Low Oxygen Modulates Multiple Signaling Pathways, Increasing Self-Renewal, While Decreasing Differentiation, Senescence, and Apoptosis in Stromal MIAMI Cells

Carmen Rios,1,2,* Gianluca D’Ippolito,2–6,* Kevin M. Curtis,1,2 Gaëtan J.-R. Delcroix,2,3,5 Lourdes A. Gomez,2 Jimmy El Hokayem,1 Megan Rieger,1 Ricardo Parrondo,2 Alicia de las Pozas,2 Carlos Perez-Stable,2,7 Guy A. Howard,1,2,7 and Paul C. Schiller1–5

Human bone marrow multipotent mesenchymal stromal cell (hMSC) number decreases with aging. Subpopulations of hMSCs can differentiate into cells found in bone, vasculature, cartilage, gut, and other tissues and participate in their repair. Maintaining throughout adult life such cell subpopulations should help prevent or delay the onset of age-related degenerative conditions. Low oxygen tension, the physiological environment in progenitor cell-rich regions of the bone marrow microarchitecture, stimulates the self-renewal of marrow-isolated adult multilineage inducible (MIAMI) cells and expression of Sox2, Nanog, Oct4a nuclear accumulation, Notch intracellular domain, notch target genes, neuronal transcriptional repressor element 1 (RE1)-silencing transcription factor (REST), and hypoxia-inducible factor-1 alpha (HIF-1α), and additionally, by decreasing the expression of (i) the proapoptotic proteins, apoptosis-inducing factor (AIF) and Bak, and (ii) senescence-associated p53 expression and β-galactosidase activity. Furthermore, low oxygen increases canonical Wnt pathway signaling coreceptor Lrp5 expression, and PI3K/Akt pathway activation. Lrp5 inhibition decreases self-renewal marker Sox2 mRNA, Oct4a nuclear accumulation, and cell numbers. Wortmannin-mediated PI3K/Akt pathway inhibition leads to increased osteoblastic differentiation at both low and high oxygen tension. We demonstrate that low oxygen stimulates a complex signaling network involving PI3K/Akt, Notch, and canonical Wnt pathways, which mediate the observed increase in nuclear Oct4a and REST, with simultaneous decrease in p53, AIF, and Bak. Collectively, these pathway activations contribute to increased self-renewal with concomitant decreased differentiation, cell cycle arrest, apoptosis, and/or senescence in MIAMI cells. Importantly, the PI3K/Akt pathway plays a central mechanistic role in the oxygen tension-regulated self-renewal versus osteoblastic differentiation of progenitor cells.

Introduction

Human multipotent mesenchymal stromal cells (hMSCs) are a heterogeneous population of cells present in bone marrow [1]. The number of hMSCs with osteogenic potential (CFU-F/ALP+) decreases during human aging [2]. Marrow-isolated adult multilineage inducible (MIAMI) cells represent a homogeneous subpopulation of developmentally immature MSCs [3] expressing sustained levels of pluripotency markers (Oct4, Nanog, Sox2, Rex1, hTeRT, and SSEA-4) under defined growth conditions [4]. Supporting the immature nature of MIAMI cells, we demonstrated in a recent proteomic analysis that MIAMI cells have more proteins in common with embryonic stem cells (ESCs) than regular MSCs do [5]. MIAMI cells expanded at low oxygen consistently respond to osteogenic stimulation independently of donor age [4], suggesting a sustained osteogenic capacity during normal aging. Physiologic oxygen tension ranges from ≤1% in cartilage and regions of the bone marrow to 10%–13% in the main arteries of most tissues [6]. Increasing evidence suggests that oxygen tension is a key regulator of cell growth, proliferation, senescence, and differentiation of stem and progenitor cells [4,7–13]. We reported that low oxygen enhances MIAMI cell pluripotency, proliferation, and maintenance of osteoblastic differentiation capability [4]. However, little is known about the influence of oxygen on the modulation of signaling pathways, particularly, those that influence the interplay between self-renewal, proliferation, differentiation, and senescence.

1Department of Biochemistry and Molecular Biology, University of Miami Miller School of Medicine, Miami, Florida.
2GRECC and Research Service, Veterans Affairs Medical Center, Miami, Florida.
3Department of Orthopaedics, Geriatrics Institute, and Interdisciplinary Stem Cell Institute, University of Miami Miller School of Medicine, Miami, Florida.
4Department of Biomedical Engineering, University of Miami, Coral Gables, Florida.
5Department of Medicine, University of Miami Miller School of Medicine, Miami, Florida.
6These authors contributed equally to this work.
Stem cell self-renewal involves several pathways, including Notch, Wnt, repressor element (RE)-1-silencing transcription factor (REST), Sirtuin, and the PI3K/Akt/mTOR pathway [13–18]. The highly conserved Notch pathway can regulate stem cell self-renewal [13,19]. Notch releases an intracellular fragment (notch intracellular domain, NICD) [20] that migrates to the nucleus where it regulates expression of target genes. NICD can also recruit HIF-1α to Notch-responsive promoters and elevate expression of Notch downstream targets, while blocking neuronal and myogenic progenitor differentiation [21]. Hypoxia may block this differentiation program by upregulating the stem cell transcription factor, Oct4. Covello et al. used a genetic knock-in strategy to demonstrate that targeted replacement of the oxygen-regulated transcription factor, HIF-1α, with its alternative isoform, HIF-2α, resulted in expanded expression of HIF-2α-specific target genes, including Oct4 [22]. Moreover, Oct4 shRNA reduced β-catenin levels in Hif-2α KI/KI tumors [22]. β-Catenin is a major mediator of the canonical Wnt signaling pathway and is also involved in self-renewal [23].

The Wnt signaling pathway [23,24] regulates downstream targets involved in modulating stem cell proliferation and self-renewal, including cell cycle regulator, Cyclin D1 [25], and pluripotency marker, Oct4 [26]. Pluripotent transcription factors, SOX2, OCT4, and NANOG, key regulators in ESCs, maintain self-renewal and multipotency of adult stem cells [27–30]. Specifically, maintenance of self-renewal and multipotency of hMSCs requires Sox2 [31,32] through mechanisms involving sirtuin [18] and Wnt [31] signaling. Wnt signaling can cross talk with the antipapoptotic/prosurvival pathway, PI3k/Akt, promoting self-renewal and expansion of stem cells [33]. Interestingly, transient hypoxia increases mouse ESC proliferation through activation of the PI3K/Akt pathway [17]. Moreover, proapoptotic protein BAD, an upstream regulator of cell self-renewal, including cell cycle regulation, apoptosis, and/or senescence gene expression. Furthermore, we used specific inhibitors to determine a mechanistic role of these pathways. This may help develop molecular approaches aimed at promoting stem cell self-renewal, thus preventing or decreasing the observed age-related decline of stem cells and tissue function in vivo [2].

### Materials and Methods

**Antibodies**

Oct4 was from Abcam (Cambridge, MA). FITC/rhodamine secondary for Oct4 was from Invitrogen (Carlsbad, CA). Mounting solution with DAPI was from Vector Laboratories (Burlingame, CA). Antibodies for western blotting, Nanog, hTeRT, Sox2, Oct4a, Oct4b, Lrp5, p53, AIF, Bak, p21, p27, EndoG, Bax, Bid, Bcl-xI, CIAP-2, Mcl-1, XIAP, HIF-1α, NICD, REST, Akt, and α-tubulin were from Abcam.

### Isolation and culture of human MIAMI cells

We previously reported the isolation of MIAMI cells in detail [3,4]. Briefly, we isolated MIAMI cells from commercially available human whole bone marrow (BM) (Lonza, Walkersville, MD, BM from a 20-year-old male, a 3-year-old male, or a 7-year-old male donor). For expansion, we plated MIAMI cells at a density of 100 cells/cm² in expansion medium in a 100% humidified atmosphere of 3% O₂,5% CO₂, and 92% N₂ (low oxygen tension) and one-half medium changed twice a week. We used only MIAMI cells from passages 3 and 4. Cells receiving exogenous treatments with recombinant mouse Wnt3a underwent a 24-h period of serum starvation before addition of 5 ng/mL of recombinant mouse Wnt3a in the medium. To test Akt activation (phospho-Akt), we plated MIAMI cells in expansion medium and exposed them to atmospheres of 21% or 3% O₂. We measured phospho-Akt and total Akt after 30 and 60 min of incubation. We inhibited the PI3K/Akt signaling pathway using different concentrations of wortmannin, 20, 50, 100, 200, and 300 nM, on MIAMI cells incubated for 30 min at 3% O₂ in expansion medium.

### Quantitative real-time PCR

We performed quantitative real-time PCR (RT-qPCR) as previously described and normalized the fluorescence data to the housekeeping genes, eukaryotic translation elongation factor 1 alpha (EF1α) and/or ribosomal protein large subunit 13a (RPL13a) [40]. Supplementary Table S1 (Supplementary Data are available online at www.liebertpub.com/scd) lists the DNA oligonucleotide sequences used for RT-qPCR.

### Western blot

We prepared protein samples using NP-40 lysis buffer, separated them by sodium dodecyl sulfate–polyacrylamide gel electrophoresis (SDS-PAGE) on 10% polyacrylamide gels, and then transferred them onto a PVDF membrane (electrophoresis and transfer apparatus from Bio-Rad; Hercules, CA) for approximately 60–90 min at 100V. We incubated blots with primary antibodies at 1:1,000 overnight at 4°C with gentle rocking, rinsed them with TBS, and incubated them with secondary antibodies at 1:5,000 for 45 min at room temperature with gentle rocking. We normalized protein levels to α-tubulin and/or Coomassie blue-stained membranes. We performed quantitation of blots using ImageJ software from NIH. We normalized data (using either Coomassie blue total protein or β-catenin) to correct for variations in protein loading.

### Immunoprecipitation

We aliquoted 25 μL of protein A/G Plus agarose beads (Santa Cruz Biotech., Inc., Santa Cruz, CA) with 2 μg of primary antibody to TCF4 and incubated them for 1 h at room temperature on a rotator. We washed the protein A/G agarose beads 3× with 1 mL PBS (Invitrogen, Carlsbad, CA) and then centrifuged them at 3,000 × g for 2 min at 4°C. We adjusted the concentration to 200–400 μg of protein sample to 200 μL PBS.
and incubated the protein/bead solution overnight at 4°C on a rotator. The next day, we washed the protein/bead solution 3× with PBS before centrifugation at 3,000 g for 2 min at 4°C. We saved the supernatants and separated the proteins using PAGE. Samples run without primary antibody or with isotypic nonimmune (normal rabbit) serum served as negative controls. We incubated the blots in primary antibodies (diluted in either 0.1% BSA or 5% nonfat milk) overnight at 4°C and secondary antibodies conjugated with horseradish peroxidase (diluted in either 0.1% BSA or 5% nonfat milk) for 45 min at room temperature. We finally incubated the blots with ECL chemiluminescence solution (GE Life Sciences) as per the manufacturer’s instructions. Final levels of proteins were normalized to Coomassie-stained blots for total protein.

siRNA transfections

We transfected MIAMI cells 1 day after seeding with 50, 100, or 200 nM siRNA ON-TARGET plus SMART pool (siLrp5: CGUCAAAGCCAUCGCAUUUU, CGUCAUGG GUGGUGUCUAUUU, GGACGGAUCCACGGAGAUAU, GUACAGGC CUUCAUCAUUUU; siLrp6: GCAGAUAU CAGA GAUUUUU, CAGAUGAACUUGAUGUAAUU, CCACAGAAGGCUACCAUAUUU, GCUCAAACGGUGAAG UUAUUU) or siCONTROL Non-Targeting pool (combination of four siRNA sequences that are not targeted to Lrp5 or 6 and found to have little off-target effects, all from Dharmacon, Chicago, IL). We performed the transfections with the Nucleofector electroporator (Amaxa; Walkersville, MD). We trypsinized 1 to 2×10⁶ cells for each transfection procedure. We used Nucleofector solution as our transfection reagent according to the Amaxa protocol for the Nucleofector Kit. Immediately following transfections, we placed the cells dropwise into incubated six-well plates containing MIAMI expansion media. Twenty-four hours after the transfections, we replaced all media with fresh MIAMI expansion media. Cells were left undisturbed for 3 days before trypsinization for a second round of transfection using the same procedure stated above. We collected the cells for RNA and protein analysis 3 days after the second transfection. To assess transfection efficiency, we transfected MIAMI cells with 1 μg GFP (green fluorescent protein) plasmid pEGFP-C1 (Clontech, Mountain View, CA) and visualized them on a Nikon fluorescence microscope.

Enzyme-Linked Immunosorbent Assay

We measured Wnt3a concentrations in the media of MIAMI cells cultured in 3% and 21% O₂ using enzyme-linked immunosorbent assay (ELISA) as follows: we coated a 96-well plate with capture antibody anti-Wnt3a (R&D Systems, Minneapolis, MN) at a 0.5 μg/mL dilution with PBS. We added media from MIAMI cells grown at 3% and 21% oxygen to the capture antibody preparation. We added fresh media as a negative control. We carefully washed the medium from the plate and added the detection biotinylated anti-Wnt3a (0.5 μg/mL). We finally used a horseradish peroxidase-conjugated streptavidin solution and stopped the colorimetric reaction with 2 M sulfuric acid. Optical density was measured at 450 nm on a microplate reader (BioRad, Hercules, CA).

Oct4 immunofluorescence

For nuclear proteins, we fixed cells with 4% PFA at 4°C for 10 min and permeabilized them with 0.1% Triton X-100 for 10 min. Blocking and diluent solution consisted of PBS with 1% BSA. We blocked the fixed cells for 30 min and incubated them sequentially for 1 h with the primary antibody to Oct4, followed by a 1-h incubation of the fluorescein- or rhodamine-conjugated secondary anti-goat IgG antibody. We used PBS plus 0.3% BSA for the washes between each step. We used isotypic antibodies as controls. Imaging was performed using a Nikon Eclipse TI microscope as well as a Nikon C2 confocal microscope (Nikon Instruments, Melville, NY).

Microarray analysis

We used the Oligo GEArray® System (SuperArray). The array requires biotin-labeled cRNA to hybridize to the membrane. We synthesized the labeled cRNA needed for hybridization to the array with the TrueLabeling-AMP™ 2.0 (SuperArray) system. We used a total of 3 μg of RNA in the system to optimize for the final cRNA yield. We quantified the cRNA and used the same initial amount for both the 21% and 3% O₂ samples. We performed hybridization until chemiluminescent detection performed on the ChemiGenius (Syngene, Frederick, MD) imager and analysis with the GEArray Analysis Suite.

Senescence-associated β-galactosidase activity

We assayed the cells for senescence-associated β-galactosidase (SA-β-Gal) activity using the Cell Signaling kit #9860 (Cell Signaling Technology, Danvers, MA) as per the manufacturer’s instructions. In brief, we plated the MIAMI cells (passage 4–6) at 200 cells/cm² in six-well plates in either 3% or 21% oxygen tension for up to 1 or 3 weeks. To assess for SA-β-Gal, we removed the medium and washed the cells with PBS before fixation for 10–15 min at room temperature. We rinsed the cells again with PBS, followed by application of the β-galactosidase staining solution and incubation at 37°C overnight in a dry incubator in the absence of CO₂. Development of a blue color indicated SA-β-Gal activity. Ten fields of cells were counted per sample to determine the percentage of cells in a senescence state.

Osteoblastic differentiation conditions

We plated MIAMI cells at 2,000 cells/cm² in 24-well plates in α-MEM supplemented with 10% FBS (HyClone, Logan, UT), 100 U/mL penicillin, 1 mg/mL streptomycin, 10 nM dexamethasone, 100 μM ascorbic acid 2-phosphate, and 10 nM β-glycerophosphate [2].

Alkaline phosphatase histochemical analysis

MIAMI cells in osteogenic medium at 3% and 21% pO₂ were rinsed in PBS, fixed with a solution of 2% cold-buffered formaldehyde and 0.2% gluteraldehyde for 30 min at 4°C, rinsed with distilled H₂O, and then stained for alkaline phosphatase activity at the times indicated in the corresponding Figure legend. Briefly, we dissolved 8 mg of naphthol AS-TR phosphate in 0.3 mL of N, N-dimethylformamide. Simultaneously, we dissolved 24 mg of fast blue BB in 30 mL of 100 mM Tris-HCl, pH 9.6. We mixed these two solutions, added 10 mg of MgCl₂, and then adjusted the pH to 9.0 with 1N
HCl. We filtered the final solution through a 0.2-μm filter. We finally incubated the cells at 37°C for 30 min and rinsed them with distilled H2O [2].

Statistical analysis

Results are presented as the mean of three independent experiments ± standard deviations (SD). We tested the differences by one-way ANOVA and calculated the P-values using the Newman–Keuls post hoc test. We considered a threshold P-value ≤0.05 as significant and depicted it with an * in the Figures.

Results

Low oxygen stimulates the expression of transcription factors associated with stem cell self-renewal

Previously, we reported that low oxygen (3%) upregulated mRNAs for total Oct4 (isoforms a and b), Rex-1, hTeRT, SSEA-4, and HIF-1α under expansion and osteogenic conditions [4]. We hypothesized that low oxygen may also stimulate the expression of other known mediators of self-renewal, such as Nanog, Sox2, and Oct4a (the active Oct4 splice variant, which localizes to the nucleus [41]). In cells grown at 3% versus 21% oxygen for 7 days, we found increased levels of transcripts for these genes and Lrp5 (Fig. 1A), a key coreceptor and modulator of the Wnt signaling pathway implicated in the maintenance of mouse embryonic stem cell (mESC) pluripotency. To determine whether low oxygen tension treatment would acutely increase the protein level, we exposed to 21% O2 overnight cells previously expanded at 3% O2 and then switched to 1% O2 for 2 h (acute setting). Western blot analysis showed that protein levels of Oct4a/b, Nanog, and, more dramatically, Sox2 increased after acute low oxygen (Fig. 1B).

Low oxygen increases the nuclear levels of Oct4a

To determine whether Oct4 localized to the nucleus and to distinguish between the Oct4a/b isoforms, we examined cytoplasmic and nuclear fractions and used immunofluorescence
to assess subcellular Oct4 localization. Western blot analysis demonstrated increased levels of nuclear Oct4 after 1 week at 3% versus 21% O2 (Fig. 1C). We further confirmed nuclear localization by immunofluorescence and confocal studies that showed stronger Oct4 nuclear staining in cells at low (3%) versus ambient (21%) oxygen (Fig. 1D).

**Low oxygen increases expression of Notch signaling pathway mediators**

To examine potential signaling pathways mediating the low oxygen-stimulated self-renewal program of MIAMI cells, we performed a signaling pathway array on cells cultured for 7 days at 3% versus 21% O2 (Supplementary Fig. S1). Genes upregulated at 3% O2 are listed in Supplementary Table S2. The data revealed upregulation of key Notch mediators among others. RT-qPCR confirmed gene expression stimulation of key genes: Notch2, STAT6, and Jagged 1 (Fig. 2A). To determine involvement of Notch signaling activation, we examined low oxygen effects on NICD nuclear levels. Western blot analyses demonstrated that NICD protein levels increased (3.8-fold) in the nucleus at low oxygen along with a simultaneous nuclear increase of HIF-1α (3.2-fold), a low-oxygen transcriptional mediator and NICD cofactor for the transactivation of specific target genes [21] (Fig. 2B).

**Low oxygen increases REST/NRSF levels**

MIAMI cells can be induced to differentiate into neuron-like cells [3,42–45]. REST/NRSF repressed neuronal differentiation in mouse ESCs [36]. Thus, we sought to examine whether REST/NRSF may contribute to MIAMI stem cell self-renewal by inhibiting neuronal differentiation. RT-qPCR showed that MIAMI cell mRNA levels of REST and co-REST increased in 3% versus 21% O2 conditions (Fig. 2C). Furthermore, the REST protein accumulated at 3.6-fold higher levels in the nucleus of cells under low oxygen conditions (Fig. 2D).

**Low oxygen regulates expression of specific apoptosis/senescence pathway genes**

We examined whether the expression of cell cycle control, apoptosis, and senescence pathway-associated genes was affected by low oxygen. The tumor suppressor p53 is a master regulator of these programs by differentially modulating

---

**FIG. 2.** Low oxygen activates Notch signaling in MIAMI cells and increases REST and co-REST expression. (A) Relative fold change of Notch mediator (Notch2, Stat6, and Jagged1) mRNA levels after 7 days at 3% compared with 21% O2 measured by RT-qPCR and normalized to housekeeping genes, EF1α and RPL13a. (B) Western blot analysis: lane 1 shows protein lysates from MIAMI cells grown at 21% O2, lane 2 cells were grown at 3% O2. Top panel shows HIF-1α protein, center panel shows NICD protein. (C) mRNA levels of REST and co-REST as measured by RT-qPCR from MIAMI cells grown at high and low oxygen for 7 days. (D) Protein levels of REST and co-REST were measured by western blot from the nuclear fraction of MIAMI cells grown at high and low oxygen for 7 days (top panel). Coomassie blue staining used as loading control (bottom panel). REST, repressor element-1-silencing transcription factor; RT-qPCR, quantitative real-time PCR. *P ≤ 0.05.
target genes controlling cell cycle arrest, apoptosis, or senescence [46,47]. We found that low oxygen decreased protein expression of p53 by approximately fourfold (Fig. 3A), while p21 and p27 levels did not appear to be affected. Furthermore, expression of the proapoptotic proteins, apoptosis-inducing factor (AIF) and Bak, was reduced five- and threefold, respectively (Fig. 3A). Expression of endoG, Bax, Bid, Bcl-xL, and cIAP-2 was not affected, while the inhibitors of apoptosis, XIAP and Mcl-1, appeared to increase slightly (Fig. 3A). These results are consistent with an overall cell prosurvival program at low oxygen. To further assess an effect on cell senescence, we quantified the number of cells with detectable senescence-associated β-galactosidase (SA-βgal) activity and determined that 3% O₂ dramatically decreased positive cells versus 21% O₂ (5% ± 2% vs. 48% ± 8% at 3 weeks, respectively, P ≤ 0.01; Fig. 3B).

**Low oxygen stimulates the canonical Wnt pathway**

Since we observed 3% O₂ stimulation of Lrp5 (Fig. 1A) and frizzled 1 and 2 (Supplementary Fig. S1B), we further examined canonical Wnt pathway activation. Western blot analysis of nuclear and cytoplasmic extracts of MIAMI cells demonstrated that 3% oxygen stimulated cytoplasmic β-catenin (1.5-fold) and, to a larger extent, nuclear β-catenin (2.3-fold, Fig. 4A). Once in the nucleus, β-catenin binds a member of the high mobility group (HMG) DNA-binding protein family of transcription factors. We investigated canonical Wnt signaling specific protein–protein interaction in MIAMI cells. We pulled down proteins bound by T-cell Factor 4 (TCF4) antibody with protein-A/G beads. Subsequently, we size separated, blotted, and probed the immunoprecipitated proteins with β-catenin antibody. Samples from cells at 3% O₂ displayed more than twofold greater association between β-catenin and TCF4 versus 21% O₂ (Fig. 4B). Subsequently, we examined downstream targets of canonical Wnt activation. The cell cycle protein and direct Wnt target, Cyclin D1, was increased by 3% O₂ about 2.2-fold in the cytoplasm and by more than 80-fold in the nucleus (Fig. 4C).

To determine the magnitude of the canonical Wnt signaling activation by low oxygen, we compared β-catenin nuclear localization at 3% and 21% O₂ in the absence and presence of recombinant Wnt3a. Both 3% O₂ and Wnt3a increased to similar levels of cytoplasmic (1.5 vs. 1.4-fold) and nuclear (2.3-fold) β-catenin, while the combination did not have an additive effect (Fig. 4D). To determine the effect of Wnt3a on the nuclear localization of a different low oxygen-sensitive protein, we examined the levels of Oct4 in the cytoplasm and nucleus at 3% and 21% O₂ in the presence and absence of Wnt3a. Surprisingly, we found that in contrast to low oxygen that stimulated nuclear Oct4...
Localization by 1.5-fold (Fig. 1C), Wnt3a increased Oct4 nuclear translocation by 8.6-fold at 21% O2 and by 6.2-fold at 3% O2 (Fig. 4D).

Since low oxygen induced canonical Wnt pathway stimulation, we sought to determine whether this was a paracrine effect accomplished through a low oxygen-stimulated secretion of canonical Wnt signaling activators. We found that 3% O2 significantly stimulated Wnt3a mRNA (20-fold), while Wnt7 and Wnt11 mRNAs were only minimally stimulated (Supplementary Table S2). Interestingly, low oxygen failed to stimulate Wnt3a secretion, quantified by ELISA (Fig. 4E). Furthermore, we demonstrated that either low oxygen or Wnt3a stimulation increased cell numbers to similar extents; however, when combined, the effect was additive (Fig. 4F).

Lrp5-mediated inhibition of the canonical Wnt signaling pathway decreases self-renewal capacity of MIAMI cells, self-renewal marker mRNAs, and Oct4a nuclear accumulation

Canonical Wnt activity requires receptors, Lrp5/6. Basal mammary cells express both Lrp5 and Lrp6 similarly. Lrp5-mediated inhibition of the canonical Wnt signaling pathway decreases self-renewal capacity of MIAMI cells, self-renewal marker mRNAs, and Oct4a nuclear accumulation.

Interestingly, high Lrp5 expression identifies the stem cell-enriched population and is required for maintenance of the basal cell layer. The absence of Lrp5 specifically depletes adult regenerative stem cell activity to less than 1% [48]. It has been shown that an increased expression of Lrp5 is sufficient for activation of the canonical Wnt signaling pathway [49]. In light of our data showing that low oxygen increased Lrp5 mRNA transcript levels (Fig. 1A) and our ELISA data showing equal secretion of Wnt3a at low and high oxygen tension (Fig. 4E), we sought to examine the possibility that Lrp5 could be the rate-limiting step in the low oxygen-stimulated Wnt activity we observed in our cells and that it may be essential for MIAMI cell self-renewal.

We found that exposure of cells to siLrp5 (100-nM) decreased Lrp5 protein levels (Fig. 5A) as well as Oct4a, Sox2, and Lrp5 mRNA levels, although only the last two with statistical significance (Fig. 5B, *P ≤ 0.05 and **P ≤ 0.01, respectively). The number of cells was decreased by 50% after comparing the cell number of cells seeded at 100 cells/cm2 and grown at 3% or 21% oxygen with or without rmWnt3a treatment for 7 days. **P ≤ 0.01 and ***P ≤ 0.001.

FIG. 4. Low oxygen tension stimulates the canonical Wnt pathway. Low oxygen increases (A) nuclear accumulation of β-catenin. (B) β-Catenin/TCF4 interactions: immunoprecipitation of 200 μg protein samples with TCF4, which were size separated and probed with β-catenin Abs in MIAMI cells grown at 3% oxygen (lane 1) and 21% oxygen (lane 2). Isotypic negative controls (normal rabbit serum); 3% samples (lane 3) and 21% (lane 4) Coomassie blue stain (bottom panel) as loading control. (C) CyclinD1 cytoplasmic (lanes 1–2) and nuclear proteins (lanes 3–4) at 3% and 21% oxygen. Coomassie stain is loading control. (D) Changes in the levels of cytoplasmic and nuclear β-catenin in response to oxygen and Wnt3a stimulation. Changes in the levels of cytoplasmic and nuclear Oct4 in response to oxygen and Wnt3a stimulation. (E) Low oxygen fails to stimulate Wnt3a secretion: ELISA from MIAMI cell-conditioned media taken from cells grown for 7 days at 3% and 21% oxygen. A media-only sample was run to determine baseline levels of Wnt3a. (F) Exogenous Wnt3a increased the cell number at low and high oxygen tension: total cell numbers of cells seeded at 100 cells/cm2 and grown at 3% or 21% oxygen with or without rmWnt3a treatment for 7 days. **P ≤ 0.01 and ***P ≤ 0.001.
nuclear localization of Oct4 in siControl cells (Fig. 5C) consistent with untreated cells in Fig. 1D. In siLrp5-treated cells, Oct4 protein did not appear to accumulate in the nucleus or cytoplasm (Fig. 5C); an indication that Oct4a-specific nuclear levels decrease in the absence of Lrp5. We sporadically observed a diffuse area of Oct4 protein outside the nucleus in cells transfected with siLrp5 compared with the negative siControl (Fig. 5C) with an overall reduced Oct4 signal similar to that seen in untreated cells grown at 21% O2 in Fig. 1D.

Low oxygen transiently upregulates Akt expression

Akt is a cytosolic serine/threonine kinase critical for cell survival under adverse conditions [34]. Akt blocks apoptosis induced by many death stimuli and is required for growth factor-mediated cell survival in different cell types [34]. Akt may therefore play a key role in promoting self-renewal, and thus we assessed whether oxygen tension affects Akt signaling. We incubated cells at 3% and 21% O2 for 30 and 60 min and observed strong Akt activation (phospho-Akt) by low oxygen after a 30-min transient exposure, which then returned to basal levels. Total Akt was unaffected (Fig. 6A). Furthermore, we assessed the effect of PI3K/Akt inhibitor, wortmannin, on MIAMI cells. We tested different concentrations of wortmannin, 20, 50, 100, 200, and 300 nM, and demonstrated that 100 nM is most effective to inhibit phospho-Akt protein (Fig. 6B) without cell damage (not shown).

Blocking the Akt signaling pathway reverses the osteogenic inhibition induced by low oxygen tension

Akt plays central mechanistic roles in the regulation of proliferation, differentiation, and survival, in addition to cell survival, in several systems. This is achieved, in part, through functional cross talk with other signaling pathways (including canonical Wnt). We assessed whether the Akt pathway played a mechanistic role in maintaining the oxygen-dependent balance between MIAMI cell self-renewal and differentiation. To accomplish this, we assessed its role in the osteoblastic differentiation program of MIAMI cells. We previously demonstrated that MIAMI cells fail to progress toward the osteoblastic differentiation program under low oxygen tension conditions [4]. Thus, we sought to determine if blocking Akt activation reversed the effect of low oxygen tension on osteoblastic differentiation. We grew cells at 21% and 3% O2 in osteogenic differentiation medium for 14 days in the presence or absence of wortmannin (100 nM). Cells grown at 21% O2 showed high levels of alkaline phosphatase activity (Fig. 6Ca). We did not detect alkaline phosphatase activity in cells grown at 3% O2 (Fig. 6Cb). However, when we added 100 nM wortmannin to cells growing at low oxygen, blocking Akt activation reversed the low oxygen-dependent osteogenic inhibition and alkaline phosphatase activity was high (Fig. 6Cd) with no detectable effect on cells grown at 21% O2. To further confirm the Akt/PI3K functional role, we analyzed the expression of key osteoblastic gene markers on cells grown for 1, 3, and 7 days under osteogenic conditions at 21% and 3% O2 in the presence or absence of wortmannin. Osteoblastic marker (Runx-2, collagen type I-α1, and osteopontin) expression increased in cells at 21% O2, while in cells at 3% O2, the markers did not increase (Fig. 6D, left panels). Blocking Akt activation using 100-nM wortmannin in cells grown at low oxygen reversed the inhibition of Runx-2, collagen I-α1, and osteopontin genes induced by 3% O2 (Fig. 6D, right panels).
Stemness of a cell can be defined as the capability of self-renewal and developing into more differentiated cell types \[50\]. This distinctive state of the cell is assigned to various cell types, including ESCs, cancer stem cells, and adult stem cells such as the stromal MIAMI cells studied here. Decline in tissue homeostasis and repair during adult life may indeed be associated with a decrease in the number and/or function of stem cells \[51\]. Thus, a better understanding of the molecular drivers regulating the balance between self-renewal and proliferation of primitive MSCs can help design strategies to maintain a stem cell pool capable of tissue repair as we age as well as implement needed improvements in current stem cell culture techniques for proper ex vivo expansion and subsequent clinical use. MSCs are multipotential cells giving rise to various differentiated cells \[52–57\] and display in vivo functional reconstitution of injured tissues \[58–60\]. The development of strategies to prevent decline of endogenous MSCs in the bone marrow during adult life will have a positive impact in delaying age-related tissue function decline.

Oxygen is a key regulator of gene expression programs and a potential modulator of the balance between stemness and differentiation \[7\]. Oxygen homeostasis is maintained by complex mechanisms that have evolved to sense oxygen levels and alter physiological responses accordingly \[61\]. Distribution of hematopoietic progenitor cells is not random throughout the marrow space as many interrelated spatial and temporal cues regulate their proliferation and differentiation \[62\]. Committed precursors localize in close proximity to oxygen-rich blood vessels (vascular niche), while...
less mature progenitors preferentially reside in areas of lower oxygen tension [63]. A model suggesting that human stem cells can self-renew without engaging in differentiation when located in a specific anatomical area with a unique architecture defines what is known as the stem cell niche [64].

We here demonstrated that Nanog, Sox2, Oct4a, and Wnt coreceptor Lrp5 increased at low oxygen (Fig. 1A). Our results suggest that low oxygen maintains MIAMI cell stemness, in part, by increasing Sox2 expression and Oct4a nuclear localization. Additionally, the data suggest that both transcriptional and post-transcriptional events may be mediating these low oxygen-stimulated effects, and transient exposure (≤2 h.) to atmospheric oxygen appears to be at least partially reversible (Fig. 1B–D). Moreover, we found that multiple pathways appear to mediate the low oxygen-stimulated self-renewal maintenance. Notch pathway activation is suggested by increased nuclear NICD levels and target genes (ie, Hes1). Additionally, HIF-1α costimulation raises possible HIF-1α/NICD interactions in targeting gene promoters, which may mediate self-renewal [21].

It is generally accepted that self-renewal mechanisms require increased expression of stemness genes with repression of differentiation programs. MIAMI cells develop an immature neuronal phenotype in response to neurotrophic factors at higher oxygen levels [43], while at low oxygen, neuronal differentiation is inhibited. REST appears to block neuronal differentiation of stem cells as a component of the transcriptional network maintaining stem cell self-renewal [36]. We observed a low oxygen-dependent stimulation of both REST/NRFS transcripts and nuclear localized protein (Fig. 2), suggesting that low oxygen activates pathways stimulating repressors of differentiation.

Furthermore, low oxygen appears to contribute to a pro-survival program in MIAMI cells, evidenced by the activation of the prosurvival Akt pathway and the downregulation of proapoptotic factors, p53, AIF, and Bak, with a concomitant dramatic decrease in SA-βgal activity, increasing self-renewal, while decreasing cell cycle arrest, senescence, and apoptosis. Low oxygen simultaneously activates the canonical Wnt pathway through increased Lrp5 expression, which in turn

FIG. 7. Oxygen tension modulates multiple signaling pathways regulating self-renewal, differentiation, senescence, and apoptosis in MIAMI cells. Diagrammatic representation of the pathways, targets, and biological responses modulated by oxygen tension in MIAMI stem cells, leading to functional changes. Color images available online at www.liebertpub.com/scd
may play a key role in self-renewal. Other studies provide precedence for an Lrp5-directed activation of the canonical Wnt signaling [49], suggesting that the low oxygen-stimulated Lrp5 gene expression may be sufficient for the observed activation of this pathway in MIAMI cells. Wnt signaling is particularly complex, involving a large number of mediators, some of which may be directly involved in different aspects of cellular physiology, including self-renewal and differentiation. Thus, the specific mechanistic context of these specific molecular mediators, together with their interaction with other simultaneously activated signaling pathways, will ultimately result in the final phenotypic or functional outcome.

An intracellular regulatory protein central to mediating the effect of complex extracellular signaling pathways (ie, canonical Wnt), self-renewal, and differentiation, p53, senescence, and apoptosis is the serine/threonine kinase, Akt. The strong transient activation of Akt in response to low oxygen tension (Fig. 6A) provides evidence that it may be playing a central mechanistic role. Having established that low oxygen tension inhibited osteoblastic differentiation under strong cytokine-mediated stimulatory conditions [4], it sets the stage to test such a role. We found that inhibition of Akt activation released the low oxygen-mediated block of the osteoblastic gene expression program and phenotypic determination (Fig. 6C, D). Thus, these results place Akt as a central mechanistic player in regulating the oxygen tension-controlled balance between self-renewal and osteoblastic differentiation in MIAMI cells.

Importantly, a recent study performed in the context of cancer stem cells suggests that nuclear localization of Akt is a key factor promoting cell stemness [65]. Our study corroborates this observation, and future studies will examine the distribution of Akt in MIAMI cells when exposed to low oxygen tension. Future studies will extend our analysis to differentiation pathways other than the osteoblastic lineage and will examine the role of Akt in mediating other processes involved in maintaining a stem cell pool during adult life. It has recently been reported that PFKFB3 plays a critical role in controlling the cell cycle entrance decision [66], and it would therefore be interesting to assess changes of PFKFB3 expression in MIAMI cells when exposed to various levels of oxygen.

In summary, we demonstrate that low oxygen stimulates the activity of the Notch, Akt/P3K, and canonical Wnt pathways. The latter appears to be mechanistically dependent on Lrp5 function. Low oxygen also increased the nuclear localization of the transcriptional repressor REST/NRSF. We have identified key pathways activated in MIAMI cells that are similar to those responsible for ESC self-renewal and stemness. Collectively, Notch and canonical Wnt pathways, in association with REST, may lead to increased self-renewal and concomitant decreased differentiation, cell cycle arrest, apoptosis, and/or senescence in MIAMI cells in which Akt appears to play a convergence role, setting the stage for further analyses of mechanistic roles and potential cross talk of these pathways (Fig. 7). Importantly, we demonstrated that the P3K/Akt pathway plays a central mechanistic role in the oxygen tension-regulated self-renewal versus osteoblastic differentiation of MIAMI cells.

Acknowledgments

The authors thank David Vazquez and Blanca Rodriguez for their assistance throughout this study, as well as the entire GRECC staff.

This study was supported by a Department of Veterans Affairs, Veterans Health Administration, Office of Research and Development (Biomedical Laboratory Research and Development) Merit Review award (BX000952) to P.C.S and by the James and Esther King Biomedical Research Program from the State of Florida to G.D.I.

Author Disclosure Statement

No competing financial interests exist.

References

LOW OXYGEN MODULATES MULTIPLE SIGNALING PATHWAYS

cells and is required to maintain the basal lineage. PLoS One 4:e6594.


Address correspondence to:
Paul C. Schiller, PhD
GRECC and Research Service
Veterans Affairs Medical Center
1201 NW 16th Street
Miami, FL 33125
E-mail: p.schiller@miami.edu

Received for publication November 24, 2015
Accepted after revision April 8, 2016
Prepublished on Liebert Instant Online April 8, 2016