The Primary Cilium as a Biomarker in the Hypoxic Adaptation of Bone Marrow-Derived Mesenchymal Stromal Cells: A Role for the Secreted Frizzled-Related Proteins

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Abstract: A pivotal role in guiding mesenchymal stem cell (MSC) differentiation has recently been attributed to the primary cilium. This solitary, non-motile microtubule-based organelle emerging from the cell surface acts as a sensorial membrane structure reflecting developmental and adaptive processes associated with pathologies including human cystic kidney disease, skeletal malformations, obesity and cancer. Given that the intrinsic hypoxic adaptation of MSC remains poorly understood within ischemic tissues or hypoxic tumours, we questioned whether the hypoxia inducible factor-1α (HIF-1α) might be a downstream effector regulating cilium maintenance. We show that murine bone marrow-derived MSC cultured under hypoxic conditions (1.2% O2) lose their primary cilia in a time-dependent manner. Gene silencing of HIF-1α prevented cilia loss in hypoxic cultures, and generation of MSC expressing a constitutively active HIF-1α (MSC-HIF) was found to decrease primary cilium formation. A Wnt pathway-related gene expression array was also performed on MSC-HIF and indicated that the secreted Frizzled-related proteins (sFRP)-1, -3 and -4 were down-regulated, while sFRP-2 was up-regulated. Overexpression of recombinant sFRP-2 or gene silencing of sFRP-1, -3 and -4 in MSC led to primary cilium disruption. These results indicate a molecular signalling mechanism for the hypoxic disruption of the primary cilium in MSC involving an HIF-1α/sFRP axis. This mechanism contributes to our understanding of the adaptive processes possibly involved in the oncogenic transformation and tumour-supporting potential of MSC. Our current observations also open up the possibility for the primary cilium to serve as a biomarker in MSC adaptation to low oxygen tension within (patho)physiological microenvironments.

Keywords: cilia, hypoxia, mesenchymal stem cells, hypoxia-inducible factor (HIF)
Introduction

Most commonly isolated from the bone marrow, mesenchymal stem cells (MSC) represent a population of pluripotent adult stem cells that can differentiate into many mesenchymal phenotypes, and that can adapt to low oxygen environments such as those encountered within ischemic tissues or hypoxic tumours. This adaptive property has been exploited to study the therapeutic efficacy of genetically-modified MSC. Homing of MSC to tumours was recently reported in a mouse model where injected human MSC could be found preferentially migrating to implanted human melanoma tumours. In fact, recruitment of MSC by experimental vascularizing tumours also resulted in the incorporation of MSC within the tumor architecture which, combined with intrinsic immunomodulatory mechanisms, suggests that they must also respond to tumour-derived growth factor cues. Consequently, their potential contribution to tumour development implies that MSC must metabolically adapt to the low oxygen environment and nutrient deprivation that characterizes hypoxic tumours. Moreover, the sum of this evidence, in line with their increased ability to migrate under an atmosphere of low oxygen, suggests that MSC may be active participants in the development of hypoxic solid tumours.

In order to survive within the stressful hypoxic microenvironment, cells have developed a coordinated set of responses orchestrating their adaptation to hypoxia. In cancer cells, the resultant of such cellular responses to hypoxia is often associated with aggressive disease and resistance to therapy. A critical mediator of the hypoxic response is the transcription factor hypoxia inducible factor 1 (HIF-1) which upregulates expression of proteins that promote angiogenesis, anaerobic metabolism, and many other survival pathways. Regulation of HIF-1α, a component of the HIF-1 heterodimer, occurs at multiple levels including translation, degradation, and transcriptional activation, and serves as a testimony to the central role of HIF-1. More recently, the canonical Wnt pathway was shown to be activated in stem cells under low oxygen culture conditions via HIF-1α. How such HIF/Wnt signalling affects the MSC adaptive mechanisms remains poorly documented.

Stem cell differentiation and proliferation are among the important processes regulated through the Wnt pathway. Its activation in hematopoietic stem cells and in MSC enhances cell proliferation, maintains pluripotency and prevents induction of apoptosis. Moreover, expression of Dickkopf (DKK), a Wnt inhibitor, prevented osteogenic differentiation of cultured human MSC. Among the Wnt molecular players, secreted Wnt antagonists were also found to be important in stem cell homeostasis in an in vivo gastro-intestinal cancer model. Autocrine Wnt signaling also operates in MSC populations to regulate mesenchymal lineage specification. The molecular mechanisms that regulate self-renewal, lineage-specific differentiation and/or adaptation still remain to be linked to specific biomarker expression.

The primary cilium is a sensory membrane structure which transduces surrounding mechanical and chemical signals and which serves as a control center for many protein signalling complexes. Despite its purpose in development where a role in guiding lineage commitment was reported, the primary cilium mainly serves as a cell surface biomarker associated with a growing number of pathologies, including human cystic kidney disease, skeletal malformations, obesity and cancer. Recent evidence links the tumour suppressor pVHL, a protein involved in the nuclear translocation of HIF-1α, to cilium integrity maintenance in cystic kidney disease. We therefore examined the biomarker potential of the primary cilium, as well as the impact of Wnt signalling in MSC adaptation to hypoxic cues.

Experimental Procedures

Materials

Sodium dodecylsulfate (SDS) and bovine serum albumin (BSA) were purchased from Sigma (Oakville, ON). Cell culture media was obtained from Invitrogen (Burlington, ON). All other reagents were from Sigma-Aldrich Canada.

Cell culture and experimental hypoxic conditions

Bone marrow-derived MSC were isolated from the whole femur and tibia bone marrow of C57BL/6 female mice; cells were cultured and characterized as previously described. Analysis by flow cytometry, performed at passage 14, revealed that MSC expressed CD44 yet were negative for CD45, CD31, KDR/flk1 (VEGF-R2), flt-4 (VEGF-R3) and Tie2 (angiopoietin receptor) (data not shown). Hypoxic
culture conditions were attained by incubation of 65%–80% confluent cells in an anaerobic box. The oxygen was maintained at 1%, as described by others, using a compact gas oxygen controller Proox model 110 (Reming Bioinstruments Co., Redfield, NY) with a residual gas mixture composed of 94% N₂ and 5% CO₂.

cDNA construct generation and transduction of the MSC-HIF-1α

The human full-length HIF-1α cDNA construct was generously provided by Dr Gregg L. Semenza (Johns Hopkins University, Baltimore, MD, USA), and was used as a template for generating an HIF-1α mutant which lacked its oxygen-dependent degradation domain (ODD 401–603). The deletion mutant (HIF-1α ΔODD) was constructed by overlap extension using PCR. The deletion was confirmed by DNA sequencing, and the 1.95 kb HIF-1α ΔODD cDNA was subcloned into pcDNA3.1. For generation of retroviral particles, the HIF-1α ΔODD construct was digested out of the pcDNA3.1 vector using BamHI and HpaI restriction enzymes and subcloned into the multiple cloning site of the bicistronic retrovector pIRES-GFP. 293-GP2 viral packaging cells were transfected with either the HIF-1α ΔODD-pIRES-GFP or null-pIRES-GFP plasmids and the viral supernatant was collected at 48 and 72 hours post-transfection. MSC were subjected to 8 rounds of viral transduction. Following viral transduction each GFP(+)MSC (AP2-MSC) and HIF-1α-ΔODD-GFP(+) MSC population were each subjected to high speed cell sorting using a BD FacsAria flow cytometer to obtain polyclonal pooled clones of retrovirally-transfected MSC that were 100% GFP(+) and similar in regards to GFP signal intensity.

Total RNA isolation, cDNA synthesis and real-time quantitative RT-PCR

Total RNA was extracted from cell monolayers using TriZol reagent (Invitrogen). For cDNA synthesis, 2 µg of total RNA were reverse-transcribed into cDNA using a high capacity cDNA reverse transcription kit (Applied Biosystems, Foster City, CA). cDNA was stored at −80 °C prior to PCR. Gene expression was quantified by real-time quantitative PCR using iQ SYBR Green Supermix (Bio-Rad, Hercules, CA). DNA amplification was carried out using an iCycler iQ5 (Bio-Rad, Hercules, CA) and product detection was performed by measuring binding of the fluorescent dye SYBR Green I to double-stranded DNA. The QuantiTect primer sets were provided by Qiagen (Valencia, CA): sFRP-1, -2, -3, -4 (Mm_Sfrp1_1_SG QT00167153, Mm_Sfrp2_1_SG QT00101759, Mm_Frzb_1_SG QT00169232, Mm_Sfrp4_1_SG QT00120491) and β-actin (Hs_Actb_2_SG QT01680476). The relative quantities of target gene mRNA against the internal control β-actin RNA were measured by following a ΔCt method employing an amplification plot (fluorescence signal vs. cycle number). The difference (ΔCt) between the mean values in the triplicate samples of target gene and those of β-actin mRNAs were calculated by iQ5 Optical System Software version 2.0 (Bio-Rad, Hercules, CA) and the relative quantified value (RQV) was expressed as 2^−ΔCt.

Mouse Wnt signaling pathway PCR array

The Mouse Wnt Signaling Pathway RT² Profiler PCR Arrays (PAMM-043, SA Biosciences, Frederick, MD) were used according to the manufacturer’s protocol. Quantitative RT-PCR was performed and relative gene expressions were calculated using the 2^−ΔΔCt method, in which Ct indicates the fractional cycle number where the fluorescent signal reaches detection threshold. The ‘delta–delta’ method uses the normalized ΔCt value of each sample, calculated using a total of five endogenous control genes (B2M, HPRT1, RPL13A, GAPDH, and ACTB). Fold change values are then presented as average fold change = 2 (averageΔΔCt) for genes in MSC-HIF relative to control MSC. Detectable PCR products were obtained and defined as requiring <35 cycles. Using real-time PCR, we reliably analyzed expression of a focused panel of genes related to Wnt-mediated signal transduction with these arrays. The resulting raw data were then analyzed using the PCR Array Data Analysis Template (http://www.sabiosciences.com/pcrar-raydataanalysis.php). This integrated web-based software package automatically performs all ΔΔCt based fold-change calculations from our uploaded raw threshold cycle data.
Immunofluorescent microscopy

Cells were seeded on 1.5-mm thick glass coverslips in 6-well culture plates. After hypoxic treatment, media were removed and cells fixed in 10% formalin phosphate buffer (Fisher Scientific, Ottawa, ON) for 20 min, permeabilized in 0.5% Triton X-100/PBS for 5 min, then blocked 1 h in 1% BSA/PBS. Immunostaining was performed for 1 h with a monoclonal anti-acetylated tubulin antibody (clone 6-11B-1) 1:200 in 1% BSA/PBS (Sigma, St-Louis, MO), followed by 1:200 anti-mouse-RedX (Invitrogen). Nuclei were stained using 5 µg/mL DAPI and then glass coverslips were mounted on slides using Pro-Long Gold Antifade reagent (Invitrogen, ON) before fluorescence was examined by microscopy, using a Nikon Eclipse TE2000-U microscope coupled to a QImaging Retiga 1300 camera.

Transfection method and RNA interference

Cells were transiently transfected with 20 nM siRNA (Qiagen) against HIF-1α (Mm_Hif1a_2 Flexitube siRNA, SI00193018), sFRP-1, -2, -3, -4 (Mm_Sfrp1_1 SI01415855, Mm_Sfrp2_1 SI00182567, Mm_Frzb_1 SI01005991, Mm_Sfrp4_1 SI00209818) or scrambled sequences (AllStar Negative Control siRNA, 1027281) using Lipofectamine 2000 (Invitrogen, ON). Specific gene knockdown was evaluated by qRT-PCR as described above. Small interfering RNA and mismatch siRNA were synthesized by Qiagen and annealed to form duplexes.

Morphological analysis of apoptotic and necrotic cells

To visualize nuclear morphology and chromatin condensation by fluorescence microscopy, cells were stained with 0.06 mg/mL Hoechst (33258, blue fluorescence) for apoptotic cells or with 50 µg/mL propidium iodide (red fluorescence) for necrotic cells.

Statistical data analysis

Data are representative of three or more independent experiments. Statistical significance was assessed using Student’s unpaired t-test. Probability values were considered significant when an asterisk identifies such significance in the figures. * = P < 0.05; ** = P < 0.01; and *** = P < 0.001.

Results

The primary cilium is expressed in MSC and hypoxic culture conditions diminish ciliogenesis

We first wished to monitor whether any primary cilia were expressed in MSC. MSC were seeded into petri dishes and cultured under normoxic conditions. Immunofluorescent staining was performed as described in the Methods section with the anti-acetylated tubulin antibody. We found that approximately 70% of the cells expressed a single plasma membrane protrusion attributable to the primary cilium (Fig. 1A). When MSC were cultured under hypoxic conditions for 24 and 48 hours, ciliogenesis was time-dependently and significantly decreased in hypoxic MSC cultures (Fig. 1B, closed circles) when compared to normoxic MSC cultures (Fig. 1B, open circles). Hoechst-33258 and propidium iodide staining only revealed basal levels of respective apoptotic and necrotic cells (<10%, data not shown) in accordance to previous studies, and suggests that the effects observed were not due to hypoxia-increased cell death. Significant decrease in ciliogenesis was also observed at 48 hours under normoxic culture conditions and may, although speculative, be attributable to low nutrient supply. Collectively, these observations still confirm that MSC express the primary cilium and support some hypoxia-mediated signaling events that would regulate such expression.

Hypoxia inducible factor-1α expression is crucial in the hypoxic downregulation of ciliogenesis

In order to assess the impact of the hypoxia inducible factor-1α (HIF-1α) in the hypoxia-mediated decrease in MSC ciliogenesis, MSC were transiently transfected with a scrambled siRNA sequence (Fig. 2A, upper panels; Mock) or with an siRNA designed to downregulate HIF-1α (Fig. 2A, lower panels; siHIF-1α) as described in the Methods section, then put in culture under normoxic (white bars) or hypoxic (black bars) conditions. Typical knockdown in HIF-1α gene expression was over 80% (not shown) and in accordance with our previous studies. Cilium staining was performed for each experimental condition and cilia were decreased, in agreement with the data in Figure 1, in hypoxic Mock-transfected cells (Fig. 2B, black bars).
This decrease in cilium expression in hypoxic MSC was prevented from diminishing in siHIF-1α-transfected cells (Fig. 2B, black bars). These observations prompt for a crucial involvement of HIF-1α-mediated regulation of MSC ciliogenesis.

HIF-1α stable expression dowregulates ciliogenesis
In order to rule out the sole effect of hypoxia and to firmly establish the direct HIF-1α’s impact on MSC ciliogenesis, cells were engineered to stably express a ΔODD HIF-1α mutant (MSC-HIF) as described in the Methods section. MSC and MSC-HIF were then cultured under normoxic conditions for 24 hours and cilia staining was performed (Fig. 3A). Cilium staining quantification confirmed the significant decrease in primary cilia expression in the MSC-HIF, therefore establishing HIF-1α as a major molecular actor in cilia downregulation.

Gene array analysis reveals involvement of members from the secreted frizzled-related proteins in the hypoxic adaptation of MSC
Given that the Wnt signalling pathway has been reported to regulate ciliogenesis,30,31 we performed a Wnt pathway-related gene expression array. Total RNA was isolated from MSC and from MSC-HIF, and cDNA was synthesized as described in the
Methods section. A schematic representation of the genes which were up-regulated and downregulated (threshold in dotted lines) is depicted (Fig. 4A). A value between $-3.00$ and $3.00$ was considered not significant. While most of the Wnt family gene members were unaffected (Wnt-1 to -16; Fig. 4B), Wnt-9a gene expression was the only one to be upregulated in MSC-HIF. Interestingly, gene expression of four members of the secreted Frizzled-related proteins (sFRP) were found modulated. While sFRP-2 was upregulated, the expression of sFRP-1, -3, and -4 was significantly downregulated (Fig. 4B). Collectively, these observations prompted us to explore whether sFRP transcriptional regulation may effectively occur within MSC.

sFRP family members possess putative hypoxia responsive elements within their promoter region and are downregulated in hypoxic MSC

We next needed to validate the gene expression array data obtained previously. The data from Figure 4B were first used to generate a representative histogram of the sFRP-1, -3, and -4 gene down-regulation and of the sFRP-2 gene up-regulation (Fig. 5A). Specific involvement of HIF-1α in directing sFRP-1 to -4 gene regulation was next explored through sequence promoter analysis. 9,000 bp sequences upstream of the ATG coding sequence of the murine sFRP-1, -2, -3, and -4 gene promoter sequences was analyzed for HIF putative transcription factor binding sites with PROMO 3.0 (http://alggen.lsi.upc.es/) using version 8.3 of the TRANSFAC database. Several core consensus sequences of the hypoxia responsive elements (HRE) $(A_G)CGT(G_C)C$ were found in the murine sFRP sequences analyzed (Fig. 5B). Quantitative qRT-PCR validation of the sFRP gene expression levels was finally performed in order to validate the gene array data and confirmed those gene expression data characterizing sFRP levels in MSC-HIF compared to MSC (Fig. 5C).

Overexpression of sFRP-2 or gene silencing of sFRP-1, -3, -4 family members dowregulates ciliogenesis

We next proceeded to validate the gene expression array significance of sFRP-2 gene upregulation and of sFRP-1, -3, and -4 gene dowregulation in MSC-HIF. We performed transient gene silencing of all sFRP members (Fig. 6A), and transiently transfected sFRP-2 cDNA (Fig. 6B) in MSC. We next went on to immunostain and assess the primary cilium expression in each of these conditions. We found that overexpression of recombinant sFRP-2 led to a significant decrease in transfected MSC (Fig. 6C). In parallel experiments, gene silencing
of sFRP-1, -3, or -4 led to decreased ciliogenesis, while that of sFRP-2 did not change primary cilium expression (Fig. 6D).

**Discussion**

Hypoxia is thought to be a tissue-specific condition that could promote oncogenic processes. In this study we have demonstrated that hypoxia significantly impacts ciliogenesis via a HIF-1α/Wnt signalling axis. While the Wnt family of proteins is known to influence the MSC phenotype through both canonical and non-canonical signalling pathways, our data support the importance of the Wnt pathway in the maintenance of cilium integrity. Accordingly, we also identified the crucial joint action of several sFRP members on MSC ciliogenesis. Our current observations also open

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**Figure 3.** HIF-1α overexpression downregulates ciliogenesis. MSC stably expressing a ΔODD HIF-1α mutant (MSC-HIF) were generated as described in the Methods section. (A) Cells were then cultured under normoxic conditions for 24 hours and cilia staining performed. (B) Quantification of ciliogenesis in MSC (open bars) or MSC-HIF (black bars) was performed. Data are representative of three independent cell cultures. Probability values of less than 0.05 were considered significant, and an asterisk (*) identifies such significance between MSC and MSC-HIF.

**Figure 4.** Gene array analysis reveals sFRP family involvement in hypoxic adaptation of MSC. Total RNA was isolated from MSC and from MSC-HIF and cDNA was synthesized as described in the Methods section. (A) Graphic representation of the array gene expression levels (thresholds in dotted lines). (B) Representative values of the sFRP1-4 and Wnt family gene expression. Values between −3.00 and 3.00 are not considered significant.
up the possibility for the primary cilia to serve as a biomarker for MSC adaptation to low oxygen tension within (patho) physiological microenvironments.

The complex nature of the hypoxic tumor microenvironment may result in pro-oncogenic conditions for MSC. As such, mobilization and migration of MSC to the peripheral blood, and tumors in response to hypoxic cues have recently been evidenced. These events are in part explained by the fact that hypoxic cancer cells secrete various cytokines including IL-6, VEGF, PDGF and FGF, that attract and promote MSC proliferation and differentiation into tumor-supporting cells. Recently, chemosensory response to PDGF-AA in fibroblasts was shown to require the primary cilium. In contrast, in the bone marrow, the hypoxic niche is important for maintaining stemness, cell cycle, cell survival and metabolism of MSC and HSC. The function of MSC in tumors is likely to parallel the role of MSC in wound healing. Accordingly, MSC derived from bone marrow are thought to endogenously support wound healing and hematopoiesis, but many of their native functions under hypoxic conditions remain poorly understood. Under such conditions the low oxygen tension is believed to protect the genomic integrity of stem cell populations by

Figure 5. sFRP family members possess putative Hypoxia Responsive Elements within their promoter regions. (A) Histogram of the sFRP-1, -3, and -4 genes downregulation and the sFRP-2 gene upregulation. (B) Sequences of the putative HRE found within the promoter regions of the four sFRP genes. NCBI sources: sFRP-1, NC_000074.5, chromosome 8; sFRP-2, NC_000069.5, chromosome 3; sFRP-3, NC_000068.6, chromosome 2; sFRP-4, NC_000079.5, chromosome 13. (C) qRT-PCR validation of sFRP gene expression levels.
sFRP regulate ciliogenesis in hypoxic MSC

Limiting the production of reactive oxygen species by mitochondrial respiration.\(^{42}\)

Primary cillum identification as a biomarker may allow efficient monitoring of hypoxic MSC adaptation processes and help increase our understanding of the mechanisms involved in their oncogenic regulation. In combination with other MSC cell surface markers, hypoxic disruption of the primary cilium may also represent a useful marker for clinical purposes. In support of this, the recent discovery of the importance of primary cilia in a variety of cell functions raises the possibility that this structure may indeed have a role in a variety of cancers. Recently, the formation of the primary cilium was disrupted in cells derived from astrocytoma/glioblastoma tumors.\(^{43}\) This observation is among the first evidence that altered primary cilium expression and function may be part of some malignant phenotypes.\(^{44,45}\) MSC were also shown to integrate, engraft and differentiate within hypoxic brain tumors,\(^{46}\) which further highlights the fact that the primary cilium could serve as a diagnostic tool and provide new insights into the mechanism of tumorigenesis.

Whether MSC are pro- or anti-tumorigenic is a subject of controversial reports that is in part explained by the complexity of their homing, engraftment, and differentiation mechanisms within the tumor microenvironment.\(^{47}\) The differentiation of MSC into lineage-specific cells is controlled by external factors in the environment, including cell–cell and cell–ECM adhesion and cytokine, chemokine, and growth factor availability.\(^{48,49}\) Several signalling pathways have recently been identified in MSC proliferation and differentiation control including canonical and non-canonical Wnt, RhoA/ROK, and Erk.\(^{50–52}\) Our current study highlights the increased levels of sFRP-2 in hypoxic MSC. Accordingly, while sFRP-1 and sFRP-2 are

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**Figure 6.** Overexpression of sFRP-2 or gene silencing of sFRP-1, -3, -4 family members downregulates ciliogenesis. (A) Gene silencing was performed using specific siRNA for each of the sFRP-1, -2, -3, or -4 members as described in the Methods section. Total RNA was extracted and qRT-PCR performed in order to evaluate efficiency and validate gene expression. (B) Cells were transfected with Mock or sFRP-2 cDNA plasmid, and sFRP-2 gene expression validated by qRT-PCR. (C) Primary cilium staining was performed in Mock- (white bars) and in sFRP-2- (black bars) transfected cells. (D) Gene silencing was performed with a scrambled sequence (siScr; Mock), or with the respective sFRP-1, -2, -3, -4 siRNA sequences. Primary cilium staining was performed in Mock- (white bars) and in si-sFRP-transfected cells (black bars).
produced by the majority of long-term and ex vivo malignant glioma cell lines, only sFRP-2 was shown to strongly promote the growth of intracranial glioma xenografts in nude mice. We also report that sFRP-1, -3, and -4 downregulation correlates with the hypoxic MSC phenotype that leads to decreased ciliogenesis. Intracellular Wnt effectors, such as Fuzzy and Inturned, were found to disrupt the primary cilium structure in Drosophila melanogaster embryos. Interestingly, sFRP-1 is frequently silenced in many types of cancer leading to aberrant activation of Wnt signaling. Given they all possess HRE in their promoter region, it remains to be clarified how different sFRPs may have opposing effects on the same process and how they affect cell cycle progression to release the centrosome from the ciliary basal body, which makes it available for duplication and organization of the mitotic spindle.

Our current observations may also impact on the potential of MSC to promote tissue repair in a diverse array of diseases, including ischemic heart disease, diabetes, and Parkinson’s disease. When engrafted at sites of tissue injury, MSC differentiate into connective tissue elements, support vasculogenesis, and secrete cytokines and growth factors that facilitate healing. Accordingly, MSC were confirmed as promising tools for cell therapy, as proven effective in US FDA-approved clinical trials for myocardial infarction, stroke, meniscus injury, limb ischemia, graft-versus-host disease and autoimmune disorders. The hypoxia/ischemia alteration of MSC within these pathological states still remains poorly understood. Clinical trials for MSC injection into the CNS to treat traumatic brain injury and stroke are also ongoing. One may envision that ischemic/hypoxic regulation of primary cilium expression may help monitor intravenous infusion of MSC to in vivo cerebral ischemia tissues. Neuroprotective, trophic support and therapeutic efficacy through MSC mobilization was also demonstrated in several ischemia/reperfusion models. In conclusion, our study enabled us to identify secreted Wnt-related proteins that contribute to hypoxic regulation of ciliogenesis in MSC. A schematic summary of the hypoxia-mediated regulation of ciliogenesis in MSC under (patho)physiological conditions is provided (Fig. 7). Altogether, we also provide new insight into stem cells biology and stem cell adaptation to hypoxia.
torship, and compliance with ethical requirements in respect to treatment of human and animal test subjects. If this article contains identifiable human subject(s), author(s) were required to supply signed patient consent prior to publication. Author(s) have confirmed that the published article is unique and not under consideration nor published by any other publication and that they have consent to reproduce any copyrighted material. The peer reviewers declared no conflicts of interest.

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