Erythropoietin guides multipotent hematopoietic progenitor cells toward an erythroid fate

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The erythroid stress cytokine erythropoietin (Epo) supports the development of committed erythroid progenitors, but its ability to act on upstream, multipotent cells remains to be established. We observe that high systemic levels of Epo reprogram the transcriptomes of multi- and bipotent hematopoietic stem/progenitor cells in vivo. This induces erythroid lineage bias at all lineage bifurcations known to exist between hematopoietic stem cells (HSCs) and committed erythroid progenitors, leading to increased erythroid and decreased myeloid HSC output. Epo, therefore, has a lineage instructive role in vivo, through suppression of non-erythroid fate options, demonstrating the ability of a cytokine to systematically bias successive lineage choices in favor of the generation of a specific cell type.

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preceding effect on erythroid progenitor numbers, we isolated the bone marrow Lin−Sca-1−c-Kit+ population, which contains the myelo-erythroid progenitors (Akashi et al., 2000), including CFU-E and proerythroblasts (Pronk et al., 2007), at 2 d after injection, and plated these cells under conditions permissive for both myeloid and erythroid differentiation. Staining of the developed colonies using 2,7-diaminofluorescein (DAF) to detect hemoglobin showed an increase in the proportion of erythroid colonies (Fig. 1 A). However, this was accompanied by a decrease in the total colony number (Fig. 1 B).

Plating Lin−Sca-1−c-Kit+ progenitor cells under conditions specifically promoting erythroid (E), megakaryocyte (Mk), B-lymphoid (B), or granulocyte/macrophage (GM) differentiation showed a significant decrease in GM, Mk, and B colony–forming cells, whereas E colony numbers were unchanged (Fig. 1 C). The increase in circulating Epo levels obtained through hydrodynamic injection were comparable to those observed in anemic patients (Jelkmann and Wiedemann, 1990), whereas no change was observed in other lineage-specific cytokines (Thrombopoietin, G-CSF; Fig. 1 D).
decreased (Fig. 1, G and H). Similar skewing of myelo-erythroid progenitor numbers was observed upon injection of recombinant Epo (unpublished data). To determine if lineage skewing occurred in the upstream Lin-Sca-1-c-Kit+ (LSK) hematopoietic stem cell (HSC)/multipotent progenitor (MPP) compartment, this population was analyzed for Flt3 expression, which is normally associated with loss of Mk/E potential and increased lymphoid potential (Adolfsson et al., 2005). We observed that the proportion of Flt3hi lymphoid-primed MPPs (LMPPs; Adolfsson et al., 2005) was decreased upon Epo exposure (Fig. 1 I), consistent with the observed suppression of lymphoid...
were isolated and gene profiled. This showed that CD45.1 on a per-cell basis. 6 wk later, CD45.1 LSKFlt3 deficient CD45.2 HSCs to generate LMPPs remains to be determined. Lineage programming of previous studies have shown that the transcriptional priming of HSCs is an indicator of their lineage potential, and that the priming pattern is regulated by the transcription factors required for subsequent lineage commitment (Bereschkenko et al., 2009; Mancini et al., 2012). To determine whether Epo exposure alters the priming pattern of stem and progenitor cell populations, we mined existing global gene profiling data (Prunk et al., 2007) to identify gene sets highly and selectively expressed in preGMs, preCFU-Es, MkPs, and common lymphoid progenitors (CLPs), representing the first GM, E, M, and lymphoid committed progenitors, respectively (Fig. S1). Gene profiling of in vivo Epo-exposed LSKFlt3− cells showed that genes associated with E lineage commitment were up-regulated, whereas those associated with M and GM lineage commitment were down-regulated (Fig. 2 A). This occurred without any change to the size or composition of the HSC compartment (Fig. 2, B and C), to the cell cycle status of HSC/MPPs (Fig. 2 D), or to HSC repopulating activity (not depicted). Downstream of HSCs, bipotent preMegEs had up-regulated preCFU-E genes, whereas most MkP gene expression was suppressed (Fig. 2 E), consistent with the observed erythroid bias at the MkP/preCFU-E bifurcation. LMPPs showed general down-regulation of CLP and preGM genes (Fig. 2 F). The decrease in LMPP lymphoid gene expression was accompanied by suppressed LMPP B cell differentiation potential (Fig. 2 G), and a decreased number of downstream CLPs (Fig. 2 H). The effect of Epo on GM and T cell potential of LMPPs remains to be determined. Lineage programming of stem/progenitor cells was therefore regulated by Epo, and correlated with altered lineage potential and progenitor output.

To address if Epo acts directly and instructively on the HSC/MPP compartment, we isolated LSKCD150+Flt3− cells and treated these with recombinant Epo in vitro. We observed a virtually identical pattern of regulation of lineage programming as seen upon in vivo Epo exposure (Fig. 3 A). To determine if endogenous erythropoietic stress would cause similar transcriptional HSC reprogramming, CD45.2 Gata1 conditional null (or control) cells (CD45.2) were co-transplanted with wild-type CD45.1 cells into lethally irradiated CD45.1/2 hosts at a 10:1 ratio, followed by Gata1 deletion through poly(I-C) induction of the Mxi1-Cre transgene. This selectively blocks erythropoiesis at the preMegE stage (Mancini et al., 2012), leading to a selective increase in circulating Epo levels (Fig. 3 B) and forcing the wild-type CD45.1 HSCs present along Gata1-deficient CD45.2 HSCs to generate ~10× more erythrocytes on a per-cell basis. 6 wk later, CD45.1 LSKFlt3− HSC/MPPs were isolated and gene profiled. This showed that CD45.1 LSKFlt3− cells present along Gata1-deficient CD45.2 cells up-regulated erythroid genes and concomitantly suppressed preGM and MkP genes (Fig. 3 C). This divergent regulation was disrupted in the Gata1-deficient CD45.2 cells exposed to the same Epo levels (unpublished data). Transcriptional HSC reprogramming therefore occurs in response to exogenous Epo, to endogenous stress erythropoiesis, and to direct in vitro Epo stimulation and is at least partially GATA-1-dependent.

There are several signaling pathways that emanate from the Epo receptor, including the phosphoinositol-3 (PI3) kinase/ Akt, Erk/MAPK, and JAK–STAT pathways (Richmond et al., 2005). To determine their individual contributions to Epo-induced transcriptional reprogramming, LSKFlt3−CD150+ HSCs were exposed to Epo in vitro in the presence of inhibitors specific for these pathways (Fig. 3 D). We observed that up-regulation of preCFU-E genes was consistently dependent on PI3 kinase activation, whereas JAK/STAT and Erk activation was involved in a gene-specific manner. In contrast, preGM gene expression was not significantly affected by PI3 kinase inhibition, whereas, remarkably, Erk inhibition converted the normal inhibition by Epo to robust up-regulation. The major Epo-activated signaling pathways therefore play distinct and lineage-specific roles in reprogramming of the HSC transcriptome. We have previously shown that loss of C/EBPα function leads to loss of myeloid programming in LSKFlt3−CD150+ HSC/MPPs (Bereschkenko et al., 2009). Direct regulation of myeloid genes by C/EBPα may therefore explain the highly coordinated regulation of preGM genes upon inhibitor treatment.

To further investigate the cellular mechanism underlying Epo-induced transcriptional reprogramming, we performed single cell gene expression profiling of control and Epo exposed LSKFlt3− HSCs. We investigated the expression of GM (Mpo, Egr1, Gfi1, Csf1r, and Csf2rl)-, E (Gata1 and Klf5)-, and HSC (Hoxb4, Bmi1, Hes1, and Cxcr4)-associated genes. We observed a 50% increase in proportion of cells expressing one or more E genes (Fig. 3 E). In addition, the average expression of E genes in these cells was increased (1.8-fold, P = 0.10), with a similar decrease in the average expression of GM genes (1.7-fold, P = 0.008; Fig. 3 F). Importantly, we consistently observed coexpression of E genes with one or more GM genes, as well as HSC-associated genes, consistent with the increase in E programming occurring in multipotent stem/progenitor cells, rather than through the expansion of a subpopulation of committed erythroid progenitors.

To establish if Epo-induced reprogramming of stem/progenitor cells resulted in skewing of their lineage output, we transplanted control and Epo-exposed HSCs (defined as LSKFlt3−CD150+; CD45.2) into sublethally irradiated CD45.1 recipients. Transplanted HSCs were isolated from CD45.2 miR144/451GFP/+ mice. This strain contains an EGFP knockin into a microRNA locus uniformly expressed in E lineage cells, including circulating mature erythrocytes, allowing for EGFP-based tracking of E lineage output (Rasmussen and O’Carroll, 2011; Mancini et al., 2012; Fig. 4 A). Analysis of peripheral blood showed that Epo-exposed HSCs generated significantly higher numbers of erythrocytes (Fig. 4, B and C) and fewer myeloid cells (Fig. 4 D). Overall, a fourfold increase of the erythroid/myeloid output ratio was observed.
Figure 3. Cellular and molecular mechanism of Epo action on HSCs. (A) LSKFlt3−CD150+ cells were cultured in the presence (Epo) or absence (Control) of Epo. After 24 h, cells were harvested and analyzed for gene expression. Data are represented as in Fig. 2 A. Values are means, n = 2, from 2 experiments with triplicate measurements. (B) Bar graph showing different serum cytokines levels in CD45.1/2 mice competitively transplanted with 10^6 CD45.1 wild-type cells and either 10^6 CD45.2 Gata1^fl/fl; Mx1-Cre^tg/+ (cKO) or 10^6 Gata1^fl/fl BM cells (Con). Values are mean ± SD, n = 9 (control) and 8 (cKO), from 2 experiments. (C) CD45.1/2 mice were competitively transplanted with 10^5 CD45.1 wild-type and either 10^6 CD45.2 Gata1^fl/fl; Mx1-Cre^tg/+(cKO) or 10^6 Gata1^fl/fl BM cells (Con). Cre recombination was induced by three poly(I-C) injections at 2-d intervals, and CD45.1+CD45.2−LSKFlt3−CD150+ cells were isolated 6 wk after the first poly(I-C) injection and subjected to gene expression profiling. Data are represented as the log2 of the ratio between gene expression in WT/cKO and WT/Control co-transplanted mice after normalization to Hprt. Values are means, n = 2, from 2 experiments with triplicate measurements. (D) LSKFlt3−CD150+ cells were cultured in the presence (Epo) or absence (Control) of Epo. For both conditions, individual cultures were supplied with either vehicle (DMSO) or inhibitors of one of the following kinases: 25 µM Jak2 (AG490), 25 µM PI3K (LY294002), or 25 µM ERK1/2 (PD98059). After 24 h, cells were harvested and analyzed for gene expression. Values are means, n = 2, from 2 experiments with triplicate measurements. (E) Heat map of gene expression in single sorted LSKFlt3− HSCs analyzed by microfluidics-based real time PCR. Expression values were normalized using Kit expression and are shown as deviation from the mean expression value of each individual gene. n = 3 from 3 experiments. (F) Bar graph of mean expression of GM−, E−, and HSC-associated genes in single cells from E. Values represent the mean ΔCt(gene)−ΔCt(Kit) for each of the genes indicated, including only cells where expression was detected. ***, P < 0.0005.
Exposure of HSC/MPPs to Epo is therefore sufficient to bias lineage output away from the myeloid lineage and toward erythropoiesis, even in the absence of exogenous Epo during their further lineage commitment and differentiation.

These results challenge our current view of lineage-specific cytokine action in two ways. First, they show that Epo can act on HSC/MPPs, in addition to E lineage–committed cells, to simultaneously increase E and decrease GM output. Second, we find that this occurs through transcriptional reprogramming, leading to increased E lineage priming and decreased GM, Mk, and lymphoid priming, and the systematic suppression of alternative lymphoid and myeloid lineage options at all lineage bifurcations examined. Up-regulation of erythroid gene expression showed a strong dependence on PI3 kinase signaling. In contrast, Erk signaling was critical for myeloid gene suppression; if Erk activity was blocked, Epo stimulation led to a general up-regulation of myeloid gene expression. This dissociation of lineage reprogramming is consistent with an instructive, rather than selective, effect of Epo on the HSC compartment; Epo exposure could lead to the selective expansion or survival of cells expressing high levels of preCFU-E and low levels of MkP and preGM genes. However, this model is not readily reconciled with the ability of simultaneous Epo stimulation and Erk inhibition to up-regulate CFU-E and preGM genes and suppress MkP genes. Our results therefore show that the high systemic levels of Epo associated with severe anemia can act in a direct and instructive manner on the HSC/MPP compartment, leading to a reduction of GM, Mk, and lymphoid phenotypic progenitors and colony-forming cells. The overall result is the creation of enhanced throughput from HSCs to the E compartment, an “erythroid superhighway,” allowing the focused generation of E-lineage progenitors (Fig. 4E), upon which Epo may then further act through its well-characterized proliferative and pro-survival effects.

**MATERIALS AND METHODS**

**Mouse lines.** C57BL/6J (CD45.2+) mice and congenic B6.SJL–PtprcaPep3b/BoyJ (CD45.1+ congenic C57BL/6J) mice were bred at the University of Edinburgh. The Gata1 conditional allele (Lindeboom et al., 2003), Mx1-Cre transgenic mice (Kühn et al., 1995), and mir144/451-eGFP reporter mice (Rasmussen and O’Carroll, 2011) were previously described. Mouse strains were backcrossed to C57Bl6/J for six or more generations before use, and mice were aged 8–12 wk at the start of experiments. All animal procedures were approved by the University of Oxford Clinical Medicine Ethical Review Committee and were maintained in accordance with Institutional and UK Home Office guidelines.

**Hydrodynamic gene transduction and ELISA.** Mice were tail vein–injected, as described previously (Hirai et al., 2006), in Krebs-Ringer buffer with 1 µg of a pCMV6-Entry vector containing Epo cDNA (OriGene) or empty pCMV6-Entry vector. Epo, Thrombopoietin, and G-CSF serum levels were determined using Quantikine immunoassays (R&D Systems).
Flow cytometry, cell separation, and cell sorting. Flow cytometry was performed essentially as previously described (Bereschkenko et al., 2009). A list of antibodies used is provided in Table S3. Cells were sorted using FACSaria II cell sorters and analyzed using LSRFortessa (BD). All FACS data were analyzed with Flowjo software (Tree Star).

In vitro culture assays. Media for colony-forming assays were from STEMCELL Technologies and used according to the manufacturer’s instructions. 500 cells were seeded per culture in M334, M3434, and Megacult media and 1,000 cells in M3436 and M3630 media. The colonies generated in methylcellulose were scored as BFU-E, CFU-GM, and CFU-MK by morphology 7–10 d after seeding, and as CFU-Mk (in MegaCult assay) by acetylcholinesterase staining. To score colonies in multilineage culture (M3434), colonies were stained with 2,7-diaminofluorene (DAF) staining solution (10 mg/ml DAF in 90% glycerol-acetic acid, 50% H2O2, and 200 mM Tris-HCl, pH 7.5) and DAF-positive cells were identified as cells with intracellular blue granules. Cells were plated in duplicate or triplicate for each assay. B cell potential of LMPPs was evaluated as previously described (Adolfsson et al., 2005) with B-lineage cells defined as Mac-1 assay. B cell potential of LMPPs was evaluated as previously described (Bereshchenko et al., 2009). The donor and competitor HSC/MPPs (LSKFlt3(CD45.1)) were transplanted in 10:1 ratio in recipients (CD45.1/CD45.2). For single cell LSKFlt3 cells were incubated with 200 ng/ml of recombinant murine Epo (R&D Systems) for 24 h, followed by gene expression analysis. Where indicated, cells were treated with 25 μM Jak2 inhibitor (AG490; EMD Millipore), PI3-kinase inhibitor (LY294002; EMD Millipore), Erk1/2 inhibitor (PD98059; Sigma-Aldrich), or DMSO vehicle.

Multiplex specific target amplification and qPCR analysis. Multiplexing of specific target amplification of the cDNA samples from various BM populations and qPCR analysis using UPL assays (Roche) were performed as described previously (Sanjuan-Pla et al., 2013). For each gene, the chosen qPCR assay was the most highly ranked by UPL Assay Design Center.

Single cell gene expression analysis. For single cell LSKFlt3 gene expression analysis, cell capture and target preamplification were performed using the C1 system (Fluidigm) according to the manufacturer’s instructions. Real-time PCR was performed using the BioMark 96.96 Dynamic Array platform (Fluidigm) and DELTAgene assays (Fluidigm; Table S2). In total, >60 single cells from both Epo-exposed and control mice (~20 cells from each of 3 biological replicates) were analyzed, and those scoring positive for Kit mRNA expression (>85% of cells analyzed) were included in the analysis. ΔCt values relative to Kit (Ct(Kit) – Ct(Gene)) were zero centered for each gene by subtraction of the mean value of all positive cells for the gene. These normalized values were used to generate heat maps, and to calculate the cumulative significance of Epo regulation of E and GM genes (i.e., all E genes without Epo against all E genes with Epo, and the same comparison for GM genes) using Student’s t test.

Bone marrow transplantation. To evaluate in vivo capacities to generate erythrocytes and myeloid cells, 500 Lin- Sca1+c-Kit+ CD150+ cells from mIR144/151* arms (CD45.2 allotype) after 2 d of Epo exposure (or control) were transplanted into sublethally irradiated CD45.1 recipients. At the indicated time point, blood samples were evaluated by flow cytometry for the levels of GFP^Ter119^ erythrocytes that were identified by their distinct scatter profile. Peripheral blood analysis was performed as described previously (Kirstetter et al., 2006). For Gata1 conditional knockouts, lethal irradiation, bone marrow transplantation, poly(I-C)-induced recombination, and peripheral blood analysis were performed as described previously (Mancini et al., 2012). To evaluate the effect of E stress on lineage priming of HSCs, Gata1-null BM cells (alloytotype CD45.2) and competitor cells (alloytotype CD45.1) were transplanted in 10:1 ratio in recipients (CD45.1/CD45.2). The donor and competitor HSC/MPPs (LSKFlt3^–/) were sorted for gene expression analysis after 6 wk of first poly(I-C) injection.

Lineage-specific gene sets. Previously published microarray data (Prong et al., 2007) were normalized and analyzed as previously described (Bereschkenko et al., 2009). Genes differentially expressed between progenitor subsets (CLP, preGM, preMegE, MKP, and preCFU-E) were manually curated to identify those selectively and highly expressed in each of the following populations: CLP, preGM, MKP, and preCFU-E (Table S1).

Statistical analyses. Significance for all the data was calculated using Student’s t test.

Online supplemental material. Lineage-specific genes and their normalized expression values are found in Fig. S1 and Table S1. Gating strategies for OP9 cultures and CLPs are shown in Fig. S2. Table S2 contains DELTAgene assays used in single-cell PCR, and Table S3 the list of antibody conjugates used. Online supplemental material is available at http://www.jem.org/cgi/content/full/jem.20131189/DC1.

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Figure S1. Identification of progenitor-specific gene sets. Genes selectively expressed in early lineage-committed progenitors of the erythroid (pre-CFU-E), Mk (MkP), myeloid (preGM), and lymphoid (CLP) lineages. Each value is the mean of at least three individual Affymetrix microarray-based determinations. Differentially expressed genes were manually curated to identify those with specific, high-level expression in each particular progenitor population. The underlying microarray data are listed in Table S1.
Figure S2. Gating strategy for the identification of CLPs and their B cell progeny. (A) Identification of CLP B cell progeny in OP9 single cell cultures. Cells were identified as live singlets based on scatter and 7-AAD staining, and B cells identified as Mac-1⁻NK1.1⁻CD19⁺. (B) Gating strategy for CLPs. For each gate the percentage of the gated population relative to the parental population is indicated. CLPs were defined as Lin⁻/7AAD⁻B220⁻IL7Rα⁻Flt3⁻Sca-1⁻c-Kit⁻ singlets.

Tables S1, S2, and S3 are provided as Excel files.