ARAP3 Functions in Hematopoietic Stem Cells

Yiwen Song¹,², Jing Jiang², Sonja Vermeren³, Wei Tong¹,²*

¹. Department of Pediatrics, University of Pennsylvania Perelman School of Medicine, Philadelphia, Pennsylvania, United States of America, ². Division of Hematology, Children’s Hospital of Philadelphia, Philadelphia, Pennsylvania, United States of America, ³. MRC Centre for Inflammation Research, The Queen’s Medical Research Institute, The University of Edinburgh, United Kingdom

* tongw@email.chop.edu

Abstract

ARAP3 is a GTPase-activating protein (GAP) that inactivates Arf6 and RhoA small GTPases. ARAP3 deficiency in mice causes a sprouting angiogenic defect resulting in embryonic lethality by E11. Mice with an ARAP3 R302,303A mutation (Arap3KI/KI) that prevents activation by phosphoinositide-3-kinase (PI3K) have a similar angiogenic phenotype, although some animals survive to adulthood. Here, we report that hematopoietic stem cells (HSCs) from rare adult Arap3KI/KI bone marrow are compromised in their ability to reconstitute recipient mice and to self-renew. To elucidate the potential cell-autonomous and non-cell-autonomous roles of ARAP3 in hematopoiesis, we conditionally deleted Arap3 in hematopoietic cells and in several cell types within the HSC niche. Excision of Arap3 in hematopoietic cells using Vav1-Cre does not alter the ability of ARAP3-deficient progenitor cells to proliferate and differentiate in vitro or ARAP3-deficient HSCs to provide multi-lineage reconstitution and to undergo self-renewal in vivo. Thus, our data suggest that ARAP3 does not play a cell-autonomous role in HSPCs. Deletion of Arap3 in osteoblasts and mesenchymal stromal cells using Prx1-Cre resulted in no discernable phenotypes in hematopoietic development or HSC homeostasis in adult mice. In contrast, deletion of Arap3 using vascular endothelial cadherin (VEC or Cdh5)-driven Cre resulted in embryonic lethality, however HSCs from surviving adult mice were largely normal. Reverse transplantsations into VEC-driven Arap3 conditional knockout mice revealed no discernable difference in HSC frequencies or function in comparison to control mice. Taken together, our investigation suggests that despite a critical role for ARAP3 in embryonic vascular development, its loss in endothelial cells minimally impacts HSCs in adult bone marrow.
Introduction
Hematopoietic stem cells (HSCs) are the critical source of all blood cells. Their potential for self-renewal and multi-lineage repopulation sustains the rapid turnover of the blood system throughout life. The first HSC arises from the hemogenic endothelium in the Aorta-Gonad-Mesonephros (AGM) region of the embryo and subsequently colonizes the fetal liver [26]. In the adult mouse, HSCs reside in complex bone marrow (BM) niches that are not mutually exclusive. Extensive research has shown that HSC perivascular and osteoblastic niches are comprised of endothelial cells, mesenchymal stromal cells, osteoblasts, sympathetic nerves and non-myelinating Schwann cells [1–3].

HSC functions are tightly regulated by a plethora of extrinsic and intrinsic regulatory pathways. One such family of regulators is the Rho family of GTPases, molecular switches that cycle between an active guanosine triphosphate (GTP)-bound form and an inactive guanosine diphosphate (GDP)-bound form [4, 5]. Rho GTPases play pivotal roles in hematopoietic stem and progenitor cell (HSPC) actin cytoskeletal reorganization [6–13], with recent genetic studies expanding our knowledge of their roles to include HSC self-renewal, multi-lineage differentiation, homing/migration, proliferation, cytokinesis and survival [14–19]. GTPase-activating proteins (GAPs) stimulate the hydrolysis of bound GTP to GDP, thereby inactivating GTPases. GAPs, such as p190B RhoGAP, have been shown to be important regulators of HSC engraftment and interaction with its microenvironment [20, 21].

ARAP3 is a dual Arf and Rho GTPase-activating protein that was first identified in porcine leukocytes for its ability to bind to phosphatidylinositol (3,4,5)-triphosphate (PIP$_3$) [22]. ARAP3 contains two distinct GAP domains that accelerate the rate of GTP hydrolysis to attenuate Arf6 and RhoA signaling [23, 24]. Previous in vitro studies found that either exogenous ARAP3 expression in epithelial cells or RNAi-mediated ARAP3 depletion in endothelial cells disrupts F-actin or lamellipodia formation, respectively, resulting in a cell rounding phenotype and failure to spread [25, 26]. This implies that ARAP3 controls Arf6 and RhoA in a tightly regulated fashion, and that maintaining precise regulation of ARAP3 activity is crucial to actin organization in the cell. RhoA has been characterized in vivo to regulate migration and chemotaxis of mature hematopoietic cells [27, 28], as well as HSPC engraftment, multi-lineage repopulation and cell survival [9, 14, 15], while the role of Arf6 in hematopoiesis is largely unknown.

In mice, ARAP3 is most highly expressed in the endothelium and bone marrow, and has been found to be critical to vascular development [29, 30]. Germline deletion or Tie2-Cre-mediated deletion of Arap3 in mice leads to embryonic lethality by E11 due to defects in sprouting angiogenesis of the endothelium [29]. Since HSCs arise from the hemogenic endothelium during embryonic development around E10.5 [31], and give rise to all subsequent hematopoietic cells in the fetal liver and in the adult BM, this genetic model precludes further studies of ARAP3 function in definitive hematopoiesis and HSC function. Conditional
Arap3 deletion in neutrophils has been shown to alter their adhesion-dependent functions [32, 33], but the role of ARAP3 in HSPCs has yet to be defined.

ARAP3 is a phosphoinositide 3-OH kinase (PI3K)- and Rap-regulated GAP that is recruited to the plasma membrane in a PIP_3-dependent fashion. PI3K-dependent activation of ARAP3 involves binding of its two most N-terminal pleckstrin homology (PH) domains to PIP_3, a lipid second messenger generated downstream of PI3K. This drives recruitment of ARAP3 to the plasma membrane to facilitate interaction with its GTPase substrates [34]. PIP_3 binding is ablated when a tandem arginine to alanine mutation is introduced at residues R302, R303 in the first PH domain of ARAP3, preventing ARAP3 activation and recruitment to the plasma membrane [29]. Arap3^{R302,303A/R302,303A} knock-in mutant mice (here referred to as KI/KI) phenocopy Arap3 null mice, suggesting an essential role for PI3K-dependent activation of ARAP3 [29].

In this study, we first investigate ARAP3 function in adult hematopoiesis using KI/KI mice, since about 2% of KI/KI mice are viable [29, 33]. We report that KI/KI HSCs are compromised in their ability to repopulate and self-renew in serial transplantation assays. To elucidate potential cell-autonomous and non-cell-autonomous roles for ARAP3 in HSC function, we selectively delete Arap3 in the hematopoietic compartment, and in endothelial and stromal cells of the HSC niche, respectively, using Cre driven by suitable promoters (Vav1, Prx1, Cdh5/VE-cadherin). Using these genetic models, we report that ARAP3 does not play a major role in regulating HSPC functions.

Results

Arap3 R302,303A mutation impairs HSC functions
To study whether ARAP3 function affects adult hematopoiesis and HSC function, we studied KI/KI mice expressing mutant ARAP3 R302,303A. This point mutation interferes with the ability of ARAP3 to bind PIP_3 and its subsequent activation by PI3K [29, 33]. Most KI/KI mutant mice die embryonically at E11 [29], but a small subset (~2%) was viable and fertile when the expected birth ratio was 25% (Table 1). By 8–12 weeks of age, these mice were indistinguishable from their littermate controls in gross appearance as well as by phenotypic characterization of their peripheral blood (Fig. 1A). KI/KI BM also showed normal progenitor cell numbers as determined by colony-forming cell (CFC) assays (Fig. 1B), and normal HSC frequencies as determined by flow cytometry using SLAM family surface markers, CD48^−CD150^+LSK (Lin^−Sca1^+cKit^+)] [35] (Fig. 1C).

To study the function of these mutant HSPCs, purified LSK cells from KI/KI mutant mice or control mice were injected with competitor bone marrow cells into each irradiated recipient mouse. Reconstitution in individual recipient mice was followed every 4 weeks post-transplant. We found that KI/KI LSKs displayed a significantly lower donor chimerism in all lineages of the peripheral blood from recipient mice (Figs. 1D and 1E). Donor-derived cells in the BM, LSK, and SLAM
LSK (CD48^+CD150^+LSK) compartments were significantly lower in mice transplanted with KI/KI cells, in comparison to control cells (Fig. 1F). Four months after the primary transplant, BM cells were harvested and $2 \times 10^6$ total BM cells were injected into secondary irradiated recipient mice. Tertiary transplants were performed similarly. Peripheral blood and BM HSC reconstitution after each transplant was analyzed by flow cytometry. We found that the defects in reconstitution of KI/KI cells were exacerbated upon serial transplantations, indicating compromised HSC self-renewal (Figs. 1G–1I). Interestingly, a myeloid bias in the multi-lineage reconstitution of serially transplanted recipients arose within KI/KI reconstituted mice (Figs. 1J and 1K), reminiscent of aged HSCs [36–38].

ARAP3 is dispensable for steady-state hematopoiesis

Our data from the knock-in mice prompted us to further elucidate the role of ARAP3 in regulating HSC function using ARAP3 conditional knockout (CKO) mice. To study if ARAP3 plays a cell-autonomous role in hematopoietic cells, we crossed Arap3^{flox/flox} (f/f) mice with mice expressing a Vav1 promoter-driven Cre. Vav1 expression begins around E11.5, and is fully turned on and expressed in greater than 99% of hematopoietic cells by E13.5, thereby excising floxed alleles in most, if not all, fetal liver and adult hematopoietic cells [39].

To measure the deletion efficiency at the DNA level, we cultured $1.5 \times 10^4$ unfractionated BM cells from either Arap3^{flox/flox};Vav1-Cre^{R26S} (f/f;Vav) CKO or f/f control mice in semi-solid methylcellulose cultures. Each progenitor cell gives rise to an individual colony, from which DNA was isolated to measure Arap3 deletion efficiency on a clonal basis. Clonal analysis by polymerase-chain reaction (PCR) genotyping showed near 100% Vav1-Cre-mediated excision of Arap3 (S1A Fig.). We also measured Arap3 deletion efficiency at the transcript level using quantitative real-time PCR (qRT-PCR). f/f;Vav showed a greater than 95% deletion of Arap3 transcripts in the BM when compared to f/f control mice (S1C Fig.). In contrast, the transcript levels of the other ARAP family members, Arap1 and Arap2, remained unchanged in the BM (S1D Fig.). To ensure a more complete deletion of ARAP3 in hematopoietic cells, we generated Arap3^{flox/};Vav1-Cre^{R26S} (f/--;Vav) mice. These mice showed 100% deletion in all f/--;Vav mice we examined at both DNA and RNA levels (Figs. 2A and 2B). We found that both

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**Table 1. KI/KI live birth rates.**

| Breeding pair | KI/KI | KI/+ | +/+ |
|---------------|-------|------|-----|
| KI/+ x KI/+   | 2/110 (1.8%) | 68/110 (61.8%) | 40/110 (36.4%) |
| Expected ratios | 25% | 50% | 25% |
| KI/+ x KI/KI | 6/36 (16.7%) | 30/36 (83.3%) | – |
| Expected ratios | 50% | 50% | – |

Breeding pair genotypes are displayed in the left column. Genotypes of expected pups are labeled at the top of each column. Birth rates of each genotype are displayed as a percentage and fraction of total pups born alive. Expected Mendelian ratios of each genotype are listed below each cross.

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Fig. 1. ARAP3 R302,303A mutation impairs HSC function. (A) Complete blood counts (CBC) of f/f control mice (white bars) and KI/KI mutant mice (dotted black bars), n=4. (B) Colony-forming cell (CFC) assays of f/f and KI/KI bone marrow cells enumerated after 11 days in culture. Colonies identified as multipotent myeloid progenitors (GEMM), erythroid (BFU-E), granulocyte-monocyte progenitors (GM), granulocyte (G), or monocyte (M). n=3. (C) The percentage of SLAM LSK population enriched for long-term hematopoietic stem cells in f/f and KI/KI bone marrow, gated on the Lin- c-Kit^+ Sca1^+ (LSK) population, followed by SLAM markers CD48^+ CD150^- (LSK). (D–K) Transplantation of LSK cells from f/f and KI/KI donor mice. Results were pooled from two separate experiments. (D) Peripheral blood in the recipients was assessed at 4, 8, and 12 weeks after primary transplant by flow cytometry for the percentage of donor-derived leukocytes. (E,H) Donor contribution to myeloid, B-cell, and T-cell compartments in the peripheral blood at the end of the primary (E) or secondary
Adult f/f;Vav and f/−;Vav mice showed normal peripheral blood composition (S2A and 2C Figs.). The lineage distribution of hematopoietic tissues as analyzed by flow cytometry in Arap3 CKO mice was comparable to that of their littermate controls (S2B and 2D Figs.). Our data suggest that ARAP3 is dispensable for steady-state hematopoiesis in the adult mouse.

ARAP3 does not play a cell-autonomous role in HSCs
We next examined if ARAP3 affects primitive hematopoietic compartments. We found that f/−;Vav mice had a normal distribution of phenotypic HSPCs within the LSK compartment, as determined by SLAM markers (Fig. 2E). Furthermore, Arap3 deficiency in BM cells did not affect hematopoietic progenitor cell proliferation or differentiation in CFC assays (Fig. 2F and S2C Fig.). Of note, when purified f/f;Vav LSKs were plated, they exhibited abilities to form various types of colonies with normal frequencies (S2D Fig.) and morphology (not shown). In contrast, Arap3−/− neutrophils from the CKO mice exhibited enhanced polyRGD-induced adhesion (S2E Fig.), as previously published [32]. These data together suggest the cell-intrinsic functions of ARAP3 are limited to more differentiated myeloid cells, rather than in the immature HSPC populations.

We next assessed whether ARAP3 plays a role in HSC function in vivo using competitive bone marrow transplantation (BMT) assays. LSK cells sorted from f/f;Vav mice or f/f littermate controls were transplanted into lethally-irradiated recipient mice and peripheral blood reconstitution was evaluated. There were no significant differences in multi-lineage reconstitution (S2H Fig.) or donor chimerism, either in the total blood cell population (S2F Fig.) or in myeloid/T-cell/B-cell lineages after primary BMT (S2G Fig.). Secondary transplants also showed normal multi-lineage reconstitution (S2I–S2K Figs.). Furthermore, serial transplantation of f/−;Vav LSKs showed no discernable differences in donor contribution (Figs. 2G and 2I) or multi-lineage reconstitution (Figs. 2H and 2J) when compared to littermate control donors. Together, our data firmly established that ARAP3 does not cell-autonomously impact HSC homeostasis or function.
Fig. 2. Hematopoietic-specific deletion of Arap3 using Vav1-Cre does not affect HSC repopulation or self-renewal. (A) Representative PCR genotyping results of individual colonies from CFC assays of f/-;Vav and f/f;Vav CKO mice. Controls are tail DNA isolated from f/f, f/- and f/-;Vav mice, showing the Arap3 floxed band (top band) and the Arap3 deleted band (bottom band). (B) Arap3 RNA transcript levels from BM of f/-, f/f;Vav, and f/-;Vav mice were assayed by qRT-PCR and first normalized to Gapdh levels. The graph shows the relative Arap3 transcripts remaining in the CKO mice compared to that in f/f controls. Each symbol
ARAP3 is dispensable in Prxl-expressing stromal and osteoblastic cells for supporting steady-state hematopoiesis

We next investigated whether ARAP3 plays a role in the HSC niche, acting non-cell-autonomously to regulate HSCs. To study this, we deleted Arap3 using Prxl-Cre, shown to induce excision in nearly all osteoblasts and 95% of perivascular stromal cells but not in endothelial cells [2, 40]. Arap3{flox/flox};Prxl-Cre{f/f};Prxl{f/f} mice were born alive in expected Mendelian ratios. Hematopoietic development of f/f;Prxl mice appeared normal as determined by CBC and lineage distribution in hematopoietic tissues, as well as by progenitor numbers and function in CFC assays (data not shown). f/f;Prxl mice also exhibited normal frequencies of phenotypic HSCs and progenitors as characterized by flow cytometric analysis (Fig. 3A). Using competitive BMT assays, we found that f/f;Prxl LSKs repopulated comparably to control f/f LSK donor mice. Data pooled from 3 independent experiments. n = 8–15. (G) Peripheral blood in the recipients was assessed at 4, 8, and 12 weeks after primary transplant by flow cytometry for the percentage of donor-derived cells. White bars represent control donors and black bars represent CKO donors. Data represent an individual mouse; horizontal lines indicate mean ± SEM levels. (C) CBC of f/f (white bars), f/− (crosshatch bars), and f/−;Vav (black bars) mice. n = 8. (D) Percentage of various cell populations in the BM and spleen of f/f, f/−, and f/−;Vav mice was analyzed by flow cytometry, as defined by the surface markers indicated. n = 5. (E) Percentage of various HSPC populations in the BM from f/f, f/−, and f/−;Vav mice was quantified using flow cytometry, defined by Lin−Sca1+c-Kit+ (LSK) and SLAM markers CD48 and CD150. n = 5. (F) CFC assays of f/f, f/−, and f/−;Vav BM cells were enumerated after 11 days in culture. n = 8. (G–J) Primary and secondary transplantation of LSK cells from control (CTL = f/f and f/−) and knockout (KO = f/f;Vav and f/−;Vav) donor mice. Data pooled from 3 independent experiments. n = 8–15. (G) Peripheral blood of recipients assessed at the end of the secondary transplant for donor chimerism by flow cytometry. (H, J) Bars show lineage distributions within donor-derived cells of individual recipient mice at the end of the primary (H) and secondary (J) transplants. T: CD3+; B: CD19+; M: Mac1+. Graphs show mean ± SEM. P-values determined by Student’s t-test.

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ARAP3 expression in endothelial cells is important for embryonic development but not adult HSC functions

Arap3 deletion in endothelial cells using Tie2-Cre resulted in embryonic lethality due to a cell-autonomous angiogenesis defect, though only with more robust excision on the f/f−, but not f/f, background [29]. Since Tie2-Cre also shows some expression in stromal cells [41, 42], we utilized the vascular endothelial cadherin (VEC or Cdh5) promoter-driven Cre (VEC-Cre) that is found to be more endothelial cell-specific than Tie2-Cre [43, 44]. Arap3{flox/flox};VEC-Cre{f/f};VEC{f/f} mice were born at expected Mendelian ratios (Table 2).

To obtain a more complete deletion of Arap3 in endothelial cells, we generated Arap3{flox/flox};VEC-Cre{f/f−};VEC{f/f−} mice. These CKO mice were born at a significantly reduced ratio (Table 2). The surviving CKO mice appeared grossly normal, and showed an approximate 88% excision efficiency (Fig. 4A), ranging from 80% to 96%, compared to an average 80% excision rate in f/f;VEC mice (S1B Fig.). By qRT-PCR, an approximate 9% Arap3 transcripts remained in f/f−;VEC BM (Fig. 4B), in contrast to 15% in f/f;VEC BM (S1C Fig.). In agreement with
previously published data, this indicates that ARAP3 function in endothelial cells is essential for embryonic development.

Both \( f/f;VEC \) and surviving \( f/-;VEC \) adult mice had similar peripheral blood composition to their control littermates (S3A Fig. and Fig. 4C) and normal lineage distribution in hematopoietic tissues (S3B Fig. and Fig. 4D). These mice also exhibited a normal distribution of immature HSPC populations, with expected numbers of CD48\(^-\)CD150\(^+\)LSK cells (Fig. 4E). Hematopoietic progenitor numbers were normal as determined by CFC assays (Fig. 4F and S3C Fig.). We next examined HSC functions in vivo using competitive BMT assays. \( f/f;VEC \) LSKs showed comparable multi-lineage reconstitution to that of \( f/f \) controls in the primary (S3D–S3F Figs.) and secondary BMTs (S3G–S3I Figs.). Similarly, \( f/-;VEC \) LSK cells showed largely normal long-term reconstitution upon primary (Figs. 4G–4H) and secondary transplantation (Figs. 4I–4J) compared to \( f/- \) LSK cells.

Fig. 3. ARAP3 is dispensable in Prx1-expressing bone marrow niche cells for steady-state hematopoiesis. (A) BM cells from \( f/f \) control and \( f/-;Prx1 \) KO mice were quantified for the percentage of cells in various HSPC populations. Bars show mean ± SEM, \( n=3 \). (B) LSK cells from \( f/f \) and \( f/-;Prx1 \) donor mice were transplanted into irradiated recipient mice. Peripheral blood of individually reconstituted mice was assessed by flow cytometry at 12 weeks post-transplant for the distribution of cell lineages within donor-derived cells (bars, left Y-axis) and the percentage of total donor leukocytes (black diamonds, right Y-axis). P-values determined by Student’s t-test.

Table 2. \( f/f;VEC \) and \( f/-;VEC \) live birth rates.

|                  | \( f/f;VEC \) | \( f/-;VEC \) | \( f/f \) | \( f/- \) |
|------------------|-------------|-------------|-----------|-----------|
| \( f/f \times f/-;VEC \) | 30/142 (21.1%) | 37/142 (26.1%) | 36/142 (25.3%) | 39/142 (27.5%) |
| Expected ratios  | 25%         | 25%         | 25%       | 25%       |

|                  | \( f/-;VEC \) | \( f/f;VEC \) | \( f/- \) | \( f/f \) |
|------------------|-------------|-------------|-----------|-----------|
| \( f/- \times f/f;VEC \) | 14/136 (10.3%) | 30/136 (22.1%) | 45/136 (33.1%) | 47/136 (34.6%) |
| Expected ratios  | 25%         | 25%         | 25%       | 25%       |

Breeding pairs are labeled in the left column. Genotypes of expected pups are labeled at the top of each column. Birth rates are displayed as a percentage and fraction of total pups born alive. Expected Mendelian ratios are listed below each cross breeding.

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Fig. 4. Arap3^floxed^;VEC-Cre mice show largely normal HSC homeostasis and function. (A) Representative PCR genotyping results of individual colonies from CFC assays of f^−^;VEC CKO BM cells are shown. Controls are tail DNA isolated from f/f and f^−^ control mice as well as f^−^;VEC CKO mice, showing the Arap3 floxed band (larger band) and the Arap3 deleted band (smaller band). (B) Arap3 RNA transcript levels from BM of f^−^ and f^−^;VEC mice were assayed by qRT-PCR. The graph shows the relative Arap3 transcript level to that in f/f controls. Each symbol represents an individual mouse; horizontal lines indicate mean ± SEM levels. (C) CBC of f/f (white bars) and f^−^ (crosshatch bars) control mice and f^−^;VEC (gray bars) CKO mice. n=5. (D) Percentage of various cell populations in the BM and spleen of f/f, f^−^, and f^−^;VEC mice was analyzed by flow cytometry. n=5. (E) Quantification of the percentage of various HSPC populations in the BM from f/f, f^−^, and f^−^;VEC mice using flow cytometry. n=5. (F) CFC assays of f/f, f^−^, and f^−^;VEC BM cells
To assess whether ARAP3 expression in BM endothelial cells is important to support HSC functions, we performed reverse transplantation of wild-type bone marrow cells into f/f, f/−, and f/−;VEC donor mice. Data pooled from 4 independent experiments. n=18–24. (G,l) Peripheral blood in the recipients was assessed every 4 weeks post-primary transplant (G) and at the end of the secondary (l) transplant for the percentage of donor-derived cells. (H,j) Multi-lineage reconstitution was analyzed by flow cytometry. Bars show lineage distributions within donor-derived cells of individual recipient mice at the end of the primary (H) and secondary (J) transplantsations. (K) ARAP3 expression in BM endothelial cells is not required to support HSC homeostasis. Reverse transplantation of wild-type CD45.1+ bone marrow cells into f/f and f/− control (SJL:CTL) or f/f;VEC and f/−;VEC CKO (SJL:VEC-CKO) mice were performed. Percentage of CD45.1+ LSK and SLAM LSK cells in the recipient bone marrow was quantified by flow cytometry 8 weeks post-transplantation. n=9–15 pooled from two independent experiments. Graphs show mean ± SEM. P-values determined by Student’s t-test.

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To assess whether ARAP3 expression in BM endothelial cells is important to support HSC functions, we performed reverse transplantation of wild-type bone marrow cells into f/f and f/− control (SJL:CTL) or f/f;VEC and f/−;VEC CKO (SJL:VEC-CKO) recipient mice. HSC homeostasis or engraftment was not altered, and multi-lineage reconstitution of the hematopoietic compartment was unaffected by Arap3 deletion in the host endothelial HSC niche (Fig. 4K and data not shown). Taken together, our investigation suggests that in spite of a critical role for ARAP3 during embryonic vascular development [29, 30], loss of ARAP3 in endothelial cells minimally impacts HSCs in the adult BM.

Discussion

In the present study, we utilize KI/KI mice and generate three Arap3 CKO mouse models to study ARAP3 functions in hematopoiesis of the adult mouse. We show that KI/KI mice exhibit defective HSC functions upon transplantation, indicating that PI3K-mediated ARAP3 function is important for HSCs. To elucidate the potential cell-autonomous and non-cell-autonomous roles of ARAP3 in HSCs, we conditionally deleted Arap3 in hematopoietic cells and in several cell types in the HSC niche. Arap3 null HSCs from f/−;Vav mice were functionally competent to repopulate the HSC pool and self-renew upon serial transplantation, revealing ARAP3 is not required in regulating HSC homeostasis or function in a cell-autonomous manner. Furthermore, ablation of ARAP3 in perivascular stromal cells, osteoblastic cells, and endothelial cells of the HSC niche did not alter HSC function or maintenance in f/−;Prx1 and f/−;VEC mice. Reverse transplantation experiments strongly suggest that ARAP3 expression in the BM endothelial niche is not required to support HSC functions; however, future investigation is needed to assess the potential role of ARAP3 in BM mesenchymal and osteoblastic niches. Our study demonstrates that f/−;VEC mice displayed partial embryonic lethality, indicating a critical role for ARAP3 in endothelial cells during embryonic development but not in the adult BM niche for HSCs.

ARAP3 was first purified from porcine leukocytes for its PIP₃ binding ability and is highly expressed in hematopoietic tissues, however our studies show that ARAP3 is not a critical cell-intrinsic regulator of hematopoiesis. ARAP3 does not play a cell-autonomous role in regulating HSC homeostasis or function. However,
our observation does not rule out a cell-autonomous role for ARAP3 under stress conditions. This phenomenon has been seen in genetic studies of other proteins, such as SIRT1 [45, 46]. It is possible that a physiological challenge to the mice is necessary to elicit a specific function for ARAP3 in hematopoiesis. In our model, Arap3 is excised in hematopoietic cells during early development using Vav1-Cre, and compensatory mechanisms may account for normal hematopoiesis in adult mice. It would be interesting to study whether acute loss of ARAP3 in hematopoietic cells of the adult mouse, such as with an inducible Cre system [47], would reveal a role for ARAP3 in hematopoiesis. Additionally, ARAP3 is part of a dual-GAP family that also includes ARAP1 and ARAP2. Although each family member targets different G-protein substrates and has different active functional domains [48–50], these three proteins may have overlapping and redundant roles in hematopoiesis, and may work in conjunction to regulate HSPC function. Thus, genetic studies of mice deficient in Arap1 or Arap2, or with combination deletions of multiple ARAP proteins would clarify whether there is a significant role for the ARAP family of proteins in hematopoiesis.

We find that deletion of Arap3 in stromal and osteoblastic cells using Prx1-Cre does not affect steady-state hematopoiesis or HSC homeostasis. Furthermore, deletion of Arap3 in endothelial cells using VEC-Cre does not impact its ability to support wild-type HSCs. However, high efficiency of Arap3 deletion in f/−;VEC mice results in embryonic lethality, suggesting ARAP3 plays an important role in developing endothelial cells, in agreement with previously published data [29, 30]. Since ARAP3 deficiency in embryonic endothelial cells disrupts angiogenic sprouting and vascular structure, this developmental-related dysregulation may indirectly affect HSC emergence, development, and maintenance in the adult mouse. However, our studies do not exclude the possibility that compensatory mechanisms may be upregulated in surviving f/−;VEC mice sometime between HSC emergence and adulthood. One approach to answer this question would be to induce the excision of Arap3 in endothelial cells when ARAP3 is no longer required for vasculature development. Future investigations are warranted to examine the emergence of definitive hematopoietic progenitors and HSCs from the endothelium in the AGM region of Arap3 null embryos and further our understanding of the role of ARAP3 in endothelial cells.

Our results demonstrate that the ARAP3 R302,303A mutation (KI/KI) that disrupts ARAP3 recruitment to the plasma membrane or activation by PI3K markedly impairs HSC function, while loss of ARAP3 does not. This could be due to compensatory mechanisms or changes in gene expression and signaling pathways of rare surviving KI/KI mice. Another possibility for compromised HSCs in KI/KI mice might be that the ARAP3 R302,303A mutant acts in a dominant negative manner to prevent translocation of other interacting players [51–55] to the plasma membrane, which may affect HSC function. For example, ARAP3 can bind the phosphatase SHIP2, a negative regulator of PI3K signaling [53, 54, 56], as well as CIN85 and Odin, both shown to be involved in receptor endocytosis and motility [51, 55, 57–59]. Although ARAP3 has not been shown to act as a scaffold for the plasma membrane translocation of any interacting
proteins, and the in vivo relevance of these interactions remains to be tested, the fact that we see a cell-intrinsic defect with KI/KI HSCs but not with f/−;Vav HSCs in the BMT assays speaks in favor of this possibility. Lastly, it is possible that ARAP3 plays an important role in a cell type other than hematopoietic, endothelial, stromal, or osteoblastic cells to impact HSC function. One way of determining whether the phenotype of KI/KI HSCs is due to a cell-autonomous effect of the R302,303A mutant in HSCs would be to generate a Vav1-Cre-driven conditional KI/KI mouse to genetically study this question.

RhoA has been characterized in vivo to regulate migration and chemotaxis of mature hematopoietic cells [27,28], as well as HSPC engraftment, multi-lineage repopulation and cell survival [9,14,15]. ARAP3 is not the only GTPase-activating protein that targets RhoA, and our data suggest it is not a major regulator of RhoA activation in HSPCs. Other GAPs, such as p190B RhoGAP, play important roles in mediating HSPC function through its inactivating activity on RhoA [20,21]. Due to the large number of regulators for RhoA, it is likely that each acts in its own individual temporal- and spatial-specific manner. The possibility of redundancies between the various GAPs also exists [60], such that changing the dynamic by ablating one GAP is not enough to alter the process of normal hematopoiesis. It would be interesting to investigate whether deletion of multiple GAPs in hematopoietic cells would result in greater deficiencies than migration or engraftment alone.

As a dual GTPase-activating protein, ARAP3 targets Arf6 as well as RhoA. Arf6 has mostly been studied in non-hematopoietic cells with regard to its role in membrane trafficking and the cell actin cytoskeleton [61–63]. Like RhoA, it is actively involved in cell migration, adhesion, proliferation and cytokinesis [64–66]. However, the potential role of Arf6 in HSPCs and hematopoiesis has not been well established. One study showed that the decrease of active Arf6-GTP in platelets is critical to the activation of Rho GTPases that is necessary for cytoskeletal rearrangements preceding full platelet function [67]. It is important to further investigate and understand the role of Arf6 in hematopoiesis, particularly in HSCs.

ARAP3 has been implicated in the regulation and progression of several human diseases, including defense against bacterial infection, diabetes and gastric carcinoma, by capitalizing on the ability of ARAP3 to manipulate vesicle internalization and cell invasion [68–70]. Dysregulation of Rho family GTPases and their regulators have also been correlated with human blood disorders and tumorigenesis [71–75]. While aberrant expression of ARAP3 has not yet been found in blood disorders, its ability to regulate the actin cytoskeleton makes it a potential target for the dysregulation of homeostatic cell functions. Thus, continued study of ARAP3 in normal and abnormal hematopoiesis will be important to elucidate a more comprehensive understanding of its role in the blood system.
Methods

Generation of Arap3 transgenic mice

Arap3\textsuperscript{flox/flox} and Arap3\textsuperscript{KI/KI} mice were generated as described previously \[29\]. Vav1-Cre mice were originally generated by Dr. Thomas Graf \[39\] and backcrossed to C57Bl/6J background for 8 generations. VEC-Cre mice were kindly provided by Dr. Nancy Speck \[44\] and backcrossed to C57Bl/6J background for 8 generations. These two strains of mice were crossed to generate Arap3\textsuperscript{flox/+};Cre\textsuperscript{g} mice, which were then crossed to Arap3\textsuperscript{flox/} mice to generate Arap3\textsuperscript{flox/+};Cre\textsuperscript{g} conditional knockout mice. Initial studies presented in S1–S3 Figs of \textsuperscript{f/f};Vav and \textsuperscript{f/f};VEC mice were done in mixed Bl6/129 background, while studies in main figures were later performed on a pure C57Bl/6J background following backcrossing for 8 generations. Arap3\textsuperscript{flox/+} mice on the pure C57Bl/6J background were crossed with CMV-Cre mice on the C57Bl/6J background (The Jackson Laboratory) to generate Arap3\textsuperscript{flox/+} mice. These mice were used to generate Arap3\textsuperscript{flox/+};Vav1-Cre\textsuperscript{g} mice, Arap3\textsuperscript{flox/+};VEC-Cre\textsuperscript{g} and Arap3\textsuperscript{flox/+};Prx1-Cre\textsuperscript{g} conditional knockout mice (all on a pure Bl6 background) that will ensure a more complete deletion efficiency. Prx1-Cre mice on a C57Bl/6J background were purchased from The Jackson Laboratory.

The animal studies were carried out in strict accordance with the recommendations in the Guide for the Care and Use of Laboratory Animals of the National Institutes of Health. The protocol was approved by the Institutional Animal Care and Use Committee of the Children’s Hospital of Philadelphia.

PCR genotyping and qRT-PCR

For genotyping and clonal PCR analysis, primers were generated to detect wildtype, floxed and deleted alleles of Arap3 using DNA isolated from CFU-C assays or mouse tissues. PCR reactions were performed on a BioRad thermal cycler using the following primers for Arap3: 5’-AGAGGCTCAGGACTAGAAGGACTA-3’ (Arap3_461F) and 5’-GGGCTGAGTAGAGACTGACGCGCC-3’ (Arap3_EcorV_F) and 5’-GAGGCCAGCCTGAGATAGATGAAACCC-3’ (Arap3_EcorV_R).

For quantitative real-time PCR, total RNA was isolated from FACS-sorted bone marrow cells, CFC assays or hematopoietic tissues using Trizol Reagent (Invitrogen Life Technologies) followed by isolation with the RNeasy Mini kit (Qiagen). cDNAs were produced using BioRad iScript kit and qRT-PCR reactions were performed on an Applied Biosystems 7900HT real-time PCR system using Sybr-Green detection with the following primers: Arap3: 5’-CCCTCTGACTCAGACTAGAAGGACTA-3’ and 5’-ATTCCAGGTATTACCGGCCAGC-3’; Arap1: 5’-GATGCCGCACTGTCTGTAGCT-3’ and 5’-CTGCTCAAGAGTGCAGGCAC-3’; Arap2: 5’-CAGGACGAGATGAGTTATTAG-3’ and 5’-TCTCGCCCTGAACTGAAAGA-3’; Gapdh: 5’-GGAGCGAGACCCACTAACA-3’ and 5’-TTCACACCCATCACAAACAT-3’. The transcript levels in CKO mice were first
normalized to Gapdh levels, then expressed as a percentage of normalized levels in control f/f mice.

**Complete blood counts**
Peripheral blood was collected by retro-orbital bleeding into capillary blood collection tubes with EDTA (BD). CBC analysis was performed using the mouse setting on a HemaVet 950 machine (Drew Scientific).

**Cell sorting and flow cytometry**
Cells from either peripheral blood or hematopoietic tissues were lysed of red blood cells, then stained with surface markers on ice and washed in PBS with 2% bovine calf serum. Surface markers used to identify cell populations are: CD3e-PE for T-cells, CD19-APC for B-cells, Gr1-PE and Mac1-APC for myeloid cells (eBiosciences), and propidium iodide for viability. This method was also used for peripheral blood analysis of transplanted mice with the addition of CD45.1-PE-Cy7 and CD45.2-FITC antibodies. Flow cytometry was performed on a FACS Canto analyzer (BD).

For HSPC analysis, total bone marrow cells were flushed from femurs, tibias and iliac crests of mice. Single-cell suspensions were lysed of red blood cells and stained with the following primary antibodies: biotinylated lineage cocktail (B220, CD4, CD5, CD8, CD19, IL-7R, Gr1, Mac1, Ter119), Sca1-PerCP-Cy5.5, cKit-APC-Cy7, CD48-FITC, and CD150-PE-Cy7 (eBiosciences). Cells were then washed in PBS with 2% bovine calf serum and stained with streptavidin-PE-TexasRed secondary antibodies. DAPI was used for viability. Immunophenotypes were defined by signaling lymphocytic activation molecule (SLAM) family markers [35] CD48^+^CD150^+^LSK (SLAM LSK) as a population enriched for long-term HSCs. CD48^−^CD150^−^LSK and CD48^−^CD150^−^LSK populations are enriched for HPCs. Flow cytometry was performed on a LSR Fortessa analyzer (BD).

For cell sorting, bone marrow cells were first depleted of lineage-positive cells using biotinylated lineage cocktail and streptavidin-coupled Dynabeads (Invitrogen) per manufacturer’s protocol. Lineage-negative (Lin^-_)_ cells were then stained for LSKs as described above and sorted on a FACS Aria Cell Sorter (BD). All analysis of FACS data was performed using FlowJo software (TreeStar).

**Colony-forming cell (CFC) assays**
Unfractionated bone marrow cells or sorted LSK cells were plated at a concentration of 15,000 or 200 cells per plate, respectively, in duplicate using semisolid methylcellulose (Methocult M3434, StemCell Technologies) containing SCF, IL-3, IL-6 and erythropoietin. Cells were incubated at 37°C, 5% CO₂ with high humidity, and colonies were enumerated after 10–12 days in culture.
Competitive bone marrow transplantation (BMT), serial BMTs, and reverse BMTs

Bone marrow cells (CD45.2+) were harvested and sorted for LSK surface markers as described above. 500–1000 LSK cells were mixed with 350,000 competitor bone marrow cells (CD45.1+) and injected into each lethally-irradiated (a split dose of 10Gy, 137Cs source) F1 recipient mouse (CD45.1+CD45.2+). Donor-derived reconstitution in the periphery was measured by flow cytometry every 4 weeks post-transplant. At 12–16 weeks, recipient mice were sacrificed and analyzed for donor HSPCs, as described above with the addition of CD45.1-PE-Cy7 and CD45.2-FITC antibodies. For secondary transplantation, $2 \times 10^6$ unfractionated bone marrow cells from pooled primary recipient mice were transplanted into lethally irradiated F1 recipients. Reconstitution was again measured every 4 weeks post-transplant and data of secondary transplant endpoints are shown where mentioned. Tertiary transplants were performed similarly. In reverse BMTs, $1 \times 10^6$ total bone marrow cells from CD45.1+ wild-type donor SJL mice were transplanted into lethally irradiated f/f and f−/− control or f/f;VEC and f−/−;VEC (CD45.2+) CKO recipients. Reconstitution and the percentage of CD45.1+ LSK and SLAM LSK cells were measured 8 weeks after transplantation.

Supporting Information

S1 Fig. Deletion efficiency of Arap3flox/flox;Vav-Cre and Arap3flox/flox;VEC-Cre mice. (A,B) Genotyping of individual colonies from CFC assays of f/f;Vav and f/f;VEC BMs. Representative PCR genotyping results from f/f;Vav (A) and f/f;VEC (B) colonies are shown. Controls are tail DNA isolated from control and CKO mice. The top band is floxed Arap3 while the bottom band reflects deleted Arap3. (C) Arap3 RNA transcript levels from f/f;Vav and f/f;VEC mice were assayed by qRT-PCR. The graph shows relative Arap3 transcript levels remaining in the CKO mice compared to that in f/f controls. (D) Relative Arap1 transcript levels (left) and Arap2 transcript levels (right) to that in f/f mice. Each symbol represents an individual mouse; horizontal lines indicate mean ± SEM levels. doi:10.1371/journal.pone.0116107.s001 (TIF)

S2 Fig. Arap3flox/flox;Vav1-Cre mice maintain steady-state hematopoiesis and normal HSC function. (A) CBC of f/f control (white bars) and f/f;Vav CKO mice (black bars). n=17. (B) Flow cytometric analysis of the percentage of various cell populations in the BM, spleen, and thymus of f/f and f/f;Vav mice. n=17. (C,D) CFC assays of f/f and f/f;Vav BM cells (C) and LSK cells (D) enumerated after 11 days in culture. n=17 and 3, respectively. (E) Poly-RGD induced adherence of bone marrow-derived neutrophils from f/f and f/f;Vav mice was quantified in triplicate as the number of cells per field of view at 20 × magnification. (F–K) Serial transplantation of LSK cells from f/f and f/f;Vav donor mice. Results were pooled from three separate experiments. (F,I) Peripheral blood of recipient mice assessed at 4, 8, and 12 weeks after primary (F) or secondary (I) transplants for the percentage of donor-derived leukocytes. (G,J) Donor contribution to myeloid, B-
cell, and T-cell compartments in the peripheral blood at the end of the primary (G) or secondary (J) transplants was analyzed by flow cytometry. (H,K) Bars show lineage distributions within donor-derived cells of individual recipient mice (left Y-axis), while diamond symbols indicate total donor leukocyte percentages (right Y-axis) in the peripheral blood, at the end of the primary (H) or secondary (K) transplants. T: CD3⁺; B: CD19⁺; M: Mac1⁺. Graphs show mean ± SEM. P-values determined by two-tailed Student’s t-test.
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S3 Fig. Arap3flox/flox;VEC-Cre mice show normal hematopoiesis and HSC functions. (A) CBC of f/f control (white bars) and f/f;VEC CKO mice (gray bars). n=12. (B) Percentage of various cell populations in the BM, spleen, and thymus of f/f and f/f;VEC mice. n=12. (C) CFC assays of f/f and f/f;VEC BM cells enumerated after 11 days in culture. n=12. (D–I) Serial transplantation of LSK cells from f/f and f/f;VEC donor mice. (D,G) Peripheral blood of recipient mice assessed every 4 weeks during the primary (D) or secondary (G) transplants for the percentage of donor-derived cells. (E,H) Donor contribution to myeloid, B-cell, and T-cell compartments in the peripheral blood after the primary (E) or secondary (H) transplants. (F,I) Bars show lineage distributions within donor-derived cells of individual recipient mice (left Y-axis), while diamond symbols indicate total donor leukocyte percentages (right Y-axis) in the peripheral blood, at the end of the primary (F) or secondary (I) transplants. Graphs show mean ± SEM. P-values determined by two-tailed Student’s t-test.
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Author Contributions
None. Conceived and designed the experiments: WT YS. Performed the experiments: YS JJ. Analyzed the data: WT YS. Contributed reagents/materials/analysis tools: SV. Wrote the paper: WT YS SV.

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