Runx3 inhibits endothelial progenitor cell differentiation and function via suppression of HIF-1α activity

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Abstract. Endothelial progenitor cells (EPCs) are bone marrow (BM)-derived progenitor cells that can differentiate into mature endothelial cells, contributing to vasculogenesis in the blood vessel formation process. Runt-related transcription factor 3 (RUNX3) belongs to the Runt domain family and is required for the differentiation of specific immune cells and neurons. The tumor suppressive role of RUNX3, via the induction of apoptosis and cell cycle arrest in a variety of cancers, and its deletion or frequent silencing by epigenetic mechanisms have been studied extensively; however, its role in the differentiation of EPCs is yet to be investigated. Therefore, in the present study, adult BM-derived hematopoietic stem cells (HSCs) were isolated from Runx3 heterozygous (Rx3+/−) or wild-type (WT) mice. The differentiation of EPCs from the BM-derived HSCs of Rx3+/− mice was found to be significantly increased compared with those of the WT mice, as determined by the number of small or large colony-forming units. The migration and tube formation abilities of Rx3+/− EPCs were also observed to be significantly increased compared with those of WT EPCs. Furthermore, the number of circulating EPCs, defined as CD34+/CD133− cells, was also significantly increased in Rx3+/− mice. Hypoxia-inducible factor (HIF)-1α was upregulated in Rx3+/− EPCs compared with WT EPCs, even under normoxic conditions. Furthermore, in a hindlimb ischemic mouse models, the recovery of blood flow was observed to be highly stimulated in Rx3+/− mice compared with WT mice. Also, in a Lewis lung carcinoma cell allograft model, the tumor size in Rx3+/− mice was significantly larger than that in WT mice, and the EPC cell population (CD34+/VEGFR2+ cells) recruited to the tumor was greater in the Rx3+/− mice compared with the WT mice. In conclusion, the present study revealed that Runx3 inhibits vasculogenesis via the inhibition of EPC differentiation and functions via the suppression of HIF-1α activity.

Introduction

The Runt-related transcription factor (RUNX) family comprises context-dependent transcriptional regulators including RUNX1, RUNX2 and RUNX3 in humans. The RUNX family has a highly conserved Runt-homology domain that mediates site-specific DNA binding and interactions with various proteins (1,2). RUNX1 is essential for hematopoiesis; RUNX2 is involved in osteogenesis during bone formation, and RUNX3 is expressed in epithelial, neuronal and hematopoietic stem cells, and is involved in the transforming growth factor (TGF)-β signaling pathway (3), in which it functions to control the lineage differentiation of hematopoietic and neuronal progenitor cells (4). RUNX3 has been documented to function as a tumor suppressor in numerous types of tumors (5). Under hypoxic conditions, RUNX3 is silenced by histone deacetylation and methylation via histone deacetylase 1 and G9a histone methyltransferase, respectively (6).

Blood vessel formation occurs via two distinct mechanisms, namely vasculogenesis and angiogenesis; angiogenesis comprises endothelial sprouting and intussusception from pre-existing vessels, while vasculogenesis initiates from vascular stem cells or hemangioblasts in the developing embryo or adult bone marrow (BM) (7). Endothelial progenitor cells (EPCs), mainly derived from BM hemangioblasts and hematopoietic stem cells (HSCs), possess an ability to self-renew, mobilize into the circulatory system, migrate to sites of neovascularization and differentiate into mature endothelial cells contributing to neonatal vasculogenesis as well as tumor vasculature (8-10). Generally, EPCs can be obtained by isolating a heterogeneous subpopulation of BM-derived mononuclear cells (MNCs) that respond to stresses, such as tissue ischemia and vascular-damaging signals (11,12). In a previous study,
an EPC colony-forming unit (CFU) assay was used to discriminate the hierarchical lineage between primitive (small colony) and definitive (large colony) CFUs in vitro (13). These EPC CFUs take up 3,3'-dioctadecyldiacarbocyanin-labeled acetylated low-density lipoprotein (DiI-Ac-LDL), bind with isolecitin B4, and express vascular endothelial growth factor (VEGF) receptor 2 (VEGFR2) and endothelial nitric oxide synthase (eNOS) independent of BM-MNC isolation markers (13,14). Using a short-term protocol (4-7 days culture), it is possible to obtain cells with myeloid/hematopoietic and typical endothelial cell markers that function as therapeutics in ischemic diseases (15,16).

Cord blood-derived EPCs enhance tissue neovascularization via physical incorporation into the ischemic area and the perivascular paracrine effect (17). Hypoxia or ischemic stress is a stimulator of EPC function through the activation of hypoxia-induced signaling molecules, such as signal transducer and activator of transcription 3 in the preconditioning to hypoxia (18). A number of studies have demonstrated that EPCs promote the angiogenic switch in solid tumors, thereby increasing tumor growth and metastasis (10). Likewise, the genetic suppression of EPC mobilization from BM inhibits tumor development and colonization to remote organs (19). These findings indicate that EPCs may be used as cell therapeutics in injured or regenerative tissues, and also as therapeutic targets for the inhibition of tumor growth or metastasis.

In a previous study, the present research team identified a new function of RUNX3 that facilitates the degradation of hypoxia-inducible factor (HIF)-α by promoting the interaction of proline hydroxylases (PHDs) with HIF-α, thus inhibiting hypoxia-induced VEGF secretion and angiogenesis in vitro and in vivo (20). Hypoxia is a critical microenvironment for the differentiation and maintenance of hematopoietic stem cells (18,21) and leukemia development (22). However, the role of Runx3 in the differentiation and function in EPCs has not yet been investigated.

In the present study, using a Runx3 heterozygous deletion mutant mouse model, it was demonstrated that the mouse homolog Runx3 inhibits vasculogenesis via the suppression of EPC differentiation and function. The mobilization of EPCs into peripheral blood (23), their differentiation into neovessels and homing to an ischemic limb region was significantly increased in Runx3 heterozygous mice. Signaling pathways involved in the maintenance or function of EPCs, as well as HIF-1α were also significantly upregulated by Runx3 heterozygous deletion. These results suggest that Runx3 suppresses EPC function and differentiation from BM-derived MNCs by the suppression of HIF-1α.

Materials and methods

Mice. The animal experiments were approved according to the guidelines for care and use of laboratory animals by the Kyungpook National University Institutional Animal Care and Use Committee (Daegu, Korea). FVB/N male mice (n=10, 4-6 weeks old, 10-12 g; Japan SLC, Inc., Hamamatsu, Japan) were maintained under specific pathogen-free conditions with a constant humidity and temperature at 26˚C, filtered air and a 12/12-h light/dark cycle. Mice were treated with CO₂ by inhalation in a chamber for anesthesia, immediately prior to sacrifice. Runx3 heterozygous mice (Rx3+/-) were generated as previously reported (24).

Isolation of mouse MNCs from BM and PB, and hypoxic conditions. MNCs were isolated from the BM or PB of wild-type (WT) and Rx3+/- mice using a Histopaque-I1083 density gradient centrifugation method (Sigma-Aldrich; Merck KGaA, Darmstadt, Germany) according to the manufacturer's instructions. Briefly, BM or PB cells were centrifuged at 400 x g for 30 min at room temperature following the addition of Histopaque-1083 and MNC-containing layers were isolated. The freshly isolated MNCs were washed with 5 mM EDTA in PBS and suspended in 4 ml of 0.8% ammonium chloride (StemCell Technologies, Inc., Vancouver, BC, Canada) for 5 min at 4˚C. The MNCs were then resuspended in Endothelial Cell Growth Medium (EGM)-2 (Lonza Group Ltd., Basel, Switzerland) supplemented with 5% fetal bovine serum (FBS; HyClone; GE Healthcare Life Sciences, Logan, UT, USA), basic fibroblast growth factor (bFGF; R&D Systems, Wiesbaden, Germany), VEGF (PeproTech, Inc., Rocky Hill, CT, USA), insulin-like growth factor-1 (R&D Systems), epidermal growth factor (EGF; PeproTech, Inc.), ascorbic acid and heparin (Sigma-Aldrich; Merck KGaA), and then seeded onto 60-mm dishes coated with 2% gelatin (2x10⁵ cells/dish). After culture for 3 days at 37˚C, the non-adherent cells were removed by washing with PBS, and the attached cells were cultured for a further 4 days. These cells are designated 'early EPCs' (15,16). For exposure of the early EPCs to hypoxic conditions, hypoxic chambers (Thermo Fisher Scientific, Inc., Waltham, MA, USA and Astec Co., Ltd., Kasuya, Japan) were used for maintaining low oxygen tension (1% O₂ and 5% CO₂, balanced with N₂).

EPC colony-forming assay. The isolation of c-kit+Sca-1+Lin⁻ (KSL) cells from mouse BM-MNCs was performed as previously described (14,25,26). The numbers of small and large colonies, i.e., primitive endothelial progenitor cell colonies with clusters of homogeneous small immature cells and definitive endothelial progenitor cell colonies with clusters of large spindle-shaped differentiated cells were counted following the culture of KSL cells (500 cells/35-mm dish) for 10 days in a methyl cellulose-containing medium (M3236; StemCell Technologies, Inc.) with 20 ng/ml stem cell factor (Sigma-Aldrich; Merck KGaA), 50 ng/ml VEGF, 20 ng/ml interleukin-3 (R&D Systems, Inc., Minneapolis, MN, USA), 50 ng/ml bFGF, 50 ng/ml EGF, 2 U/ml heparin, 30% FBS and antibiotics (100 U/ml penicillin and 100 mg/ml streptomycin; Thermo Fisher Scientific, Inc.). The CFUs were counted by blinded investigators 12 days after seeding by visual inspection with an inverted microscope at x40 magnification. The expression of additional endothelial marker genes, such as cluster of differentiation (CD)31, VEGFR2 [also known as fetal liver kinase 1 (Flk1)], von Willebrand factor, vascular endothelial cadherin and eNOS in small and large CFUs was conducted as previously described (14).

Cell counting kit-8 (CCK-8) assay. EPCs (3x10⁵) were cultured on 96-well plates. Following 24 or 36 h incubation in a hypoxic chamber, CCK-8 solution (Dojindo Molecular Technologies, Inc., Rockville, MD, USA) was added to each
well and incubated for another 2 h at 37°C. Absorbance of the plate was measured with a microplate reader (Infinite M200 Pro; Tecan Group, Ltd., Mannedorf, Switzerland) at 540 nm. Three replicate wells were used for each analysis.

Migration assay of early EPCs. A migration assay of early EPCs was performed as previously reported (27). Briefly, using a modified Boyden chamber, 1 day-starved early EPCs (1x10⁶) were seeded onto a 24-well Transwell membrane and treated with VEGF (10 ng/ml) or stromal cell-derived factor (SDF)-α (100 ng/ml) in the bottom chamber for 6 h. Migrated EPCs were stained with hematoxylin and eosin, and then counted using a CKX41SF microscope (Olympus Corporation, Tokyo, Japan).

Matrigel tube formation assay. Early EPCs (1:1; 1x10⁶ cells/100 µl 5% FBS/EGM-2) were incubated with 0.4 µg/ml Dil-Ac-LDL (Biomedical Technologies Inc., Stoughton, MA, USA) for 4 h at 37°C and then cocultured with human umbilical vein endothelial cells (HUVECs; BD Biosciences, San Jose, CA, USA) on a 96-well culture plate coated with Matrigel® (BD Biosciences) for 30 min at 37°C. Plates were examined for tube formation following incubation for 8 h. The tube length and the number of incorporated Dil-expressing EPCs per HPF were counted (n=3/group) using a microscope (Olympus Corporation) at x200 magnification.

Enzyme-linked immunosorbent assay (ELISA). The amount of VEGF protein secreted by the medium was determined using a VEGF ELISA kit (cat. no. MWV00; R&D Systems). The cells were cultured in 6-well plates until they reached 80-90% confluence and the amount of VEGF was quantified using a microplate reader (Infinite M200 Pro; Tecan Group, Ltd.)

Reverse transcription-quantitative polymerase chain reaction (RT-qPCR) and semi-quantitative PCR. Total RNA was isolated from mouse early EPCs using TRizol (Invitrogen; Thermo Fisher Scientific, Inc.) and RT-qPCR was performed as previously reported (28). The semi-quantitative PCR was performed using AccuPower PCR premix (Bioneer Corporation, Daejeon, Korea). The PCR conditions were as follows: 28 cycles of denaturation (94°C/1 min), annealing (60°C/1 min), extension (72°C/1 min) and final extension (72°C/10 min). The primer sequences for the genes tested are shown in Table I. PCR products were separated on 1.5% agarose gels in 1X TAE buffer, stained with ethidium bromide and visualized under UV light.

Western blot analysis. EPCs were lysed with ProPrep protein extraction solution (Intron Biotechnology, Inc., Seoul, Korea), and the protein concentrations of the lysates were measured using a Pierce BCA protein assay kit (Thermo Fisher Scientific, Inc.). Proteins (50 µg/lane) were separated by 10% SDS-PAGE and transferred to a nitrocellulose membrane (Whatman; GE Healthcare Life Sciences). The membrane was blocked with 5% non-fat skimmed milk in Tris-buffered saline containing 0.1% Tween-20, for 30 min at room temperature. The membrane was incubated with primary antibodies at 4°C overnight, and then with secondary antibodies conjugated with horseradish peroxidase for 1 h at room temperature. Finally, the membrane was visualized using the West Pico Chemiluminescent Substrate (Pierce; Thermo Fisher Scientific, Inc.). The quantification was performed using ImageJ software (version 1.52a; National Institutes of Health, Bethesda, MD, USA). Primary antibodies used for western blot analysis were as follows: VEGF (1:500; cat. no. sc-152; Santa Cruz Biotechnology, Inc., Dallas, TX, USA), VEGFR2 (1:1,000; cat. no. 2479), phospho (P)-VEGFR2 (1:500; cat. no. 2478), extracellular signal-regulated kinase (ERK; 1:1,000; cat. no. 9101), AKT (1:1,000; cat. no. 4691), P-AKT (1:1,000; cat. no. 9271), Jun N-terminal kinase (JNK; 1:1,000; cat. no. 9252), P-JNK (1:1,000; cat. no. 9251), eNOS (1:1,000; cat. no. 9586), P-eNOS (1:1,000; cat. no. 9571) (all from Cell Signaling Technology, Inc., Danvers, MA, USA), HIF-1α (1:500; cat. no. 610958; BD Biosciences), RUNX3 (1:500; cat. no. 40278; Abcam, Cambridge, UK) and α-tubulin (1:1,000; cat. no. 3873; Cell Signaling Technology, Inc.). HRP-linked secondary antibodies were purchased from Cell Signaling Technology Inc. (1:5000; cat. nos. 7076 and 7074).

Flow cytometric analysis. To count the number of circulating EPCs, MNCs isolated from PB were stained with fluorescein isothiocyanate (FITC)-conjugated anti-CD34 monoclonal antibody (1:100; cat. no. GTX2021; GenTex, Inc., Irvine, CA, USA) and phycoerythrin-conjugated anti-VEGFR2 monoclonal antibody (1:100; cat. no. 555308; BD Biosciences) for 1 h at room temperature. The cells were washed with PBS three times. A FACS flow cytometer (FACSAria flow cytometer; BD Biosciences) was used to detect the double-positive fluorescent cells using FACSeqExpress Software (version 4; De Novo Software, Glendale, CA, USA).

Mouse hindlimb ischemia model. A unilateral mouse hindlimb ischemia model was established as previously described (29). Briefly, WT and Rx3⁻/⁻ mice were anesthetized.

<table>
<thead>
<tr>
<th>Gene</th>
<th>Direction</th>
<th>Primer sequence</th>
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<tbody>
<tr>
<td>SDF-1α</td>
<td>Forward</td>
<td>5'-CTGTAGCGTACGGAGCAGAT-3'</td>
</tr>
<tr>
<td></td>
<td>Reverse</td>
<td>5'-CCATTCTCGAGCAGACCCAAAA-3'</td>
</tr>
<tr>
<td>CXCR4</td>
<td>Forward</td>
<td>5'-AGCCTCTGCTCATGGAGTTG-3'</td>
</tr>
<tr>
<td></td>
<td>Reverse</td>
<td>5'-GCCAGGTCCAAGGTCTGC-3'</td>
</tr>
<tr>
<td>VEGF</td>
<td>Forward</td>
<td>5'-GGGCCAGCTGATGGTTGAC-3'</td>
</tr>
<tr>
<td></td>
<td>Reverse</td>
<td>5'-TCTCCAGATCGTGACAGT-3'</td>
</tr>
<tr>
<td>Flik1</td>
<td>Forward</td>
<td>5'-TCCCCCCCTGGAAATCTC-3'</td>
</tr>
<tr>
<td></td>
<td>Reverse</td>
<td>5'-ACAGACC GGCAAACAA-3'</td>
</tr>
<tr>
<td>eNOS</td>
<td>Forward</td>
<td>5'-CCGCACTCACAGGAGAAGA-3'</td>
</tr>
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<td></td>
<td>Reverse</td>
<td>5'-CATGACGGCAGGCAGAT-3'</td>
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<tr>
<td>β-actin</td>
<td>Forward</td>
<td>5'-AACTCCTCACCTCCTCCAAAG-3'</td>
</tr>
<tr>
<td></td>
<td>Reverse</td>
<td>5'-AAAGAATCTCTGTCACCTCCC-3'</td>
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SDF, stromal cell-derived factor; CXCR4, C-X-C chemokine receptor type 4; VEGF, vascular endothelial growth factor; Flik1, fetal liver kinase (VEGF receptor 2); eNOS, endothelial nitric oxide synthase.
with 2% Avertin (250 mg/kg) by intraperitoneal injection (n=3 mice/group). The proximal and distal portions of the femoral artery and the distal portion of the saphenous artery were ligated. The arteries and all side branches were dissected free and excised. A laser Doppler perfusion imaging system (PeriScan PIM II; Perimed AB, Järfälla, Sweden) was used to measure hindlimb blood perfusion immediately following surgery and at the end of the study on day 14. To avoid the influence of ambient light and temperature, the results were expressed as the ratio between perfusion in the left (ischemic) versus right (non-ischemic) limbs.

**In vivo tumor allograft experiment.** Lewis lung carcinoma (LLC; American Type Culture Collection, Manassas, VA, USA) cells (2x10^6) were injected subcutaneously into the right and left flanks of 6-week-old WT or Rx3+/− mice on day 0 (n=4 mice/group). Tumor growth was measured with a caliper every other day from day 9, and the tumor volume was calculated using the following formula: Volume (cm^3) = height x length x depth (cm). Tumor tissues were removed at day 14 and fixed with 4% paraformaldehyde.

**Immunohistochemistry.** LLC-derived tumor tissues from the allograft experiment and hindlimb muscle tissues from the mice in the ischemia experiment were removed at day 14 and fixed with 4% paraformaldehyde. Frozen blocks were made at -26˚C and cut into 10-µm sections, blocked with 5% goat serum (Abcam) in PBS containing 0.03% Triton X-100 for 1 h, and then incubated for 3 h at room temperature with anti-CD34 (cat. no. 553731), anti-CD31 (cat. no. 553370) (both from BD Pharmingen; BD Biosciences) and anti-VEGFR2 (cat. no. 2479; Cell Signaling Technology, Inc.) primary antibodies, followed by Alexa Fluor 488 anti-rabbit IgG and Alexa Fluor 594 anti-rat IgG (Thermo Fisher Scientific, Inc.) secondary antibodies for 1 h at room temperature with protection from light. After washing with PBS, the samples were mounted with mounting medium containing DAPI (Vector Laboratories, Inc., Burlingame, CA, USA), and observed under a confocal microscope (Leica Microsystems, Inc., Buffalo Grove, IL, USA). The quantitation was acquired using ImageJ software.

**Statistical analysis.** SPSS software was used for all statistical analyses. Differences among experimental groups were compared using one-way analysis of variance with the Newman-Keuls post hoc test. P<0.05 was considered to indicate a significant difference. Results are presented as the mean± standard deviation.

**Results**

**Runx3 knockout enhances the differentiation of EPCs.** Since Runx3 homozygote knockout (Rx3−/−) mice die soon after birth (24), Rx3−/− mice were used for the isolation of MNCs from adult BM. To identify the role of Runx3 in EPC differentiation, the BM-derived MNCs isolated from Rx3−/− mice were differentiated into EPCs. Western blotting results confirmed that Runx3 was expressed in the early EPCs from WT mice but repressed in those from Rx3−/− mice (Fig. 1A). Using the EPC colony-forming assay, the self-renewal and differentiation abilities of EPCs were investigated via the detection of two differently differentiated EPCs in terms of EPC hierarchy. Colonies formed from relatively small and rounded cells are known as small EPC CFUs, which exhibit the characteristics of primitive EPCs, including high proliferative activity and immature properties, and colonies formed from spindle-shaped cells are known as large EPC CFUs, which exhibit late or mature EPC characteristics, including as improved tube formation ability and neovascularization (13). The numbers of small and large EPC CFUs were observed to be significantly increased in the Rx3−/− mice compared with the WT mice (Fig. 1B and C). These results suggest that the deletion of Runx3 is associated with the increased self-renewal and differentiation capacity of EPCs.

**In vitro vasculoegenic activity is increased in Rx3−/− EPCs.** Adherent MNCs isolated from Runx3 knockout mice cultured for 7 days, with the removal of non-adherent BM-MNCs after 3 days, changed their morphology from round to spindle-like but did not form colonies in the presence of endothelial cell growth supplements. These cells were characterized as early EPCs (15,16) and exhibited Dil-Ac-LDL uptake, isolectin B4 binding, expression of CD34 (a HSC marker) and VEGFR2 (Flk1, an endothelial cell marker) as in a previous study (21).

The growth rate of early EPCs derived from Rx3−/− MNCs was significantly increased compared with that of early EPCs derived from WT MNCs under normoxia and hypoxia (Fig. 2A). As VEGF and SDF-1α serve important roles in the regulation of various cellular functions of EPCs, such as migration and homing to injured tissues (16,30), a Boyden chamber migration assay using early EPCs in the presence of VEGF or SDF-1α was performed in the present study. VEGF or SDF-1α treatment induced the migration of EPCs, and the number of migrated cells was significantly increased in Rx3−/− EPCs compared with WT EPCs (Fig. 2B). To evaluate the ability of EPCs to differentiate into ECs and form tubes, a tube formation assay was performed on a Matrigel matrix using a 1:1 mixture of Dil-Ac-LDL-labeled EPCs and HUVECs. The tube formation ability was monitored using tube length measurements. As shown in Fig. 2C, the capillary networks developed by Rx3−/− EPCs and HUVECs were more pronounced than those formed by WT EPCs and HUVECs. The total tube length and number of EPCs incorporated into the tubes were significantly increased in the Rx3−/− group compared with the WT group. These results suggest that the absence of Runx3 increases the neovessel-forming differentiation ability of EPCs.

**Expression of growth factors and their receptors involved in EPC functions is upregulated in Rx3−/− EPCs.** To investigate whether the increased vessel formation ability of EPCs with Runx3 deletion is due to increased vasculogenic gene expression patterns in EPCs, the mRNA expression levels of growth factors and their receptors involved in vasculogenesis were analyzed. RT-qPCR revealed that the mRNA expression levels of VEGF, SDF-1α, C-X-C chemokine receptor type 4 (CXCR4) and eNOS were significantly upregulated in Rx3−/− EPCs compared with WT EPCs (Fig. 3A), while a significant increase in VEGFR2 (Flk1) mRNA expression was detected. To confirm these results, the phosphoinositide-3 kinase (PI3K) and mitogen-activated protein kinase (MAPK) signaling pathways that are activated by VEGFR2 were
VEGFR2 mRNA expression was not altered with Runx3 deletion, but the phosphorylation of VEGFR2 was observed to be significantly increased in Rx3<sup>+/−</sup> EPCs compared with WT EPCs (Fig. 3B). As PI3K and MAPKs, including JNK and ERK1/2, are downstream molecules for CXCR4/SDF-1α (31,32) as well as VEGF/VEGFR2 signaling pathways (33), the activation of Akt, JNK and ERK1/2 in Rx3<sup>+/−</sup> EPCs was examined. The phosphorylation levels of Akt, JNK and ERK1/2 were significantly increased in Rx3<sup>+/−</sup> EPCs compared with WT EPCs (Fig. 3C).
JNK and ERK1/2 were significantly increased in Rx3+/− EPCs compared with WT EPCs (Fig. 3C). The activation of eNOS is positively controlled by Akt through the phosphorylation of Ser1178 (34) and results in the generation of NO that serves a crucial role in vessel vasodilation and vasculogenesis (35). Thus, the amount of the activated form of eNOS was evaluated. The activated form of eNOS was observed to be significantly increased in Rx3+/− EPCs compared with WT EPCs (Fig. 3C). Furthermore, the secretion of VEGF from Rx3+/− EPCs was confirmed to be significantly increased compared with that from WT EPCs (Fig. 3D). These results suggest that Runx3 knockout enhances vasculogenic signaling via the activation of VEGF/VEGFR2 and CXCR-4/SDF-1α pathways.

HIF-1α is stabilized under normoxic conditions in Rx3+/− EPCs. As VEGF, VEGFR (36), SDF-1α and CXCR4 (37) are downstream targets of HIF-1α, the HIF-1α protein levels in Rx3+/− EPCs were measured under normoxic and hypoxic conditions. Compared with WT EPCs, HIF-1α expression was increased 3-fold in Rx3+/− EPCs under normoxic conditions (Fig. 4, lanes 1 and 2). Under hypoxic conditions, HIF-1α was highly stabilized by 5-fold in WT EPCs (Fig. 4, lane 1 and lane 3), whereas in Rx3+/− EPCs, HIF-1α was more increased by 2.7-fold (Fig. 4, lanes 2 and 4), suggesting that Runx3 destabilizes HIF-1α protein.

Mobilization into PB circulation is reduced in Rx3+/− EPCs. The mobilization of EPCs from the BM into the peripheral circulation, resulting in circulating EPCs (CEPCs), is a critical step during the process of neovascularization at ischemic sites (35). Therefore, the percentage and number of CEPCs in the PB were investigated via the flow cytometric detection of the EPC markers CD34 and VEGFR2. The percentage of EPCs in PB-MNCs was found to be significantly increased in Rx3+/− mice (1.1±0.12%) compared with WT mice (0.5±0.08%; Fig. 5A and B). Furthermore, the number of EPCs in the PB of Rx3+/− mice (1,603.2±38.12 EPCs/µl) was 3-fold higher than that of WT mice (497.1±32.74 EPCs/µl; Fig. 5C). The results indicate that, consistent with the expression patterns of vasculogenic signaling molecules, the mobilization of EPCs into the PB is also stimulated by the knockdown of Runx3.

Repair of blood flow in hindlimb-ischemia and tumor tissue infiltration of EPCs of Rx3+/− mice. As the aforementioned results reveal that the haploinsufficient deletion of Runx3...
Figure 4. Heterozygous deletion of Runx3 increases HIF-1α expression in cultured EPCs. Early EPCs from WT or heterozygous Runx3-deleted mice were cultured in normoxic or hypoxic conditions for 8 h. Whole cell extracts were subjected to western blot analysis of HIF-1α and RUNX3. *P<0.05, Runx3, runt-related transcription factor 3; Rx3+/−, Runx3 heterozygous; WT, wild-type; HIF, hypoxia-inducible factor; EPC, endothelial progenitor cell.

Figure 5. Heterozygous deletion of Runx3 increases the mobilization of EPCs from the bone marrow to the peripheral circulation and ischemic hindlimb. (A) Mononuclear cells were isolated using the density gradient method and stained with a FITC-conjugated anti-CD34 antibody and PE-conjugated anti-VEGFR2 antibody. Flow cytometric analysis was performed to detect CD34+/VEGFR2+ circulating EPCs. (B) Percentage and (C) total numbers of circulating EPCs were counted in 10,000 peripheral blood cells. *P<0.01 vs. WT. (D) Representative images of the ischemic (L) and non-ischemic (R) hindlimbs immediately and 6, 10 and 14 days after surgery. In the color-coded images, red indicates normal perfusion, and blue indicates a marked reduction in blood flow in the ischemic hindlimb. (E) Blood flow rate was recovered in Rx3+/− mice compared with WT mice (n=3/group). *P<0.05 vs. WT. (F) CD31+ regions (green fluorescence) in ischemic muscle detected using immunohistochemistry (left panels) were quantified, and representative images of hematoxylin and eosin stained tissue are shown (right panels). Arrow heads indicate the vessels. Scale bar, 50 µm. (G) LLC cells (2x10^5) were inoculated into bilateral flanks of WT or Runx3 heterozygous knockout mice (day 0) and the tumor size were measured at various time points. *P<0.05 vs. WT (n=4). (H) Sectioned (10 µm) tumor tissue from LLC-injected WT or heterozygous Runx3-deleted mice was immunostained with anti-VEGFR2 and anti-CD34 antibodies, which exhibit green and red fluorescence, respectively. Infiltrated EPCs (CD34+/VEGFR2+ cells) were counted. *P<0.05 vs. WT. Scale bar, 200 µm. Runx3, runt-related transcription factor 3; Rx3+/−, Runx3 heterozygous; WT, wild-type; EPC, endothelial progenitor cell; FITC, fluorescein isothiocyanate; PE, phycoerythrin; CD, cluster of differentiation; VEGFR2, vascular endothelial growth factor receptor 2; L, left; R, right; LLC, Lewis lung carcinoma.
increased EPC function (Figs. 1-4), and a previous study demonstrated that RUNX3 inhibits the stability of HIF-1α (20), a main transcription factor of angiogenesis and vasculogenesis in hypoxia, angiogenesis in Rx3+/− mice was investigated in vivo using a hindlimb-ischemia model. Compared with WT mice, blood flow was significantly increased in Rx3+/− mice from day 6 after ligation of the hindlimb artery and vein. At 10 and 14 days post-ligation, the blood flow rate in the hindlimb and foot was completely recovered in Rx3+/− mice (Fig. 5D and E). When the hindlimb muscle was analyzed by immunohistochemistry using an anti-CD31 antibody, the microvessel density was observed to be markedly increased in the Rx3+/− mice compared with the WT mice (Fig. 5F). Furthermore, in an LLC allograft model, the tumor volume in Rx3+/− mice was significantly larger than that in WT mice (Fig. 5G) and the EPC population (CD34+/VEGFR2+ cells) in the tumor mass was ~2.5-fold greater in the Rx3+/− mice compared with the WT mice (Fig. 5H). These results suggest that the mobilization of EPCs into the tumor vessels is increased in Rx3+/− mice.

Discussion

The present study suggests that Runx3 may be an inhibitor of EPC function and the differentiation of BM-MNCs into EPCs. EPCs, first identified in human PB (38), are a small population of circulating or BM-derived MNCs, which are incorporated into sites of neovascularization, and contribute to vascular repair in ischemic tissues by increasing the expression of cytokines, growth factors and/or hormones through autocrine, paracrine and/or endocrine systems (39); this is the process of postnatal vasculogenesis. Thus, EPCs are considered to have potential value in therapeutic vasculogenesis for various ischemic diseases (40). With the same properties as EPCs, they could also be incorporated into hypoxic tumor regions for neovessel formation. Thus, they may be a target for the molecular inhibition of tumor vasculature.

RUNX3, a tumor suppressor, activates the TGF-β signaling pathway and inhibits angiogenesis via the suppression of VEGF transcription (41) and by reducing HIF-1α stability (20). Since VEGF and HIF-1α are crucial factors for vasculogenesis, the present study investigated whether RUNX3 influences the differentiation of EPCs and vasculogenesis. Similar to RUNX3, RUNX1 is also known to function as a tumor suppressor because its mutation results in the development of leukemia (23). As RUNX1 is required for hematopoiesis, a RUNX1 null mutation results in defective hematopoiesis, which leads to the impairment of angiogenesis in the developing embryo (42), suggesting that RUNX1 is required for embryonic vasculogenesis via the modulation of hematopoiesis. However, a transcriptional activity assay in a previous study revealed that VEGF gene expression is inhibited by RUNX1 but increased by an acute myeloid leukemia (AML)/RUNX1 fusion protein (AML/ETO) (43). Therefore, the function of RUNX1 in vessel formation is different between embryogenesis and leukemia development. As RUNX3 and RUNX1 have similar properties in a number of biological responses, the observation that loss of Runx1 function in zebrafish embryos leads to defects in hematopoiesis and vasculogenesis (44) suggests that Runx3 deletion may affect hematopoiesis and/or vasculogenesis. However, RUNX1 cross-regulates the expression of RUNX3 in lymphoid cells (45), further suggesting that the role of Runx3 in the differentiation of BM-HSCs might be distinct from that of Runx1. A genomic study demonstrating different gene expression patterns between early EPCs (small EPCs in the present study) and outgrowth ECs (OECs having similar properties as large EPCs in the present study) revealed that Runx1 is expressed in early EPCs (46). Early EPCs (small EPCs) exhibit more premature and undifferentiated properties in a differentiation hierarchy relative to large EPCs. The present study demonstrated that in small and large EPC CFUs, Runx3 knockout cells formed a greater number of CFUs than did WT cells, suggesting that Runx3 has an inhibitory role in the early and late stages of EPC differentiation from BM stem cells. Runx3 regulates the differentiation of BM-derived dendritic cells by the TGF-β signaling pathway (47). Dendritic cell maturation is essential for the immune response; however, the differentiation process of EPCs may differ from those of other myeloid and lymphoid cells. Runx3 is required for the silencing of CD4 during T-cell lineage decisions (48,49). Human CD34+ HSCs and several normal and malignant hematopoietic cell lines express RUNX3, suggesting that it plays a role in EPCs (50,51). As the differentiation of EPCs from BM-MNCs and their mobilization or homing to injured or ischemic regions are dependent on VEGFR2 and CXCR4-SDF1α signaling pathways (52,53), Runx3 knockout may cause increased EPC differentiation, probably through VEGFR2 and CXCR4-SDF1α pathways. As Runx3 destabilizes HIF-1α and VEGF/VEGFR2 expression is induced by HIF-1α, it may be speculated that the heterozygous knockout of Runx3 increases HIF-1α accumulation and thus VEGF/VEGFR2 expression.

The observation that CXCR4 and SDF1α are also increased by HIF-1α indicates a causative role in the increased EPC function induced by the heterozygous deletion of Runx3. The increase of EPC differentiation and function in heterozygous deletion of Runx3 may be explained by the tumor suppressive function of Runx3. Our results suggest that the stabilization of HIF-1α in Rx3+/− EPCs increases VEGF/VEGFR2 and SDF-1α/CXCR4 expression and their signaling pathways to enhance the differentiation of BM stem cells into EPCs. In a previous study, it was found that RUNX3 facilitates the degradation of HIF-1α (54) by enhancing the interaction of PHDs with HIF-1α, which is a key inhibitor of hypoxia-induced VEGF secretion and angiogenesis in vitro and in vivo (20). Hypoxia downregulates RUNX3 expression and function (6), and is a strong inducer of EPC mobilization and homing via the enhanced expression of various cytokines, including eNOS and SDF1α. The present study indicates that Runx3 downregulation in ischemic regions results in hypoxia-induced cytokine expression and secretion which induces the recruitment of EPCs. Taken together, these results suggest that Runx3 functions to inhibit the differentiation of EPCs from BM-MNCs. Furthermore, Runx3 inhibits EPC functions, including self-renewal, migration, tube formation, as well as mobilization into the PB and homing to injured tissues. Therefore, Runx3 may totally inhibit vasculogenesis via the suppression of EPC differentiation and recruitment to ischemic tissues during the development of new vessels from BM. Therefore, it may be concluded that the inhibition
of EPC differentiation and function by RUNX3 recovery or overexpression is a potential therapeutic strategy for the treatment of tumor progression and metastasis, but may inhibit tissue regeneration in ischemic diseases.

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Availability of data and materials

The datasets used and/or analysed during the current study are available from the corresponding author on reasonable request.

Authors’ contributions

SYNC and SHY performed EPC isolation and analysis, cell culture, molecular analyses and in vivo experiments. DJL and SHL performed the isolation of EPCs and western blot analysis. KL, IJK and WL performed in vivo tumor allograft and IHC experiments with tumor tissue. SCB provided the Runx3−/− mice and analyzed the data. YML initiated the study, supervised the experiments, analysed data and wrote the manuscript.

Ethics approval and consent to participate

The animal experiments were by the Kyungpook National University Institutional Animal Care and Use Committee.

Patient consent for publication

Not applicable.

Competing interests

The authors declare that they have no competing interests.

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