Constitutive Expression of Human Telomerase Enhances the Proliferation Potential of Human Mesenchymal Stem Cells

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Abstract

Human mesenchymal stem cells (hMSCs) are highly desirable cells for bone engineering due to the inherent multipotent nature of the cells. Unfortunately, there is a high degree of variability, as primary hMSC cultures quickly undergo replicative senescence with loss of proliferative potential as they are continually propagated in cell culture. We sought to reduce the variability of these cells by insertion and expression of human telomerase reverse transcriptase (TERT) to immortalize the cell line. hMSCs were transduced with a lentivirus containing the human TERT gene. The resulting cell line has been propagated through more than 70 population-doubling level (PDL) to date and continues to grow exhibiting the characteristic fibroblastic hMSC phenotype. Expression of TERT mRNA and protein activity was confirmed in the TERT-transduced cells. Mock-transduced hMSCs had almost undetectable levels of TERT mRNA and protein activity and lost proliferation potential at PDL 14. The enhanced growth capacity of the hMSC TERT cells was due to increased cell proliferation and reduced cellular senescence rather than due to inhibition of apoptosis. The multipotent nature of the TERT cells was confirmed by differentiation toward the osteoblastic and adipogenic lineages in vitro. Osteoblastic differentiation was confirmed by expression of alkaline phosphate and mineral deposition visualized by Alizarin Red staining. Adipogenic differentiation was confirmed by production of lipid droplets, which were detected by Oil Red-O staining. In summary, we have generated a stable hMSC line that can be continually propagated and retains both osteoblastic and adipogenic differentiation potential.

Key words: adipogenic differentiation; human mesenchymal stem cell; osteoblastic differentiation; TERT expression

Introduction

Mesenchymal stem cells (MSCs) have the potential to differentiate into a number of cell types, sparking a great interest in their use in regenerative medicine for cartilage and bone. MSCs have been isolated from bone marrow stromal cells (BMSC) and are present in a very low frequency, whereas only 0.001% to 0.01% of nucleated cells in BMSC are actually MSCs.1 As a stem cell source, human MSCs (hMSCs) are well studied and have been shown to differentiate toward osteogenic, chondrogenic, adipogenic, and myogenic lineages.2,5 Unfortunately, hMSCs undergo replicative senescence4 with prolonged growth in cell culture, further limiting the availability of hMSCs that could be used in clinical applications. Thus, continuous passage of hMSCs in cell culture resembles an aging process with the later-passage cells being more differentiated than early-passage cells until the cells cease dividing altogether.4 Rapidly dividing cells undergo about 20–40 population doublings before succumbing to cell senescence, resulting in a shorter life span than slower growing cells.6 Replicative senescence (ageing) is associated with the loss of telomeric DNA with each cell division due to incomplete replication of the telomeric ends.7 Telomeric loss results in a variety of consequences, including inhibition of the mitotic clock, genotoxic damage due to accumulation of free radicals, and chromosomal rearrangement, which can trigger a DNA damage response and ultimately leading to senescence and cell apoptosis.8–10

The length of the telomeric ends is controlled by expression of the telomerase reverse transcriptase (TERT) gene.11 Somatic cells usually have low or undetectable levels of telomerase. Decreased TERT activity correlates with increased differentiation and passage of MSCs in cell culture. Cancer cells have been shown to have increased levels of telomerase

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activity.

Transgenic mice constitutively expressing TERT have a higher incidence of cancers, although the tumorigenicity is due to loss of tumor suppressor genes and not specifically due to expression of TERT.

Constitutive expression of TERT has been shown to prevent senescence, increase proliferation, yet maintain differentiation ability in hMSCs.

We report here the generation of an hMSC line expressing TERT that exhibits enhanced cell proliferation and stability in cell culture and retains the ability to differentiate toward both the osteogenic and adipogenic lineages.

Materials and Methods

Cell culture

hMSCs, hMSC complete growth medium (HMSCGM), and growth supplements were obtained from Lonza, Inc. (Allendale, NJ). hMSCs used in this study were from a 32-year-old Caucasian man (Lonza Lot # 4F0591). Cells were subcultured at 60%–70% confluence at 37°C under 5% air/5% CO₂ atmosphere, and infected with TERT-expressing lentivirus (LV) at passage 4. For differentiation to the osteoblastic lineage, hMSCs were treated every 3–4 days with an osteogenic medium (OGM) consisting of HMSCGM (Lonza PT-3001) supplemented with 50 mM ascorbic acid-2-phosphate, 10 mM β-glycerophosphate, and 10⁻⁷ M dexamethasone (Sigma-Aldrich, St. Louis, MO). For differentiation to adipocytes, hMSCs were treated with three cycles of an hMSC adipogenic induction medium (AIM; Lonza PT-3102B containing h-insulin, l-glutamine, MCGS, dexamethasone, indomethacin, 3-isobutyl-l-methyl-xanthine [IBMX], and GA-1000) for 3 days and an hMSC adipogenic maintenance medium for 3 days (AMM; Lonza PT-3102A containing h-insulin, l-glutamine, MCGS, and GA-1000), and then further cultured in the AMM until analysis.

Generation of telomerase-expressing LV

Plasmid pBABE-puro-hTERT (#1771) containing the TERT gene was obtained from Addgene (Cambridge, MA) and the TERT gene isolated on a 3.5-kB EcoRI/Sall fragment. PLVX-IRES-Hyg (Clontech, Mountain View, CA) was restricted with SpeI. The EcoRI/Sall/Spel 5'-overhangs were end-filled using Klenow to generate blunt ends for cloning. The blunt EcoRI-Sall fragment was ligated into the blunt SpeI-PLVX-IRES-Hyg vector and transformed into NEB 5-alpha- (New England Biolabs, Ipswich, MA) competent Escherichia coli selecting for ampicillin resistance. Transformants were screened for correct insertion/orientation of the TERT fragment by restriction analysis. In this construct (pLVX-TERT-IRES-Hyg), TERT is expressed as a bicistronic transcript with the hygromycin (hyg) gene allowing hyg resistance to be used as an indicator of transduction efficiency and as a marker for selection.

TERT-expressing LV was generated after transfection of pLVX-TERT-IRES-Hyg into HEK293 cells (American Type Culture Collection, Manassas, VA) using the Lenti-X HTX Packaging system (Clontech). Briefly, 5 x 10⁶ HEK293 cells were plated in one 10-cm dish in a tetracycline-free medium and transfected the next day with pLVX-TERT-IRES-Hyg and the Lenti-X HT Packaging Mix for 4h using Xfect Polymer (Clontech). After 4h, the medium was replaced with 10 mL tetracycline-free medium. Infectious LV was collected after 72h, centrifuged 10 min at 500 g, and stored in aliquots at −80°C. LV obtained was at a high titer and did not need to be concentrated further.

Transduction of hMSCs with TERT-expressing LV

hMSCs (passage 4) were plated in six-well dishes (10,000 cells/cm²; 2 mL HMSCGM medium/well) and allowed to attach overnight. Two cycles of LV transduction were used. LV TERT was diluted 1:2 with HMSCGM, and polybrene (Sigma-Aldrich) was added (24 µg/mL). One milliliter of this dilution was added drop-wise to the 2 mL already in the wells. This gave a final concentration of 8 µg/mL polybrene in each well. The plate was centrifuged (1200 g; 32°C) for 1 h and then incubated (37°C) for an additional 5 h before changing the medium (2 mL HMSCGM) overnight. The next day, the LV TERT transduction was repeated using a fresh dilution of virus on the same wells. The transduction medium was replaced with HMSCGM containing 25 µg/mL hyg (Sigma-Aldrich). Mock-transfected wells containing no virus (with polybrene) were treated with hyg as controls. After 6 days, LV TERT hyg-resistant wells were passaged into T₂₅ flasks with hyg. Mock-transfected wells with hyg were dead. After 1 week, the cells were passaged into T₂₅ flasks without hyg to begin cell characterization and were designated hMSCs (mock-transfected) and hMSC TERT (TERT-expressing) cells.

Analysis of cell culture growth characteristics

hMSC and hMSC TERT cells were passaged into two T₂₅ flasks each at 5000 cells/cm² without hyg for 1 week and then harvested and counted. Similar cultures were combined and passaged again into two T₂₅ flasks. Using cumulative counts, the population-doubling level (PDL) was calculated at each passage as follows:

$$ PDL_{n} = \log \left( \frac{N_f}{N_0} \right) / \log 2 $$

where $n$ = passage number, $N_f$ = final number of cells, and $N_0$ = number of cells seeded at passage. After each passage, $PDL_{n}$ was added to the previous $PDL_{n-1}$ to get the new $PDL_{n}$. Serial passage levels were expressed as PDL or as the passage number as $PDL_{n}$. At each passage, samples were taken for RNA analysis using the Qiagen RNaseasy Miniprep kit (Qiagen, Valencia, CA); cell proliferation was studied using the Cell Proliferation ELISA and BrdU (colcemid) assay using the TiterTACS™ apoptosis detection kit (Roche, Roche Applied Science, Indianapolis, IN); and telomerase activity using the TeloTAGGG telomerase polymerase chain reaction (PCR) ELISA (Roche Applied Science, Indianapolis, IN). Cell proliferation was studied using the Cell Proliferation ELISA and BrdU (colcemid) assay (Roche Applied Science); cell senescence using the Senescence Cells Histochemical Staining Kit (Sigma-Aldrich); and apoptosis using the TiterTACS™ apoptosis detection kit from R&D Systems, Inc. (Minneapolis, MN).

Quantitative real-time reverse transcription–PCR

Gene sequences for the osteogenic marker alkaline phosphatase (ALP), the telomerase (TERT), the 18S ribosomal RNA (18S rRNA), and the tumor suppressor genes p31, p53, and p16Ink4a, and klotho were designed using the National Center for Biotechnology (NCBI) gene database (Table 1). Relative mRNA levels were quantitated by real-time reverse transcription (RT)–PCR using the Opticon continuous fluorescence system (Bio-Rad Laboratories, Hercules, CA) and the SYBR Green RT-PCR kit (Qiagen). Reactions were
performed in triplicate: reverse transcription at 50°C for 30 min, 95°C for 15 min; followed by 40 cycles of 94°C for 15 sec (denaturing), 60°C for 30 sec (annealing), and 72°C for 30 sec (extension). Relative changes in gene expression were calculated in relation to 18S levels and the reference time point using the 2\(^{-\Delta\Delta C(T)}\) method.\(^{19}\)

### Chemotaxis assays

Chemotaxis assays were performed as previously described.\(^{20}\) Cells (2.5 \(\times\) 10\(^4\) in 100 \(\mu\)L) were plated in triplicate onto Costar transwell membranes in the serum-free hMSCGM with 0.1% bovine serum albumin. Lower chambers contained hMSCGM with 1% fetal bovine serum (FBS). Transwells were incubated at 37°C (4 h), fixed with 10% neutral buffered formalin, and stained with hematoxylin. Migrating cells were counted (400 x) on a Nikon Eclipse E400 microscope with five fields analyzed per membrane.

### Osteogenic and adipogenic differentiation

hMSCs and hMSC TERT cells were plated (5000 cell/cm\(^2\)) in hMSCGM in triplicate in 48-well plates and allowed to attach overnight before treatment with OGM. Total RNA was isolated at 0, 3, 7, 10, and 14 days. Differentiation was monitored by mRNA expression of the osteoblastic marker gene \(ALP\) and 18S rRNA as a housekeeping marker. For mineralization analysis, cells were plated (10,000 cell/cm\(^2\)) in 96-well plates, treated with hMSCGM or OGM for 4 weeks, and stained with Alizarin Red. For adipogenic analysis, cells were seeded in 96-well plates (5000 cell/cm\(^2\)), treated with three cycles of AIM/AMM (3 days each cycle), and then cultured in the AMM. Lipid droplets were visualized with Oil Red-O staining 3 weeks after adipogenic induction.

### Statistics

Data are reported as the mean value ±SD. Values were analyzed by nonparametric one-way analysis of variance with Bonferroni post hoc test analysis and by Student’s \(t\)-test using GraphPad Prism 5.04 (GraphPad Software, San Diego, CA). A value of \(p < 0.05\) was considered significant.

### Results

#### Construction of LV

To address the problem of variability in hMSC populations, we sought to develop a cell line of hMSCs that expresses TERT, and therefore may have the potential for continued growth in cell culture. The human TERT gene was subcloned into pLVX-ires-Hyg (Clontech) and is expressed as a bicistronic transcript with the hyg-resistant gene, allowing hyg to be used as an indicator of transduction efficiency and as a marker for selection. The bicistronic construction of this vector ensures that hyg-resistant cells must also be expressing the TERT gene.

#### Phenotype of cells

Early-passage hMSCs have a characteristic fibroblast morphology. Figure 1A shows passage P3 hMSCs detailing the long fibroblast phenotype and the characteristic alignment of the cells as the culture becomes confluent. Later-passage [PDL 25; P15(19)] hMSC TERT cells still retain the fibroblastic phenotype of younger-passage hMSCs.

#### Analysis of culture growth characteristics

Both the hMSC and hMSC TERT cell lines were propagated through serial passage in T\(_{75}\) flasks to assess the ability of the cells to continue to grow in cell culture in the absence of hyg. At early passages, the hMSCs and hMSC TERT cells grew at similar rates, but at PDL 11 [passage P6(10)], the cell yield of the hMSCs decreased steadily and reached a minimum between P13(17) and P16(20) at ~2 \(\times\) 10\(^5\) cell/flask (Fig. 1B). In contrast, hMSC TERT cells produced between 9 \(\times\) 10\(^5\) and 1.5 \(\times\) 10\(^6\) cells/flask throughout the time course of the study. The hMSC TERT cell line has produced a cumulative total of over 7.8 \(\times\) 10\(^7\) cells and shows no signs of diminished proliferation even after 70 PDLs [P40(44); Fig. 1B], whereas the

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**Table 1. Primer Sequences Used in Quantitative Real-Time Reverse Transcription–Polymerase Chain Reaction**

| Gene of interest | Oligonucleotide sequence | GenBank ID |
|------------------|--------------------------|------------|
| ALP sense        | 5'-AACGGCGTGCCTGGGTTGCGG-3' | NM_000478  |
| ALP antisense    | 5'-ACAGGAGAGTCGGCTTTCAG-3' | NM_198253.2|
| TERT sense       | 5'-CTACGGGAGCATGGAGGAAC-3' | NR_003286.2|
| TERT antisense   | 5'-GACACTTCAAGCCGAAGAC-3' | NM_004795.3|
| 18S rRNA sense   | 5'-CCGCGAGTTCACCTACTG-3' | NM_000389.4|
| 18S rRNA antisense| 5'-CGGGTCATAAGCTTGCGT-3' | NM_000546.4|
| KLOTHO sense     | 5'-GTCGGCCGTGACCGGCTACG-3' | NM_058195.3|
| KLOTHO antisense | 5'-GCCAGACTTTGCGATGAGCCAG-3' | NM_000389.4|
| p21 sense        | 5'-AGAGAAGGCGCGGTCAGGACCC-3' | NM_000546.4|
| p21 antisense    | 5'-GAATGCAGAGCGGCTACG-3' | NM_000389.4|
| p53 sense        | 5'-TAACCGAGCCTGCGCAAAACAAC-3' | NM_000546.4|
| p53 antisense    | 5'-GCTTCGGAACATCTGCGACCC-3' | NM_000546.4|
| p16\(^{INK4a}\) sense | 5'-GTGGACCTGCTGAGGAG-3' | NM_058195.3|
| p16\(^{INK4a}\) antisense | 5'-CTTTCATTGGGATGTCTG-3' | NM_058195.3|

\(ALP\), alkaline phosphatase; \(TERT\), telomerase reverse transcriptase; 18S rRNA, 18S ribosomal RNA; \(p21\), cyclin-dependent kinase inhibitor 1A; \(p53\), tumor protein 53.
mock-transduced hMSC line ceased proliferating at $1.5 \times 10^7$ cells after 14 PDLs [P16(20)].

**Analysis of TERT expression and activity in hMSC TERT cells**

RNA isolated at various passages throughout the time course was analyzed by real time RT-PCR to confirm TERT expression. hMSCs [PDL 6; P4(8)] have a low level of detectable TERT mRNA (Fig. 2A), whereas P4(8) hMSC TERT cells have over 6400 $\times$ the level of TERT mRNA as hMSCs. TERT mRNA in hMSCs decreases to almost undetectable levels by early passage P6(10). Even though the level of TERT mRNA in the hMSC TERT cells decreases slightly, there is still significantly higher expression than hMSC levels at P4(8). Telomerase enzyme activity was also measured at various points in comparison to early-passage hMSCs (Fig. 2B). At PDL 10 [P6(10)], hMSC TERT cells have significantly higher levels of telomerase activity in comparison to hMSCs at the same passage. The high level of telomerase activity was sustained at later passages [P16(20) and P26(30)]. mRNA levels of the cell cycle regulator p16$^{INK4a}$ significantly increased at P13(17) in hMSCs, but not in TERT cells (Fig. 3). Tumor suppressor genes p21, p53, and klotho were also quantitated, but were not significantly different comparing hMSC and hMSC TERT cultures at these passages (data not shown).

**Cell proliferation, senescence, and apoptosis**

To determine the nature of the enhanced growth capacity of hMSC TERT cells, we examined cell proliferation, cellular senescence, and apoptosis at various passages. Cell proliferation was assayed at both early [P8(12)] and late [P16(20)] passages. hMSC TERT cells exhibited a significantly higher rate of cell proliferation after 4 days in culture in comparison to hMSCs at a similar passage (Fig. 4A). Cell proliferation at late passage in the hMSCs was significantly reduced from that seen in early passage hMSCs, whereas cell proliferation in hMSC TERT cells was only slightly reduced when comparing early- and late-passage cells.

Cell senescence was assayed using $\beta$-galactosidase activity as a marker at four consecutive passages. The percentage of cells exhibiting $\beta$-galactosidase staining was unchanged in hMSC TERT cells even up to P40(44), whereas $\beta$-galactosidase activity was not significantly different in hMSCs at P10 and...
but significantly increased at passages P12 and P13, representing an increase in cell senescence in hMSCs after only 12 passages (Fig. 4B).

DNA fragmentation as a measure of apoptosis was analyzed in late-passage [P16(20)] cells. Although the apoptosis level was lower in hMSC TERT cultures in comparison to hMSC cultures, the results were not significantly different (Fig. 4C). Taken together, these results indicate that the enhanced growth characteristics of hMSC TERT cells are due to increased cell proliferation and inhibition of cellular senescence.

hMSCs have the ability to chemotax toward several attractants. To test if later-passage hMSC TERT cells retain this ability, chemotaxis assays were performed to compare late-passage [P8–P10] and hMSC TERT cells [P11(15)–P12(16)]. hMSCs show a low level of chemotaxis toward a medium containing 1% FBS, whereas hMSC TERT cells exhibit a significantly higher level of chemotaxis at even later passages (Fig. 5).

**Differentiation capability of hMSC TERT cells**

To test the capacity of hMSC TERT cells to undergo differentiation toward the osteogenic lineage, cells were treated for 14 days with OGM. OGM treatment of late-passage [P16(20)] hMSC TERT cells significantly induced mRNA expression of the early chondro-/osteogenic marker alkaline phosphatase (hALP) beginning at 7 days and throughout the 14-day time course of the experiment (Fig. 6A). This is consistent with our previous results for hMSC osteogenic differentiation.21 hMSC TERT cells from early and late passages were also treated for 4 weeks in OGM and then analyzed for mineral deposition using Alizarin Red stain (Fig. 6B). Alizarin Red-stained wells were only present in wells treated with OGM, indicating that the cells reached the late stage of osteoblastic differentiation and could deposit mineral. Late-passage [P22(26)] hMSC TERT cells were also treated with AIM to confirm the ability of these cells to differentiate toward the adipogenic lineage (Fig. 6C). Oil Red-O staining confirmed the presence of lipid deposits in those wells treated with AIM and absent in those treated with AMM.

**Discussion**

hMSCs have the potential to be useful in a variety of reconstructive engineering capacities due to the inherent multipotent capacity of the cells to differentiate into osteogenic, adipogenic, and chondrogenic lineages.23 A major drawback to using hMSCs is that they are only present in low numbers...
in the bone marrow, and although they propagate readily in cell culture, serial passage of the cells ultimately leads to replicative senescence limiting the number of available cells that could be used in a clinical setting. Replicative senescence in hMSCs resembles an aging process with each consecutive passage becoming slightly more differentiated. Problems arise when trying to use the cells to study developmental processes as the cultures are continually changing (aging), necessitating the need to use multiple donors to do repetitive experiments. To address the problem of replicative senescence, we sought to constitutively express the TERT gene in hMSCs, thereby immortalizing the culture, and hopefully lessening the variability of the cultures due to aging in vitro.

The TERT gene was inserted into pLVX-IRES-Hyg (Clontech) and is expressed as a bicistronic transcript with the hygromycin (hyg)-resistant gene. Even though the hMSC TERT cell line is resistant to hyg, the cultures were grown without the addition of hyg for two reasons. First, to assess the stability of the TERT expression without selective pressure, and second, since cell growth of hMSC TERT cells was slower in the presence of hyg (data not shown), we wanted to be sure that we were assessing the effect of TERT expression and not the effect of hyg in comparison to the hyg-sensitive hMSCs. This cloning strategy (hyg resistance with TERT expression) was undertaken with the knowledge that Clontech also produces another vector with puromycin (puro) resistance (pLVX-Puro) that could be used to express other inserted genes selecting for both hyg and puro, thus generating immortalized cell lines expressing both TERT and another gene of interest.

hMSC TERT cells have been serial-passaged to over 70 population doublings and still exhibit the characteristic fibroblastic cell phenotype typical of early-passage hMSCs. TERT expression is stable without the need for continued hyg selection. Previous studies have shown that constitutive expression of TERT by itself does not generate malignant conditions; although, TERT expression has been associated with cancer. We also looked at expression of the tumor suppressor genes p16INK4a, cyclin-dependent kinase inhibitor p21 (Waf-1/Cip-1), p53, and klotho. The expression of these tumor suppressor genes in the hMSC TERT cell line is not significantly different than expression in early-passage hMSCs. Additionally, both p16INK4a and klotho have been characterized as genes associated with aging. p16INK4a is frequently induced as cells age and inhibits stem cell proliferation and regenerative potential, and was induced in later passages of hMSCs, but not induced in the hMSC TERT cell line. Klotho-deficient and klotho-null mice exhibit a premature aging phenotype, whereas overexpression of klotho has been shown to extend the lifespan of such mice by 20%–30%. While it was surmised that TERT expression may increase klotho expression, we did not see any changes in klotho gene expression. TERT expression does not significantly alter expression levels of these tumor suppressor genes, and therefore the hMSC TERT cells do not appear to be neoplastic, although this would have to be verified in vivo. The multipotent nature of hMSC TERT cells is demonstrated with differentiation toward the osteogenic and adipogenic lineages.

FIG. 5. Chemotaxis toward a medium containing 1% fetal bovine serum (FBS). Data are expressed as a chemotactic index, defined as the number of cells migrating in response to the medium (1% FBS) divided by the number of cells migrating in the negative control (–). hMSC TERT cells [P12(16)] exhibit significantly higher levels of chemotaxis than hMSCs at earlier PDL (P8). *p < 0.05 compared to negative control.

FIG. 6. Differentiation of hMSC TERT cells toward the osteoblastic and adipogenic lineages. hMSC TERT cells were treated with an osteogenic medium (OGM) for 14 or 28 days. (A) Relative changes in alkaline phosphatase (ALP) expression were determined by real-time RT-PCR. ALP mRNA expression is significantly upregulated at 7, 10, and 14 days. *p < 0.05 compared to hMSC TERT levels at day 0. (B) Alizarin Red staining of early- [P5(9)] and late- [P16(20)] passage cells demonstrates that the hMSC TERT cells can undergo terminal differentiation, resulting in the deposition of mineral after OGM treatment for 28 days. (C) Oil Red-O staining of lipid droplets confirms that the hMSC TERT cells can also undergo adipogenic differentiation after induction for 21 days.
Osteogenic differentiation leads to both ALP mRNA induction and mineralization as we have reported previously in hMSC cultures, and adipogenic differentiation leads to cells filled with lipid droplets.

In summary, we have generated a hMSC line that is hyg resistant, does not exhibit replicative senescence typical of primary hMSCs, and retains the ability to chemotax and differentiate toward the osteogenic and adipogenic lineages. The hMSC TERT cell line is ideal for both basic and applied tissue engineering studies of bone development and repair due to the reduced variability offered by the enhanced growth potential in cell culture over that of hMSC primary cultures.

Acknowledgment

This work was supported by a Veterans’ Affairs National Merit Review Grant awarded to D.T.Y.

Author Disclosure Statement

The authors have no financial interest in the products mentioned in the article.

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