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What Kind of Signaling Maintains Pluripotency and Viability in Human-Induced Pluripotent Stem Cells Cultured on Laminin-511 with Serum-Free Medium?

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Abstract

Xeno-free medium contains no animal-derived components, but is composed of minimal growth factors and is serum free; the medium may be supplemented with insulin, transferrin, and selenium (ITS medium). Serum-free and xeno-free culture of human-induced pluripotent stem cells (hiPSCs) uses a variety of components based on ITS medium and Dulbecco's modified Eagle's medium/Ham's nutrient mixture F12 (DMEM/F12) that contain high levels of iron salt and glucose. Culture of hiPSCs also requires scaffolding materials, such as extracellular matrix, collagen, fibronectin, laminin, proteoglycan, and vitronectin. The scaffolding component laminin-511, which is composed of $\alpha 5$, $\beta 1$, and $\gamma 1$ chains, binds to $\alpha 3\beta 1$, $\alpha 6\beta 1$, and $\alpha 6\beta 4$ integrins on the cell membrane to induce activation of the PI3K/AKT- and Ras/MAPK-dependent signaling pathways. In hiPSCs, the interaction of laminin-511/ $\alpha 6\beta 1$ integrin with the cell-cell adhesion molecule E-cadherin confers protection against apoptosis through the Ras homolog gene family member A (RhoA)/Rho kinase (ROCK) signaling pathway (the major pathways for cell death) and the proto-oncogene tyrosine-protein kinase Fyn (Fyn)-RhoA-ROCK signaling pathway. The expression levels of $\alpha 6\beta 1$ integrin and E-cadherin on cell membranes are controlled through the activation of insulin receptor/insulin, FGF receptor/FGF2, or activin-like kinase 5 (ALK5)-dependent TGF- β signaling. A combination of growth factors, medium constituents, cell membrane-located E-cadherin, and $\alpha 6\beta 1$ integrin-induced signaling is required for pluripotent cell proliferation and for optimal cell survival on a laminin-511 scaffold. In this review, we discuss and explore the influence of growth factors on the cadherin and integrin signaling pathways in serum-free and xeno-free cultures of hiPSCs during the preparation of products for regenerative medicinal therapies. In addition, we suggest the optimum serum-free medium components for use with laminin-511, a new scaffold for hiPSC culture.

Key words: cellular biology; cell culture; extracellular matrix; growth factor; stem cells

Introduction

Initially, human pluripotent stem cell (hPSC) lines were grown in a coculture system using a layer of mitotically inactivated mouse embryonic fibroblast (MEF) feeder cells.¹⁻³ However, to be clinically compliant, human-induced pluripotent stem cells (hiPSCs) need to be cultured in a medium that is serum free, chemically defined, and free of animal products (xeno free). Serum-free medium is generally composed of minimal growth factors, bovine serum albumin (BSA) and Fraction V protein; the insulin, transferrin, and selenium (ITS) medium is

supplemented with ITS. Considerable efforts are being made to develop culture systems that do not require animal feeder cells and use xeno-free medium.⁴⁻¹¹

hPSCs cultured in MEF-conditioned medium (MEF-CM), which is based on ITS medium, can be maintained on Matrigel,⁹ a complex mixture of matrix proteins derived from Engelbreth-Holm-Swarm mouse tumors¹²⁻¹⁴; this mixture includes laminin-111, collagen IV, fibronectin, and vitronectin and supports robust hPSC growth.^{8,14-16} Moreover, a combination of collagen IV, fibronectin, laminin, and vitronectin can be used to

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replace Matrigel for deriving and expanding hPSCs under defined culture conditions.⁸ A laminin-511 scaffold can bind $\alpha 3\beta 1$, $\alpha 6\beta 1$, and $\alpha 6\beta 4$ integrins and is able to maintain hPSCs in the undifferentiated state in serum- and xeno-free medium under feeder cell-free culture conditions.^{17–21}

In this review, we consider the effects of medium components and intracellular signal transduction in cultures using serum- and xeno-free medium and a feeder cell-free culture system. Substrates obtained from recombinant human laminin-511 offer significant improvements for establishing defined conditions for adhesion cultures of hPSCs under clinically compliant settings. We suggest that choosing the components of the culture medium and the scaffolding materials can ensure maintenance of pluripotency and cell survival through PI3K/AKT and Fyn-RhoA-ROCK signaling pathways in hiPSC culture.

Signaling Pathways in Serum-Free Culture Systems

hiPSCs easily lose pluripotency due to a component in serum. Therefore, hiPSCs have to be cultured using serum-free culture medium. Initially, researchers examined serum albumin, a primary component of serum, and found that Dulbecco's modified Eagle's medium/Ham's nutrient mixture F12 (DMEM/F12) culture medium supplemented with serum albumin, FGF2, and the ITS medium components. ITS functioned similarly as serum-based culture media for hiPSCs. However, the serum-free medium used for the culture of hiPSCs initially contained a component derived from an animal. In addition, mouse feeder cells were also used.

FGF2/FGFR signaling

FGFR1 (fibroblast growth factor receptor 1) is the predominant FGFR in hiPSCs. Binding of FGF2 activates FGFR and major signal transduction pathways, such as all three branches of the PI3K and AKT pathways, the mitogen-activated protein kinase (MAPK) pathway, and the extracellular signal-regulated kinase (ERK)/MAPK pathway. FGF2 activates the MAP kinases ERK1/ERK2, and also PI3K/AKT and c-Jun, and stimulates signaling cascades by activation of activin A.^{22,23} Induction of $\alpha 2\beta 1$, $\alpha 3\beta 1$, $\alpha 6\beta 1$, and $\alpha 6\beta 4$ integrin transcription requires the FGF2-FGFR signal pathway and is reinforced by the ALK4-activin A signaling pathway. FGF2 signaling may act through the PI3K pathway to regulate the synthesis of pericellular laminin-332, -511, -521, and collagen IV isotypes and provide a framework for the basement membrane of hPSCs.²⁴ Blocking the

FGF2 pathway in hPSCs can lead to differentiation.^{6,8,25} Extracellular matrix (ECM) modulated signals may converge with FGF2 signaling and be involved in the maintenance of pluripotency (Fig. 1).

Bovine serum albumin

BSA has many roles in cell culture; it acts as a lipid carrier, provides physical protection,^{26,27} and prevents toxic effects by β -mercaptoethanol (a reducing agent normally required for culturing cells). In addition, BSA acts as a nutrient for MEFs. Many proteins secreted by MEFs have been identified recently in mass spectrometric analyses.^{28–30} MEF secreted heparan sulfate proteoglycans (HSPG), fibronectin, transforming growth factor β (TGF- β), and insulin-like growth factors 1 (IGF1) and 2 (IGF2). In addition, it has been shown that critical products produced by MEF feeder layers include activin A³¹ (Fig. 1).

Insulin/IGF-R signaling

The IGF-1R signaling axis is critical to cell proliferation and glucose metabolism and also induces E-cadherin transcription in hiPSCs. Upon binding insulin, IGF-1R is activated through autophosphorylation. PI3K activation then leads to an increase in phosphatidylinositol 3,4,5-trisphosphate (PIP3), which results in the activation of the critical protein AKT.²⁵ This signaling axis is commonly referred to as the PI3K/AKT, nuclear factor κ B (NF- κ B), and jun N-terminal kinases (JNK) pathway and as c-Jun phosphorylation of IGF-1R signaling; it is ultimately responsible for the maintenance of hPSCs and hiPSCs.^{32,33} In parallel, IGF-1R signaling also promotes cell proliferation through the Ras/MAPK pathway (Fig. 1).

Transferrin

Transferrin, a glycoprotein, is required in the culture medium to act primarily as an iron transporter and secondarily to bind endogenous iron. Transferrin enables the cells to take up oxygen and also stimulates growth-related enzyme activities. Two types of transferrin are generally added to culture media: iron-saturated transferrin (holo-transferrin) and iron-deficient transferrin (apo-transferrin) (Fig. 1).

Selenium

Selenium, in the form of selenoproteins in particular glutathione peroxidase (Glu.Px), is an essential component of the human cellular antioxidant defense system. It also has anti-inflammatory effects by its ability to scavenge free radicals.³⁴ As is evident from Figure 1, selenium

FIG. 1. Signaling pathways in serum-free on feeder culture systems. BSA acts as a nutrient for MEFs. MEFs secrete heparan sulfate proteoglycans (HSPG), fibronectin, TGF- β , IGF1, and IGF2. HSA acts as a nutrient for hiPSCs. Albumin in the culture medium promotes expression of collagen I, collagen IV, and HSPG that detoxify β -mercaptoethanol. Collagen I binds to $\alpha 2\beta 1$ integrin and transduces Ras function to activate PI3K binding to AKT; activation of the latter pathway promotes self-renewal in hiPSCs. Collagen IV binds to $\alpha 2\beta 1$ integrin and transduces a RhoGAP function to activate PI3K binding to AKT; activation of the latter pathway promotes cell growth in hiPSCs. With regard to downstream functions in the IGF1R and ALK5 pathways of hiPSCs, activation induces E-cadherin transcription that recruits the PI3K/AKT and ERK/MAPK. This suggests that both PI3K/AKT and MAPK/ERK signaling are required for insulin- and TGF- β -induced E-cadherin upregulation on the cell membrane. Glucose promotes fibronectin and TGF- β manifestation through c-Jun signaling. TGF- β is activated by $\alpha V\beta 6$ integrin. The pathways that IGF1R and ALK5 activate through unidentified growth factors, which are present in the culture medium for hiPSCs, are sufficient to maintain pluripotency in hiPSCs. Transferrin regulates cellular iron uptake and, thus, is an essential culture medium ingredient. Selenium is a component of some glutathione peroxidases; these enzymes prevent oxidative damage in cells by inducing the reduction of lipid hydroperoxides. With regard to downstream functions in the FGFR and ALK4 pathways of hiPSCs, the induction of $\alpha 2\beta 1$, $\alpha 3\beta 1$, $\alpha 6\beta 1$, and $\alpha 6\beta 4$ integrin transcription requires the FGF2-FGFR signal pathway and is reinforced by ALK4-activin A signal pathway. Collagen I and HSPG concentrate on FGF2 in the vicinity of FGFR to activate laminin-332, -511, -521, and collagen IV transcription through the PI3K/AKT signaling pathway. Fibronectin is secreted by hiPSCs and binds with $\alpha 8\beta 1$ integrin to promote cell survival of hiPSCs through PI3K/AKT signaling. In addition, fibronectin can also bind to $\alpha V\beta 1$ integrin and $\alpha 5\beta 1$ integrin to promote secretion of laminin-332, -511, -521, and collagen IV from hiPSCs. The pathways activated by fibronectin-integrin signaling are responsible for the secretion of ECM, which maintains pluripotency of hiPSCs. ALK4, activin-like kinase receptor-4; BSA, bovine serum albumin; ECM, extracellular matrix; ERK, extracellular signal-regulated kinase; hiPSCs, human-induced pluripotent stem cells; FGFR, fibroblast growth factor receptor; HSA, human serum albumin; IGF, insulin-like growth factor; IGF1R, insulin-like growth factor-1 receptor; MAPK, mitogen-activated protein kinase; MEF, mouse embryonic fibroblast; PI3K, phosphatidylinositol 3-kinase; TGF- β , transforming growth factor β .

is a component of some glutathione peroxidases; these enzymes prevent oxidative damage in cells by inducing the reduction of lipid hydroperoxides.

Glucose

In hPSCs, the c-jun N-terminal kinase (JNK) inhibitor SP600125 blocks fibronectin expression induced by high glucose and inhibits cell cycle regulatory protein expression induced by high glucose-mediated TGF- β expression.³⁵ High glucose levels (4.5 g/L) also increase

intracellular calcium (Ca^{2+}) concentration, PKC phosphorylation, and TGF- β expression in hPSCs. Thus, c-Jun is necessary for expression of fibronectin and TGF- β (Fig. 1).

Signaling Pathways in Feeder Cell-Free Culture Systems

When hiPSCs are cultured on an MEF feeder layer, the MEFs provide proteins, such as activin A, TGF- β , IGF1 and 2 (IGF2), and glycans, including HSPG, fibronectin,

collagen I, collagen IV, and laminin. These proteins and glycans are necessary for hiPSC attachment, proliferation, and self-renewal³⁶ (Fig. 1). Matrigel, which includes laminin-111, collagen IV, fibronectin, and vitronectin, can be used to replace the MEF feeder layer and support robust hPSC growth.^{12–14}

Activin A/ALK4 signaling

Activin A has a role in a wide range of cellular processes from cell proliferation and differentiation to apoptosis. Initially, activin A binds to type II activin A receptors (ActRIIA or ActRIIB) and then recruits type IB activin A receptor (ALK4). ALK4 interacts with and phosphorylates the SMAD family members 2 and 3 (SMAD2 and SMAD3). Vallier et al.^{37–39} demonstrated that the cooperation of the activin A pathway and FGF2 is necessary for the maintenance of pluripotency in hPSCs.⁴⁰ When active, PI3K/AKT establishes conditions where activin A/SMAD2/3 stimulate self-renewal in hiPSCs by activating target genes, including Nanog.^{41,42} This stimulatory effect involves the interaction of activin A and SMAD2/3, and may also require upregulation of FGF2 pathways^{39,43} (Fig. 2).

TGF- β /ALK5 signaling

TGF- β type 1 receptor, also known as ALK5, phosphorylates SMAD2/3 and forms a complex with SMAD4 that translocates to the nucleus. The complex

induces transcription of TGF- β -dependent genes that regulate signaling pathways involved in pluripotency and self-renewal, such as the MAPK and PI3K/AKT pathways. Signaling through these pathways supports proliferation and continued pluripotency in hiPSCs⁴⁴ (Fig. 2).

Signaling Pathways in Xeno-Free Culture Systems

Animal serum contains xenogeneic antigens, and cells cultured in serum have the potential to induce immune reactions in recipients and also raise the risk of transmission of animal viruses, prions, and zoonotic infections. Thus, animal serum is not desirable for clinical cell production or for transplantation therapies. Laminin-511 scaffold can bind $\alpha 3\beta 1$, $\alpha 6\beta 1$, and $\alpha 6\beta 4$ integrins and can be used to maintain hPSCs in the undifferentiated state under serum and xeno-free conditions in a feeder cell-free culture system.^{17–21}

Recombinant human albumin

Human albumin has many biological and physical roles in hiPSC culture and is necessary for StemFit medium (Ajinomoto Co., Inc.) to enable the efficient formation of colonies by hiPSCs on recombinant laminin-511.⁴⁵ However, albumin in the culture medium promotes the secretion of ECM molecules such as collagen I, collagen IV, and HSPG. Consequently, protein lytic enzymes such as the trypsin

FIG. 2. Signaling pathways in serum-free and feeder cell-free culture systems (using MEF-CM on Matrigel). Matrigel includes laminin-111, collagen IV, fibronectin, and vitronectin and supports robust hPSC growth. Collagen IV binds to $\alpha 2\beta 1$ integrin and transduces a RhoGAP function to activate PI3K binding to AKT; activation of the latter pathway promotes cell growth in hiPSCs. Vitronectin supports the maintenance of hPSCs through $\alpha V\beta 5$ integrin. $\alpha V\beta 1$ integrin is able to recognize vitronectin and fibrinogen. Fibronectin can also bind to $\alpha V\beta 1$ integrin and $\alpha 5\beta 1$ integrin to promote secretion of laminin-332, -511, -521, and collagen IV from hiPSCs. hPSC, human pluripotent stem cells; MEF-CM, MEF-conditioned medium.

are necessary to break the strong cell-to-plate adhesion (Fig. 3).

Recombinant vitronectin scaffold

Recombinant vitronectin as a glycoprotein can function as a chemically defined matrix component in hiPSCs. Vitronectin is a functionally defined substrate that supports hiPSCs self-renewal through $\alpha V\beta 5$ integrin in albumin-free medium such as Essential 8 medium (Thermo Fisher Scientific). It is possible to dislodge cells from the culture plate using only EDTA; thus, integrins on the cell surface are not damaged by trypsin treatment.

Recombinant laminin-511 scaffold

The scaffolding component laminin-511, which is composed of $\alpha 5$, $\beta 1$, and $\gamma 1$ chains, binds to $\alpha 3\beta 1$, $\alpha 6\beta 1$, and $\alpha 6\beta 4$ integrins on the cell membrane to induce activation of the PI3K/AKT- and Ras/MAPK-dependent signaling pathways. In hiPSCs, the interaction of laminin-511/ $\alpha 6\beta 1$ integrin with the cell-cell adhesion molecule E-cadherin confers protection against apoptosis through the Ras homolog gene family member A (RhoA)/Rho kinase (ROCK) signaling pathway (the major pathway for cell death) and the proto-oncogene tyrosine-protein kinase Fyn (Fyn)-RhoA-ROCK signaling pathway.

Signaling Pathways in Cultures Using Laminin-511 Cadherin

Cell adhesion molecules (CAMs), such as cadherins and integrins, stimulate intercell adhesion. The “classical” CAMs include epithelial (E-), neural (N-), and placental (P-) cadherins and, as their names suggest, they tend to be expressed in a tissue-specific pattern.^{46–52} In addition to their role in cell-cell adhesion, cadherins may also function in signaling pathways between the cell membrane and the nucleus.⁵³

E-cadherin signaling. E-cadherin, a Ca^{2+} -dependent cell-cell adhesion molecule,^{50,54} is stabilized at the cell surface by its link to the actin cytoskeleton through β -catenin and α -catenin and is essential for PSC intercellular adhesion and colony formation.^{55,56} It has been shown that functional interaction between Rac1, a small Rho family GTPase, and E-cadherin is responsible for regulating hiPSC self-renewal.^{57,58} E-cadherin-mediated adhesion is commonly associated with β -catenin signaling and also stimulates PI3K/AKT signaling.^{59,60} The ability of the PI3K/AKT pathway to promote cell survival indicates a broader role for Rho family GTPases through p120-catenin.^{61,62} The PI3K inhibitor wortmannin prevents E-cadherin-dependent Rac1 activation, but does not interfere with the intracellular distribution of Rac1

FIG. 3. Signaling pathways in serum-free and feeder cell-free culture systems (on laminin-511). Cadherin is activated by a cell junction. When N-cadherin is activated, transcriptional enhancement of FGFR occurs. As a result, the FGF2 signal pathway is activated. This signaling pathway has an important role in the expression of FGFR attachment to the cell membrane. In cultures using a laminin-511 scaffold, the E-cadherin- $\alpha 6\beta 1$ integrin signal pathway is required to control cell death in hiPSCs. PI3K signaling promotes expression of Rac1. Rac1 binds to cadherins, and cadherins prevent endocytosis. When $\alpha 6\beta 1$ integrin is combined with laminin-511, Fyn-RhoA-ROCK signaling is induced. As a result, hiPSC death caused by ROCK is repressed. Laminin-511 can directly bind to $\alpha 3\beta 1$ integrin and, thereby, activate the PI3K/AKT signaling pathway through Ras. In addition, linkage of $\alpha 6\beta 4$ activates the MAP kinase pathways directly through Ras.

or E-cadherin.⁶³ The activation of Rac1 through PI3K by E-cadherin-mediated cell–cell adhesion involves at least two steps as follows: (1) recruitment of Rac1 to E-cadherin mediates cell–cell contact sites and (2) Rac1 activation by a guanine nucleotide exchange factor.⁶⁴ E-cadherin levels at adherens junctions are regulated by endocytosis and recycling; activated Rac1 reduces E-cadherin endocytosis and, thus, increases the levels of cell surface E-cadherin and, consequently, promotes cell–cell adhesion. Ectopic overexpression of E-cadherin also increases the rate of survival in dissociated hPSCs.⁶⁵ However, when dissociated hPSCs are grown on E-cadherin-coated plates, they form membrane protrusions and have a lower rate of survival. Thus, E-cadherin interactions alone are not the only factor in cell survival.⁶⁶ Transcription factors for E-cadherin expression appear to act downstream in various signaling pathways, such as in the TGF- β , FGF2, NF- κ B, and integrin cascades⁶⁷ (Fig. 3).

N-cadherin signaling. N-cadherin induces FGFR upregulation⁶⁸ and has also been shown to stimulate FGF2 signaling.⁶⁹ N-cadherin adhesion can lead to PI3K-mediated activation of AKT,^{70,71} and activated AKT signaling can stimulate β -catenin signaling. The E-cadherin function can be replaced by N-cadherin in the rescue of the undifferentiated pluripotent iPSC state⁷² (Fig. 3).

P-cadherin signaling. P-cadherin controls cell-to-laminin adhesion by modulating the expression and activation of the laminin receptor $\alpha 6\beta 4$ integrin heterodimer.⁷³ Importantly, it has been reported that $\alpha 6\beta 4$ integrin (adhering to laminin-332-containing $\alpha 3$, $\beta 3$, and $\gamma 2$ chains) promotes survival and invasion by activating the PI3K/AKT pathway.^{74–76} $\alpha 6$ Integrin activation induces P-cadherin transcription.⁷⁷ P-cadherin results in the translocation of p120ctn from the plasma membrane to the cytoplasm, thereby activating Rac1.⁷⁸

Integrin

Integrins are transmembrane proteins that have a wide range of functions, including adhesion and cell–cell interactions.⁷⁹ They are heterodimeric molecules consisting of α and β chains. The 24 different heterodimeric combinations of integrin subunits in vertebrates have different ligand binding preferences. Based on their ligand specificities, the integrin family can be divided into three subgroups with respect to hPSC culture: collagen receptors ($\alpha 2\beta 1$ and $\alpha 11\beta 1$), laminin receptors ($\alpha 3\beta 1$, $\alpha 6\beta 1$, $\alpha 7\beta 1$, and $\alpha 6\beta 4$), and RGD (Arg-Gly-

Asp) receptors ($\alpha 5\beta 1$, $\alpha 8\beta 1$, $\alpha V\beta 1$, $\alpha V\beta 5$, $\alpha V\beta 6$, and $\alpha IIb\beta 3$).^{36,80–82}

Collagen receptors ($\alpha 2\beta 1$ and $\alpha 11\beta 1$)

Total collagen complex, including types I, II, III, and IV, can induce an increased proliferative capacity in iPSCs.⁸³ The effect of collagen I and IV on cell migration is mediated by $\alpha 2\beta 1$ integrin.^{84,85} Collagen is not included in cultures using a laminin-511 scaffold, although hiPSC lines do express both collagen I and IV.^{86,87} Albumin in the culture medium promotes collagen secretion by hiPSCs. Collagen I has been reported to stabilize FGF2, similar to HSPG, for floating cultures of hiPSCs.⁸⁸

$\alpha 2\beta 1$ integrin/collagen I/IV signaling. Collagen I-bound $\alpha 2\beta 1$ integrin stimulates the self-renewal of PSCs by increasing the PI3K/AKT signal through the GTP Ras pathway.⁸⁹ Collagen IV pretreatment enhances *in vivo* growth of undifferentiated PSCs without affecting their pluripotency through $\alpha 2\beta 1$ integrin-mediated actin remodeling and the regulatory molecules PI3K and Rho-family GTPases⁸⁴ (Fig. 3).

Laminin receptors ($\alpha 3\beta 1$, $\alpha 6\beta 1$, $\alpha 7\beta 1$, and $\alpha 6\beta 4$)

Laminins are large glycoprotein molecules consisting of α , β , and γ chains. To date, five α , three β , and three γ chains have been identified. Laminin expression is present in early embryos and in pluripotent mouse and human ESCs. Cultured pluripotent hPSCs express a laminin-511 heterotrimer ($\alpha 5\beta 1\gamma 1$), consisting of $\alpha 5$, $\beta 1$, and $\gamma 1$ laminin chains.⁹⁰

$\alpha 3\beta 1$ integrin/laminin-332, -511, -521 signaling. The role of integrin $\alpha 3\beta 1$ in cell migration is unclear. Laminin-332 drives PI3K activation and cell survival.⁷⁶ Gu et al.⁹¹ demonstrated that laminin-511 and laminin-521 directly bind to $\alpha 3\beta 1$ integrin and, thereby, activate the PI3K/AKT signaling pathway (Fig. 3). However, it is not yet clear whether laminin-511 and laminin-332 use the same or different binding sites in these receptors.

$\alpha 6\beta 1$ integrin/laminin-332, -511, -521 signaling. Laminin-511 and laminin-521 maintain hPSC pluripotency after $\alpha 6\beta 1$ integrin signaling to induce the PI3K/AKT signaling pathway.^{90,92} In cultures using a laminin-511 scaffold, the interaction between laminin-511 and $\alpha 6\beta 1$ integrin partly compensates for the ROCK inhibitor, Y-27632. The proto-oncogene tyrosine-protein kinase Fyn⁹³ associates with the p85 subunit of PI3K. The Fyn-RhoA-ROCK signaling

cascade⁹⁴ is of interest since Fyn directly associates with $\alpha 6 \beta 1$ -integrin.^{95,96} PSCs interact with laminin-511 through $\beta 1$ -integrins, mostly $\alpha 6 \beta 1$. Strong Fyn-RhoA-ROCK signaling is expected in cultures using laminin-511 as the scaffold (Fig. 3).

$\alpha 6 \beta 4$ integrin/laminin-332, -511, -521 signaling. Upon binding to laminin-332, the integrin $\alpha 6 \beta 4$ can organize hemidesmosomes, which are essential for stable cell adhesion.^{97,98} Furthermore, FGF2 upregulates the expression of $\alpha 2 \beta 1$, $\alpha 3 \beta 1$, $\alpha 6 \beta 1$, and $\alpha 6 \beta 4$ integrins in cells cultured *in vitro*.⁹⁹ Therefore, it is believed that $\alpha 6 \beta 4$ integrin in hiPSC membranes increases if concentrated FGF2 is added to the culture medium (Fig. 3).

RGD receptors ($\alpha 5 \beta 1$, $\alpha 8 \beta 1$, $\alpha V \beta 1$, $\alpha V \beta 5$, and $\alpha V \beta 6$)

The tripeptide sequence arginine-glycine-aspartic acid (RGD) has been suggested to be a recognition sequence for integrins.¹⁰⁰ This peptide sequence has been identified in many ECM proteins such as fibronectin, vitronectin, fibrinogen, von Willebrand factor, thrombospondin, laminin, entactin, tenascin, osteopontin, and bone sialoprotein.^{101,102}

$\alpha 5 \beta 1$ integrin/fibronectin signaling. Human fibronectin, which binds to integrin $\alpha 5 \beta 1$,¹⁰³ supports hPSC expansion in serum-free medium.¹⁰⁴ $\alpha 5 \beta 1$ integrin binds soluble fibronectin efficiently; first, the $\alpha 5$ integrin binds to the synergy site to cause a conformational

change in the fibronectin; second, $\beta 1$ integrin binds to the exposed RGD sequence.¹⁰⁵ Overexpression of $\alpha 5 \beta 1$ provides considerable protection against several proapoptotic stimuli. Furthermore, the protective effects of $\alpha 5 \beta 1$ are mediated through a PI3K/AKT-dependent pathway¹⁰⁶ (Fig. 3).

$\alpha 8 \beta 1$ integrin/fibronectin signaling. $\alpha 8 \beta 1$ integrin, a cell adhesion receptor that links the cytoskeleton to ECM proteins, not only mediates mechanical and chemical signals from the matrix but also regulates cell adhesion through bidirectional membrane transduction.¹⁰⁷ Under serum deprivation conditions, $\alpha 8 \beta 1$ can interact with fibronectin to increase cell survival rates so that $\alpha 8$ -mediated survival is mediated by the PI3K pathway¹⁰⁸ (Fig. 2).

$\alpha V \beta 1$ integrin/fibronectin signaling. $\alpha V \beta 1$ has been described as a receptor for fibronectin,¹⁰⁹ vitronectin,^{110,111} or collagen.¹¹² $\alpha V \beta 1$ integrin is able to recognize vitronectin and fibrinogen and also cooperate with $\alpha 5 \beta 1$ to mediate attachment to and spreading on fibronectin;^{91,113} Ras/MEK/MAPK signaling and kinases are also stimulated by integrin ligation (Fig. 2).

$\alpha V \beta 5$ integrin/vitronectin signaling. Recombinant vitronectin is a functionally defined substrate that supports hPSCs self-renewal through $\alpha V \beta 5$ integrin³⁶ (Fig. 4). $\alpha V \beta 5$ activates latent TGF- β .¹¹⁴ Chen et al.

FIG. 4. Signaling pathways in serum-free and feeder cell-free culture systems (using E8 medium on vitronectin). Recombinant vitronectin is a functionally defined substrate that supports hPSCs self-renewal through $\alpha V \beta 5$ integrin. $\alpha V \beta 1$ has been described as a receptor for vitronectin through Ras/MEK/MAPK signaling, and kinases are also stimulated by integrin ligation. This culture system does not need albumin; as collagen I and HSPG do not concentrate on FGF2, it is necessary to have a high concentration of FGF2 in the culture medium.

enhanced the attachment and survival of hPSCs on vitronectin-coated surfaces by excising the N-terminal and/or C-terminal fragments of full-length vitronectin.³² It has also been reported that vitronectin and one of its variants (VTN-NC) support the maintenance of hPSCs through $\alpha V\beta 5$ integrin.^{32,36,115}

$\alpha V\beta 6$ integrin/TGF- β signaling. $\alpha V\beta 6$ integrin has a key role in regulating TGF- β activation of ALK5, a TGF- β type I receptor.^{114,116,117} The association with $\alpha V\beta 6$ integrin expression may be linked to TGF- β activation, which has an important role in intracellular signaling. TGF- β activates c-Jun and Erk through $\alpha V\beta 6$ integrin. TGF- β is an essential additive in culture medium using a laminin-511 scaffold (Fig. 3).

Conclusion

This review highlights the components that are suitable for serum- and xeno-free media and feeder cell-free culture systems that use laminin-511 as the scaffold. The binding of $\alpha 6\beta 1$ with laminin-511 and the interaction with E-cadherin stimulate Fyn-RhoA-ROCK signaling to act as a deactivation ROCK signal. These results indicate that the signaling pathways induced by a laminin-511 scaffold depend on PI3K signaling through E-cadherin-mediated cell-cell adhesion and that the Fyn-RhoA-ROCK signaling cascade is associated with $\alpha 6\beta 1$ integrin-laminin-511 adhesion.

Author Disclosure Statement

No competing financial interests exist.

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Abbreviations Used

ALK = activin-like kinase
BSA = bovine serum albumin
CAMs = cell adhesion molecules
ECM = extracellular matrix
ERK = extracellular signal-regulated kinase
FGFR = fibroblast growth factor receptor
HSPG = heparan sulfate proteoglycans
hiPSCs = human-induced pluripotent stem cells
hPSC = human pluripotent stem cell
IGF = insulin-like growth factor
ITS = insulin, transferrin, and selenium
MAPK = mitogen-activated protein kinase
MEF = mouse embryonic fibroblast
NF-κB = nuclear factor κB
TGF-β = transforming growth factor β

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