Pak2 restraints endomitosis during megakaryopoiesis and alters cytoskeleton organization

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Key Points

- Bone marrow-specific deletion of Pak2 is associated with macrothrombocytopenia and abnormal megakaryocyte morphology and function.
- Pak2 deletion is associated with defects in megakaryocyte endomitosis and the activation of Aurora-A and LIM kinase.

Introduction

Megakaryocytes are both the largest (50-100 μm) and rarest (~0.03%-0.06%) cell type in the bone marrow.1,2 To produce sufficient numbers of platelets, these cells become polyplloid and undergo massive nuclear proliferation, together with an enlargement of the megakaryocyte cytoplasm, which becomes filled with platelet-specific granules. Megakaryocytes undergo a complex maturation process by which their cytoplasmic contents are packaged into multiple elongated proplatelet processes.3 As proplatelets extend through the sinusoid vessels, physiological shear force aids in dissociating proplatelets into circulating platelets.4,5 Megakaryocyte maturation is triggered primarily through cellular signaling events initiated by the cytokine thrombopoietin (TPO).5 TPO binds to the c-Mpl receptor on megakaryocytes to activate the Janus kinase-2 signaling pathway and stimulates the phosphatidyl inositol 3 kinase (PI3K)/Akt and mitogen-activated protein kinase (MAPK) pathways.6,7

The p21-activated kinases (Paks) are serine/threonine kinases that support cell homeostasis, contractility, and survival processes.8 After their activation by the p61 GTPases, Rac1 and Cdc42, Paks phosphorylate dozens of effector proteins to regulate the mitogen-activated protein kinase signaling and cytoskeletal remodeling.9,10,11 Accordingly, the Paks regulate numerous hematopoietic processes, including hematopoietic stem cell engraftment and homing to the bone marrow, assembly of the actin cytoskeleton, and chemotaxis.12

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Materials and methods

Generation of mice with Pak2<sup>−/−</sup> bone marrow

Pak2-deficient mice were generated by crossing Pak2-Boxed mice with mice carrying the Mx1-cre transgene (M.R. and J.C., manuscript in preparation).<sup>13,31,32</sup> In 3-month-old mice, wild-type and Mx1-cre<sup> Pak2<sup>−/−</sup></sup> mice were injected with 400 µg polynosinic-polycytidylic acid (p(3)pC) every other day. Fourteen days after the first injection, mice were euthanized for experiments.

Administration of drug treatment

Animals received vehicle or Frax1036 (30 mg/kg) via daily oral gavage, diluted in 20% (2-hydroxypropyl)-β-cyclodextrin (Sigma-Aldrich) for 3 weeks, according to a previously published protocol.<sup>33</sup> Severe combined immunodeficiency animals were used for drug treatment, as C57BL/6 animals were hypersensitive to drug treatment. All animal studies were performed according to protocols approved by the Fox Chase Cancer Center Institutional Animal Care and Use Committee.

Antibodies and reagents

All reagents were purchased from Sigma-Aldrich unless otherwise stated. Antibodies for western blotting (WB) against pLIMK (T508), pCofilin (S3), and pAurora (T288/T232/T198) were from Cell Signaling. Anti-Aurora A was from BD Biosciences. WB technique on day 5 megakaryocytes was carried out as previously described by our laboratory.<sup>13</sup> Recombinant mouse TPO was from Shennandoah Biotechnology. Interleukin 3 (IL-3) and IL-6 were purchased from PeproTech. Antibodies for fluorescence-activated cell sorting were from Ebioscience (antibody name [clone]) murine hematopoietic lineage eFluor 450 cocktail, Cat-APC (ZB8), sca-1-phycoerythrin (PE)-Cy7 (D7), CD150-FTC (BioLegend-TC15-12F12.2), CD41-APC-Cy7 and CD41-eFluor 450 (MWReg30), CD105-PE (M7/18), and FcγRII/III-PerCP-eFluor 710(clone 93). Antibodies for immunofluorescence (CD41 and β1-tubulin) were from Abcam. Frax1036 was kindly supplied by Genentech.

Analysis of platelet clearance and production

To determine platelet clearance, an in vivo biotinylation approach was used. Brieﬂy, 10 days after pPcC, mice were IV injected with sulfo-N-hydroxysulfosuccinimidobiotin (Pierce Chemical). After blood collection daily with retroorbital bleeds, the blood was diluted in phosphate-buffered saline and incubated with streptavidin-PE (BD Biosciences) to label biotinylated platelets. Thiazole orange (10 µg/mL) was then added to measure reticulated platelets, after which samples were ﬁxed and stained for acetylcholinesterase according to the manufacturer’s protocol.

Bone marrow collection and MegaCult-C assays

Bone marrow cell suspension was isolated from the tibias and femurs of pIpC-/ bone marrow, with deletion of Pak2<sup>−/−</sup> mice were injected with 400 µg polynosinic-polycytidylic acid (p(3)pC) every other day. Fourteen days after the first injection, mice were euthanized for experiments.

Measurement of hematological parameters

Mice were euthanized with CO<sub>2</sub>, and blood was extracted via the hepatic portal vein with a syringe containing acid-citrate dextrose (10% final) and added into EDTA tubes. Counts were performed using a VetScan HM5 (Abaxis).

In vitro culture of bone marrow-derived megakaryocytes

In vitro assays used CAGG-Cre-ERT2<sup>Pak2<sup>−/−</sup></sup> bone marrow, with deletion of Pak2 induced with 500 nM 4-hydroxystilbamidine. Frax1036 treatment of cultured megakaryocytes was performed at a concentration of 1 µM. Bone marrow-megakaryocytes were cultured for 5 days in Dulbecco’s modified Eagle medium, 10% fetal bovine serum, penicillin/streptomycin, and 75 ng/mL TPO and fractionated on a 1.5%/3% discontinuous bovine serum albumin gradient.

Megakaryocyte ploidy analysis

For ploidy measurements, cells were analyzed as described previously.<sup>35</sup> DNA content in CD41<sup>+</sup> megakaryocytes was determined by labeling red blood cells with anti-CD41-efuor450, followed by ﬁxation with 0.5% formalin. Cells were permeabilized with 70% ice-cold methanol and washed and incubated with 10 µg RNase A, followed by brief incubation with 1 µg/mL propidium iodide. Cells were analyzed on an LSR-II.

Proplatelet analysis

Bone marrow-derived megakaryocytes on day 5 were added to fibrinogen-coated plates and imaged at 5 and 24 hours. The amount of proplatelet extensions, relative to total megakaryocytes, was enumerated by light microscopy (×20 objective, EVOS, Invitrogen). At least 10 images per genotype (n ≥ 5 genotype) were used to evaluate the number of proplatelet-forming megakaryocytes.

Analysis of megakaryocyte progenitor cells

To analyze megakaryocyte stem cells, the protocol from Pronk et al was followed.<sup>36</sup> Bi_potential megakaryocyte/erythroid progenitor (PreMegE) cells were gated as lin<sup>−</sup> sca-1<sup>−</sup> kit<sup>−</sup> CD41<sup>−</sup> FcyRII/III<sup>−</sup> CD105<sup>−</sup> CD41<sup>−</sup> FcγRII/III<sup>−</sup> PE-Cy7<sup>−</sup> CD150<sup>−</sup> and megakaryocyte progenitors were gated as lin<sup>−</sup> sca-1<sup>−</sup> kit<sup>−</sup> CD105<sup>−</sup> CD41<sup>−</sup> PE-Cy7<sup>−</sup> CD150<sup>−</sup>. Megakaryocytes were analyzed by anti-CD41-APC-Cy7 and calculated as percentage of total bone marrow. Flow cytometric data collection was performed on an LSRII and analyzed with FlowJo software (TreeStar).

Immunofluorescence microscopy

In vitro bone marrow-derived megakaryocytes adhered to fibrinogen-coated slides (200 µg/mL) for 5 hours in the presence of TPO. Cells were ﬁxed with 2% paraformaldehyde and centrifuged onto coated slides for 5 minutes at 1000 g. Cells were permeabilized with 0.5% triton X-100 for 5 minutes before blocking for 1 hour and antibody incubation overnight. Slides were counterstained with 4′,6 diamidino-2-phenylindole (DAPI) and phalloidin and were mounted in ProLong gold before visualization. Confocal microscopy was done on an inverted Leica SP8 3 channel confocal system at ×63 magniﬁcation. Image analysis was performed with Fiji software (National Institutes of Health).

Histology

Five-µm sections of paraffin-embedded sternum and spleen were stained with hematoxylin and eosin and analyzed for megakaryocytes using an Olympus BX53 microscope (×40 objective).

Transmission electron microscopy

Bone marrow cores from mice 14 days post-pPcC were ﬁxed with 2.5% glutaraldehyde, 2.0% paraformaldehyde in 0.1 M sodium cacodylate buffer at pH 7.4 for 24 hours at 4°C. After subsequent buffer washes, the samples were postﬁxed in 2.0% osmium tetroxide for 1 hour at room temperature, washed, and placed in dH<sub>2</sub>O. After dehydration through a graded ethanol series, bone marrow was infiltrated and embedded in EMbed-812 (Electron Microscopy Sciences). Thin sections were stained with uranyl acetate and lead citrate and examined with a JEOL 1010 electron microscope ﬁtted with a Hamamatsu digital camera and AMT Advantage image capture software.

Statistical analysis

The data presented in this report are the result of at least 3 independent experiments with individual mice per genotype. Statistical analysis was performed using a 2-tailed Student t test, and a P value <.05 was considered statistically significant.
Results

Inducible deletion of Pak2 results in macrothrombocytopenia in adult mice

To study the function of the Pak2 gene in various hematopoietic-derived cells, mice carrying conditional Pak2<sup>fl/fl</sup> alleles were crossed with mice harboring the interferon-inducible Mx1-Cre transgene to produce Mx<sup>1</sup>PK2<sup>2/2</sup> (Pak2<sup>−/−</sup>) and Mx<sup>1</sup>PK2<sup>1/1</sup> controls (wild-type) (M.R. and J.C., manuscript in preparation).31 Deletion of the floxed exon 2, which encodes the start site of Pak2, was achieved by intraperitoneal administration of pIpC. Neither Pak2 protein levels in the bone marrow nor megakaryocytes were detectable 14 days post-pIpC (DPI) (Figure 1A). Deletion in megakaryocytes of Pak2<sup>−/−</sup> mice was confirmed by culturing bone marrow extracted at 14 DPI with TPO for 5 days.

Complete blood count analysis of Pak2<sup>−/−</sup> and wild-type mice displayed moderate thrombocytopenia (wild-type, 997 ± 3103/μL [698 ± 3103], n = 29; Pak2<sup>−/−</sup>, 498 ± 3103/μL [25 ± 3103], n = 52; Table 1). Platelet size was significantly increased, as determined by mean platelet volume (MPV; wild-type, 6.56 ± 0.05 fL vs Pak2<sup>−/−</sup> 7.1 ± 0.07 fL, P < .001). Additional changes in blood counts included
increased neutrophils and monocytes and decreased lymphocytes (Table 1). Deletion of Pak1 did not affect peripheral blood indices, including no changes in platelet counts or megakaryopoeisis.37 (J.C.K., unpublished data) We also tested the effects of Frax1036, a specific small-molecule inhibitor of group I Paks.33 Unlike the genetic deletion of Pak2, mice treated daily with Frax1036 for 3 weeks did not develop thrombocytopenia, nor other changes in peripheral blood (data not shown). Assuming effective inhibition of Pak2 kinase activity, these results suggest that Pak2, similar to Pak1, may have kinase-independent functions that affect certain aspects of hematopoiesis in vivo.38-40

Pak2 regulates megakaryocyte maturation

Mice with megakaryocyte-specific deletion of Pak-activating proteins, Rac1/Cdc42, are macrothrombocytopenic, with ultrastructurally abnormal megakaryocytes that fail to form proplatelets with defects in microtubule stabilization.25 As Paks are downstream effectors of these GTPases, we sought to determine whether Pak activation is required for megakaryocyte development and function, as well as to maintain platelet counts.23-25 As Pak1/2 mice do not have a notable platelet or megakaryocyte phenotype,37 we examined the effects of genetic deletion of Pak2 on megakaryocyte differentiation. We evaluated bone marrow for megakaryocyte number and morphology. Bone marrow from Pak2−/− mice had an increased number of mature megakaryocytes (CD41) with increased diameter (Figure 1B-D).

We next sought to determine whether Pak kinase activity changed during stem cell maturation into megakaryocytes. We isolated CD34+ stem cells and cultured them with TPO for 5 days. Sample lysates were collected and analyzed for activated Pak1/2/3, total Pak2, and CD41 protein expression (Figure 1E). Our results demonstrated that Pak1/2/3 activation increased during the maturation process, until decreasing at Table 1. Complete blood count profile for Pak2-deficient mice

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<th>Mx1creTgPak2fl/fl</th>
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<td>7.6 ± 0.4</td>
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<td>2.3 ± 0.13*</td>
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<td>5.4 ± 0.3</td>
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<td>9.67 ± 0.25</td>
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<tr>
<td>MPV, fL</td>
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<td>7.1 ± 0.07*</td>
</tr>
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HCT, hematocrit; MCV, mean corpuscular volume; MPV, mean platelet volume. *P < .05

![Figure 2. Clearance rate and production of platelets is increased in Pak2-null mice.](image-url)

(A) Quantification of in vivo biotinylated platelets 24, 48, 72, and 96 hours after N-hydroxysulfosuccinimidobiotin-biotin injection. Data expressed as percentage of baseline (24 hours postinjection; n = 4-8 mice/genotype per time point; mean ± SEM). (B) Quantification of the percentage of new, reticulated platelets as a percentage of total platelets (n = 5; mean ± SEM), 10 DPI. (C) MPV of Pak2+/+ and Pak2−/− platelets, 14 DPI. (D) Representative images of hematoxylin and eosin stained spleen sections from Pak2+/+ and Pak2−/− mice 14 DPI. Arrows indicate splenic megakaryocytes. Scale bar = 50 µm (40× original magnification). (E) Quantification of megakaryocytes in spleen sections counted in 10 low-power fields. (F) Spleen weight (mg) 14 DPI. All data from at least 3 mice/genotype; mean ± SEM. *P < .05.
Figure 3. Pak2 deficiency increases megakaryocyte precursors and hematopoietic stem cells. (A) Percentage of bone marrow from wild-type or Pak2−/− mice that are PreMegE identified by CD150+ and CD105− after gating for Lin−/CD150−/Scal− (P < .05). (B) Percentage of bone marrow from wild-type or Pak2−/− mice that are committed MkPs, defined by CD41+ and CD150+, after gating for Lin−/CD150+/Scal− (P < .0001). (C) MegaCult-C colony-formation assays (CFU-Mk) of 105 unsorted bone marrow cells from wild-type and Pak2−/− mice 14 DPI. Number of megakaryocyte colony-forming units (CFU-Mk) were counted after staining with acetylcholinesterase (P < .0002). (D) Hematopoietic stem cells in bone marrow of wild-type and Pak2−/− mice. LSK, Lin−/CD150−/Scal−; LK, Lin−/CD150+/Scal− (P < .05). All bone marrow measured at 14 DPI. Experiments included n > 5 mice/genotype, mean ± SEM.

the fully mature state (D5). Pak2 was substantially autophosphorylated in CD34+ stem cells (D0), but this level, similar to that of pPak1/3, also increased during maturation (Figure 1E, top panel, lower band).

Localization and expression of Pak2 in mature megakaryocytes, proplatelets, and platelets was found throughout the maturation process (Figure 1F). Together, these data demonstrate that Pak2 has a role in megakaryocyte development from the stem cell to megakaryocyte.

**Pak2 deficiency increases platelet clearance rate and thrombopoiesis**

The combination of increased bone marrow megakaryocyte size and number, along with decreased platelet counts, suggests that Pak2−/− mice produce abnormal platelets with decreased survival in the circulation. To test whether platelet clearance was altered, we measured platelet life span, as well as the production of newly synthesized platelets. Pak2−/− platelet life span was significantly reduced relative to wild-type mice (Figure 2A). These data are comparable to the shortened platelet life span in the Rac1/Cdc42−/− mice. To assess whether platelet production was altered, we labeled blood samples ex vivo with thiazole orange. As shown in Figure 2B, there was nearly double the amount of reticulated platelets in Pak2−/− mice compared with controls (wild-type, 5.8 ± 0.4% vs Pak2−/−, 11.5 ± 0.6%; mean ± standard error of the mean [SEM]) at 10 DPI. Young reticulated platelets are larger than older, nonreticulated platelets, therefore, we evaluated MPV, and it was significantly increased in Pak2−/− mice (Figure 2C).

The spleen acts as a site for platelet clearance and production after bone marrow damage. Therefore, we evaluated the spleen for extramedullary megakaryopoiesis and discovered that Pak2−/−null spleens had significantly increased mature megakaryocytes; however, overall spleen weight was only slightly decreased (wild-type, 81 ± 3.5 mg; Pak2−/−, 70 ± 4 mg) (Figure 2D-F). Similar to bone marrow megakaryocytes, splenic megakaryocytes were also increased in diameter (supplemental Figure 1A, available on the Blood Web site). Increased platelet clearance can result in splenomegaly; however, our results demonstrated decreased spleen weight. Histological evaluation of Pak2−/− spleens demonstrated a distortion in normal splenic architecture, demonstrated by reduced white pulp, indicative of decreased lymphocytes and increased hemosiderin deposition (supplemental Figure 1B). Platelet clearance could still be occurring in the spleen, but other alterations to the spleen on Pak2 deletion make splenomegaly difficult to observe. Together, these data demonstrate that Pak2−/− mice are macrothrombocytopenic as a result of decreased platelet life span.

**Megakaryocyte progenitors are increased in Pak2-deficient mice**

Extensive flow cytometry studies have identified surface markers that constitute bone marrow megakaryocyte progenitors. Because we observed significantly more mature megakaryocytes in Pak2−/− bone marrow (Figure 1D), we investigated whether there were also increased...
megakaryocyte progenitors. The gating scheme used is delineated in supplemental Figure 2A. PreMegE cells are the progenitor cells that produce megakaryocytes and erythrocytes and have the highest capacity to form megakaryocyte colonies in vitro.36 This population is defined as Lin^− c-Kit^− CD41^− FcγRII/III^− CD150^− CD105^−.

We found that Pak2^+/− bone marrow had significantly more bipotential PreMegE cells than wild-type bone marrow (Figure 3A). We next examined committed megakaryocyte progenitors (MkP) by expression of CD41 after gating for Lin^− c-Kit^− Sca1^− CD150^−. These progenitor cells represent an intermediate stage between the bipotential precursor and mature megakaryocyte. Lineage-committed MkPs were significantly increased in the Pak2^+/− bone marrow (Figure 3B).

Given the increase in megakaryocyte stem cell progenitors, we next tested whether these progenitors produced more mature megakaryocyte colonies. To address this question, we cultured an equal amount of unsorted bone marrow from wild-type and Mx1cre^+/−Pak2^fl/fl animals, 14 DPI injection, in a megakaryocyte specific colony-formation assay (CFU-MK). Analysis of the colonies with acetylcholinesterase staining after 8 to 10 days of growth with TPO, IL-3, and IL-6 demonstrated an increased number of colonies in Pak2^+/− bone marrow, relative to...
Ter119high cells were additionally analyzed for CD71 surface expression and increased mature erythrocytes in bone marrow (supplemental Figure 4A-B).

Pak2 proliferative potential of megakaryocyte progenitors are increased in (supplemental Figure 2B). These results suggest that the number and proliferative potential of megakaryocyte progenitors also affected erythroid precursors, we examined bone marrow of wild-type and Pak2−/− micrographs and (B) proplatelet structure by fluorescence microscopy of wild-type and Pak2−/− megakaryocytes adhered to collagen for 5 hours. (D) Fetal liver-derived megakaryocytes treated with Frax1036 for duration of culture (4 days) and stained for β1-tubulin (green) and DAPI (blue). (E) Western blot detection of serine 141 phosphorylated Pak1/2/3 (pPak) of fetal liver-derived megakaryocytes treated with Frax1036 for duration of culture (4 days) and stained for β1-tubulin (green) and DAPI (blue). (F) Representative Alexa Fluor 594-phalloidin staining for actin (red) and 1-tubulin (green) and DAPI nuclear (blue) analyzed by fluorescence microscopy of wild-type and Pak2−/− megakaryocytes adhered to collagen for 5 hours. Bone marrow treated with 500 nM 4-hydroxytamoxifen to induce Cag-Cre-ERT2 expression and delete Pak2fl/fl expression and deletion resulted in significantly decreased endomitosis (polyploidization) in megakaryocytes (percentage of cells containing more than 8N DNA content) (Figure 4A). To determine whether this was cell-intrinsic, we measured polyploidy with binase and found significantly increased 8N, 16N, and 32N populations (P < .03). These in vitro data suggest that increased polyploidization observed in vivo is

Because Pak2−/− mice demonstrated altered megakaryopoiesis, we next asked whether loss of the Pak2 gene was associated with abnormal hematopoiesis. Total bone marrow hematopoietic stem cells (LSK [Lin−/C-kit−/Sca1+] and LK [Lin−/C-kit−/Sca1−]) were evaluated and significantly increased in Pak2−/− bone marrow (Figure 3D). LSK cells were further evaluated for number of long-term HSCs and multipotent progenitors, and both populations were increased (supplemental Figure 3A-B). Collectively, these findings identify Pak2 as a negative regulator of megakaryocyte maturation via regulation of the bipotential precursors (PreMegE) and MkP cells in the bone marrow. The observed increase in megakaryocyte stem cells is not caused by elevated serum TPO levels induced by a reduction in circulating platelets (data not shown).

To examine whether expansion of megakaryocyte–erythroid progenitors also affected erythroid precursors, we examined bone marrow for erythroblasts (CD71+Ter119+9b) and erythrocytes (CD71+Ter119+9b) (supplemental Figure 4A-B). Pak2 deletion significantly reduced early erythroblasts and increased mature erythrocytes in bone marrow. Ter119+9b cells were additionally analyzed for CD71 surface expression and size by FSC. Pak2 deletion resulted in significant decreased bone marrow immature and intermediate erythrocytes (EryA, EryB) and increased mature erythrocytes (EryC) (supplemental Figure 4C-D). Evaluation of erythroid stem cell progenitors (proerythroblasts), identified as Lin−/ckit−/Sca1−/CD41+/CD150−/CD105−/Ter119+/CD71− was significantly decreased in Pak2-null bone marrow (supplemental Figure 4E). Together, these data demonstrate that Pak2 deletion shifts the balance in the PreMegE stem cells toward the megakaryocyte lineage, leading to a decrease in expansion of proerythroblasts and erythroblasts.

Megakaryocyte endomitosis is negatively regulated by Pak2

Megakaryocytes progress through development to an endomitotic phase. Here cells are programmed to fail cytokinesis and accumulate DNA content in a single polylobulated nucleus before a final maturation state, consisting of proplatelet formation and platelet release.24,45 Accordingly, we examined endomitosis (polyploidization) in megakaryocytes from Pak2−/− mice. We found that Pak2 deletion in vivo markedly increased polyploidization of megakaryocytes (percentage of CD41+ cells containing more than 8N DNA content) (Figure 4A). To determine whether this was cell-intrinsic, we measured polyploidy with in vitro–deleted megakaryocytes with tamoxifen-regulated Cre recombinase and found Pak2 regulated polyploidization in a cell-intrinsic manner (Figure 4B). Pak2−/− megakaryocytes demonstrated significantly increased 8N, 16N, and 32N populations (P < .03). These in vitro data suggest that increased polyploidization observed in vivo is

Figure 5. Altered cytoskeleton structure in Pak2-null megakaryocytes. (A) Analysis of β1-tubulin structure and (B) proplatelet structure by fluorescence microscopy of wild-type and Pak2−/− megakaryocytes adhered to fibrinogen for 5 hours. Bone marrow treated with 500 nM 4-hydroxytamoxifen to induce Cag-Cre-ERT2 expression and delete Pak2fl/fl deletion signifi-
from cell-intrinsic signaling, but does not rule out influence from cell-extrinsic factors.

Small-molecule inhibitors specifically targeting group I Pak are currently being developed for the treatment of tumors overexpressing Pak proteins.10,28-30,46,47 We examined whether treatment with group I-specific Pak inhibitor Frax1036 also altered megakaryocyte polyploidization or induced thrombocytopenia.33 Treatment with Frax1036 for 3 weeks increased polyploidization in megakaryocytes and increased CD41⁺ megakaryocytes (Figure 4C-D). Frax1036 treatment effectively ablated Pak1-3 phosphorylation in the bone marrow (Figure 4C, inset). Bone marrow-derived megakaryocytes treated with Frax1036 also increased polyploidization at 8N and 16N ploidy stages (P < .008 and P < .003, respectively) (Figure 4E). Collectively, these findings indicate that genetic deletion of Pak2, as well as pharmacologic kinase inhibition, causes increased megakaryocyte polyploidization in vivo and in vitro.

**Altered cytoskeleton and ultrastructure of Pak2⁻/⁻ megakaryocytes**

During maturation, megakaryocytes undergo cytoskeletal alterations critical for polyploidization and proplatelet formation.38-50 As Pak kinases are principally known for their regulation of actin and tubulin cytoskeleton networks,51 we next measured β1-tubulin expression in

![Figure 6. Decreased proplatelets and altered ultrastructure of Pak-null megakaryocytes.](image)

(A) Bone marrow imaged for megakaryocytes from wild-type and Mx1cre Pak2⁻/- mice analyzed by transmission electron microscopy, 14 DPI. Wild-type megakaryocyte (upper, left, x5000), periphery magnified (upper, right, x12000). Pak2⁻/- megakaryocyte (lower, left, x5000), periphery magnified (lower, right, x20 000). Scale bar = 2 μm. (B) Proplatelet formation imaged from cultured megakaryocytes from wild-type and Cag-Cre-ERT2 Pak2⁻/- bone marrow, induced with 500 nM 4-hydroxytamoxifen and adhered to fibrinogen. White arrows indicate proplatelet. (C) Percentage proplatelet-forming megakaryocytes expressed as a percentage of total megakaryocytes per visual field ± SEM, n = 5/genotype. Scale bar = 200 μm. *P < .05. (D) Representative image of proplatelet formation in vehicle and Frax1036-treated bone marrow-derived megakaryocytes. Scale bar = 200 μm. White arrows indicate proplatelet.
megakaryocytes adherent to fibrinogen, relative to cell area. Quantitative fluorescence microscopy analyses found that Pak2−/− megakaryocytes expressed significantly less β1-tubulin (24.1 ± 2.5%) relative to wild-type megakaryocytes (Figure 5A). Proplatelet extension structures were also altered in Pak2−/− megakaryocytes (Figure 5B). Actin polymerization of megakaryocytes adherent to collagen was also decreased with Pak2 deletion (57.6% ± 15.3% decrease compared with wild-type) (Figure 5C). Similar to Pak2−/− megakaryocytes, β1-tubulin staining decreased in Frax1036 treated-megakaryocytes, further supporting a role for Pak2 activity in megakaryocyte maturation and proplatelet formation (Figure 5D-E).

To better investigate megakaryocyte morphology in situ, we next examined wild-type and Pak2−/− bone marrow sections by transmission electron microscopy. Mature wild-type megakaryocytes displayed characteristic membrane invaginations, indicative of the invaginated membrane system (Figure 6A, top). Pak2−/− megakaryocytes demonstrated fewer invaginations in the periphery and around the nucleus (Figure 6A, bottom), supporting a role for Pak2 in proplatelet formation. Indeed, on adherence to fibronectin, Pak2−/− bone marrow-derived megakaryocytes developed significantly fewer proplatelet extensions (Figure 6B-C). Consistent with genetic deletion, Frax1036-treated bone-marrow-derived megakaryocytes phenocopied Pak2-null megakaryocytes, with no proplatelet extensions observed (Figure 6D). Together, these data support a role for Pak2 in megakaryocyte cytoskeletal dynamics and ultrastructural mechanics that orchestrate proplatelet formation.

Altered signal transduction in Pak2−/− megakaryocytes

To establish a potential mechanism underlying the effects of Pak2 deletion on megakaryocyte function, we next investigated the phosphorylation status of Pak2 effectors known to regulate cytoskeletal dynamics. We found that LIMK phosphorylation was significantly reduced (>60% reduction) and was associated with reduced phosphorylation of cofilin (>40% reduction) (Figure 7A-B) in Pak2−/− megakaryocytes. As cofilin is active in the nonphosphorylated state, these results suggest that Pak2−/− megakaryocytes undergo more rapid actin severing and depolymerization compared with wild-type megakaryocytes. Such properties have previously been associated with enhanced polyploidization in megakaryocytes.50,52

Similar to actin severing, enhanced microtubule depolymerization with colchicine, nocodazole, and vincristine have also been shown to enhance megakaryocyte ploidy.50,53,54 Microtubule dynamics are regulated by a variety of proteins, including the Pak substrate Aurora A44,55 a negative regulator of megakaryocyte polyploidization.56 We evaluated Pak2−/− megakaryocytes for Aurora A/B/C phosphorylation and found markedly decreased levels in Pak2−/− cells compared with wild-type levels (>48% reduction) (Figure 7A-B). These data strongly suggest that Pak2 regulates multiple functions in the developing megakaryocyte by regulating both the actin and microtubule networks (Figure 7C).

Discussion

Here we report a novel role for Pak2 in megakaryocyte biogenesis, cytoskeletal remodeling, proplatelet formation, and endomitosis. Using a conditional knockout mouse model, as well as a small-molecule inhibitor of group I Paks, we establish Pak2 as regulator of the megakaryocyte cytoskeleton. We demonstrate that Mx1-crePak2−/− mice develop macrothrombocytopenia, which manifests in vivo with decreased platelet lifespan, along with increased mature megakaryocytes in the bone marrow and spleen, increased level of megakaryocyte ploidy, and increased megakaryocyte bone marrow progenitor cells. Pak2 also regulates megakaryocyte cytoskeleton structure, demonstrated with transmission electron microscopy of in situ bone marrow and immunofluorescence of in vitro primary culture of megakaryocytes.

Evidence from signaling studies suggests that at least 2 signaling pathways that regulate cytoskeletal function in megakaryocytes are regulated by Pak2: Aurora A, which classically regulates microtubule dynamics and mitotic entry; and LIMK/cofilin, which regulate actin dynamics (Figure 7). Aurora A deletion increases megakaryocyte differentiation and levels of polyploidization.56 Aurora A could function to regulate polyploidization through its role in microtubule-organizing center localization, bipolar spindle formation, and inhibition of proteins that destabilize the microtubule network (ie, stathmin and mitotic centromere-associated kinesin).59,61 We evaluated the effect of Pak2 deletion on megakaryocyte Aurora activation and found that Pak2−/− megakaryocytes had decreased Aurora phosphorylation. Actin regulatory mechanisms mediated by the Pak effector LIMK and its substrate cofilin also play a role in later stages of megakaryocyte development, and in particular in proplatelet formation.62 We observed significantly decreased F-actin formation in Pak2-null megakaryocytes (Figure 5C), supporting the hypothesis that Pak2 deletion increases actin-severing activity of cofilin through LIMK inhibition. Notably, LIMK can also
directly phosphorylate Aurora to stabilize the microtubule network.\textsuperscript{53,64} Thus, loss of LIMK phosphorylation in Pak2\textsuperscript{-/-} megakaryocytes may serve to disrupt a signaling node between actin and microtubule systems, affecting both Aurora and cofilin activity (Figure 7C). However, LIMK regulation downstream of Pak is complex, as megakaryocyte double-knockout deletions of genes encoding the Pak activating proteins Rac1/Cdc42 increase LIMK and cofilin phosphorylation.\textsuperscript{25} Such findings support roles for Rac1 and Cdc42 beyond that of their effector Pak2 alone and suggest that Pak2 has a specific role in megakaryocyte endomitosis and proplatelet formation by means of regulating signal transduction cascades involved in cytoskeletal network regulation.

Although the use of the inducible Mx1-cre system for Pak2 deletion in mice makes it difficult to determine whether our observations regarding megakaryocyte morphology, function, and platelet half-life are cell autonomous, our findings with in vitro megakaryocyte culture strongly suggest a cell-autonomous effect. In vivo bone marrow megakaryocyte ultrastructure revealed disrupted megakaryocyte plasma membrane invaginations (Figure 6A) in the Mx1-cre model system. Consistent with in vivo data, in vitro megakaryocytes were also defective in their structure, mainly observed by defects in proplatelet formation, using both a genetic and pharmacological model (Figure 6B-C). Elevated levels of polyploidization were also observed both in vivo and in vitro (Figure 4). These data strongly suggest a cell-autonomous role for Pak2 in megakaryocytes. It remains possible, however, that there are also non-cell-autonomous effects related to Pak2 deletion in other hematopoietic lineages in Mx1-cre mice, especially given the known role of Pak2 in hematopoietic stem cells and their progeny.\textsuperscript{15,32,65} These issues will be addressed in future studies using transgenic mice with megakaryocyte-specific Cre drivers such as P-F4-cre.\textsuperscript{56}

In summary, our results demonstrate that Pak2 regulates megakaryocytes, in part through signaling networks that regulate actin and microtubule cytoskeleton. This multineuflence influence of Pak2, through phosphorylation of LIMK and Aurora, supports a role for Pak2 in controlling the dynamic cytoskeleton of developing megakaryocytes to play a novel role in megakaryocyte biology.

Acknowledgments

We thank J. Oesterling (Fox Chase Cancer Center Flow Cytometry Core) for flow cytometry assistance, A. Efimov (Fox Chase Cancer Center Microscopy Core), and D. Williams (University of Pennsylvania, Electron Microscopy Resource Laboratory) for imaging assistance. We also thank Dr J. Field, Dr S. Ryeeom, Dr S. Sykes, Dr A. Mazarhian, and Dr M. Larson for insightful comments.

This work was supported by grants from the National Institutes of Health, National Cancer Institute (R01-CA142928 and R01-CA148805 to J.C., and F31-CA177182 to R.E.K.), the Fox Chase Cancer Center (P30-CA006927), the National Heart, Lung, and Blood Institute (R01-HL101972 to O.J.T.M.; and R01-HL118593 and R01-HL93231 to A.P.K.), as well as by an appropriation from the State of Pennsylvania. In addition, grant funding was provided from the American Heart Association (13POST13730003 to J.E.A.; 15POST22420000 to J.C.K.; and 13EIA12630000 to O.J.T.M.).

Authorship


Conflict-of-interest disclosure: The authors declare no competing financial interests.

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Pak2 restrains endomitosis during megakaryopoiesis and alters cytoskeleton organization

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