks hair count 193, 5 month hair count 201); subject 013 (baseline hair count 266.5, 12 weeks hair count 267, 5 month hair count 294); subject 024 (baseline hair count 335.5, 12 weeks hair count 415, 5 month hair count 433). Further, 12 out of 13 patients (92.3%) administered hECM in the study showed efficacy at 12 weeks.

[00182] Additional hair parameters were analyzed throughout the study. At 12 weeks and 20 weeks relative hair count was analyzed as compared to baseline (0 weeks), terminal hair was analyzed as compared with baseline, and hair thickness was analyzed as compared with baseline. For example, in one subject, treatment increased hair count, terminal hair and follicle thickness, at 12 weeks, 22.4%, 27.8% and 23.9% respectively as compared to baseline. In another subject, treatment increased hair count, terminal hair and follicle thickness, at 12 weeks, 23.7%, 24.2% and 22.2% respectively as compared to baseline. In subject 009 (having baseline hair count 179, 12 weeks hair count 193, 5 month hair count 201, as above) treatment increased hair count, terminal hair and follicle thickness, at 12 weeks, 7.8%, 48.5% and 19.2% respectively; and, at 20 weeks, 12.9%, 33.0% and 21.1% respectively, as compared to baseline. In subject 013 (having baseline hair count 266.5, 12 weeks hair count 267, 5 month hair count 294, as above) treatment increased hair count, terminal hair and follicle thickness, at 12 weeks, 0.2%, 25.0% and 8.3% respectively; and, at 20 weeks, 10.3%, 41.4% and 23.0% respectively as compared to baseline. In subject 024 (baseline hair count 335.5, 12 weeks hair count 415, 5 month hair count 433 as above) treatment increased hair count, terminal hair and follicle thickness, at 12 weeks, 23.7%, 24.2% and 22.2% respectively; and, at 20 weeks, 29.1%, 5.9% and 17.3% respectively as compared to baseline respectively.

[00183] Of note, treated study members showed a significant increase in the number of terminal hairs and increase in thickness density at 3 months (84.6% of pts). Additionally, no adverse reactions observed, normal histology was observed and no hamartomas were observed.

[00184] The results point to use of hECM in additional applications, such as to prevent hair loss in patients post transplant and for eyebrow and eyelash growth. In hair transplant

patients, hair is known to fall out and generally take 4 to 5 months to return, thus, treatment with hECM would prevent hair loss in such individuals post transplant.

EXAMPLE 8

GENERATION OF HUMAN EXTRACELLULAR MATRIX COMPOSITIONS (HECM)

[00185] Human ECM composition was generated using newborn human fibroblasts. Fibroblasts were seeded onto beadlike structures conditioned with liquid media. Culture conditions were optimized without the need for fetal bovine serum. Within a few days, under embryonic culture conditions described herein, cells produced a dense embryonic –like ECM. Secretion of Wnt family proteins, as well as several growth factors was observed.

[00186] Cultures were grown to confluency. The cultures were subsequently exposed to sterile water to induce uniform lysing of the cells. The acellular hECM was then washed to ensure removal of all living cells and cellular debris and examined microscopically to confirm removal of cellular debris. Next, human fibroblasts were exposed to culture flasks coated with the hECM or plated onto a non-treated flask and then covered with a thick layer of matrix. The ECM proteins identified in the hECM are shown in Table 7.

Table 7. Extracellular Matrix Proteins Observed in Hecm

Matrix Protein	Function
Versican	structural, binds hyaluronic acid (HA) and collagen
Decorin	binds growth factors, influences collagen structure
Betaglycan	TGF- β Type III receptor
Syndecan	binds growth factors, enhances activity
Collagen Type I, II, III, V	major structural proteins of dermis
Fibronectin	cell adhesion, spreading, migration, motogenesis
Tenascin	induced in wound healing, control of cell adhesion

[00187] The hECM was observed to induce an increase of metabolic activity of the cells, as measured by increased enzymatic activity using the MTT assay. Human ECM, unlike mouse ECM, induced a dose-dependant increase in cellular metabolic activity as measured by MTT

assay. Cells were observed to rapidly and uniformly infiltrate the hECM overlay material. In addition, there was a dose-dependant increase in cell number in response to hECM, as measured by the Pico Green assay.

[00188] Known coatings, injectables, and implantable matrix products are typically either bovine collagens, porcine matrix proteins derived from the intestines or urinary bladder, hyaluronic acid, or human ECM derived from cadaver skin. While these products may offer benefits by creating a more physiologically equivalent environment, none are completely human and contain the entire range of matrix proteins found in young, developing tissue. The hECM produced contains the same ECM materials found in young, healthy tissue. It also was observed to support the active proliferation of human cells as well as rapid in-growth of cells. There are several advantages evident in using hECM in applications involving a human subject. For example, hECM promotes rapid host cell integration and improved healing (acts as normal scaffold for host cells and subsequent remodeling). Additionally, hECM eliminates the concern regarding viral transmission from non-human animal and human tissues (particularly BSE from bovine tissue and TSE from human tissue). Further, consistent product composition and performance is observed for hECM as compared to biologic products, particularly human dermis and fascia lata. Additionally, hECM reduces erosion of host tissues as compared to synthetic implants.

EXAMPLE 9

HUMAN FIBROBLAST DERIVED HYPOXIC CONDITIONED EXTRACELLULAR MATRIX FOR MEDICAL AESTHETIC APPLICATIONS

[00189] A double blind, randomized study of topical hECM administration post facial ablative laser surgery was conducted. The study enrolled 41 subjects between the ages of 40 and 60 years of age. All members of the study group were without prior invasive or minimally invasive surgery, or topical anti-aging treatments within the prior 12 months. The laser procedure included full fractional ablative laser procedure, peri-ocular, peri-oral and full face. A Palomar Starluz 550p laser was used (1540-non-ablative and 2940 ablative). Subjects were administered topical hECM compositions once a day (at different concentrations) or placebo vehicle for 14 days. End points of the study included clinical photography (3 blinded evaluations- dermatologists), transepidermal water loss (TEWL), punch biopsy, and evaluation of erythema, edema, and crusting.

[00190] The 10X strength hECM composition provided the most clinical improvement in symptoms as compared to the vehicle control (evaluations were conducted “blindly” by two cosmetic dermatologists, unrelated to any conduct of the clinical study). Photographic evaluation also indicated a reduction of erythema severity in several patients at days 3, 7 and 14.

[00191] Transepidermal water loss (TEWL) values were also evaluated 3, 7, and 14 days post laser treatment for all 41 subjects. The 10X strength hECM composition provided improvement in stratum corneum barrier function as noted at day 3, and day 7 as compared to the vehicle control. At day 7, the hECM composition is statistically significant at ($p < 0.05$) as compared to the vehicle control. This observation is consistent with the fact that there were subjects at day 7 post ablative fractional laser treatment that were demonstrating reepithelialization.

[00192] A double blind, randomized study of topical hECM administration for anti-aging (e.g., wrinkle reduction) was also conducted. The study enrolled 26 subjects between the ages of 40 and 65 years of age. All members of the study group were without prior invasive or minimally invasive surgery, or topical anti-aging treatments within the prior 12 months. Subjects were administered topical hECM compositions twice a day or placebo vehicle for 10 weeks. Endpoints of the study included clinical photography (2 blinded cosmetic dermatologists), corneometer-surface hydration, cutometer-elasticity, punch biopsy, molecular evaluation (Epidermal Genetic Information Retrieval (EGIR)).

[00193] Photographic evaluation of the facial area indicated a generation of lighter pigmentation, smoother skin texture, more evenly toned skin, and a reduction in the appearance of fine wrinkles and lines after 10 weeks of hECM administration.

[00194] Three dimensional profilometry image analysis of silicon replicas of the periorcular area was also performed for 22 of the 26 subjects. To perform the analysis a collimated light source was directed at a 25° angle from the plane of the replica. The replica was placed in a holder that fixed the direction of the tab position of the replica so that the replica could be rotated to align the tab direction normal or parallel to the incident light direction. The replicas were taken from the crow's feet area adjacent to each eye with the tab direction pointing toward the ear. The normal sampling orientation provided texture measurements sensitive to the major, expression-induced lines (crow's feet). The parallel sampling orientation provided texture measurements sensitive to the minor, fine lines.

[00195] A double blind, randomized study of topical hECM administration post facial ablative laser surgery was conducted. The study enrolled 49 subjects between the ages of 40 and 60 years of age. All members of the study group were without prior invasive or minimally invasive surgery, or topical anti-aging treatments within the prior 12 months. The laser procedure included full fractional ablative laser procedure, peri-ocular, peri-oral and full face. A Palomar Starluz 550p laser was used (1540-non-ablative and 2940 ablative). Subjects were administered topical hECM compositions twice a day or placebo vehicle for 14 days. End points of the study included clinical photography (3 blinded evaluations-dermatologists), mexameter and subject assessment.

[00196] Photographic evaluation of the facial area at days 1, 3, 5, 7 and 14 post surgery showed a clear reduction in erythema at every time point as compared to placebo.

[00197] The results of the studies indicated several beneficial characteristics of hECM containing topicals. Such benefits included 1) facilitating re-epithelization following resurfacing; 2) reduction of non-ablative and ablative fractional laser resurfacing symptoms (e.g., erythema, edema, crusting, and sensorial discomfort); 3) generating smooth, even textured skin; 4) generating skin moisturization; 5) reducing appearance of fine lines/wrinkles; 6) increasing skin firmness and suppleness; 7) reducing skin dyspigmentation and 8) reducing red, blotchy skin.

[00198] Although the invention has been described with reference to the above examples, it will be understood that modifications and variations are encompassed within the spirit and scope of the invention. Accordingly, the invention is limited only by the following claims.

What is claimed is:

1. A method of producing a composition comprising:
culturing cells in a suitable growth medium, wherein the cells produce and secrete a cell culture conditioned medium (CCM) composition, wherein the CCM promotes hair, lash and/or nail growth and/or promoting hair follicle development and/or activation or stimulation on an area of the skin when administered to the region of skin or tissue in need of growth or repair in a subject, thereby producing the composition.
2. The method of claim 1, wherein the cells are cultured under hypoxic or normoxic conditions.
3. The method of claim 2, wherein the hypoxic conditions comprise 1-5% O₂.
4. The method of claim 1, wherein the CCM composition is a soluble fraction.
5. The method of claim 1, wherein an ECM composition is produced as a non-soluble fraction.
6. A composition comprising growth factors or a CCM in combination with an active agent to promote hair, lash and/or nail growth in a subject.
7. The composition of claim 6, wherein the CCM is produced by the method of claim 1.
8. The composition of claim 6, wherein the active agent is a hair growth promoting agent.
9. The composition of claim 7, wherein the hair growth promoting agent is selected from the group consisting of: a potassium channel opener, an ATP-sensitive potassium channel (KATP opener), minoxidil, diazoxide, or phenytoin, a 5 α -reductase inhibitors, finasteride, dutasteride (e.g., Avodart), turosteride, bexlosteride, izonsteride, epristeride, epigallocatechin, MK-386, azelaic acid, FCE 28260, and SKF 105,1 1 1, ketoconazole, fluconazole, spironolactone, flutamide, diazoxide, 17-alpha-hydroxyprogesterone, 1 1-alpha-hydroxyprogesterone, ketoconazole, RU58841, dutasteride (marketed as Avodart), fluridil, or QLT-7704, an antiandrogen oligonucleotide, a prostaglandin F2a analogs, prostaglandin analogs, a prostaglandin, bimatoprost (e.g., Latisse, Lumigan), latanoprost (trade name Xalatan), travoprost (trade name Travatan), tafluprost, unoprostone, dinoprost (trade name Prostin F2 Alpha), AS604872, BOL303259X, PF3187207, carboprost (trade name Hemabate), kopexil (for example, the product Keranique™), CaC12, botulinum toxin A, adenosine, ketoconazole, DoxoRx, Docetaxel, FK506, GP 1 1046, GP1 151 1 1, LGD 1331, ICX-TRC, MTS-01, NEOSHIOL, HYG-102440, HYG-410, HYG-420, HYG-430, HYG-440, spironolactone, CB- 03-01 , RK-023, Abatacept, Viviscal®, MorrF, ASC-J9, NP-619, AS

101, Metron-F-1, PSK 3841, Targretin (e.g., 1% gel), MedinGel, PF3187207, BOL303259X, AS604872, THG1 1331, PF-277343, PF-3004459, Raptiva, caffeine, coffee, a herb (such as, e.g., saw palmetto, glycine soja, Panax ginseng, Castanea Sativa, Arnica Montana, Hedera Helix Geranium Maculatum), triamcinolone acetonide, a topical irritant (e.g., anthralin) or sensitizer (e.g., squaric acid dibutyl ester [SADBE] or diphenyl cyclopropanone [DPCP]), clomipramine, unsaturated fatty acids (e.g., gamma linolenic acid), a fatty acid derivative, a thickener (such as, e.g., carbomer, glycol distearate, cetearyl alcohol), a hair loss concealer, niacin, nicotinate esters and salts, adenosine, methionine, an androgen receptor inhibitor, a copper peptide, a compound with superoxide dismutation activity, an agent that increases nitric oxide production (e.g., arginine, citrulline, nitroglycerin, amyl nitrite, or sildenafil (Viagra)), a compound that mobilizes bone marrow-derived stem cells (e.g., growth factors such as G-CSF and/or chemical agents such as plerixafor (Mozobil®)), a compound that regulates the differentiation of stem cells into gender-specific specialized human hair follicles (e.g., finasteride, fluconazole, spironolactone, flutamide, diazoxide, 17 β -alpha-hydroxyprogesterone, ketoconazole, RU58841, dutasteride, fluridil, or QLT-7704, an antiandrogen oligonucleotide, cyoctol, topical progesterone, topical estrogen, cyproterone acetate, ru58841, combination 5 α -reductase inhibitors, oral contraceptive pills), an antiestrogen, an estrogen, or estrogenlike drug, an anti-oxidants (e.g., glutathione, ascorbic acid, tocopherol, uric acid, or polyphenol antioxidants), inhibitors of reactive oxygen species (ROS) generation (e.g., superoxide dismutase inhibitors; stimulators of ROS breakdown, such as selenium; mTOR inhibitors, such as rapamycin; or sirtuins or activators thereof, such as resveratrol, or other SIRT1, SIRT3 activators, or nicotinamide inhibitors), an agent that induces an immune response or causes inflammation (e.g., tetanus toxoid, topical non-specific irritants (anthralin), or sensitizers (squaric acid dibutyl ester [SADBE] and diphenyl cyclopropanone [DPCP]), and an antiapoptotic compound.

10. The composition of claim 6, wherein the composition comprises growth factors or CCM and Minoxidil.

11. The composition of claim 10, wherein minoxidil is present at a concentration from about 0.5% to about 5% by weight.

12. The composition of claim 11, wherein minoxidil is present at a concentration from about 1% by weight.

13. The composition of claim 11, wherein minoxidil is present at a concentration from about 2% by weight.
14. The composition of claim 11, wherein minoxidil is present at a concentration from about 5% by weight.
15. The composition of claim 11, wherein minoxidil is in the form of a pharmaceutically acceptable derivative.
16. The composition of claim 15, wherein the pharmaceutically acceptable derivative comprises pharmaceutically acceptable salts, solvates, hydrates, isomers, esters, tautomers, anhydrates, enantiomers, complexes, polymorphs or prodrugs.
17. The composition of claim 6, wherein the composition comprises growth factors or CCM and bimatoprost.
18. The composition of claim 17, wherein bimatoprost is present at a concentration from about 0.01% to about 5% by weight.
19. The composition of claim 18, wherein bimatoprost is present at a concentration of about 0.1% by weight.
20. The composition of claim 18, wherein bimatoprost is present at a concentration of about 1% by weight.
21. The composition of claim 18, wherein bimatoprost is present at a concentration of about 3% by weight.
22. The composition of claim 18, wherein bimatoprost is present at a concentration of about 5% by weight.
23. The composition of claim 18, wherein bimatoprost is provided as a pharmaceutically acceptable salt.
24. The composition of any of claims 6-23 further comprising penetration enhancers.
25. The composition of any of claims 6-24 further comprising a pharmaceutically acceptable excipient.
26. The composition of claim 6, wherein the composition is adapted for topical application to mammalian skin as a foam, wherein said foam comprises bimatoprost and/or minoxidil, and at least one surfactant, wherein the surfactant optionally includes a foam stabilizer; an aqueous-alcohol solvent, and wherein said aqueous-alcohol solvent comprises water and an alcohol.
27. A composition produced by the method of claim 1.

- 28.** A method of stimulating hair, nail or lash growth and/or promoting hair follicle development and/or activation or stimulation on an area of the skin of a subject comprising contacting the hair (scalp), nail or lash or adjacent areas thereof with a composition of any of claims 6-24 under conditions that allow for hair, nail or lash growth and/or promoting hair follicle development and/or activation or stimulation on an area of the skin in the subject.
- 29.** The method of claim 28, wherein the scalp, dermis or hair follicle is contacted with the composition.
- 30.** A topical pharmaceutical composition comprising minoxidil, growth factors and/or a CCM and at least one or more pharmaceutically acceptable excipients.
- 31.** The topical pharmaceutical composition of claim 30, wherein the CCM is derived from the method of claim 1.
- 32.** The topical pharmaceutical composition of claim 30, wherein minoxidil is in the form of a pharmaceutically acceptable derivative.
- 33.** The topical pharmaceutical composition of claim 32, wherein the pharmaceutically acceptable derivative comprises pharmaceutically acceptable salts, solvates, hydrates, isomers, esters, tautomers, anhydrides, enantiomers, complexes, polymorphs or prodrugs.
- 34.** The topical pharmaceutical composition of claim 30, wherein minoxidil is present at a concentration from about 0.5% to about 5% by weight.
- 35.** The topical pharmaceutical composition of claim 30, wherein minoxidil is present at a concentration of about 1%, 2% or 5% by weight.
- 36.** The topical pharmaceutical composition of claim 30 further comprising bimatoprost.
- 37.** A topical pharmaceutical composition comprising bimatoprost, growth factors and/or a CCM and at least one or more pharmaceutically acceptable excipients.
- 38.** The topical pharmaceutical composition of claim 37, wherein the CCM is derived from the method of claim 1.
- 39.** The composition of claim 37, wherein bimatoprost is present at a concentration from about 0.01% to about 5% by weight.
- 40.** The composition of claim 39, wherein bimatoprost is present at a concentration of about 0.1%, 1%, 3% or 5% by weight.
- 41.** The composition of claim 37, wherein bimatoprost is provided as a pharmaceutically acceptable salt.
- 42.** The method of claim 1, wherein the cells are fibroblasts.

- 43. The method of claim 1, wherein the cells are grown under 1-, 2-, or 3-dimensional conditions.
- 44. The method of claim 43, wherein the cells are grown as monolayers.
- 45. The method of claim 43, wherein the cells are grown on beads.
- 46. The method of claim 43, wherein the cells are grown on mesh.
- 47. The composition of claims 30 or 37, further comprising vitamins or other supplements.
- 48. The composition of claim 47, wherein the vitamins or other supplements are selected from Vitamin D, niacin, biotin, Vitamin A, Vitamin C, B Vitamins including Vitamin B12 or other supplements.

INTERNATIONAL SEARCH REPORT

International application No.

PCT/US2019/044405

A. CLASSIFICATION OF SUBJECT MATTER

IPC(8) - A61K 35/33; A61K 35/00; A61K 35/12; A61Q 7/00 (2019.01)

CPC - A61K 35/33; A61K 35/00; A61K 35/12; A61Q 7/00; C12N 2502/1323 (2019.08)

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

See Search History document

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

USPC - 435/41; 435/70.1; 435/70.3 (keyword delimited)

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

See Search History document

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	US 2010/0047305 A1 (NAUGHTON et al) 25 February 2010 (25.02.2010) entire document	1-5, 27, 42-46
A	US 5,130,142 A (WONG et al) 14 July 1992 (14.07.1992) entire document	1-5, 27, 42-46
A	US 2007/0036742 A1 (ROUFS et al) 15 February 2007 (15.02.2007) entire document	1-5, 27, 42-46

☐ Further documents are listed in the continuation of Box C.☐ See patent family annex.

* Special categories of cited documents:

"A" document defining the general state of the art which is not considered to be of particular relevance

"E" earlier application or patent but published on or after the international filing date

"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)

"O" document referring to an oral disclosure, use, exhibition or other means

"P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art

"&" document member of the same patent family

Date of the actual completion of the international search

08 November 2019

Date of mailing of the international search report

27 NOV 2019

Name and mailing address of the ISA/US

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INTERNATIONAL SEARCH REPORT

International application No.

PCT/US2019/044405

Box No. II Observations where certain claims were found unsearchable (Continuation of item 2 of first sheet)

This international search report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☐ Claims Nos.:
because they relate to subject matter not required to be searched by this Authority, namely:
2. ☐ Claims Nos.:
because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out, specifically:
3. ☒ Claims Nos.: 7, 9, 24, 25, 28, 29, 31, 38
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box No. III Observations where unity of invention is lacking (Continuation of item 3 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

See extra sheet(s).

1. ☐ As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims.
2. ☐ As all searchable claims could be searched without effort justifying additional fees, this Authority did not invite payment of additional fees:
3. ☐ As only some of the required additional search fees were timely paid by the applicant, this international search report covers only those claims for which fees were paid, specifically claims Nos.:
4. ☒ No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:
1-5, 27, 42-46

Remark on Protest

- ☐ The additional search fees were accompanied by the applicant's protest and, where applicable, the payment of a protest fee.
- ☐ The additional search fees were accompanied by the applicant's protest but the applicable protest fee was not paid within the time limit specified in the invitation.
- ☐ No protest accompanied the payment of additional search fees.

INTERNATIONAL SEARCH REPORT

International application No.

PCT/US2019/044405

Continued from Box No. III Observations where unity of invention is lacking

This application contains the following inventions or groups of inventions which are not so linked as to form a single general inventive concept under PCT Rule 13.1. In order for all inventions to be examined, the appropriate additional examination fees need to be paid.

Group I: claims 1-5, 27, 42-46 are drawn to method of producing a compositions and compositions thereof.

Group II: claims 6, 8, 10-23, 26, 30, 32-37, 39-41, 47, and 48 are drawn to compositions and topical pharmaceutical compositions.

The inventions listed in Groups I and II do not relate to a single general inventive concept under PCT Rule 13.1, because under PCT Rule 13.2 they lack the same or corresponding special technical features for the following reasons:

The special technical features of Group I, method of producing a compositions and compositions thereof, are not present in Group II; and the special technical features of Group II, compositions and topical pharmaceutical compositions, are not present in Group I.

Additionally, even if Groups I and II were considered to share the technical features of a composition comprising growth factors or a CCM in combination with an active agent to promote hair, lash and/or nail growth in a subject, and at least one or more pharmaceutically acceptable excipients, these shared technical features do not represent a contribution over the prior art as disclosed by US 5,130,142 A to Wong et al.

US 5,130,142 A to Wong et al. teach a composition (Abstract and Claim 1, A composition for regulating hair growth...) comprising growth factors or a CCM (Claim 1, ...a. a safe and effective amount of a supernatant derived from a culture of epithelial cells which comprises a growth stimulating factor...; Col. 4, Lns. 1-43, Example 1, Preparation of Epidermal Cell Derived Growth Stimulating Factor, The epidermal cells are incubated in this medium for up to 48 hours at which time the culture medium is decanted. This medium ("Conditioned Defined Medium") is then centrifuged at 100,000xG for 30 minutes. The medium is fractionated and concentrated by ultrafiltration...; Col. 8, Lns. 37-50, Hair Grown Regulating Compositions,... comprising a safe and effective amount of an epithelial cell derived hair growth factor...and a pharmaceutically acceptable carrier...; Col. 17, Example XIX, see table for ingredients.) in combination with an active agent to promote hair, lash and/or nail growth in a subject (Claim 9, The composition of claim 1 which additionally comprises a safe and effective amount of minoxidil...; Col. 17, Example XIX, see table for ingredients...; Col. 13, Lns. 55-63, The composition according to the invention can also optionally comprise other hair growth stimulants capable of functioning in different ways to enhance the benefit of the growth stimulating factors. Examples of other substances which themselves possess the ability to regulate hair growth include, but are not limited to, minoxidil...), and at least one or more pharmaceutically acceptable excipients (Col. 8, Lns. 37-50, Hair Grown Regulating Compositions,... comprising a safe and effective amount of an epithelial cell derived hair growth factor... and a pharmaceutically acceptable carrier...).

The inventions listed in Groups I and II therefore lack unity under Rule 13 because they do not share a same or corresponding special technical feature.