A mouse model of hereditary folate malabsorption: deletion of the PCFT gene leads to systemic folate deficiency

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The human proton coupled folate transporter (PCFT) is involved in low pH-dependent intestinal folate transport. In this report, we describe a new murine model of the hereditary folate malabsorption syndrome that we developed through targeted disruption of the first 3 coding exons of the murine homolog of the PCFT gene. By 4 weeks of age, PCFT-deficient (PCFT−/−) mice developed severe macrocytic normochromic anemia and pancytopenia. Immature erythroblasts accumulated in the bone marrow and spleen of PCFT−/− mice and failed to differentiate further, showing an increased rate of apoptosis in intermediate erythroblasts and reduced release of reticulocytes. In response to the inefficient hematologic development, the serum of the PCFT−/− animals contained elevated concentrations of erythropoietin, soluble transferrin receptor (sCD71), and thrombopoietin. In vivo folate uptake experiments demonstrated a systemic folate deficiency caused by disruption of PCFT-mediated intestinal folate uptake, thus confirming in vivo a critical and nonredundant role of the PCFT protein in intestinal folate transport and erythropoiesis. The PCFT-deficient mouse serves as a model for the hereditary folate malabsorption syndrome and is the most accurate animal model of folate deficiency anemia described to date that closely captures the spectrum of pathology typical of this disease. (Blood. 2011;117(18):4895-4904)

Introduction

Folate is an essential cosubstrate of many biochemical reactions, such as the de novo synthesis of purines and pyrimidines, methionine, and deoxythymidylate monophosphate.1,2 Mammals cannot synthesize folate; therefore, inadequate dietary supply or folate malabsorption results in defective DNA synthesis. One of the first manifestations of a folate deficiency is in the rapidly proliferating cells of the hematopoietic system, leading to pancytopenia and anemia of the megaloblastic type. Patients with megaloblastic anemia demonstrate ineffective erythropoiesis by harboring large immature red blood cell (RBC) precursors that fail to survive to the postmitotic, terminal stages, undergoing premature apoptosis.2

Three mammalian folate transporter systems have been described to date in a variety of tissues: (1) the bidirectional reduced folate carrier 1 (RFC1), also known as SLC19A1,3,4 (2) the glycosyl-phosphatidylinositol-anchored folate receptors (FOLR1, FOLR2, and FOLR4) and one secreted receptor in humans without a mouse homolog (FOLR3),5 and (3) the human proton coupled folate transporter (PCFT).6-8 The RFC1 transporter is expressed ubiquitously, including the brush-border membrane of epithelial cells in the small intestine.9 Although RFC1 is necessary for folate transport in erythroid cells, its involvement in intestinal folate uptake has not been confirmed.6 Inactivation of RFC1 in mice by homologous recombination led to either embryonic lethality or defective erythropoiesis in pups born to mothers who were supplemented with 1 mg daily subcutaneous doses of folic acid.10,11 Because this transporter functions at a neutral pH optimum whereas the majority of intestinal folate transport occurs in an acidic luminal milieu, RFC1 is an unlikely candidate for an intestinal folate transporter.7 The role of FOLR1 in folate absorption and transport has also been demonstrated, where a 60% to 70% reduction was observed in plasma folate of FOLR1−/− mice fed low folate and normal chow.12 FOLR1 also regulates folate homeostasis via endocytotic mechanisms during embryonic development, and mice rendered null for this receptor display severe morphogenetic abnormalities and die in utero unless provided supraphysiologic concentrations of either folic acid or 5-methyltetrahydrofolate.5,13

The PCFT transporter is highly expressed in tissues involved in folate and heme transport, including the duodenum and liver.6,14 Initially, PCFT was identified as a low-affinity, pH-dependent heme transporter14 and then later described to function as a low pH-dependent folate transporter in intestinal cells.6 The latter role of the transporter was confirmed by the identification of loss-of-function mutations in the human PCFT gene in persons diagnosed with hereditary folate malabsorption syndrome.6 Studies have also indicated that PCFT facilitates folate transport during folate receptor-mediated endocytosis, where FOLR1 binds folate, and its export into the cytosol is driven by PCFT activity as the vesicle undergoes endocytosis and becomes acidified.15 In this report, we describe a new murine model of hereditary folate malabsorption syndrome and folate deficiency anemia, which we developed through targeted deletion of the PCFT gene, and confirm in vivo a critical and nonredundant role of the PCFT transporter in folate metabolism and erythropoiesis.
Methods

Generation of PCFT mutant mice

The targeting vector was derived using long-range polymerase chain reaction (PCR) to generate the 5′ and 3′ arms of homology using 129/SvEv<sup>thd</sup> embryonic stem cell (ESC) Lex-1 DNA as a template. The 2388-bp 5′ arm was generated using primers PCFT-3 [5′-GGATCCGGATGGGCTGACAGCCAACACGGAGGGGCTAGCAC-3′] and PCFT-2 [5′-GGCCGCTATGGGCTGACAGCCAACACGGAGGGGCTAGCAC-3′] and cloned using the TOPO cloning kit (Invitrogen). The 3722-bp 3′ arm was generated using primers PCFT-4 [5′-GGCCAGCGAGGCCGGATCAGTTTGTCATCTGAAACTACCTG-3′] and PCFT-6 [5′-CTCGAGCATGCCCTTGCTCTGTAACCTCTG-3′] and cloned using the TOPO cloning kit (Invitrogen). The 5722-bp 5′ arm was generated using primers PCFT-7 [5′-GGCCAGCGAGGCCGGATCAGTTTGTCATCTGAAACTACCTG-3′] and PCFT-8 [5′-CTCGAGCATGCCCTTGCTCTGTAACCTCTG-3′] and cloned using the TOPO cloning kit (Invitrogen). The 3′ arm was excised from the holding plasmid using BamHI and SfiI. The 3′ arm was excised from the holding plasmid using SfiI and XhoI. The arms were ligated to a SfiI prepared selection cassette containing a neomycin fusion marker (B-Geo) along with a PGK promoter-driven porcymycin resistance marker and inserted into a BamHI/XhoI cut pKOScrambler vector (Stratagene) to complete the PCFT targeting vector. The NotI linearized targeting vector was electropropagated into 129/SvEv<sup>thd</sup> (Lex-1) ESCs. G418/FI.AU-resistant ESC clones were isolated, and correctly targeted clones were identified and confirmed by Southern analysis using a 291-bp 5′ selection cassette probe (8/9), generated by PCR using primers Ires-8 [5′-AGGAAAGCAGTGCTCTCTGGAA-3′] and Ires-9 [5′-CACATGAAAGCAGTGCTCTCTGGAA-3′] and a 441-bp 3′ external probe (20/21), amplified by PCR using primers PCFT-20 [5′-GGCCAGCGAGGCCGGATCAGTTTGTCATCTGAAACTACCTG-3′] and PCFT-21 [5′-CTCGAGCATGCCCTTGCTCTGTAACCTCTG-3′]. Southern analysis using probe 8/9 detected a 9.3-kb mutant band in ApalI digested genomic DNA, whereas probe 20/21 detected a 15.5-kb wild-type (WT) band and 12.3-kb mutant band in EcoRI digested genomic DNA. Six targeted ESC clones were identified and microinjected into C57BL/6 (albino) blastocysts to generate chimeric animals, which were bred to C57BL/6 (albino) females. The resulting heterozygous (PCFT<sup>+/−</sup>) offspring were interbred to produce homozygous PCFT<sup>−/−</sup> mice. Determination of the genotype of mice at the PCFT locus was performed by screening DNA from tail biopsy samples using quantitative PCR for the Neo cassette. This strategy enabled discrimination of zero, one, or 2 gene disruptions representing WT, PCFT<sup>+</sup>−, and PCFT<sup>−/−</sup> mice, respectively.

All experiments were performed on mice of mixed genetic background (129/SvEvBrd and C57BL/6J) representing both sexes of littermate mutant and WT animals. Procedures involving animals were conducted in conformity with the Institutional Animal Care and Use Committee guidelines that are in compliance with the state and federal laws.

Complete blood cell count and flow cytometry (FACS) analysis

Peripheral blood was collected from the retro-orbital plexus of mice anesthetized with isoflurane, using heparinized microhematocrit capillary tubes. Complete blood cell count was obtained using a hematologic analyzer (Cell-Dyn 3500R, Abbott Diagnostics), according to the manufacturer’s instructions. Reticulocytes and reticulated platelets (rPLT) were quantitated on a FACScan flow cytometer (BD Biosciences Immunocytometry Systems) after staining whole blood samples with RET-COUNT (Thiazole Orange) reagent (BD Biosciences), according to the manufacturer’s instructions. Reticulocyte production index (RPI) in PCFT<sup>−/−</sup> mice was calculated by multiplying the percentage of reticulocytes in whole blood by the ratio of peripheral blood hematocrit (HCT) to the mean HCT in the WT group (% reticulocyte × [HCT/mean WT HCT]). The RPI was further adjusted for the longer life span of reticulocytes in mice with low HCT by dividing RPI values by the reticulocyte maturation index, which was arbitrarily set to 1 for blood samples with HCT values in the 41% to 50% range, and 1.5 for blood samples with HCT values in the 30% to 40% range.

For analysis of bone marrow (BM), cells were harvested by flushing femurs and tibia with 5 mL of Hanks balanced salt solution supplemented with 5% fetal bovine serum (fluorescence-activated cell sorter [FACS] buffer) using a 27-gauge needle. Single splenocyte suspensions were prepared by passage through a 70-μm cell strainer in the presence of FACS buffer. For FACS analysis, cells were washed 2 times in FACS buffer, incubated with 1 μg of monoclonal antibodies to CD16/CD32 molecules (Fc Block, BD Biosciences Pharmingen) for 15 minutes at 4°C, and stained for 30 minutes at 4°C in the dark with fluorescein-isothiocyanate (FITC)-conjugated anti-CD119 monoclonal antibodies (all purchased from BD Biosciences Pharmingen). The cells were washed twice in FACS buffer before analysis.

Anucleated erythrocytes with low forward scatter were not included in the analysis.

Histology

Freshly isolated mouse tissues were fixed in buffered neutral 10% formalin. Paraffin-embedded tissue was sectioned at 4 to 6 μm, mounted, and stained with hematoxylin and eosin or Gomori iron stain by standard methods, or...
modified von Kossa silver nitrate staining for Ca salts. Cytospin (Shandon Southern Products) preparations of BM cells and splenocytes were stained by the May-Grünwald-Giemsa standard technique. Sections and cytospin preparations were observed and photographed using the SPOT InSight 3.2.0 camera (Diagnostic Instruments Inc) attached to the Olympus BX51 microscope (Olympus), controlled by SPOT 4.0.9 software (Diagnostic Instruments Inc).

**Enzyme-linked immunosorbent assays**

Erythropoietin (EPO) and thrombopoietin serum levels were quantified using commercial enzyme-linked immunosorbent assay kits (R&D Systems). For measurement of soluble transferrin receptor (sCD71) concentration in mouse sera, Nunc Maxi-Sorp 96-well plates (VWR) were coated overnight at 4°C with 100 μL/well of a 5-μg/mL solution of rat antianti CD71 IgG (Southern Biotechnology Associates). The plates were washed 3 times with phosphate buffered saline (PBS) containing 0.05% Tween-20 (PBS/Tween-20) and blocked with 200 μL/well of 10% fetal bovine serum in PBS for one hour at room temperature. After 5 washes in PBS/Tween-20, the plates were incubated with mouse serum (diluted 1:20 in PBS) for 2 hours at room temperature. Plate-bound sCD71 was detected by incubating the plates for one hour at room temperature with biotinylated rat antimouse CD71 IgG (2 μg/mL; BD Biosciences Pharmingen), followed by streptavidin-alkaline phosphatase conjugate (30 minutes at room temperature; Southern Biotechnology Associates). Colorimetric detection was performed by adding 3,3',5',5'-tetramethylbenzidine to the plates and reading absorbance at 450 nm wavelength.

**Iron status, heme, folate, and homocysteine content measurements**

Total serum iron (Fe^{2+} and Fe^{3+}) and tissue folate concentrations were measured using a colorimetric assay (QuantiChrom Iron Assay Kit, BioAssay Systems) and enzyme-linked immunosorbent assay (TECRA International), respectively, according to the protocols supplied by the manufacturers. For measurement of tissue folate, liver, kidney, and splenic tissues were weighed, homogenized, lysed by 3 cycles of freeze-thaw in folate enzyme-linked immunosorbent assay sample buffer, and briefly sonicated. Extracts were then centrifuged at 10,000g at 4°C, and the supernatant was retained.

Plasma folate was determined by modification of the inhibition of folate binding assay. Briefly, bovine folate binding protein is immobilized onto glass slides and sample folates compete with horseradish peroxidase-labeled folic acid for binding to bovine folate binding protein. Standards of unlabeled folic acid (0.244-250 ng/mL) in PBS-0.1% Tween-20 (PBS-T, pH 7.3) were used to determine concentrations. Standards and plasma (10 μL) were diluted to 50 μL in PBS-TA (PBS-T with 1% weight/volume ascorbate, pH 3.0) and placed in boiling water for 10 minutes. Samples were neutralized with 1 μL of 10% NaOH, checked for neutral pH on test strips, and spun for one minute at 14,000g. Supernatants were mixed 1:4 with horseradish peroxidase-labeled folic acid (Ortho-Clinical Diagnostics) and incubated with immobilized bovine folate binding protein for 2 hours. Slides were washed 5 times with PBS-T, signal amplification is performed with the biotin-labeled tyramide (PerkinElmer Life and Analytical Sciences), streptavidin-alkaline phosphatase conjugate (Invitrogen), and enzyme-labeled fluorescein alkaline phosphatase substrate (Invitrogen). Slides were imaged using an 8-bit UV photography workstation (Kodak). Fluorescent signal intensities were determined using ImageJ 1.43 (National Institutes of Health, Bethesda, MD), and foreground signals were fit with a sigmoid, regression curve using GraphPad Prism 4 Software.

Total homocysteine (tHcy) concentrations were determined in plasma and tissue homogenates using high-performance liquid chromatography according to the previously described procedure. tHcy concentrations in tissue homogenates were further normalized to total protein content.

**Analysis of in vivo folate uptake**

Analysis of in vivo folate uptake was performed as previously described. Briefly, a single intragastric dose of tritiated [3',5',7,9-3H] folic acid (34.9 Ci/mmol; GE Healthcare) was given to mice by oral gavage (20 μCi/mouse). Control mice received a single oral dose of cold folic acid. Serum was collected by retroorbital bleed 0.5 and one hour after folate administration. Liver, kidney, proximal small intestine (duodenum and proximal jejunum), and spleen tissue were collected 16 hours after dosing, weighed, and homogenized in ice-cold 2 times radioimmunoprecipitation assay buffer (Thermo Fisher Scientific Inc) using a tissue homogenizer.
Protein concentration in the tissue lysates was analyzed by the Bradford assay. Folate-associated radioactivity in serum and tissue lysates containing equal amounts of protein was measured by β-scintillation counting and normalized to the mouse weight.

Analysis of in vitro uptake of folate and zinc protoporphyrin

The full-length coding sequences of human and mouse PCFT were cloned into the expression vector pIRESpuro2 (Clontech). The plasmids were transfected into HEK293 cells with Lipofectamine 2000 (Invitrogen), and uptake of folate was measured 24 hours later as described previously.6 Zinc protoporphyrin uptake assays were carried out in HEK293 cells transiently transfected with human PCFT essentially as described previously.14

Supplementation studies

PCFT−/− mice were given daily supplementation containing iron or folate. The volumes administered were 0.10 mL per 10 g of body weight. Hemin (ferrous sulfate (FeSO4) was dissolved in sterile reverse osmosis purified water and given by gavage (1, 10, and 40 mg/kg). Folate 5-formyltetrahydrofolate (folinic acid) and 5-methyltetrahydrofolate (5-MTHF) were dissolved in distilled water and given by gavage or injected subcutaneously at 40 mg/kg dose. Folic acid was administered daily by subcutaneous injection at 0.1 mg/kg dose, or given by gavage (0.1 and 2 mg/kg body weight). All reagents were from Sigma-Aldrich, unless indicated otherwise.

Figure 3. PCFT deficiency leads to impaired erythropoiesis and thrombopoiesis. (A) Representative May-Grünwald-Giemsa-stained cyt centrifuge preparations show accumulation of proerythroblasts and large basophilic erythroblasts (yellow arrows) in 4- to 6-week-old PCFT−/− mice. Periodic acid Schiff-stained BM sections (original magnification 10×/0.3 NA dry objective) contain reduced number of megakaryocytes (green arrows) in PCFT−/− mice than in WTs. (Inset) Megakaryocytes are less differentiated in the PCFT−/− BM, as evidenced by marked reduction in cytoplasmic volume and nuclear lobulation. (B) Abnormal neutrophil precursors (giant metamyelocytes, yellow arrows) in the BM of PCFT−/− mice (May-Grünwald-Giemsa staining of cyt centrifuge preparations). (C) Multilobed polymorphonuclear neutrophils in a 4- to 6-week-old PCFT−/− mouse (top panels) (original magnification 40×/0.75 NA dry objective). No evidence of deep intracerebral hemorrhage and calcification was found in PCFT−/− brain sections by hematoxylin and eosin (bottom panel) and von Kossa staining (not shown). The hematoxylin and eosin low power image (original magnification 2×/0.08 NA dry objective) corresponds to the location of the insets at the right (original magnification 20×/0.75 NA dry objective; sagittal fissure above corpus callosum and ventrolateral cortex, respectively).

Statistical analysis

The normality of results was evaluated by the Kolmogorov-Smirnoff test. Statistical significance of group differences between WT and PCFT−/− mice was analyzed using the 2-sample Student t test, and P < .05 was considered statistically significant.

Results

PCFT is essential for normal growth and survival

We generated PCFT-deficient mice by targeted disruption of the first 3 exons of the murine homolog of the PCFT gene (GenBank accession no. NM_026740) by homologous recombination in ESCs,20 followed by germline transmission of the mutant gene into chimeric, PCFT+/−, and PCFT−/− mice (Figure 1A). Southern hybridization analysis demonstrated the targeted mutation in ESCs (Figure 1B). PCFT transcripts were detected by reverse-transcribed PCR in all tissues of WT (PCFT+/+) mice, except for adipose tissue and total bone tissue including BM (Figure 1C).

Mating of PCFT−/− mice generated pups of the 3 possible genotypes with ratios that fit well within normal Mendelian frequencies. However, PCFT−− mice failed to thrive (body weight 21 ± 4 g, PCFT+/+, n = 8 vs 15 ± 2 g, PCFT−/+ , n = 8, at 6 weeks of age, P = .004), and more than 90% of PCFT−/− mice died by 10 to 12 weeks of age. Growth rate and survival of PCFT−/− mice were normal; therefore, they were not routinely included in additional characterization studies.
Blood profile of PCFT−/− mice demonstrates pancytopenia and ineffective erythropoiesis

Hematologic analysis of the peripheral blood of PCFT−/− mice revealed decreased cellularity in all leukocyte populations as well as erythrocytes and platelets (Figure 2A). RBC parameters in the mutant mice were consistent with a diagnosis of macrocytic normochromic anemia, showing lower RBC counts and decreased HCT and hemoglobin concentration compared with WT controls, but having normal values for mean corpuscular hemoglobin concentration. In addition, increased distribution width of RBC indicated anisocytosis, which was accompanied by an increase in the mean corpuscular volume to the level of macrocytosis (mean ± SEM; n = 10 mice each cohort). BM and splenic cellularity and the percentages of erythroid progenitor subsets in the BM and spleen, including the percentages of annexin V− intermediate erythroblasts, are summarized in panel B (mean ± SEM; n = 10 mice each cohort). Data are from one of 2 independent experiments showing similar results. *P < .05 compared with WT.

Absence of PCFT leads to impaired erythropoiesis in the BM and spleen

The hematologic profile of PCFT-deficient mice closely resembled that of human folate deficiency anemia. This observation prompted
us to characterize maturation and survival of RBC progenitors in PCFT−/− BM and spleen. Histopathologic examination revealed signs of marked erythroid cell hyperproliferation in the BM and spleens of PCFT−/− mice, leading to accumulation and over-representation of proerythroblasts and large basophilic erythroblasts (Figure 3A-B). However, both the reticular chromatin pattern and nucleocytoplasmic dissociation typically observed in human megaloblasts were infrequent findings in the PCFT−/− BM and splenic erythroblasts. The PCFT−/− BM also harbored less mature megakaryocytes with abnormal morphology. These findings are all consistent with a histopathologic profile observed in patients with folate deficiency anemia.

To delineate the erythroid maturational stages affected by the PCFT mutation, we performed phenotypic characterization of BM (Figure 4) and splenic (Figure 4B) subpopulations by FACS analysis using markers for erythroid cell differentiation and apoptosis. Differential expression of Ter119 and CD71 characterizes erythroid maturation, proceeding from proerythroblasts (Ter119(lo)CD71(hi) cells) and early basophilic precursors (Ter119(hi)CD71(hi)), through basophilic and chromatophilic erythroblasts (Ter119(hi)CD71(int) intermediate erythroblasts), to orthochromatophilic erythroblasts and reticulocytes (Ter119(hi)CD71(low) cells). In PCFT-deficient mice, this maturational process appeared to be blocked at the intermediate erythroblast stage because the mutant BM showed accumulation of Ter119(hi)CD71(mod) intermediate erythroblasts compared with WT BM. Moreover, the proportion of annexin V-positive apoptotic cells was significantly higher in the intermediate erythroblast population of PCFT−/− BM than in the WT BM (22% ± 1.4% vs 16% ± 0.7%; P = .006), indicating that erythroid differentiation in the mutant BM was impaired as a result of decreased survival of erythroblasts at the later stage of erythroid maturation. Ineffective erythropoiesis and elevated EPO levels in PCFT−/− mice also led to the rapid expansion of stress erythroid progenitors in the spleen, the primary site of extramedullary erythropoiesis in mice (Figure 4B). However, the expansion of TER119+ erythroid progenitors and accumulation of Ter119(hi)CD71(int) intermediate erythroblasts in the spleen of PCFT−/− mice was accompanied by elevated apoptosis in the intermediate erythroblast subset of PCFT−/− splenocytes, resulting in only a modest increase in splenic cellularity (Figure 4B).

**PCFT is indispensable for normal intestinal uptake and tissue accumulation of folate**

Previous studies have described PCFT as a low pH-dependent folate transporter contributing to dietary folate absorption in humans. The presence of hematologic abnormalities in PCFT-deficient mice was also consistent with systemic folate deficiency; therefore, we measured folate concentrations in the peripheral blood and tissues that are known to have high folate content. Plasma, liver, and kidney folate concentrations were all significantly decreased in PCFT−/− and PCFT+/− mice compared with WT littermates (Figure 5A). In contrast, serum concentration of total iron was increased in the mutant mice, which is in agreement with previous reports of elevated systemic iron levels observed during folate deficiency. Liver sections of the mutant mice also showed increased iron accumulation (Figure 5B). On the other hand, PCFT−/− mice registered normal serum and plasma heme concentration (Figure 5A), suggesting that deficiency in heme is not a contributor to the anemia in PCFT−/− mice. Folate depletion is also associated with hyperhomocysteinemia; therefore, PCFT−/− mice were expected to have elevated plasma homocysteine levels. In PCFT−/− mice, plasma homocysteine (Figure 5A) and liver and kidney homocysteine concentrations (Figure 5B) were significantly increased compared with WT mice. Liver sections from WT animals.

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**Figure 5. Absence of PCFT decreases systemic folate content and leads to iron and homocysteine buildup.** (A) Concentration of folate, iron, heme, and tHcy in the indicated tissues was determined in 4- to 6-week-old mice of indicated genotype. Data are presented as in Figure 2; n = 12 mice in each cohort. (B) Gomori iron staining of liver sections (original magnification 40×/0.75 NA dry objective). Iron is detected in periportal hepatocytes of PCFT−/− mice mainly as fine cytoplasmic stippling and rarely as cytoplasmic granules on a diffuse light blue cytoplasmic background. Iron is not detected by Gomori staining in liver sections from WT animals.
tissue content of homocysteine. Observations confirmed increased tHcy concentration in plasma, brain, heart, lungs, liver, and kidney of PCFT−/− mice (Figure 5A).

The main source of folate in uptake is through intestinal absorption. To investigate the role of PCFT in the intestinal transport of folate in vivo, we analyzed the kinetics of folate uptake into various tissues in WT and PCFT-deficient mice after oral gavage administration of radioactively labeled folic acid. As shown in Figure 6A, both homozygous and heterozygous deletion of the PCFT gene led to a dramatic decrease in duodenum-jejunum folate one hour after the oral folate, supporting the essential role of PCFT in the intestinal uptake of folate. In agreement with the deficient intestinal uptake, systemic accumulation of folate in serum, liver, kidney, and spleen was dramatically lower in PCFT−/− and PCFT+/− mice compared with that in WT mice.

To further confirm the function of PCFT as a folate transporter, we measured folate uptake in HEK293 cells transiently transfected with either human or mouse PCFT. PCFT of either species facilitated robust cellular uptake of radioactively labeled folic acid, whereas cells transfected with vector alone showed no significant uptake of folate (Figure 6B). Of note, uptake of zinc protoporphyrin, a measure of heme transport activity, was not affected by expression of the human PCFT in the same cell line.

Folates rescue PCFT−/− mice

To further investigate the role of PCFT in the intestinal transport of folates in vivo, we tested various folate supplementation strategies to demonstrate the biologic function of PCFT. PCFT−/− mice develop folate deficiency anemia and die prematurely because of these conditions, so rescue strategies were developed to increase the length of survival of null pups resulting from PCFT’s function as a folate or heme transporter. Subcutaneous injections of 5-formyltetrahydrofolate (folic acid), 5-MTHF, or folic acid promoted survival in PCFT−/− mice and significantly improved RBC and platelet parameters, including RBC and platelet counts, HCT, and hemoglobin concentration (Figure 7). The results confirmed that folates are the essential nutrients deficient in the PCFT−/− mice, and their replacement via the parenteral route is sufficient to rescue PCFT−/− mice from premature death. Oral supplementation with 5-MTHF at 10 to 40 mg/kg per day also rescued PCFT−/− mice from anemia and thrombocytopenia, indicating sustained transport of 5-MTHF across the gastrointestinal epithelia in the absence of PCFT. In contrast, daily oral supplementation with folic acid at a therapeutic dose (0.1 mg/kg) failed to rescue PCFT−/− mice from anemia and premature death. However, when PCFT−/− mice were treated orally with high doses of folic acid (2 mg/kg), folic acid therapy significantly increased survival and improved RBC and platelet parameters, suggesting the involvement of alternative transport systems in the intestinal uptake of folic acid given at supraphysiologic concentrations. No increases in survival rates were observed in PCFT−/− mice supplemented with oral betaine, folic acid, FeSO4, or hemin injections (data not shown).

Discussion

The PCFT knockout model presented here provides in vivo evidence for a pivotal and nonredundant role played by PCFT in intestinal folate absorption and erythropoiesis. Inactivation of the PCFT gene resulted in a marked decrease of folate uptake in the duodenum-jejunum, leading to folate deficiency. The phenotype of PCFT-mutant mice presented in this report largely recapitulated the hematologic phenotype of dietary folate deficiency in mice and closely resembles the type of anemia reported in folate-deficient humans. In mice, the key features of folate deficiency are growth retardation, anemia associated with a marked, progressive macrocytosis, and pancytopenia accompanied by ineffective hematopoiesis in the BM and spleen. In humans, the disease of folate malabsorption is characterized as macrocytic/megaloblastic anemia because of the accumulation of large immature erythroid progenitors, which undergo apoptosis before complete maturation resulting from impaired DNA synthesis. Likewise, in the absence of the PCFT-mediated intestinal folate transport, mice developed systemic folate deficiency and severe macrocytic normochromic anemia with normal corpuscular hemoglobin and elevated corpuscular volume. PCFT−/− BM showed accumulation of immature
intermediate erythroblasts, which failed to differentiate further and showed increased rate of apoptosis similarly to proerythroblasts isolated from mice fed an amino acid-based, folate-free diet. Marked expansion of erythroid PCFT/−/− precursors and elevated apoptosis of intermediate PCFT/−/− erythroblasts were also evident in the spleen, the primary site of extramedullary erythropoiesis in mice. Consistent with ineffective erythropoiesis and enhanced intramedullary destruction of erythroblasts, we observed an increase in soluble transferrin receptor serum levels in PCFT/−/− mice. Serum soluble transferrin receptor, a truncated monomer of cell surface-expressed transferrin receptor, is primarily released from erythroblasts, and its serum level reflects the erythropoietic activity of BM and spleen. We also observed multilobed polymorphonuclear neutrophils in the PCFT/−/− blood smears and giant metamyelocytes in the PCFT/−/− BM, consistent with abnormal myelopoiesis often observed in patients with folate deficiency-induced anemia. Internal bleeding in the cerebral vasculature and bloody stools were also noted in PCFT/−/− mice, which too is often associated with folate deficiency-induced anemia. Therefore, PCFT deficiency in mice presents a bona fide model of hereditary folate malabsorption syndrome and folate deficiency anemia, recapitulating all the key indicators of the disease. It is also noteworthy that deletion of a single allele of PCFT leads to haploinsufficiency and phenotypic abnormalities in the intestinal transport of folate in vivo, resulting in a systemic folate deficiency. These findings are in agreement with the previous studies in humans diagnosed with hereditary folate malabsorption, indicating that heterozygosity for loss-of-function PCFT mutations results in a clinically significant decrease in folate absorption.

It was noted that the majority of PCFT-mutant mice appeared normal before they were weaned. We therefore hypothesized that forms of folate in the dam’s milk were able to maintain early development, but the synthetic folic acid in adult chow was unable to cross the intestinal epithelia. The folate form found in milk tends to be 5-MTHF, which will dissociate with FOLR1 during gastric passage, but would be expected to bind RFC-1 with relatively high affinity (2 M), compared with folic acid (200 M) in the lower intestinal tract. We therefore tested whether 5-MTHF or folinic acid (40 mg/kg) could rescue PCFT-mutant mice from early death. Oral supplementation with 5-MTHF rescued the majority of PCFT-mutant mice from early death, anemia, and thrombocytopenia, as did injections with folic acid, 5-MTHF, or folinic acid. Increased survival and improved hematopoietic competency in PCFT/−/− mice treated with parenteral folic acid are consistent with the biotransformation of folic acid and sufficient uptake of reduced folic acid derivatives in tissues. Although oral supplementation with pharmacologic doses of folic acid failed to rescue PCFT/−/− mice from premature death and impaired hematopoiesis, except for a slight increase in platelet counts, PCFT/−/− mice supplemented orally with high doses of folic acid displayed a significant increase in survival accompanied by improved RBC and platelet parameters. These findings confirm earlier studies indicating that relatively low folate levels are sufficient to correct hematopoietic abnormalities in patients with hereditary folate malabsorption, whereas
repeated administration of high doses of folic acid is required to increase folate levels in the cerebrospinal fluid and compensate for impaired folate transport into the brain. Direct analysis of the cerebrospinal fluid/serum ratio in PCFT-mutant mice will be required to validate this hypothesis and is currently underway.

Finally, after maturity, supplementation was reduced to weekly doses, and PCFT-mutant mice were noted to live in excess of 300 days, but all PCFT-mutant mice redeveloped folate deficiency anemia 4 to 6 weeks after experimental supplementation was discontinued (R.M.C. and R.H.F., unpublished data, April 6, 2010).

Without proper supplementation, the PCFT-mutant mice displayed several additional abnormal blood parameters that are all typical diagnostic biomarkers for human megaloblastic anemia and reflect the response of BM to anemia and inefficient hematopoietic cell maturation. These included: (1) reduced release of reticulocytes from the BM and spleen resulting from a block in erythroid cell maturation, (2) thrombocytopenia accompanied by abnormal thrombopoiesis and underproduction of platelets, (3) increased serum thrombopoietin concentration as a response to thrombocytopenia, and (4) elevated Epo, sCD71, and iron concentration in the serum resulting from the increased but inefficient erythropoietic activity. Greater iron concentration was also observed in animals fed with folate-deficient diet and could be explained by diminished use of iron for synthesis of hemoglobin in the deficient animals. The elevated iron content and normal heme levels rule out both iron and heme deficiency as a probable cause of anemia in the PCFT-/− mice. Even though PCFT has been originally described as a heme carrier protein, we, like others, could not detect PCFT-related heme transport in vitro, and the normal heme content of blood in the absence of PCFT indicates that this transporter does not play a significant role in heme uptake in vivo. Furthermore, our folate supplementation studies indicate that the heme transport function of PCFT is not essential for normal erythropoiesis as oral supplementation with 5-MTHF in PCFT-mutant mice resulted in the complete rescue from anemia and normalized the length of survival of the nullizygous pups. These findings underline the critical role of PCFT in folate uptake both in vitro and in vivo and provide in vivo evidence for a redundant function of PCFT in heme uptake and metabolism.

Systemic folate deficiency in patients with hereditary folate malabsorption is accompanied by a severe progressive neurologic disorder and intracranial calcifications. However, no evidence of calcification was identified in any sections of PCFT-/− brain tissue by hematoxylin and eosin and von Kossa staining. It is probable that calcification appears in PCFT-/− animals in an age-dependent manner and premature death precludes detection of intracranial calcifications by standard histologic techniques. Finally, our findings suggest that severe thrombocytopenia may lead to subdural or subarachnoid hemorrhages observed in some PCFT-/− mice macroscopically. However, histologic examination failed to detect deep intracerebral hemorrhages in PCFT-/− brain sections. Hemorrhages associated with thrombocytopenia may develop in patients with megaloblastic anemia, and this clinical manifestation of folate deficiency is recapitulated in the PCFT knockout model.

Existing genetic models of folate transport and deficient erythropoiesis only partially replicate common clinical manifestations of human megaloblastic anemia. Targeted disruption of the other folate transporters, RFC1 and FOLR1, results in embryonic lethality in mice. Inactivation of the methionine synthase gene (MS), a B12-dependent enzyme required for demethylation of 5-MTHF to tetrahydrofolate, would be expected to produce a functional folate deficiency by trapping folate as 5-MTHF. However, MS deletion leads to embryonic lethality because of the methyl trap, whereas heterozygous MS+/− mice fail to develop anemia. Megaloblastic erythroid development was described in a mouse model of thiamine-responsive megaloblastic anemia syndrome. Erythroid cells from mice deficient for the high-affinity thiamine transporter, Slc19a2, demonstrated altered thiamine uptake. Accordingly, Slc19a2−/− mice showed some distinctive features of megaloblastic anemia with erythroid hyperplasia in the BM, ineffective erythropoiesis with a dys-synchrony of nuclear and cytoplasmic maturation, increased lobulation of megakaryocytes, and accumulation of giant metamyelocytes. Although these changes affecting erythroid, myeloid, and megakaryocyte lineages are reminiscent of megaloblastic erythroid maturation, it remains undocumented whether Slc19a2−/− mice develop pancytopenia as a result of dysregulated erythropoiesis, thrombopoiesis, and myelopoiesis in the BM, and whether the ineffective erythropoiesis in Slc19a2−/− mice stems from increased apoptosis of late erythroblasts, which is the hallmark of megaloblastic erythropoiesis. Moreover, the phenotypic expression of Slc19a2 homozygosity in the erythroid lineage was documented in the absence of systemic folate deficiency and only in Slc19a2−/− mice challenged with thiamine-deficient diet, thus limiting the relevance of the model to folate deficiency-dependent megaloblastic erythropoiesis. Finally, Fleming et al found no evidence of megaloblastosis in the BM of Slc19a2−/− mice.

In conclusion, the PCFT-deficient mouse line is the most accurate animal model that closely captures the spectrum of pathology typical of human hereditary folate malabsorption syndrome and human folate deficiency anemia. The primary cause of the disease is a disruption of PCFT-mediated intestinal folate uptake, yet the results presented here do not rule out the possibility that PCFT may control folate uptake in other tissues as well. Future studies using PCFT KO mice will help determine the physiologic function of this transporter in more detail and facilitate identification of factors involved in the pathologic mechanisms leading to ineffective hematopoiesis.

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Authorship

Contribution: K.V.S. designed experiments, analyzed data, and wrote the manuscript with extensive contributions from R.M.C., K.A.P., Q.L., R.R., W.C.C., and P.V.; T.O. and R.H.F. edited the manuscript and provided conceptual advice; T.O., R.H.F., K.V.S., and R.M.C. supervised the project; and W.S., W.C.C., C.L., and L.D. performed the experiments.
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A mouse model of hereditary folate malabsorption: deletion of the \textit{PCFT} gene leads to systemic folate deficiency

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