Heterogeneous Differentiation of Human Mesenchymal Stem Cells in 3D Extracellular Matrix Composites

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Abstract

Extracellular matrix (ECM) proteins are structural elements of tissue and also potent signaling molecules. Previously, our laboratory showed that ECM of 2D coatings can trigger differentiation of bone marrow-derived mesenchymal stem cells (MSCs) into mesodermal lineages in an ECM-specific manner over 14 days, in some cases comparable to chemical induction. To test whether a similar effect was possible in a 3D, tissue-like environment, we designed a synthetic-natural biomaterial composite. The composite can present whole-molecule ECM proteins to cells, even those that do not spontaneously form hydrogels ex vivo, in 3D. To this end, we entrapped collagen type I, laminin-111, or fibronectin in ECM composites with MSCs and directly compared markers of mesodermal differentiation including cardiomyogenic (ACTC1), osteogenic (SPP1), adipogenic (PPARG), and chondrogenic (SOX9) in 2D versus 3D. We found the 3D condition largely mimicked the 2D condition such that the addition of type I collagen was the most potent inducer of differentiation to all lineages tested. One notable difference between 2D and 3D was pronounced adipogenic differentiation in 3D especially in the presence of exogenous collagen type I. In particular, PPARG gene expression was significantly increased ~16-fold relative to chemical induction, in 3D and not in 2D. Unexpectedly, 3D engagement of ECM proteins also altered immunomodulatory function of MSCs in that expression of IL-6 gene was elevated relative to basal levels in 2D. In fact, levels of IL-6 gene expression in 3D composites containing exogenously supplied collagen type I or fibronectin were statistically similar to levels attained in 2D with tumor necrosis factor-α (TNF-α) stimulation and these levels were sustained over a 2-week period. Thus, this novel biomaterial platform allowed us to compare the biochemical impact of whole-molecule ECM proteins in 2D versus 3D indicating enhanced adipogenic differentiation and IL-6 expression of MSC in the 3D context. Exploiting the biochemical impact of ECM proteins on MSC differentiation and immunomodulation could augment the therapeutic utility of MSCs.

Key words: biomaterials; extracellular matrix; stem cells

Introduction

Utilization of mesenchymal stem cells (MSCs) in the clinic and attempts to drive mesodermal differentiation of MSCs in vitro would benefit from enhanced understanding of the impact of 3D, exogenous extracellular matrix (ECM) on MSC state. Indeed MSCs are a heterogeneous lot, primed to mature to several distinct mesodermal states or to remain multipotent depending on the local environment (e.g., native tissue, recipient tissue or tissue culture scenario). The local environment contains soluble signaling factors, other cells or cell types, and ECM, and the bulk of past literature reports pertain to soluble factor stimulation with respect to MSC state. More recently, an appreciation of the role of ECM in MSC specification has emerged including work of our laboratory in 2009 showing the presentation of ECM
proteins on 2D culture plastic to MSCs is capable of trig-
nering differentiation into multiple lineages at levels
comparable to soluble factor induction.\(^1\) This result is
perhaps not now surprising as the multifaceted func-
tions of ECM continue to emerge.\(^2\)–\(^4\)

ECM proteins, aside from providing support for cells
and tissues, are important modulators of cell signaling
during development, in the stem cell niche, and in the
tumor microenvironment. ECM proteins are large and
complex, and contain highly conserved regions.\(^5\) ECM
proteins act as effective ligands to initiate cell signaling
usually via integrin receptors. They may also act as a
depot to controllably bind or release growth factors
that drive a multitude of cell signaling events. Despite
the multifunctionality of ECM proteins in guiding cell
behavior such as proliferation and migration, systematic
understanding of the role of ECM in stem cell differen-
tiation is somewhat limited in conventional pseudo-3D
culture platforms.\(^6\) Covalent conjugation\(^7\) or adsorption \(^8\)
of ECM proteins on 2D surfaces have been simple and
convenient platforms to create microenvironments for
stem cell differentiation. Efforts to create more physio-
logically relevant 3D microenvironments\(^9\) were carried
out by controlling matrix mechanics,\(^10\) porosity,\(^11\) or
cell aggregates\(^12\) and by forming scaffolds with collagen,
Matrigel\(^13\), decellularized ECM, and ECM-mimicking
oligopeptides.\(^13\) Although the distinction between stem
cell differentiation in 2D versus 3D was not ignored,\(^14\)–\(^16\)
a direct comparison of ECM presentation in 2D versus
3D has been rarely evaluated.\(^17\) Here, we utilized a re-
cently developed 3D biomaterial platform\(^18\) to compare
MSC differentiation to cardiomyogenic, osteogenic, adi-
pogenic, and chondrogenic lineages in 2D versus 3D.

We created “composites” by incorporating ECM pro-
tein(s) and stem cells in poly(ethylene glycol) (PEG)
hydrogels, which are cross-linked via formation of native
amide bonds.\(^19\) This chemoselective cross-linking allows
the presentation of ECM proteins without any modifica-
tion or conjugation to PEG chains and maintains consis-
tent overall stiffness and diffusion in the presence of
exogenous ECM proteins. This platform enables bio-
chemical engagement of ECM proteins to MSCs in 3D,
largely excluding other biophysical stimuli. Here, human
embryonic stem cell (hESC)-derived MSCs (hMSCs)\(^20\),\(^21\)
were encapsulated in ECM composites with 2 mg/mL of
either collagen type I, laminin-111, or fibronectin (Fig. 1).
MSC functionality was assessed via gene and protein ex-
pression of lineage markers associated with cardiomyo-
genesis, adipogenesis, osteogenesis, and chondrogenesis
and of cytokines associated with immunomodulation.

**Materials and Methods**

**Detailed methods are provided in the Supplementary
Materials and Methods section.**

**Culture of human mesenchymal stem cells and
formation of ECM composites and ECM films**

Mesenchymal stem cells were derived from H1 (hESC, a
gift from Dr. Peiman Hematti, University of Wisconsin-
Madison)\(^20\) and markers indicative of the MSC pheno-
types (CD73, CD90, and CD105) were probed by flow
cytometry (Supplementary Fig. S1). Human MSC
(hMSC) were encapsulated in ECM composites\(^18\)
or cultured on 2D ECM films\(^1\) using protocols previously
published (Supplementary Materials and Methods).

**Imaging ECM composites using multi-photon
laser scanning microscopy**

At days 1, 14, and 28, ECM composites were fixed and
stained with previously published protocols (Supple-
mentary Materials and Methods).\(^18\) ECM proteins,
nuclei, and actin filaments were visualized with a
40x (NA = 0.8) objective lens by using a mode-locked
Ti:Sapphire laser on a multi-photon laser scanning mi-
croscope (MPLSM; Bruker Nano). 3D reconstructions
were generated using Imaris 7.5.2 software (Bitplane,
Inc.). To further confirm the expression of FABP4
and PPAR-\(\gamma\) proteins from H1 MSCs after 28 days,
ColI composites were incubated overnight at 4°C
with rabbit anti-FABP4 (cat# PA5-30591; Thermo Sci-
entific, 1:100 dilution in the blocking solution) or
rabbit-PPAR-\(\gamma\) (cat# PA3-821A; Thermo Scientific, 1:100
dilution in blocking solution) antibodies, which
were probed by goat anti-rabbit FITC (cat# 31583;
Pierce, 1:200 dilution in blocking solution) for 45 min.

**Quantitative real-time polymerase chain reactions**

RNAs were extracted from hMSC culture or ECM com-
posites using a standard TRizol\(^10\) method (Supple-
mentary Materials and Methods). Complementary DNA
cDNA) was synthesized following the instructions
from the Maxima First Strand cDNA Synthesis Kit
(cat# K1642; Thermo Scientific). Each lineage primer se-
quence is provided in Table 1. Immunomodulatory gene
primers (IL-6 and IDO) were purchased from Biorad
(cat# 10025636). Primer efficiencies were extracted from
RealPlex\(^2\) software and verified with melting curves.
Each lineage standard curve was calculated using termi-
nally differentiated cells and corresponding primers.
The standard curve method was employed to determine copy
numbers of each lineage and GAPDH gene. The lineage
gene expression level was determined as the copy number of lineage gene of interest normalized to that of GAPDH at each time point.

hMSC chemical induction

Standard chemical reagents were used for cardiomyogenic, osteogenic, adipogenic, and chondrogenic differentiation (Supplementary Materials and Methods).

Stimulation of hMSCs with IFN-γ and TNF-α

hMSCs suspended in α-MEM medium were plated to six-well plates at a density of $1.4 \times 10^4$ cells/mL and were kept at 37°C/5% CO₂ for 20 h. Stimulation media were prepared with 100 ng/mL of IFN-γ (cat# 300-02; Peprotech) or 50 ng/mL of TNF-α (cat# 68-8786-82; eBioscience) in α-MEM medium. After 20 h of culture, the medium in the six-well plates was replaced with stimulation media, which was added to the experimental wells and α-MEM medium in the control wells. The plates were kept at 37°C/5% CO₂ and hMSCs were collected at 4 and 24 h.

Statistical analyses

For comparison of expression levels of various differentiation or immunomodulatory markers of hMSCs from ECM composites, one-way ANOVA with the Tukey’s HSD post hoc test, Fisher’s LSD post hoc test, or Student’s t-test ($\alpha = 0.05$) was performed. All tests were performed using Statistical Analysis System (SAS Institute, Inc.).

Results

hMSCs in ECM composites exhibit morphologies influenced by ECM engagement in 3D

ECM proteins, provided exogenously at the time of composite formation, distinctively impacted the distribution and morphology of hMSCs in 3D. After 3 weeks in culture, hMSCs in laminin (LN) composites exhibited

Table 1. qRT-PCR Primer Sequences for hMSC Lineage Markers

| Target gene | Primer sequence (5’ → 3’) | Amplicon size (bp) |
|-------------|--------------------------|-------------------|
| SPP1        | FOR: TTCGCAGACCTGACATCCAGTACC  | 70                |
|             | REV: TCTCGCCCTTCCATGTTGAGG     |                   |
| PPARG       | FOR: AGCCTCATGAAGAGGCTTTCCAAC  | 122               |
|             | REV: TCTCGGGAGAAACCTTTGG CATC  |                   |
| ACTC1       | FOR: GCTTCGAGCTGTCGAGA         | 61                |
|             | REV: ATGCCAGCAGATTCCTAC         |                   |
| SOX9        | FOR: TACCCGGAAGTGGACAT         | 77                |
|             | REV: TCAAACTCGATGGACATCGAAGG   |                   |
| GAPDH       | FOR: TTAAAAAGCAGCCCTGGTGAC     | 144               |
|             | REV: CTCTGCCTCCTGCTGGCAC       |                   |

hMSC, human mesenchymal stem cells; qRT-PCR, quantitative real-time polymerase chain reaction.
FIG. 2. Morphology and engagement of human MSCs (hMSCs) with ECM of 3D composites over time. Multiphoton laser scanning microscopy (MPLSM) was used to optically section (at 1 μm interval) microenvironments of LN composites containing hMSCs to convey the three-dimensionality of the microenvironments within the composite (a–c, 161 planes). Z stacks of images were reconstructed from raw images (a, b) to solid objects using Imaris software to best view cell–ECM interactions (c); scale bar 80 μm. Single optical sections of ECM composites at days 1, 14, and 28 after composite formation. Coll (collagen type I) composites (d–f), LN (laminin-111) composites (g–i), and FN (fibronectin) composites (j–l). ECM proteins (green), nuclei (blue), and actin filaments (red); Scale bar 50 μm.
reticulate structures and were found to adhere to multiple dimensions to the network architecture of LN (Fig. 2a–c). The 3D surface rendering of z stacks at over 162 μm showed that hMSCs were engaged with LN throughout the PEG-entrapped microenvironments of composites (Fig. 2b, c). Similar 3D engagement with ColI was observed at this point time, but with cells assuming an elongated morphology and cell-ECM structure, usually globular in nature as previously reported from our laboratory.\textsuperscript{18} We further probed the engagement of hMSCs with surrounding ECM at 1, 14, and 28 days of culture utilizing MPLSM. Since each field of view was not taken from a flat, 2D region of interest, some optical sections show narrower aggregate structures (e.g., Fig. 2e, f or l) than others (e.g., Fig. 2i or j). All composites showed globular microenvironments at day 1 with high hMSC to ECM ratios (Fig. 2d, g, j). At later time points, composites evolved to contain a slightly lower hMSC to ECM ratio likely due to accumulating endogenous ECM and associated decrease in hMSC proliferation.\textsuperscript{18} By day 28, distinct hMSC morphologies and corresponding microenvironments emerge. ColI composites showed globular aggregation with significant hMSC spreading (Fig. 2f). LN, as noted above, assumes a reticulate or mesh-like hMSC-ECM distribution (Fig. 2i). A higher number of hMSCs attached to fibronectin (FN) than other ECM, perhaps attributed to increased cell-binding domains\textsuperscript{24} relative to ColI or LN. FN composites showed fewer numbers of cells per microenvironment and smaller total volume per microenvironment (Fig. 2l) than LN composites. Collectively, ECM proteins exogenously supplied in 3D composites were able to modulate hMSC-ECM engagement. Indeed, cell–ECM interactions appeared quite similar between groups at day 1, but quickly changed over time in distinctive ways according to ECM type.

**Differentiation of MSCs, especially adipogenesis, is initiated in 3D ECM composites**

Previously, our laboratory reported that 2D ECM coatings alone were able to initiate differentiation of bone marrow-derived MSCs into cardiomyogenic, osteogenic, and adipogenic lineages at levels comparable to chemical induction.\textsuperscript{3} Unclear at that time was whether or not differentiation in the context of ECM was an artifact of the unnatural spreading and associated strains of the 2D tissue culture scenario or whether the same could occur in a 3D, tissue-like context. Appropriate technology to directly test this question took several years to develop since many ECM types extracted from the body do not form 3D hydrogels; but we have recently validated such a system.\textsuperscript{18} We use it here to determine whether ECM composites, each containing a single ECM at a fixed concentration of 2 mg/mL and with maintained stiffness\textsuperscript{18} and diffusivity (Supplementary Table S1) could modulate the functional capacity of hMSCs both in terms of differentiation and immunomodulatory potential. For the purpose of comparison, we repeated the 2D studies\textsuperscript{1} here using hMSCs and reexamined the differentiation profile (Fig. 3a–d). In 2D, hMSCs cultured on ECM films of LN and FN showed increased expression of *ACTC1* (i.e., cardiomyogenic differentiation) at day 14 relative to day 1, but only FN stimulated gene expression levels significantly exceeding those of the no construct (NC) control and chemical induction (Fig. 3a). Osteogenic differentiation (as discerned by *SPP1* expression) of hMSCs on 2D ECM films generally increased from day 1 to 14 (with the exception of ColI; Fig. 3b) to levels comparable to chemical induction at day 14. None of the 2D ECM films promoted *PPARG* expression associated with adipogenic differentiation. Surprisingly, only the NC control yielded *PPARG* expression slightly, but significantly higher than chemical induction at day 14 (Fig. 3c). Similarly, none of the 2D ECM films promoted SOX9 expression associated with chondrogenic differentiation to levels approaching that of chemical induction. It should be noted that the chemical induction protocol for chondrogenesis requires a cell pellet that essentially creates a 3D environment, and so perhaps is not the most robust 2D control.

In 3D ECM composites, we found Coll significantly promoted the expression of *SPP1, PPARG*, and SOX9 genes of hMSCs at 28 days relative to 2D chemical induction (Fig. 3e–h). The average expression of *ACTC1* in ColI composite was relatively high while its variation was also large and thus not significantly different from 2D chemical induction at day 28 (Fig. 3e). Osteogenic and adipogenic differentiation in the Coll composite were significantly higher than chemical induction and other ECM composites at day 28 (Fig. 3f, g). In addition, chondrogenic differentiation in Coll composite was significantly increased from day 14 to 28, and was also significantly higher than chemical induction and other ECM composites at day 28 (Fig. 3h). To distinguish the contribution of the 3D environment only (PEG) from the 3D environment plus exogenous ECM on hMSC differentiation, we normalized each expression level of lineage genes to that of the PEG control at each time point (Fig. 3i–l). Coll composites significantly upregulated...
expression of each lineage gene at day 28 (Fig. 3j–l), except ACTC1 (cardiomyogenic) (Fig. 3i). Adipogenic chondrogenic gene expression of hMSCs in Coll composites was also significantly increased from day 14 to 28, respectively (Fig. 3k, l). Taken together, differentiation of ESC-derived MSCs in 3D Coll composites was significantly more effective than 2D chemical induction, while chemical induction of hMSCs was more efficient than 2D ECM films only in chondrogenic induction by 3D cell mass.
The most outstanding difference in hMSC differentiation progress between 2D and 3D exogenous ECM engagement was adipogenic differentiation. Minimal PPARG expression was detected in 2D ECM films (Fig. 3c), while 3D ECM composites all upregulated PPARG gene expression at both 14 and 28 days, especially in composites with Coll (Fig. 3g). The significant upregulation of PPARG gene led to query of the expression of adipogenic proteins, FABP4 and PPAR-γ in 3D Coll composites. After 28 days of hMSC culture in Coll composite, hMSCs were stained with antibodies against FABP4 and PPAR-γ using MPLSM (Fig. 4). Over 28 days, hMSCs in Coll composites showed clusters of hMSCs positive for both FABP4 (Fig. 4c) and PPAR-γ (Fig. 4f) protein expression. Staining of both adipocyte-associated proteins was more intense in 3D composites containing collagen type I than that observed in all 2D cultures with ECM films and even with chemical induction where only limited, faint staining was observed. Due to the limited protein staining in the 2D condition, especially with chemical induction, we also exposed an alternative ESC-derived MSC line (H9 MSCs) to chemical induction for adipogenesis (Supplementary Fig. S2a, e).21 We found a moderate level of staining for PPAR-γ and FABP4, definitive Oil Red O staining of fat droplets (Supplementary Fig. S2i) and significant upregulation of PPARG gene (Supplementary Fig. S2j) suggesting the chemical induction protocol was effective, and that the H1 MSC line is less prone to adipogenesis than other MSC populations such as H9 MSCs. Given this result, it is perhaps even more surprising that adipogenesis of H1 MSC was significantly promoted in 3D composites and was further enhanced in the composite containing Coll.

**ECM composites attenuate IL-6 gene expression**

To determine whether immunomodulatory function of MSCs might also be altered with 3D exposure to ECM, we again capitalized on our composite system. hMSCs were entrapped in composites containing Coll, LN, or FN and cultured for 28 days and then probed for transcript expression of indoleamine 2,3-dioxygenase (IDO) and interleukin 6 (IL-6), two cytokines frequently associated with MSC immunomodulation.25–28 Both IDO and IL-6 expression were inducible by IFN-γ and TNF-α stimulation in 2D culture, respectively (Fig. 5a, b). In
3D, expression of IDO gene was undetectable, similar to unstimulated MSC expression in 2D (data not shown). The expression of IL-6 on 2D ECM films was significantly increased relative to unstimulated hMSC for 14 days except NC control (Fig. 5c). However, the expression of IL-6 in 3D ECM composites was significantly increased relative to unstimulated hMSC expression in 2D. And in fact, expression was modulated in an ECM-specific manner. At 28 days, composites containing Coll or FN maintained levels of IL-6 gene expression statistically similar to that of 2D hMSC cultures with TNF-α stimulation (24 h, Fig. 5d). At the same time point, composites without ECM and those containing LN exhibited IL-6 expression above that of 2D unstimulated MSCs, but less than levels with TNF-α stimulation. Collectively, the expression of IL-6 was elevated in 3D spaces for several days and the elevation of IL-6 production in 3D was modulated to varying degrees by the type of ECM supplied exogenously at day 0.

Discussion

Here, we show for the first time the impact of individual, intact ECM proteins presented in 3D on MSC functional potential both in terms of differentiation and immunomodulation. This work is important for those attempting to develop 3D ECM-containing delivery vehicles for MSC transplantation and for predicting behavior of transplanted cells to ECM-rich microenvironments. Previously, our laboratory showed that 2D ECM coatings modulated the differentiation of bone marrow-derived MSCs into different lineages in an ECM-specific manner over 14 days, comparable to chemical induction.1 But extension of these studies to
3D has been challenging because, with the exception of type I collagen, hyaluronan, and fibrin, ECM proteins isolated from tissues cannot reassemble reliably into robust 3D scaffolds capable of supporting cellular populations. Type I collagen has been utilized as a supporting scaffold for other ECM proteins purified from tissue or lyophilized ECM mixtures extracted from decellularized tissue. However, in these cases it is challenging to distinguish biochemical effects of the added ECM components from that of the supporting ECM (i.e., type I collagen) scaffold. Recently, our laboratory designed a synthetic-natural biomaterial composite that can present whole-molecule ECM proteins to cells without chemically cross-linking ECM proteins to the 3D biomaterial platform, preserving native secondary and tertiary structures necessary for appropriate biochemical signaling. In addition, since the impact of substrate stiffness on MSC differentiation is now well appreciated, the composite platform was designed to maintain consistent macroscale storage modulus (0.9 kPa) even with alterations in ECM type. Finally, the 3D scaffold was verified to maintain diffusion of well-characterized bovine serum albumin molecules in the same order of magnitude (Supplementary Table S1). These controls allowed us to compare functional outcomes of MSC exposure to individual ECM in 3D and relate those outcomes to findings in 2D. Like 2D, ECM proteins alone could stimulate multi-lineage differentiation in 3D at levels approaching and sometimes exceeding those attained with standard chemical differentiation protocols. Unlike 2D, adipogenic differentiation was favored with ECM serving to refine this response.

Adipogenic differentiation of MSCs and other multipotent cells has been a great interest to endocrinologists, stem cell biologists, and tissue engineers. The adipose tissue is an endocrine/paracrine regulator of energy metabolism and dysregulation can lead to type II diabetes and cardiovascular disorders among others. From a tissue engineering standpoint, burn patients suffering from soft tissue atrophy or facial lipodystrophy from antiviral medications for human immunodeficiency virus would benefit soft tissue reconstruction. Despite the importance of adipose tissue model systems and regeneration, mechanistic understanding of adipogenesis is still developing and methods to derive adipocytes from stem cells are quite variable. Further, the role of ECM proteins and three dimensionality has only recently been appreciated. Indeed, human mesenchymal progenitor cell spheroids formed in 2–3 μm honeycomb scaffolds significantly upregulate PPARG, CEBPA (CCAAT-enhancer-binding protein α, C/EBP-α), ADIPOQ (adiponectin), and ALBP (fatty acid-binding protein 4, FABP4, or aP2) gene expressions over 2D culture scenarios after only 3 days. This example highlights the importance of three dimensionality for adipogenesis, but exploits chemical induction without exogenous ECM proteins.

When ECM proteins are added the induction protocol for adipogenesis, augmented differentiation is observed. For example, human MSC spheroids cultured in rotating wall vessel bioreactor in combination with adipogenic induction showed significantly enhanced PPARG, CEBPA, ALBP, and LPL (lipoprotein lipase) 7 days after culture initiation compared with 2D culture. Further, covalent conjugation of FN on 2D polyacrylamide soft (~0.6 kPa) substrate also significantly improved the expression of CEBPA and LPL over 10 days with microscale circular patterns (~1000 μm²) relative to 2D polyacrylamide substrate without patterns. This is an interesting result as our composites are on the order of 0.9 kPa in stiffness, suggesting the possibility that 3D mechanical properties in addition to ECM exposure further enable adipogenesis. The impact of mechanical properties on MSC differentiation is well characterized in 2D and 3D, where adipogenic or neurogenic commitment occurs in softer environments at levels dependent on substrate/scaffold materials. Also of note, intramuscular adipocytes (BIP) have been reported to produce collagens type I, V, VI, and fibronectin at high levels in vitro, while type IV collagen and laminin were produced at lower levels. The ECM-niche of fat then may promote adipogenesis and explain why composites with LN saw lower levels of differentiation.

Composites without ECM proteins were unable to maintain adipogenic differentiation after 14 days, supporting a necessity of not only the third dimensionality and associated mechanical properties but also ECM proteins for adipogenesis. Like other connective tissue cell types with mesenchymal origin, adipocytes are surrounded by collagens and basal lamina containing ECM proteins. Since only a lipid monolayer forms the boundary between stored fat and cytosol, transfer of mechanical stress from the outside to the inside of the adipocyte can be decreased by adipocyte ECM proteins and also redistribute forces over a larger area of tissue. These findings reflect the critical requirement of adipocyte ECM proteins for the
development of preadipocyte to fat-storing adipocytes, in part explaining the attenuation of PPARG expression from composites with no ECM. To guide interpretation of these results, it should be noted that composites without exogenous ECM were less able to sustain cell viability long term. We previously reported a reduction of viability to as low as 60% over the course of several weeks in PEG (no ECM control) and thus differences in differentiation relative to the no ECM controls might relate to factors associated with cell death instead of or in addition to lack of signaling associated with exogenous ECM. For this reason we emphasized comparison to chemical induction and between ECM-containing composites.

In addition to multipotency of hMSCs, the immunomodulatory properties of MSCs have been exploited to treat multiple diseases including myocardial infarction or graft versus host disease to name a few. Although the exact mechanistic details are unclear, MSCs have been reported to be hypoinmunogenic, enabling MSC transplantation across major histocompatibility barriers and creating off-the-shelf therapies with allogenic MSCs. It should be noted that hESC-derived MSCs (both H1 and H9 MSCs) exhibit very similar immunosuppressive properties and multipotency relative to bone marrow-derived MSCs including the three main characteristics of MSCs including adherence to plastic, positive expression of CD29, CD44, CD54, CD73, CD90, and CD105, and negative expression of CD34, CD45, and CD31 and their ability to differentiate into osteogenic, adipogenic, and chondrogenic lineages. Here, we selected a small subset of cytokines to determine whether MSCs, by the engagement of ECM proteins in 3D, will modulate the expression of two immunomodulatory genes, IDO and IL-6. IDO (indoleamine-2,3-dioxygenase) is an enzyme that catalyzes the degradation of tryptophan along the metabolic pathway to kynurenine. MSC-mediated immunosuppression is believed to result from the depletion of tryptophan and the local accumulation of tryptophan metabolites, suppressing T cell and natural killer cell proliferation. As observed in Figure 5 and according to results using single cell RNA-sequencing of murine MSCs from our laboratory, MSCs normally express minimal levels of IDO while the IDO mRNA levels are found to be significantly elevated upon stimulation with inflammatory cytokines, primarily IFN-γ. IL-6 is known as a pleiotropic cytokine suppressing T-cell proliferation and local inflammation. While MSC-secreted IL-6 is reported to be proinflammatory, IL-6 can also be anti-inflammatory to modulate other proinflammatory cytokines. Although the inflammatory nature of IL-6 in 3D ECM composites was hard to define, the upregulation of IL-6 gene in 3D is significant and is surprisingly sustained over multiple weeks (Fig. 5). Aggregates of MSCs showed a similar result after 7 days in culture and MSCs encapsulated in 3D hydrogels also modulated IL-6 secretion at day 14. In this case an association between IL-6 and hydrogel stiffness was observed within the softest hydrogel (1.5 kPa) showing the most significant upregulation of expression of IL-6. Future studies would be wise to assess how immunomodulation of MSCs changes in the face of distinct ECM of different tissue types following transplantation.

Conclusion
These results further elucidate the importance of ECM in guiding differentiation of stem cells in tissue and more surprisingly, associated immunomodulatory cytokine production. These data set the stage for synthetic biomaterial development for a multitude of applications including tuning of differentiation and immunomodulation to enable proactive delivery vehicles for stem cell transplantation.

Acknowledgments
The authors thank Brian Freeman for helping characterize H1 MSCs by flow cytometry and Quyen Tran for immunohistochemical staining of H9 MSCs, respectively. Epifluorescence microscopy was performed in the Alford lab. 3D surface rendering was performed at the University Imaging Center of the University of Minnesota–Twin Cities. This work was supported by the National Cancer Institute (R01CA181385, to PPP).

Author Disclosure Statement
No competing financial interests exist.

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Cite this article as: Jung JP, Bache-Wiig MK, Provenzano PP, Ogle BM (2016) Heterogeneous differentiation of human mesenchymal stem cells in 3D extracellular matrix composites, BioResearch Open Access 5:1, 37–48, DOI: 10.1089/biores.2015.0044.

Abbreviations Used

- BSA = bovine serum albumin
- cDNA = complementary DNA
- DABCO = 1,4-diazabicyclo[2.2.2]octane
- DAPI = 4',6-diamidino-2-phenylindole
- ECM = extracellular matrix
- FN = fibronectin
- hESCs = human embryonic stem cells
- hMSC = human mesenchymal stem cells
- IDO = indoleamine 2,3-dioxygenase
- IL-6 = interleukin 6
- LN = laminin
- MSCs = mesenchymal stem cells
- MPLSM = multi-photon laser scanning microscopy
- NC = no construct
- PBS = phosphate buffered saline
- qRT-PCR = quantitative real-time polymerase chain reaction
- RT = room temperature
- TNF-α = tumor necrosis factor-α

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