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Activated N-Ras signaling regulates arterial-venous specification in zebrafish

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Abstract

Background: The aberrant activation of Ras signaling is associated with human diseases including hematological malignancies and vascular disorders. So far the pathological roles of activated Ras signaling in hematopoiesis and vasculogenesis are largely unknown.

Methods: A conditional Cre/loxP transgenic strategy was used to mediate the specific expression of a constitutively active form of human N-Ras in zebrafish endothelial and hematopoietic cells driven by the zebrafish lmo2 promoter. The expression of hematopoietic and endothelial marker genes was analyzed both via whole mount in situ hybridization (WISH) assay and real-time quantitative PCR (qPCR). The embryonic vascular morphogenesis was characterized both by living imaging and immunofluorescence on the sections with a confocal microscopy, and the number of endothelial cells in the embryos was quantified by flow cytometry. The functional analyses of the blood circulation were carried out by fluorescence microangiography assay and morpholino injection.

Results: In the activated N-Ras transgenic embryos, the primitive hematopoiesis appeared normal, however, the definitive hematopoiesis of these embryos was completely absent. Further analysis of endothelial cell markers confirmed that transcription of arterial marker ephrinB2 was significantly decreased and expression of venous marker flt4 excessively increased, indicating the activated N-Ras signaling promotes the venous development at the expense of arteriogenesis during zebrafish embryogenesis. The activated N-Ras-expressing embryos showed atrophic axial arteries and expansive axial veins, leading to no definitive hematopoietic stem cell formation, the blood circulation failure and subsequently embryonic lethality.

Conclusions: Our studies revealed for the first time that activated N-Ras signaling during the endothelial differentiation in vertebrates can disrupt the balance of arterial-venous specification, thus providing new insights into the pathogenesis of the congenital human vascular disease and tumorigenic angiogenesis.

Keywords: Vasculogenesis, Arteriogenesis, N-Ras

Background

RAS proteins are small GTPases that have been proved to be essential for the control of cell proliferation, survival, and differentiation [1,2]. Ras genes are evolutionally and functionally conserved in all eukaryotic organisms. In mammals, the 3 Ras genes encode 4 highly homologous proteins: H-, N-, and K-RAS 4A and 4B. The four RAS proteins interact with a common set of activators and effectors, and thus share many biochemical and biological functions [1]. Both Ras genes and components of Ras signaling pathways, including membrane receptor tyrosine kinases (RTKs), Ras-GEFs, Ras-GAPs, and downstream cytoplasmic kinases (RAFs), are reported to be frequently mutated in diverse human cancers and congenital developmental diseases [3]. In particular, activated mutations of N-Ras gene have been detected in human myeloid leukemias with high frequency [3], ranging from 15% to 60%, and several myeloid leukemia-like mice models were established by
inducing the expression of oncogenic N-Ras gene in bone marrow transduction/transplantation [4], transgenic [5] and knock-in [6,7] model systems, suggesting that the oncogenic N-Ras can initiate myeloid leukemias, nevertheless, the long latency and the requirement of cooperative genetic lesions in those models imply that the activated N-Ras signaling alone is insufficient to transform the normal hematopoietic cells into leukemic cells. So far the direct effect of oncogenic N-Ras signaling in hematopoietic cells remains mostly unknown.

Abnormal activation of Ras signaling has been reported to be involved in tumorigenic angiogenesis [8] and congenital vascular developmental diseases [3]. Capillary Malformation-Arteriovenous Malformation (CM-AVM) is a cutaneous congenital vascular disease that is compound of Capillary Malformation (CM) and Arteriovenous Malformation (AVM), and in which the arterial and venous vessels in the skin are connected directly to one another without an intervening capillary bed [9]. Rasa1 gene encodes p120ras-GAP which functions by inhibiting the activity of RAS proteins, and heterozygous inactivating rasa1 mutations were detected and proved to be the causal genetic lesion for CM-AVM [9], suggesting an important role of hyperactive Ras signaling in this disease. So far whether the rasa1 mutations function through an endothelial cell-autonomous manner is still unknown. In addition, the reciprocal signaling between notch pathway mutant (dll4 and Rbpsuh genes) mice embryos exhibit defects in arterial specification of nascent blood vessels and develop Arteriovenous Malformations [12]. However, the pathogenesis of CM-AVM or AVM is largely unknown.

In human N-RAS protein, a point mutation resulting in the substitution of glycine to aspartic acid at codon 12 (G12D) has been proved to impair GAP-stimulated GTP hydrolysis, making the N-Ras signaling constitutively activated [13], hereafter referred as hNRASD12. Lmo2 is a transcriptional factor specifically expressed in the hemangioblasts (the bipotential hemato-endothelial progenitor cells) and hemangioblasts-derived primitive hematopoietic and endothelial cells [14,15]. We recently established and functionally characterized the transgenic Tg(lmo2:Cre) fish expressing the Cre recombinase under the control of zebrafish lmo2 promoter [16], and further showed that Tg(lmo2:Cre) mediated expression of dominant negative Cε/IPα and Bmi1 in zebrafish lmo2+ cells each extended short-lived hematopoietic stem/progenitor cell life span and induced lethal dyserythropoiesis [17]. To explore the roles of aberrant activation of N-Ras signaling in hematopoiesis and vasculogenesis, we used the Tg(lmo2:Cre) to mediate the expression of hNRASD12 in lmo2+ cells of zebrafish embryos, and showed that the activated N-Ras signaling in the hNRASD12 transgenic embryos didn’t induce overt hematopoietic or leukemic phenotypes, while caused severe defective vasculogenesis and subsequently embryonic lethality. The venous cell fate determination was abnormally enhanced at the expense of the arteriogenesis, indicating an endothelial cell-autonomous role of Ras signaling in the arterial-venous specification.

Results

Establishment of transgenic zebrafish lines with hemogenic and endothelial cell-specific expression of human oncogenic N-Ras

To address the roles of disease-associated activation of N-Ras signaling in hematopoietic and endothelial systems in vivo, we induced specific expression of human NRASD12 (hNRASD12) in those tissues using the Cre-loxP system under the control of lmo2 promoter in the zebrafish.

We first established the Tg(β-actin:LDL-hNRASD12) lines (Figure 1A) and obtained 7 heterozygous stable lines that ubiquitously expressed the fluorescent DsRed protein during embryonic development (data not shown). All 7 lines were crossed to the heterozygous Tg(lmo2:Cre) fish individually to generate four different genotypes in the progeny: hNRASD12 Cre− (wild-type), hNRASD12 Cre+, hNRASD12 Cre− β-actin:LDL-hNRASD12:wild-type and hNRASD12 Cre+ β-actin:LDL-hNRASD12; lmo2:Cre). In particular, the β-actin:LDL-hNRASD12;lmo2:Cre double transgenic embryos derived from line 4 showed robust expression of hNRASD12 transcripts in the intermediate cell mass (ICM) region at 22 hours post fertilization (hpf) (Figure 1B), which mostly resembled the expression pattern of endogenous lmo2 transcripts in wild-type embryos at the same stage (Figure 1D). And all the β-actin:LDL-hNRASD12;wild-type embryos, hereafter referred as control embryos, showed no detectable expression of hNRASD12 transcripts in the ICM region (Figure 1C). Consistently, the expression of hNRAS protein was only detected in the β-actin:LDL-hNRASD12;lmo2:Cre embryos at 28 hpf (Figure 1E), hereafter referred as hNRASD12 embryos.

Elevated Ras signaling by overexpression of hNRASD12 has no impact on primitive hematopoiesis

After confirmation of the hNRASD12 expression in the transgenic embryos, we examined the effect of activated N-Ras signaling on the specification and differentiation of blood cells in the hNRASD12 embryos. Like mammalian, zebrafish hematopoiesis includes two consecutive waves of hematopoiesis: primitive and definitive
hematopoiesis [18]. Primitive hematopoiesis originates from the anterior and posterior lateral plate mesoderm and produces erythrocytes, macrophages, and granulocytes, providing blood cells for early embryogenesis [18]. The definitive hematopoietic stem cells (HSCs) emerge from the hemogenic endothelium at the ventral wall of the dorsal aorta from 30 hpf onward, called endothelial hematopoietic transition (EHT) [19,20], and generate all types of blood cells in adulthood [18]. To characterize the hematopoietic phenotypes in the hNRASD12 embryos, we performed whole mount in situ hybridization (WISH) analysis to examine the hematopoietic markers expression [21,22] in the control and hNRASD12 embryos at 18 and 22 hpf, including lmo2 (hemangioblasts), gata1 (primitive erythroid progenitors), αe1 globin (mature erythrocytes), pu.1 (primitive myeloid progenitors), l-plastin (primitive macrophages), mpo (neutrophils), cmyb and runx1 (HSCs). The results showed that expression of those primitive hematopoietic markers is not affected in the hNRASD12-expressing embryos (Figure 2A & B). Consistent with the WISH results, the real-time quantitative PCR (qPCR) analysis showed no significant difference at the expression level of these markers between the hNRASD12 and control embryos.

Figure 1 Validation of hNRASD12 expression in the β-actin:LDL-hNRASD12;lmo2:Cre transgenic embryos. (A) Diagrammatic scheme of the β-actin-LDL-hNRASD12 transgenic construct. (B-D) The expression analyses of the hNRASD12 transcripts via whole mount in situ hybridization (WISH) in the β-actin:LDL-hNRASD12;lmo2:Cre embryo (hereafter referred as hNRASD12 embryo) (B) and β-actin:LDL-hNRASD12 wild-type embryo (hereafter referred as control embryo) (C) respectively at 22 hours post fertilization (hpf), and lmo2 transcripts in the wild-type embryo at 22 hpf (D). Black arrows denote the location of the intermediate cell mass (ICM) region. (E) Confirmation of the expression of hNRAS protein in the hNRASD12-embryos at 28 hpf via western blotting analysis.

Figure 2 The primitive erythropoiesis and myelopoiesis are not affected by the expression of hNRASD12. (A-B) WISH analysis of primitive hematopoietic markers expression in the control embryos and hNRASD12 expression embryos respectively at 18 (A) and 22 hpf (B). (C-D) Verification of the expression level of the hematopoietic markers in the control or hNRASD12 embryos at 18 (C) and 22 hpf (D) via real-time quantitative PCR. Data shown are means ± SEM of at least three independent experiments.
indicating the activated N-Ras doesn’t alter the development of primitive blood cells (Figure 2C & D). However, the definitive HSCs marked by runx1 and cmyb couldn’t be detected in the aorta-gonad-mesonephros (AGM) region of the hNRASD12 embryos (Figure 3A-D). According to the defective blood circulation and dorsal aorta formation in the hNRASD12 embryos described below, the lack of definitive hematopoiesis is likely the secondary effect of the vascular defects, as both the normal blood circulation and dorsal aorta formation have been proven to be indispensable for the emergency of definitive HSCs in the AGM region [23,24], thus preventing from directly dissecting the role of activated N-Ras signaling in definitive hematopoiesis. Taken together, these data show that the oncogenic N-Ras signaling in the zebrafish lmo2 cells doesn’t induce directly hematopoietic or leukemic phenotypes in the transgenic embryos.

Vascular morphogenesis defects in hNRASD12-expressing transgenic embryos

Expression of hNRASD12 under the lmo2 promoter will lead to expression both in the hematopoietic and endothelial lineages, so we then examined the circulatory system and found the hNRASD12 embryos at 28 hpf lacked blood circulation both in head and trunk vessels, and the blood cells were accumulated in the trunk axial vessels and heart chambers (Figure 4B & D, and Additional files 1 and 2: movie S3 and S4) in comparison of normal circulation in the control embryos (Figure 4A & C, and Additional files 3 and 4: movie S1 and S2). The hNRASD12 embryos died between 5 and 8 days post fertilization (dpf) with the persistent lack of blood circulation. To validate the blockage of circulation was specifically induced by the expression of human NRASD12, we designed a rescue experiment by using an antisense ATG morpholino (MO) targeting the translational initiation site of this transgenic hNRASD12. Firstly, the efficiency of the hNRAS MO was confirmed by efficiently blocking both the translation of the GFP mRNA reporter (Additional file 5: Figure S1A-D) in wild-type embryos and the expression of hNRASD12 protein in the hNRASD12 embryos (Additional file 5: Figure S1E). Compared with the control MO, the hNRAS MO that was injected into one-cell stage embryos derived from the crossing between the heterozygous Tg(lmo2:Cre) and Tg(β-actin:LDL-hNRASD12) fish restored the blood circulation of hNRASD12 embryos to a comparable level as that of control embryos (Figure 4E), indicating the blocked blood circulation is specifically induced by hNRASD12 ectopic expression. Considering the primitive hematopoiesis of hNRASD12 embryos is normal, we hypothesized that the congestion of circulation resulted from the defective cardiovascular development. We then checked the cardiovascular structure and function by using the fluorescent microangiography assay. The fluorescein-coupled latex beads were injected into the inflow tract of the atrium of the control and hNRASD12 embryos respectively, and were observed to perfuse the whole vascular system of the control embryos in five minutes (Additional file 5: Figure S2A, upper panel). However, it took over 30 minutes for injected beads to passively diffuse into the anterior trunk of the hNRASD12 embryos (Additional file 5: Figure S2A, lower panel), indicating the activated Ras signaling disrupts the circulation. In particular, the consecutive accumulation of fluorescent beads in the trunk of hNRASD12 embryos (Additional file 5: Figure S2A, lower panel) suggested the existence of only one axial vascular tube. We further examined the function of the heart of hNRASD12 embryo by counting the rate of heartbeats, and the result showed that the average rate of heartbeats in the hNRASD12 embryos at 30 hpf was only slightly slower than that of control embryos (Additional file 5: Figure S2B), suggesting the phenotype of heartbeat contributes little to the circulation blockage.

To further analyze the vascular defect in the hNRASD12 embryos, we crossed the Tg(flk1:GFP) fish [25], an endothelial reporter transgenic line, with Tg(β-actin:LDL-hNRASD12) fish to obtain the double transgenic progeny, which were raised to adulthood and subsequently crossed to Tg(lmo2:Cre) fish to generate the β-actin:LDL-hNRASD12/flk1:GFP/lmo2:Cre (referred as hNRASD12; flk1:GFP) embryos. In the trunk of β-actin:LDL-
hNRASD12;flk1:GFP;wild-type (referred as control;flk1:GFP) embryos at 28 hpf, the primary blood vessels consisted of the dorsal aorta (DA) and posterior cardinal vein (PCV), and the intersegmental vessels (ISVs) formed by sprouting from the dorsal aorta (Figure 5A) [26]. However, hNRASD12;flk1:GFP embryos failed to form morphologically distinct DA or PCV, and the ISVs were also defective (Figure 5B). The results of immunofluorescence analysis on transverse sections showed the presence of only one vascular tube in the trunk of hNRASD12-expressing embryo (Figure 5C & D). Furthermore, the morphology of head vasculature was also examined, in the control;flk1:GFP embryos at 28 hpf, the paired bilateral lateral dorsal aortas (LDAs) merged backward into single dorsal aorta at the boundary between head and trunk, and the posterior cardinal vein (PCV) split forward into a pair of vessels in the cranial trunk, then the branches of PCV emptied into the common cardinal veins (CCVs) on the either side of yolk, which came into the sinus venous of the heart (Figure 5E & G) [26]. Unlike that of control;flk1:GFP embryos, the LDAs of hNRASD12-expressing embryos were evidently atrophic while CCVs and the bifurcation of PCV were greatly expanded, however, the primordial hindbrain channels (PHBCs), the primary veins in the head at this stage, seemed normal (Figure 5F & H). In addition, basilar artery (BA), the main artery in the hindbrain of the zebrafish embryo formed by endothelial sprouting of PHBCs [27], was lost in the hNRASD12-expressing embryo (Figure 5I & J). Taken together, our observations suggested that the activated N-Ras signaling in the zebrafish lmo2+ cells caused the abnormal assembly of the axial blood vessels and the defective arteries formation and expansion of veins in the head vasculature system of the hNRASD12 embryos.

**hNRASD12 promotes venous fate specification at the expense of arterial fate in zebrafish**

As Ras signaling has been proved to be important for cell proliferation [1,2], the frequencies of endothelial cells in the hNRASD12;flk1:GFP and control;flk1:GFP embryos at 28 hpf were respectively quantified by flow cytometry analysis, and the result showed no significant difference in the frequencies of GFP+ endothelial cells between hNRASD12-expressing and control embryos (Figure 6A). We further characterized the arterial and venous cells lineages respectively in the hNRASD12 embryos by performing WISH analysis. EphrinB2, hRT, dll4 and grl genes are specifically expressed in the arterial cells of wild-type embryo’s trunk region [23,27], while ephB4, fli4 and dab2 mark the venous cells [28]. The result of WISH showed that the arterial cells were evidently reduced in hNRASD12-expressing embryos compared with control embryos (Figure 6B, B’, D, and D’), and failed to form an intact dorsal aorta (Figure 6C, C’, E, and E’); On the contrary, the venous cells in the hNRASD12 embryos were greatly expanded and organized to be an enlarged vein-like vascular tube (Figure 6F-I and F’-I). To confirm that expression of hNRASD12 induced an unbalanced arterial-venous specification in the same embryo, we performed 2-color WISH analysis with digoxigenin-labeled ephrinB2 and fluorescein-labeled fli4 antisense probes, and observed the expansion of venous cells and reduction of arterial cells occurring in a single hNRASD12-expressing embryo (Figure 6J’ & K’).
to that in control embryo (Figure 6J & K). These results from WISH with dll4, grl, and dab2 probes also supported the above observations (Additional file 5: Figure S3 A-F) that almost the entire vascular tube in the trunk of hNRASD12 embryo consists of venous endothelial cells.

Vasculogenesis is referred to as the de novo formation of the blood vessels, involving the differentiation, proliferation and migration of endothelial cells [29]. In zebrafish, the angioblasts at the bilateral posterior lateral plate mesoderm migrate medially to form the vascular cord between 14 and 18 hpf, meanwhile the arterial-venous specification is established [23,25]. Then the venous cells sprout ventrally to separate from the arterial cells between 21 and 24 hpf, called arterial-venous segregation [30]. Subsequently, the dorsal arterial cells organize to be dorsal aorta by cord hollowing [31], while the posterior cardinal vein forms by embrace of the hematopoietic cells [30].

To further confirm that the defects in the assembly of the blood vessels and the arterial and venous fate decisions are specifically induced by the transgenic expression of hNRASD12, we knocked down the hNRASD12 expression using hNRAS MO in the hNRASD12 embryos. The trunk vasculature was imaged under a confocal microscopy and arterial and venous cells in the region were evaluated by WISH analysis of arterial-venous cell fate is just established [25], indicating N-Ras signaling regulates the specification of arterial and venous cells before the arterial-venous segregation and onset of circulation. In addition, previous studies have revealed an important role of the angioblasts migration between 14 and 18 hpf in the arterial-venous specification [23,32]. To examine whether the constitutively activated N-Ras signaling affects the medial migration of angioblasts, we used flk1 as a probe to trace this process. The result showed the medial migration began at 14–15 hpf, and nearly all angioblasts arrived at the midline at 17–18 hpf in the control embryo (Figure 8A-E), and hNRASD12-expressing embryo showed a similar pattern (Figure 8F-J), indicating the activated N-Ras signaling specifically affects the arterial-venous specification in an angioblasts-medial-migration-independent manner.

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venous cells specific markers. Compared to the control MO, the hNRAS MO efficiently rescued both the defective assembly of blood vessels (Additional file 5: Figure S4A) and the unbalanced arterial-venous fates of the endothelial cells (Additional file 5: Figure S4B) in the trunk region of hNRASD12 embryo, indicating the defective vasculogenesis in the transgenic embryos is specifically caused by hNRASD12 expression.

Thus we conclude that the activated N-Ras signaling by expressing hNRASD12 in lmo2+ cells disrupts the balance of the arterial-venous specification, where the venous fate decision is enhanced while the arterial specification is repressed, which in turn causes abnormal assembly of main blood vessels and the defect in the development of intersegmental vessels, leading to blockage of blood circulation and embryonic lethality. Nevertheless, expression of hNRASD12 affects neither the endothelial specification from mesoderm cells, the medial migration of angioblasts, nor the proliferation of endothelial cells.

The hierarchical signaling molecules, including sonic hedgehog (Shh), vascular endothelial growth factor A (VEGFA), phospholipase C gamma-1 (Plcg1), and notch have been shown to be essential for the arterial cell fate decision [33]; while chicken ovalbumin upstream promoter-transcription factor II (COUP-TFII), also known as Nuclear Receptor 2F2 or NR2F2, is indispensable for the venous specification [33]. To figure out whether hNRASD12 regulates the transcriptional expression of those genes, we performed WISH analysis with riboprobes of these arterial-venous regulatory genes, and found no significant difference between the control and hNRASD12 embryos (Additional file 5: Figure S5A-F and S5A’-F).

Previous studies have shown that forced expression of vegfa121 in wild-type embryo could transform all the endothelial cells into ephrinB2+ arterial cells [34], however, we observed the failure of hNRASD12 embryos to respond to exogenous expression of vegfa121 (Additional file 5: Figure S6A-E), indicating that N-Ras signaling functions downstream of vegfa to negatively regulate the vegfa induced arterial development.

Considering notch signaling has been proved to act downstream of shh-vegfa signaling axis to regulate the arterial endothelial differentiation in zebrafish [35], we tried to rescue the hNRASD12-induced defective arteriogenesis by injecting a transient transgenic construct into the hNRASD12 embryos, in which a constitutively activated form of notch1 containing only the intracellular domain (referred as NICD1) was driven by the zebrafish flk1 promoter, however, no recovery of arteriogenesis was observed in those injected embryos (data not shown), implying defects caused by activated Ras is not through disrupting the notch signaling.

Figure 6 Expression of hNRASD12 promotes venous specification at the expense of arterial fate during embryonic endothelial differentiation. (A) Quantitation of total endothelial cells by flow cytometry analysis on the control;flk1:GFP and hNRASD12;flk1:GFP embryos at 28 hpf. Representative results obtained from three independent experiments. 2 tailed t-test of frequency of endothelial cells in hNRASD12;flk1:GFP embryos vs control;flk1:GFP embryos shows no significant difference. (B-E) WISH analysis of ephrinB2+ and hRT+ arterial cells in the trunk of control (B-E, blue triangles) and hNRASD12 embryos (B’-E’, red triangles) at 28 hpf. (F-I) WISH analysis of flt4+ and ephB4+ venous cells in the trunk of control (F-I) and hNRASD12 embryos (F-I) at 28 hpf. The yellow square brackets mark the width of the venous strip in the trunk. (J-K) 2-color WISH analysis with digoxigenin-labeled ephrinB2 (white arrows) and fluorescein-labeled flt4 (white triangles) in control (J, K) and hNRASD12 embryos (J’, K’) respectively at 28 hpf.
Functional conservation of human and zebrafish N-Ras signaling in vasculogenesis

To assess the conservation between the human and zebrafish N-Ras signaling, we compared the protein sequences of the human, mouse and zebrafish N-RAS, and found that there is a high degree of sequence similarity between the zebrafish and mammalian N-RAS with a high identity of 95% in the critical H_N_K_Ras_like domain (G domain) (Additional file 5: Figure S7). To further determine whether the activated zebrafish N-Ras signaling has a conserved function in the vasculogenesis as human N-Ras, we generated both lmo2:EGFP-2A-hNRASD12 and lmo2:EGFP-2A-zNRASD12 transgenic constructs, in which the zebrafish lmo2 promoter was used to transiently direct the specific expression of activating mutant of human and zebrafish N-Ras in blood and vasculature respectively. The WISH analysis showed that both the hNRASD12 and zNRASD12 transcripts were appropriately expressed in the ICM region (Additional file 5: Figure S8A), similar to the expression pattern of endogenous lmo2 gene (Figure 1D), and the expression of the human and zebrafish NRASD12 proteins was also confirmed via western blotting assay (Additional file 5: Figure S8B). The trunk vasculature of embryos expressing the hNRASD12 or zNRASD12 was characterized under a confocal microscopy. And compared to the embryos injected with lmo2:EGFP-2A, both the hNRASD12 and zNRASD12-expressing embryos had only one vascular tube in the trunk (Additional file 5: Figure S8C). Additional WISH analysis of the arterial marker ephrinB2 and venous marker flt4 genes showed large expansion of venous cells population and nearly lack of the arterial cells in the trunk region (Additional file 5: Figure S8C). These results strongly suggest a conserved function between activated human and zebrafish N-Ras and that the N-Ras signaling participate in regulating the zebrafish vascular development.

Figure 7 Loss of arterial specification by hNRASD12 expression before arterial-venous segregation and onset of circulation. (A-H) WISH analysis of arterial (ephrinB2+, hRT+) and venous (flt4+, dab2+) cells in the trunk of control (A, C, E, G) and hNRASD12 embryos (B, D, F, H) at 20 hpf respectively. (I-L) 2-color WISH analysis with digoxigenin-labeled ephrinB2 (white arrows) and fluorescein-labeled flt4 (white triangles) in control (I, J) and hNRASD12 embryos (K, L).
Discussion

In this study, we showed that transgenic overexpression of activated human \textit{NRAS} (\textit{hNRASD12}) in zebrafish \textit{lmo2}\(^+\) cells caused un-circulated vasculature, but had no impact on the primitive hematopoiesis during embryogenesis. In particular, the expression of \textit{hNRASD12} promoted the venous fate at the expense of arterial fate during arterial-venous specification, indicating an important novel role of activated Ras signaling in vascular morphogenesis.

Our whole mount \textit{in situ} hybridization (WISH) and quantitative RT-PCR analysis of the expression of the hematopoietic markers showed the expression of \textit{hNRASD12} in the \textit{lmo2}\(^+\) hemangioblasts neither affected the specification or differentiation of the primitive hematopoietic cells, nor induced any leukemic phenotypes. However, the mice models expressing \textit{hNRASD12} exhibited leukocytosis and progressed to myeloid leukemias [4-7]. The discrepancy between our zebrafish model and those mice models is likely due to the difference in the targeting cell type and the developmental window of the \textit{NRASD12} expression. In the mice models, the expression of the \textit{NRASD12} was induced in the adult bone marrow cells, the adult hematopoietic organs, and caused dysplastic myelopoiesis/leukemias with long latencies (> 4 weeks), which highlight the possible requirement of a definitive hematopoietic environment and the length of time to accumulate cooperative genetic mutations for the development of \textit{NRASD12}-associated leukemia [36]. In our study, the expression of \textit{hNRASD12} was activated during the transient primitive hematopoietic stage and embryos failed to develop through the definitive hematopoietic stages due to lack of definitive hematopoiesis as a result of the defective vasculogenesis, and early embryonic lethality (5–8 days post fertilization), preventing the occurrence of the myeloid disorders and leukemias. Our results supported the concept that leukemogenesis is a multistep process involving cooperating genetic mutations [37], the elevation of RAS signaling by the expression of \textit{hNRASD12} alone in the short-lived primitive hematopoietic cells was insufficient for leukemogenesis.

Whether the hyperactive Ras signaling caused by inactivating \textit{RASA1} mutations in Capillary Malformation-Arteriovenous Malformation (CM-AVM) functions through an endothelial cell-autonomous manner is still unknown [9]. Here we showed that the expression of oncogenic human N-Ras in the \textit{lmo2}\(^+\) cells disrupted arterial-venous specification and angiogenesis in zebrafish embryos, indicating an endothelial cell-autonomous role of oncogenic ras signaling in vasculogenesis. Consistently, the \textit{rasa1}-deficient mice embryos are embryonic lethal and exhibit atrophic dorsal aorta and disorganized patterns of the intersegmental arteries [38]. The EphrinB2/EphB4 signaling [10,11] and notch pathway [12] are involved in the pathogenesis of Arteriovenous Malformation (AVM), suggesting the direct genetic connection between the arterial-venous specification and CM-AVM/AVM. In our study, the transgenic expression of \textit{hNRASD12} induces defective arterial-venous specification in zebrafish, and this vertebrate genetic model may help us to further dissect the pathogenesis of CM-AVM or AVM.

In mice, it has been demonstrated that the germ line oncogenic Ras expression leads to embryonic lethality [6,39], where the \textit{Protamine-Cre} (PrmCre) mice mediates constitutive and ubiquitous expression of oncogenic \textit{KRASD12} (floxed knock-in) [39], and \textit{mox2-Cre} mice mediates germ line expression of endogenous oncogenic \textit{NRASD12} [6], however, the cellular and pathological mechanisms for the embryonic lethality in those studies are unknown. Here we show that the expression of oncogenic \textit{hNRASD12} during zebrafish embryogenesis induces embryonic lethality through disrupting the arterial-venous cell fate decision, offering a possible mechanistic explanation for the oncogenic Ras signaling induced embryonic lethality. Nevertheless, the phenotypes of endothelial development need to be further examined in those mice models [6,39].
The zebrafish has emerged as the first vertebrate model organism that is suitable for large-scaled whole-animal small chemical molecule screening, contributing to several aspects of the drug development process, including target identification, disease modeling, lead discovery and toxicology [40]. The oncogenic NRASD12-expressing mice develop leukemias in adult stage [4-7], and are not suitable for drug screening due to the cost and the large size of an individual, while the NRASD12 transgenic zebrafish embryos may be helpful for us to perform large-scaled whole-animal small chemical molecule screening to find oncogenic NRASD12-associated modulators using a developmental stage-controlled expression of oncogenic NRASD12.

Conclusions
In conclusion, we report firstly the constitutively activated N-Ras signaling regulates the arterial-venous specification in vertebrate, and also established an oncogenic hNRASD12 induced animal disease model that is feasible for high-throughput whole-animal drug screening and validating the functionality of NRASD12 modulators as potential therapeutics for a variety of human diseases caused by oncogenic mutant RAS.

Methods
Fish care
Zebrafish maintenance, breeding, and staging were performed as previously described [41]. The zebrafish experiments were approved by the Animal Experimentation Committee of Shanghai Institutes for Biological Sciences and institutional review board of Institute of Health Sciences.

Generation of Tg(β-actin:LDL-hNRASD12) transgenic line
Human NRASD12 (hNRASD12) was obtained by PCR from a myc-tagged NRASD12 in a retroviral vector [4] and cloned into pCS2+ vector at the ClaI and Xhol sites to get the hNRASD12-pCS2+ construct. Then the hNRASD12-SV40 fragment was excised from hNRASD12-pCS2+ plasmid by ClaI and Kpnl and cloned into the same sites of β-actin-loxP-DsRed-SV40-loxP-PSBK-1-SceI plasmid [16] to get the β-actin-loxP-DsRed-SV40-loxP-hNRASD12-PSBK-1-SceI transgenic construct. Then this construct was co-injected with I-SceI meganuclease (New England Biolabs) into the fertilized zebrafish eggs at one cell stage. The injected embryos were raised to adulthood to establish the stable transgenic lines.

Transient transgenic constructs
The zebrafish lmo2 promoter of 2.5 kbp was obtained by PCR amplification from the lmo2-Cre-PSBK-I-SceI plasmid [16] and cloned into the EGFP-2A-pDestTol2 plasmid (Z.W. Dong et al., unpublished data) at the Xhol and BamHI sites to get the lmo2-EGFP-2A-pDestTol2 construct. Then the human NRASD12 (hNRASD12) and zebrafish NRASD12 (zNRASD12) fragments were obtained by PCR from the β-actin-locP-DsRed-SV40-locP-hNRASD12-PSBK-1-SceI plasmid and cDNA from the wild-type embryos respectively, and cloned into the lmo2-EGFP-2A-pDestTol2 construct at the Xmal and Sall sites to get the lmo2-EGFP-2A-hNRASD12/ zNRASD12-pDestTol2 constructs. Note of, the forward primer for the amplification of zebrafish NRASD12 contains mutations at the codon 12 from GGA to GAT, resulting in the substitution of glycine (G) to aspartic acid (D). Then the lmo2-EGFP-2A-pDestTol2, lmo2-EGFP-2A-hNRASD12-pDestTol2 and lmo2-EGFP-2A-zNRASD12-pDestTol2 transgenic plasmids (25 ng/ul) were individually co-injected with KCl (0.2 M) and Tol2 transposase mRNA (25 ng/ul) as previously described [42] into the Tg(fkl: mCherry) or wild-type zebrafish embryos at one cell stage.

To generate the construct of the zebrafish flk1 promoter driven NICD1, the HA-tagged NICD1 fragment was obtained by PCR from the PME-NICD1 (zebrafish) plasmid and cloned into the flk1-PolyA-psBK1 plasmid at the BamHI and EcoRI sites to get the flk1-HA-NICD1-PolyA-pBK12 construct. Then the construct was co-injected with I-SceI meganuclease into the hNRASD12 transgenic embryos at one cell stage.

Morpholino (MO)
Embryos were injected with 2 nl human NRAS ATG MO (Gene Tools): 5’-ACCAGTTTGTACTCAGTCAT CGA-3’. The efficiency of hNRAS MO was confirmed both by western blotting and co-injection with hNRAS1-60-GFP mRNA reporter containing the targeting sites of hNRAS MO.

Preparation of mRNAs and antisense probes, whole-mount in situ hybridization (WISH)
A 60 bp fragment of human NRAS that contains the hNRAS ATG MO binding sites was fused to the N terminal of GFP in the pCS2+ vector to get the hNRAS1-60-GFP-pCS2+ construct. The hNRAS1-60-GFP, zebrafish vegfaa121, [35] and Tol2 transposase [43] mRNAs were synthesized with MESSAGE mMACHINE Kit (Ambion).

The digoxigenin- or fluorescein-labelled antisense probes used for WISH analysis in this study includes lmo2 [16], (shh, notch1) [17], vegfaa121 [35], (plcg1, pu.1, l-plastin, mpo, gata1, ace globin, flk1) [44], (cmyb, runx1) [45], (ephrinB2, hRT, gcl, notch3) [23], (dab2, ephB4 and flt4) [28]. And the hNRASD12, zNRASD12,
$dll$ and $nr2f2$ were cloned into pCS2+ plasmid and their digoxigenin-labelled antisense probes were synthesized with T3 polymerase (Ambion). Single- and 2-color WISH were performed as described previously [17]. Embryos were mounted in 3% methylcellulose and captured under the Nikon SMZ1500 microscope equipped with a Nikon Dxm1200F digital camera and ACT-1 software.

Genomic DNA isolation and genotyping
Embryos were incubated in lysis buffer (Tris–HCl 1 M, pH 8.3; KCl 1 M; Tween 20 10%; NP40 10%) and protease K (10 mg/ml) at 55°C overnight. Then 3700 rpm for 10 min at 4°C, and the supernatant was subject to genomic PCR with specific primers for $Cre$ and $hNRASD12$.

Real-time Quantitative PCR
Total RNAs were extracted from 20 zebrafish embryos using Trizol reagent (Invitrogen). RNA was reverse-transcribed using random hexamers and SuperScript III Reverse Transcriptase (Invitrogen). 2×PCR Mix (TaKaRa, Premix Ex Taq”) containing SYBR Green I was used for the real-time quantitative PCR analysis with the Applied Biosystems 7900HT Fast Real-Time PCR System. The relative expression values were normalized against the internal control $gapdh$ (qPCR primer sequences were listed in Additional file 5: Table S1).

Western blotting and immunofluorescence
The western blotting assay was performed as described previously [17] with antibodies: human NRAS (Santa Cruz), GAPDH (Cell Signaling Technology). Immunohistochemistry was performed as described previously [25], and the trunk of embryo was transversely sectioned with a Leica VT1000S vibratome into 50–100 μm sections, and F-actin was stained with Phalloidin-TRITC (0.007 μM, Sigma), and GFP antibody (Invitrogen, A6455) was used. Sections were mounted with Dako-fluorescent mounting media (Dako North America) prior to imaging with an Olympus FV 1000 confocal microscopy.

Fluorescence microangiography
The fluorescent-coupled latex beads (Molecular Probes) were injected into the inflow tract of the atrium of the zebrafish embryo at 30 hpf with a dose of 2 nl. The beads flowed with blood circulation and labeled all the vascular vessels of wild-type embryos in 5 minutes.

Flow cytometry analysis
Embryos at 28 hpf were dissected in DPBS (Invitrogen) after dechorionation and deyolking [46], and digested with 1x trypsin/EDTA (Life Technologies) for 30 min at 28.5°C. Single cell suspension was obtained by centrifugation at 400 g for 5 min, washed twice with 0.9XPBS/5%FBS, and passed through a 40 μm nylon mesh filter. Fluorescence-activated cell sorting was performed with FACS Aria (BD Biosciences) to quantify the frequency of the GFP$^+$ endothelial cells.

Image analysis
Embryos were imaged under an Olympus FV 1000 confocal microscopy equipped with the FV10-ASW version3 software.

Statistical analysis
The values were represented as mean ± SEM. Two-tailed students’-t-tests were used for two groups comparison analysis, and P < 0.05 was considered to be significant. The bar charts were used to reflect the alterations of experimental data.

Additional files

Additional file 1: Movie S3. The blood circulation in the trunk of $\beta$-actin:LDL-hNRASD12; $lmo2$:Cre embryo at 28 hpf.

Additional file 2: Movie S4. The blood circulation in the head of $\beta$-actin:LDL-hNRASD12; $lmo2$:Cre embryo at 28 hpf.

Additional file 3: Movie S1. The blood circulation in the trunk of $\beta$-actin:LDL-hNRASD12; wild-type embryo at 28 hpf.

Additional file 4: Movie S2. The blood circulation in the head of $\beta$-actin:LDL-hNRASD12; wild-type embryo at 28 hpf.

Additional file 5: Supplementary figures and table.

Competing interests
The authors declare that they have no competing interests.

Authors’ contributions
CGR performed experiments and analyzed data. LW, XEJ, YJ, YC, and MD assisted with experiments. TXL, WJP, YZ(Yong Zhou), YJ, YC, and MD assisted with experiments. TXL, WJP, YZ(Yong Zhou), RBB, and CGR designed the research plan and wrote the paper. All authors were involved in the writing and final approval of the manuscript.

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