**Mds1CreERT2**, an inducible Cre allele specific to adult-repopulating hematopoietic stem cells

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**SUMMARY**

Hematopoietic ontogeny consists of two broad programs: an initial hematopoietic stem cell (HSC)-independent program followed by HSC-dependent hematopoiesis that sequentially seed the fetal liver and generate blood cells. However, the transition from HSC-independent to HSC-derived hematopoiesis remains poorly characterized. To help resolve this question, we developed Mds1CreERT2 mice, which inducibly express Cre-recombinase in emerging HSCs in the aorta and label long-term adult HSCs, but not HSC-independent yolk-sac-derived primitive or definitive erythromyeloid (EMP) hematopoiesis. Our lineage-tracing studies indicate that HSC-derived erythroid, myeloid, and lymphoid progeny significantly expand in the liver and blood stream between E14.5 and E16.5. Additionally, we find that HSCs contribute the majority of F4/80+ macrophages in adult spleen and marrow, in contrast to their limited contribution to macrophage populations in brain, liver, and lungs. The Mds1CreERT2 mouse model will be useful to deconvolute the complexity of hematopoiesis as it unfolds in the embryo and functions postnatally.

**In brief**

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**AUTHOR CONTRIBUTIONS**

Y.Z., K.E.M., E.A., P.D.K., H.Y., K.F., K.A.M., S.R., and A.S.P. performed experiments and analyzed data. Y.Z., K.E.M., J.P., and A.S.P. wrote the manuscript.

**DECLARATION OF INTERESTS**

The authors declare no competing interests.

**SUPPLEMENTAL INFORMATION**

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Zhang et al. develop $Mds^{CreERT2}$ mice, which inducibly express Cre-recombinase in emerging fetal hematopoietic stem cells (HSCs). Lineage tracing reveals increasing HSC contributions to hematopoietic cells in E14.5–E16.5 embryos, as well as the majority of macrophages in adult spleen and marrow. $Mds^{CreERT2}$ mice will help deconvolute the complexity of hematopoietic ontogeny.

**Graphical Abstract**

**INTRODUCTION**

The emergence of the hematopoietic system during embryonic and fetal development has been the focus of extensive study, both for its paradigmatic value but also vis-à-vis its relevance to human diseases. Studies primarily in the murine model system indicate that the ontogeny of the hematopoietic system consists of two broad programs, an initial HSC-independent program followed by the emergence and establishment of an HSC-dependent program that arises from arterial vessels within the body of the embryo. HSC-independent hematopoiesis is complex, consisting of overlapping waves of primitive hematopoietic progenitors, erythro-myeloid progenitors (EMPs), and small numbers of lymphoid progenitors (Böiers et al., 2013; McGrath et al., 2015; Palis et al., 1999; Yoshimoto et al., 2011, 2012). Primitive hematopoiesis is derived from progenitors that emerge in the yolk sac beginning at embryonic day 7.25 (E7.25), producing circulating maturing primitive erythroid, megakaryocyte, and macrophage cells by E10 (Palis et al.,...
Lineage-tracing experiments indicate that these first macrophages persist as adult self-renewing tissue-resident microglia (Hoeffel et al., 2015; Palis et al., 1999; Stremmel et al., 2018). Multilineage EMPs emerge from hemogenic endothelium in the yolk sac from E8.25 through E10.5 and seed the fetal liver as early as E10 to generate definitive erythroid, megakaryocyte, granulocyte, and monocyte lineage cells (Frame et al., 2016; Gomez Perdigueró et al., 2015; Lux et al., 2008; McGrath et al., 2015; Palis and Yoder, 2001). EMPs also provide tissue-resident macrophages that persist into adulthood in multiple organs including the liver, lungs, and skin (Gomez Perdigueró et al., 2015). Rare B cell and T cell progenitors arise independently of HSCs in the yolk sac and embryo proper by E9.5 (Yoshimoto et al., 2011, 2012), and lymphomyeloid-restricted progenitors (LMP) are already present in the fetal liver by E10.5, before HSC activity is detectable there (Boiers et al., 2013; Luis et al., 2016).

Small numbers of HSCs capable of long-term reconstitution of adult mouse recipients begin to emerge at E10.5 from the ventral aspect of the dorsal aorta (Yokomizo and Dzierzak, 2010). At E11.5–12.0, HSCs are found in the fetal liver where they subsequently expand in numbers (Ema and Nakauchi, 2000; Kumaravelu et al., 2002). However, the transition from HSC-independent to HSC-derived hematopoiesis in the murine fetus remains poorly characterized, particularly since the maturing erythroid, megakaryocytic, and myeloid progenies of EMPs and HSCs are not currently easily distinguishable. Several different inducible lineage-tracing approaches have been used to distinguish HSC-independent versus HSC-derived contributions based on promoters active both in HSCs and in HSC-independent progenitors, including Cdh5 (Zovein et al., 2008), CSF1R-Cre (Qian et al., 2011), Kit-Cre (Stremmel et al., 2018), Runx1-Cre (Samokhvalov et al., 2007), Tie2-Cre (Gomez Perdigueró et al., 2015), and TLR2-Cre (Balounová et al., 2019). The efficacy of these CreER alleles relies on the timing of induction during the staggered emergence of HSC-independent progenitors and of HSCs. However, the significant temporal overlap in the emergence of HSC-independent progenitors and of HSCs, coupled with the broad cellular expression of these promoters, and the imprecision of in vivo CreER induction often results both in incomplete and in mixed labeling (Gentek et al., 2018; Gomez Perdigueró et al., 2015; Hoeffel et al., 2015; Soares-da-Silva et al., 2021). An alternative approach is to specifically label emerging HSCs. Studies of emerging HSCs have identified several genes preferentially or specifically expressed in HSCs, including Hlf (Yokomizo et al., 2019), Ly6A (Sca-1; (Ma et al., 2002), Procr (Iwasaki et al., 2010), and Mecom (Yuasa et al., 2005).

The Mecom or Mds1-Evi1 complex locus was first identified as a site of proviral insertion in murine myeloid leukemias (Morishita et al., 1988; Mucenski et al., 1988) and was also found to be activated via chromosomal rearrangements in human AML (Bitter et al., 1985; Rubin et al., 1990). Importantly, the Mds1-Evi1 gene harbors two widely separated transcription start sites (TSSs) (Fears et al., 1996) that encode a series of different isoforms, with the upstream TSS encoding MDS1-EVI1, a 160 kDa isoform harboring an N-terminal PR domain, thus placing this isoform in the PR/SET family of transcriptional regulators and chromatin modifiers (Hohenauer and Moore, 2012). The downstream TSS encodes at least three zinc finger protein isoforms of EVI1 (135/145, 123, and 103 kDa). Roles for Mecom in HSCs during embryogenesis have been documented by several groups. In the mouse,

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homozygous loss of all but the 103 kDa isoform results in loss of competitive repopulating ability within the para-aortic splanchnopleura and lethality by E10.5 (Hoyt et al., 1997; Yuasa et al., 2005). Knockout of all isoforms is associated with HSCs in the fetal liver that lack competitive repopulation ability and with lethality at E13.5 (Goyama et al., 2008). In contrast, the elimination of the 160 kDa MDS1-EVI1 isoform results in viable mice with diminished numbers of adult HSCs (Zhang et al., 2011) and with skeletal defects (Juneja et al., 2014). Using a zebrafish model, Konantz et al. (2016) documented expression of Evi1 in the ventral aspect of the dorsal aorta, and downregulation of Evi1 led to impaired HSC emergence from hemogenic endothelium.

To more clearly define the role of the different Mecom isoforms in the mammalian hematopoietic system, we introduced a lacZ reporter into the upstream TSS of murine Mecom (Mds1\textsuperscript{lacZ}) facilitating the examination of its expression histologically within the murine embryo. Mds1\textsuperscript{lacZ} is expressed in Runx1\textsuperscript{+} clusters in the ventral aspect of the dorsal aorta of the mouse but is not expressed in the fetal liver (Zhang et al., 2011). To probe this further, we developed a Cre-ER knockin into the Mds1 locus (Mds1\textsuperscript{CreERT2}). Lineage-tracing studies using this mouse model revealed that the progeny of Mds1-expressing cells are subsequently found in the fetal liver as immunophenotypic and functional LT-HSCs. In contrast, neither EMPs themselves nor their progeny were labeled, nor were the progeny of primitive hematopoietic progenitors. Lineage tracing with Mds1\textsuperscript{CreERT2} demonstrated that HSC-derived hematopoietic progeny are found in increasing numbers in the livers of E14.5 to E16.5 murine embryos and contribute significantly to circulating erythroid, myeloid, and lymphoid cells by E16.5. In addition, while varying degrees of tissue-resident macrophages in the brain, liver, and lung of adult mice originate from HSC-independent sources, the large majority of F4/80\textsuperscript{+} macrophages in the spleen and bone marrow of adult mice are lineage labeled, consistent with their derivation from HSCs. The Mds1\textsuperscript{CreERT2} model will be useful to address multiple questions regarding the complexity of the hematopoietic system as it unfolds in the murine embryo and continues to function postnatally.

RESULTS

Embryonic expression of Mecom

We reported previously the expression pattern of Mecom transcripts during embryogenesis using in situ hybridization probes that hybridized to both Evil and Mds1 transcripts. These studies revealed Mds1-Evili transcripts in the developing limb bud, lung epithelium, and developing renal system among other locations, with no discernable expression in the fetal liver (Perkins et al., 1991). In an effort to understand more precisely where Mds1 is expressed in the embryonic hematopoietic system, we created a nuclear localized NLS-lacZ knockin allele that puts the lacZ gene under transcriptional control of the Mecom upstream promoter (termed Mds1) (Fears et al., 1996; Zhang et al., 2011). Using this allele, termed Mds1\textsuperscript{lacZ} (referred to as Mds1\textsuperscript{m1} in Zhang et al., 2011), we conducted whole-mount labeling of E8.5 to E14.5 embryos with 5-bromo-4-chloro-3-indolyl-\textbeta-D-galactopyranoside (X-gal). \textbeta-galactosidase staining was evident in the limb buds, mesonephros, and vertebral bodies (Figures 1A–1E), overlapping considerably with the expression of Mds1-Evii described previously by in situ hybridization (Perkins et al., 1991).
Coronal sections of E9.5, E10.5, and E11.5 embryos (Figures 1F–1H; Figures S1A and S1C) revealed LacZ staining of endothelial cells on the ventral aspect of the dorsal aorta, as well as in intra-aortic hematopoietic clusters (IAHC). Importantly, these LacZ-positive aortic endothelial cells were found to co-express Runx1, consistent with a hemogenic endothelial identity (Figure 1I; Figures S1B and S1D) (Yzaguirre et al., 2017). Given that HSCs are known to seed and expand within the fetal liver, we expected to detect β-galactosidase-staining cells in the fetal liver. Paradoxically, we did not find evidence of Mds1-Evi1 expression in the fetal liver (Figure 1J), consistent with previous findings (Perkins et al., 1991). These data suggest that the expression of Mds1-Evi1 in the fetal hematopoietic system is remarkably transient, in contrast to the expression of Mds1-Evi1 in adult LT- and ST-HSCs (Zhang et al., 2011).

**Generation of a mouse model of TAM-inducible Cre recombinase knocked into the Mds1 locus**

To assess whether the β-gal-positive cells identified at E9.5 in the dorsal aorta of Mds1lacZ mice represent functional hematopoietic cells, we performed lineage-tracing experiments. We constructed an Mds1CreERT2 allele in which a tamoxifen (TAM)-inducible Cre recombinase was inserted into the Meconn locus so it would be expressed by the upstream TSS (Figures 2A and 2B). We first performed lineage tracing with the Mds1CreERT2 allele in adult mice by combining it with the ubiquitously transcribed, Cre-dependent Rosa26loxP-STOP-foxP-YFP (Rosa26LSL-YFP) reporter. Based on the restricted expression of Mds1-Evi1 to adult LT- and ST-HSCs, as well as a small percentage of MPPs (Christodoulou et al., 2020; Zhang et al., 2011), we expected that activation of Mds1CreERT2 via TAM treatment would result in increasing and broad YFP positivity in hematopoietic cells over time. Mds1CreERT2 Rosa26LSL-YFP mice received five treatments of TAM over 5 days (Figure 2C). Analysis of bone marrow at 4, 10, and 15 days (relative to the first treatment) revealed a time-dependent increase in YFP+ cells (Figure 2D). By 15 days, up to 40% of marrow cells were YFP+ and they consisted of myeloid, erythroid, and lymphoid cells (data not shown). These labeled cells in the marrow contributed to YFP+ progeny detected in the peripheral blood at 30 days (Figure 2E).

**Lineage-tracing preHSCs and fetal liver cells with Mds1CreERT2**

To test whether the Mds1lacZ-expressing cells present in the dorsal aorta might contain precursors of fetal liver hematopoiesis, we conducted lineage-tracing studies using the Mds1CreERT2 allele crossed with a Rosa26SL-lacZ reporter. Timed pregnant mice were treated with a single dose of TAM either at E7.5 or at E9.5, and embryos were harvested and stained for β-gal at E13.5. Labeling at E9.5 with TAM lineage traced small numbers of β-gal-positive cells in the fetal liver at E13.5, which were not seen when TAM was delivered at E7.5 (Figures 2F and 2G). While fetal liver cells themselves do not express Mds1, the fetal liver contains a population of cells derived from earlier Mds1-expressing cells.

We next investigated the specificity of the Mds1CreERT2 lineage-tracing mouse model by analyzing E11.5 fetuses. Pregnant Mds1CreERT2/wt Rosa26dSL-YFP/LSL-YFP mice were treated with TAM at E9.5 and cells from aorta-gonad-mesonephros (AGM), yolk sac, blood, and liver in Mds1CreERT2/wt Rosa26dSL-YFP/LSL-YFP (referred to as Mds1CreERT2 Rosa26dSL-YFP)
fetuses were analyzed by flow cytometry. Cells from littermate negative control Mds1WT/WT Rosa2dSL-YFP/wt mice were used to determine bone fide YFP+ expression in all experiments (data not shown). Consistent with the staining seen with Mds1lacZ, we found YFP+CD31+kit+VE-cadherin+ pre-HSCs ((Azzoni et al., 2018; Gao et al., 2020; Yokomizo and Dzierzak, 2010) in the AGM and in the circulation. In contrast, we found no labeling of CD41+kit+CD16/32+ EMPs in the yolk sac, Ter119+ primitive erythroid cells (EryP) in the bloodstream (Kingsley et al., 2004), or EMP-derived definitive erythroid cells (EryD) in the liver (McGrath et al., 2015; Gomez Perdiguero et al., 2015) (Figures 2H and 2I; Figures S2A and S2B). Other HSC-independent CD45+ early myeloid cells and CD41+ megakaryocytes in the bloodstream and in the liver were also not labeled (Figure 2I). The Mds1CreERT2 promoter did not label HSC-independent hematopoietic cells in E11.5 fetuses even when activated at E8.5 (Figure S2C). TAM treatment at E8.5 also did not label pre-HSCs, despite labeling a subset of cells in the paw, consistent with Mds1 expression in limb buds (Figure S2C; Figure 1).

We next asked whether the Mds1CreERT2 can label functional HSC in the murine embryo. As E12.5 is the first time that a significant number of transplantable HSCs are found in the liver (Ema and Nakauchi, 2000; Kumaravelu et al., 2002), pregnant Mds1CreERT2 Rosa2dSL-YFP mice were treated with TAM at E9.5, and blood and liver were analyzed at E12.5. As seen at E11.5, primitive erythroid cells in the blood stream and definitive erythroid and granulocyte progeny of EMPs in the liver were not labeled (Figure 2I; Figures S2D and S2E). In contrast, YFP+ cells in the E12.5 liver were found to constitute a subset of Kit+ cells, suggesting that Mds1CreERT2 might be lineage-labeling HSCs. To directly test this hypothesis, we transplanted 2 × 10⁶ E12.5 Mds1CreERT2 Rosa2dSL-YFP liver cells (induced with TAM at E9.5) into irradiated syngeneic recipient mice along with 2 × 10⁵ wild-type bone-marrow cells. At 12 weeks post-transplant, peripheral blood from these recipients revealed a high percentage of YFP+ T cells, B cells, and myeloid cells (Figure 2K; Figure S2F). Taken together, these data demonstrate that Mds1CreERT2, combined with TAM induction at E9.5, lineage-traces fetal HSCs but not primitive erythropoiesis or EMP-derived hematopoiesis.

Mds1CreERT2 lineage tracing of later fetal hematopoiesis

We utilized the Mds1CreERT2 lineage-tracing mouse model to examine the transition from HSC-independent to HSC-derived hematopoiesis in livers of E14.5 and E16.5 mouse embryos (Figure 3). The level of labeling of HSC-enriched LSK (lineage−, Sca1+, Kit+) cells was used to normalize for different efficiencies of excision. In contrast to E12.5 livers, YFP+ Ter119+ (definitive erythroid), Ly6G+ (granulocyte), Ly6C+ (monocyte), and CD19+ (lymphoid) cells were readily evident at E14.5. This suggests that HSCs have begun to contribute to multiple hematopoietic lineages by E14.5 (Figure 3A; Figures S3A and S3B). Furthermore, at E14.5, the higher contribution of Mds1CreERT2-labeled cells to the hematopoietic progenitor (LK, Lineage−, Sca1−, Kit+) compartment compared to the downstream lineage-positive cells suggests that HSC contribution to fetal hematopoiesis is expanding (Figure 3A). Indeed, this is evident at E16.5, with the increased contribution of HSCs to the LK hematopoietic progenitor pool, as well as downstream to maturing erythroid-, granulocyte-, and monocyte-lineage cells (Figure 3B, left panel). Consistent with
the progressive contribution of HSCs to fetal hematopoiesis, labeled myeloid cells and red blood cells are now also found in the bloodstream (Figure 3B, right panel, Figure S3C). Since the Rosa26 locus is progressively downregulated in maturing erythroid cells (Asari et al., 2004; Zambrowicz et al., 1997), the YFP+ fraction of circulating erythrocytes may underrepresent the full contribution from HSCs. Further analysis of E16.5 hematopoietic stem and progenitor cells indicated that fetal HSCs contribute equally to all lineage-biased MPP subsets, and to a lesser, but uniform degree to lineage-committed lymphoid (CD127+), myeloid (CD16/32+), and erythroid/megakaryocyte (CD16/32−) progenitors (Figure 3C; Figure S3D; Pronk et al., 2007). Taken together, these data indicate that HSCs provide an increasing contribution to hematopoiesis in the fetal liver between E14.5 and E16.5, the time points associated with significant expansion of HSC numbers and differentiating hematopoietic progenitors and precursors in the fetal liver (Ema and Nakauchi, 2000; Kumaravelu et al., 2002).

Several tissue-resident macrophage populations, established from HSC-independent sources, persist through embryogenesis as self-renewing populations (Gomez Perdiguero et al., 2015; Hoeffel et al., 2015; Liu et al., 2019). This is particularly evident for microglia, which are seeded early in development and not normally replaced by later fetal or adult myeloid cells (Gomez Perdiguero et al., 2015; Hoeffel et al., 2015). Consistent with these findings, no YFP labeling of F4/80+ macrophage cells in the E14.5 or 16.5 brain was evident in \textit{Mds1}^{CreERT2} fetuses (Figure 3D; Figure S3E). In contrast, a high proportion of macrophages in the fetal liver are HSC-derived by E16.5 consistent with contribution of HSCs to the monocyte lineage (Figures 3B and 3D).

**Lineage tracing of hematopoiesis in adult mice**

TAM administration at E9.5 clearly activated Cre-ER in fetal cells capable of long-term engraftment when transplanted into adult recipients (Figure 2K). However, differences in the regulation of Flt3 in subsets of fetal HSCs has indicated that there is heterogeneity in fetal transplantable HSCs, including subsets that may not normally contribute to adult HSCs (Beaudin et al., 2016). To directly test whether precursors of adult HSCs were being labeled in the \textit{Mds1}^{CreERT2} mouse model, E9.5 TAM-treated pregnancies containing \textit{Mds1}^{CreERT2} \textit{Rosa}^{LSL-YFP} fetuses were allowed to come to term. After 1 year, these mice were analysed for YFP+ cells both in blood and in the bone marrow. YFP+ cells constituted nearly a third of the leukocytes in the blood, and similar proportions of bone-marrow immunophenotypic LT-HSCs, MPPs, more restricted hematopoietic progenitors, as well as kit-CD45+ cells (Figures 4A and 4B; Figure S4A). Treatment with TAM at E8.5 did not lead to the labeling of hematopoietic cells in the adult, consistent with lack of preHSC labeling at E11.5 (Figure S2E).

We also examined the contribution of HSCs to F4/80+ tissue-resident macrophage populations in the adult. As expected, there was no HSC contribution to microglia, and lower but increasing contribution to macrophages in liver, lung, and kidney (Figure 4C; Figure S4B). Surprisingly, a high proportion of F4/80+ macrophages in the spleen and bone marrow were labeled, indicating that they are predominately HSC-derived. \textit{Mds1}^{CreERT2} \textit{Rosa26}^{SL-LSL-YFP} activated by TAM at E8.5 did not lead to the labeling of any of these
DISCUSSION

HSCs, which transiently emerge in the embryo from hemogenic endothelial cells in large arterial vessels, serve as the foundational source of all circulating blood cells in the postnatal organism. A number of genes, including Mecom, Hlf, Procr, and HoxA family members, have been specifically associated with HSC emergence in mice (Iwasaki et al., 2010; Ng et al., 2016; Yokomizo et al., 2019). Mecom is one of the most differentially expressed genes present in hemogenic endothelium of the aorta compared both to hemogenic endothelium in the yolk sac and to non-hemogenic endothelial cells (Baron et al., 2018; Gao et al., 2018; Solaimani Kartalaei et al., 2015; Yokomizo et al., 2019). Recently, Mecom expression has been identified in primary murine aortic arterial-identity cells, as well as in their subsequent transition in pseudotime to hemogenic endothelium and preHSC identity cells (Hou et al., 2020; Zhu et al., 2020). The MECOM locus is complex, with multiple isoforms expressed from two TSS. Here, we report the development of an Mds1\textsuperscript{CreERT2} mouse model that contains a TAM-inducible Cre cassette within the first TSS of the MECOM locus. Utilizing an Mds1\textsuperscript{lacZ} mouse (Zhang et al., 2011), we found expression of the LacZ reporter in a subset of cells of the E9.5 aorta that also express Runx1 (Figures 1G–1I), consistent with the reported expression of MECOM in aortic endothelial cells as they transition to hemogenic endothelium (Konantz et al., 2016; Yokomizo et al., 2019; Zhu et al., 2020, Hou, 2020 #42). Our data suggest that the MECOM locus is transiently expressed during HSC emergence and subsequently downregulated in the fetal liver, unlike its expression in LSK cells in adult mice (Zhang et al., 2011). Our RNA expression studies in adult tissues revealed that the two TSSs are coordinately regulated (Zhang et al., 2011), and analysis of publicly available transcription factor binding data reveals coordinate binding of transcription factors to the two TSSs, as well as similar histone markings and DNaseI hypersensitive sites (unpublished findings). However, the factors regulating transcription from the Mds1 and Evi1 promoters in adults or during development are poorly characterized and are an avenue for future research.

Subsets of myeloid and specialized lymphoid cells are formed in the fetus and self-renew postnatally with little input from adult HSCs (Gentek et al., 2018; Ghosn and Yang, 2015; Hashimoto et al., 2013). The overlapping emergence and lineage output of HSC-independent and HSC-derived hematopoiesis raise important questions regarding the identity and potential functional differences of their mature progeny. Exploring these questions will require ways to distinguish and isolate cells derived from different developmental sources. Current lineage-tracing approaches rely largely on the temporal induction of promoters, including Cdh5 (VECadherin), Runx1, Csf1r, and Kit, that are expressed both in HSC-independent progenitors and also in HSCs (Gomez Perdiguero et al., 2015; Hoeffel et al., 2015; Soares-daSilva et al., 2021; Stremmel et al., 2018; Zovein et al., 2008). An alternative approach would be to use promoters for genes such as Hlf, Procr, or Mecom that are expressed more specifically in fetal HSCs (Iwasaki et al., 2010; Yokomizo et al., 2019; Yuasa et al., 2005). Although a caveat common to all of these systems, including Mds1\textsuperscript{CreERT2}, is that it remains unclear to what degree rare lymphoid
or lympho-myeloid progenitors might also be labeled. Our studies reveal the usefulness of the \textit{Mdx^{CreERT2} Rosa26^{LSL-YFP}} mouse model that labels HSCs as they first emerge from hemogenic endothelium, but does not label the major HSC-independent waves of primitive hematopoietic progenitors and EMPs (Figure S4C).

EMP's begin to colonize the fetal liver between E9.5 and E10.5 and rapidly provide differentiating progeny, particularly the first circulating definitive erythrocytes (McGrath et al., 2015; Palis et al., 1999). Transplantable HSCs subsequently start to seed the liver between E11.5 and E12.5; however, the transition between these two sources of blood cells has only recently begun to be scrutinized (Ema and Nakauchi, 2000; Kumaravelu et al., 2002; Soares-da-Silva et al., 2021). Our analysis of TAM-treated \textit{Mdx^{CreERT2} Rosa26^{LSL-YFP}} embryos indicates that HSCs have begun to generate small numbers of differentiating erythroid, myeloid, and lymphoid progeny between E12.5 and E14.5. By E16.5, a significant proportion of differentiating erythroid, myeloid, and B cell lineage cells in the liver are HSC-derived, and HSCs have now begun to contribute erythroid and myeloid cells to the rapidly expanding pool of circulating blood cells. A similar result was reported in a recent paper that found increased HSC contribution to hematopoiesis from E14.5 to E16.5 (Soares-da-Silva et al., 2021). These authors also observed a strong lineage bias, concluding that erythropoiesis is HSC independent until birth, which we did not observe. This could possibly be due to the challenges of using different lineage-tracing mouse models. Our data support the conclusion that HSCs contribute increasing proportions of erythroid, myeloid, and lymphoid cells to the rapidly developing late gestation murine fetus, supplementing the robust blood cell output of EMPs prior to birth.

Many tissue-resident macrophage populations in adult organs contain contributions of fetal cells that persist by self-renewal. These self-renewing macrophage populations were thought to derive primarily from HSC-independent sources. Here, we used the \textit{Mdx^{CreERT2} Rosa26^{LSL-YFP}} model to examine the contribution of HSCs to F4/80$^+$ macrophage populations in several organs over developmental time. As expected, we found no evidence of HSC contribution to microglia, but increasing contributions of HSCs to macrophages in liver, lung, and kidney (Gomez Perdiguero et al., 2015; Hoeffel et al., 2015; Liu et al., 2019). Interestingly, the majority of F4/80$^+$ cells in the adult spleen and bone marrow, though also self-maintaining (Hashimoto et al., 2013), appear to be predominantly derived from fetal HSCs (Figure 4C). Macrophage cells in the marrow perform multiple functions, including regulation of the HSC niche, support of erythroid precursor maturation in erythroblastic islands, and clearance of pyrenocytes. The surprisingly high proportion of HSC-derived F4/80$^+$ cells in the fetal compared to the adult liver may reflect the need for those macrophage cells to also fulfill these specific functions in the fetus.

**STAR METHODS**

**RESOURCE AVAILABILITY**

**Lead contact**—Further information and requests for resources should be directed to Archibald Perkins (archibald_perkins@urmc.rochester.edu).
Material availability—Mds1\(^{lacZ}\) mice (also referred to as Mds1\(^{m1}\) mice) have been submitted to MMRRC (Mds1-lacZ, Stock No. MMRRC:68126); please contact the MMRRC regarding availability. Mds1\(^{CreERT2}\) mice (also referred to as Mds1\(^{m2}\) mice) have been submitted to Jackson Labs for distribution (Mds1-CreERT2, Stock No. 032863). Please contact Jen Merriman regarding availability (Jen.Merriam@jax.org). There are no restrictions on the distribution of these lines.

Data and code availability

- Any data reported in this paper will be shared by the lead contact upon request.
- This paper does not report any original code.
- Any additional information requires to reanalyze the data reported in this paper is available from the lead contact upon request.

EXPERIMENTAL MODEL AND SUBJECT DETAILS

Mice—Mds1\(^{lacZ}\) mice have been described previously (Zhang et al., 2011). Mds1\(^{CreERT2}\) (also referred to as Mds1\(^{m2}\)) mice were generated as described below. Rosa26-YFP reporter mice (Srinivas et al., 2001) Gt(ROSA)26Sor\(^{tm1(EYFP)Cos/J}\), Stock No: 006148, abbreviated Rosa26\(^{YFP}\) and C57BL/6 were obtained from Jackson Labs. Both female and male embryos and adults were analyzed in these experiments. For timed pregnancies, dams were aged 6–16 weeks; sires were aged 7–60 weeks. For bone marrow transplantation, recipient mice were aged 8–12 weeks. All animals were housed and cared for in facilities operated by the Division of Animal Care at the University of Rochester Medical Center and used in accordance with the guidelines of the University of Rochester School of Medicine and Dentistry Institutional Animal Care and Use Committee.

METHOD DETAILS

Construction of Mds1\(^{CreERT2}\) allele—Plasmid construction was carried out using BAC recombineering (Liu et al., 2003). Bacterial artificial chromosome for Mecom (clone RP24–120B18) was introduced via electroporation into bacterial strain DY380 to create GS1650. The ApSacB cassette was then inserted into exon 1 of Mecom destroying the exon’s splice donor to create GS1656. The CreERT2 cassette (Metzger et al., 1995), derived from plasmid pXL119 (Zhao et al., 2007), together with a FRT-neo-FRT cassette, was inserted into GS1656, replacing the ApSacB cassette, to create GS1690. The PciI site of plasmid pL253 (Liu et al., 2003) was changed to an NsiI site, creating p1342. The NotI site of p1342 was altered to a FseI site, creating p1343. The NsiI fragment of GS1690 was then transferred into p1343 to create the final plasmid, p1347, which was linearized with FseI and submitted to the Genome Editing Facility at the University of Rochester, which created embryonic stem cell clones that were screened by Southern blotting, cutting DNAs with EcoRV for both the 5° and 3° hybridization probes. Ten positives were identified from 192 clones, one of which was used to create mice. These were then crossed with a FLPE deleter strain (Gt(ROSA)26Sor\(^{tm1(FLP1)DymRainJ}\)) to delete the Frt-neo-Frt cassette; the resultant allele was designated Mds1\(^{CreERT2}\).
Time pregnancy and competitive BM transplantation—Time pregnancy experiments were setup using male Mds1+/CreERT2 Rosa26<sup>dSL-YFP</sup>LSL-YFP crossed with Rosa26<sup>dSL-YFP</sup> or C57BL/6 females. Mice were mated overnight and the morning of vaginal plug detection was considered embryonic day (E) 0.3. Pregnant mice were treated with tamoxifen (TAM) freshly dissolved in corn oil at 10mg/ml and 1mg/mouse delivered by gavage at E7.5, E8.5 or E9.5, and harvested at E11.5, E12.5, E14.5 or E16.5 for tracing embryonic hematopoiesis, or harvested 12 months after birth for long term tracing experiments. For competitive repopulation assays, C57BL/6 recipient mice were given 10 Gy of radiation, split in two 5-Gy doses separated by 24 hours and injected i.v. with 2 × 10<sup>6</sup> E12.5 liver cells mixed with 100,000 C57BL/6 control bone marrow cells.

Tissue collection and processing—For complete blood count (CBC) values, blood was collected by capillary action by holding a heparinized 147 mm glass blood collection tube to a razor-pricked mouse tail. CBC measurements on fresh blood samples were taken using a scil Vet ABC Plus+ hematology analyzer.

Marrow cells were collected from mouse tibias and femurs as follows. Mice were euthanized with carbon dioxide and cervical dislocation, and tibia and femurs were dissected free of muscle, tendons, and connective tissue (Amend et al., 2016) and placed into room temperature sterile phosphate-buffered saline (PBS). Marrow contents were flushed from femurs with 1 mL of PBS by cutting the bones at the distal ends of the central marrow cavity. Pooled marrow contents in PBS were washed once with PBS and used in subsequent hematopoietic analyses (see section below: Flow Cytometry).

Embryonic tissues were isolated from staged embryos were dissected from dams as follows. Dams were euthanized with carbon dioxide and cervical dislocation. Uteri were removed from the peritoneum and washed with several changes of phosphate-buffered saline (PBS). Embryos were dissected free of decidual tissues and Reichert’s membrane in IMDM/PBS with 0.2% delipated BSA using #5 watchmakers forceps (Dumoxal; Electron Microscopy Science, Ft. Washington, PA). Vestiges of maternal blood cells were removed by multiple transfers of tissues into fresh medium during the dissection process. Somite stage embryos (E8.25-E11.5) were grouped according to somite number. Most of the blood was derived from the larger vessels of the embryo proper. At E > 11.5, the liver was isolated separately from the remainder of the embryo proper. Embryonic livers were dissociated with Type I Collagenase (Stem Cell Technologies, diluted 1:3 with Dulbecco’s Phosphate Buffered Saline (DPBS or PB2), with 0.1 g/L CaCl2, 0.3% BSA (Gemini Bio-Products) and 0.1% glucose with gentle pipetting (McGrath et al., 2011). Cells were counted and used for Flow Cytometry (see below) or transplantation into irradiated recipient mice (see Competitive BM transplantation section above).

Histochemical and β-galactosidase staining—Timed pregnancies between dams and sires of appropriate genotypes. Embryos at different stages were harvested and dissected clear of yolk sac, amniotic tissue, and placenta. β-galactosidase staining (Loughna and Henderson, 2007; Nagy et al., 2007; Sanes et al., 1986) of whole-mount embryos was performed as follows: Embryos were fixed in 0.4% paraformaldehyde (in PBS, pH 7.4) on ice for 15 minutes for embryos up to E12.5 dpc and for 30 min for E13.5-E14.5.
Following fixation, embryos were rinsed 3 times in PBS. Embryos were then stained in 2 mM MgCl₂, 0.02% IGEPAL, 0.1 M phosphate buffer pH 7.2, 5 mM potassium ferricyanide, 5 mM potassium ferrocyanide, 0.01% sodium deoxycholate, and 1 mg/ml X-gal in a 24-well tissue culture plate, incubating at 37°C in a humidified tissue culture incubator until color develops (2–8 hr). After the staining is complete, embryos were rinsed 4–5 times in PBS, then post-fixed in 2% paraformaldehyde/PBS overnight at 4°C. Embryos were then stored in 70% ethanol. Whole-mount photography was performed on a Zeiss 9901 Stereo Zoom microscope. Embryos were then dehydrated, embedded in paraffin, sectioned, and counterstained with eosin. Slides were then dehydrated and coverslipped.

**Runx1 Immunohistochemistry**—β-galactosidase-stained sections were hydrated, treated with 3% hydrogen peroxide, and subjected to HIER with 10mM Citric acid pH6, 0.05% Tween. Sections were blocked in PBST with 5% BSA and 5% goat serum, incubated overnight at 4°C with Rabbit antiRunx1 (Abcam Ab92336), washed and incubated with an AF647 secondary antibody (Thermofisher A-21244). Sections were counterstained with DAPI and mounted in antifade Prolong Gold.

**Flow cytometric analysis**—All antibody used in this study are listed in Key Resources Table and combinations and gating are shown in Supplemental Figures. Flow cytometry was performed on a LSRii or LSR Fortessa X20 (BD Biosciences) and analyzed with FlowJo v10.3 software (TreeStar).

**QUANTITATION AND STATISTICAL ANALYSIS**

Differences between groups were compared with unpaired two-tailed Student’s t test using Prism 7.0 (GraphPad Software, San Diego, CA). A two-tailed paired Student’s t test and a Two-Way ANOVA analyses with Bonferroni correction made when appropriate and p values indicated on figures. All experiments are representative of at least three independent experiments, unless otherwise noted.

**Supplementary Material**

Refer to Web version on PubMed Central for supplementary material.

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Highlights

- HSCs express *Mds1* as they emerge in the aorta of the murine fetus
- Inducible *Mds1Cre-ERT2* mice facilitate the lineage tracing of HSC-derived hematopoiesis
- HSC increasingly contribute to hematopoiesis in the fetal liver after E12.5
- The majority of macrophages in adult spleen and marrow are derived from HSCs
Figure 1. Mds1 is expressed in aortic hemogenic endothelium

Embryo-fetal expression of Mds1 documented using the Mds1$^{\text{lacZ}}$ allele (Zhang et al., 2011).

(A–E) Whole-mount β-galactosidase staining of heterozygous Mds1$^{\text{lacZ}/\text{wt}}$ embryos at various stages of development as indicated. Mean crown-rump length for C57 E8.5 = 1.88, E9.5 = 3.47, E11.5 = 6.83, E12.5 = 8.60, E14.5 = 12.29 mm (Mu et al., 2008).

(F) Photomicrograph of a X-gal-stained section of an E9.5 embryo with plane of section indicated by the white line in (B). Scale bar, 100 μM.

(G) Close-up of boxed region in (F), showing the dorsal aorta, left being ventral, right being dorsal, with β-galactosidase-positive cells seen budding from the endothelium (arrows). Scale bar, 25 μM.

(H and I) Photomicrographs of E10.5 aorta (left is ventral) β-galactosidase-positive cells (H) that also express Runx1 detected by immunocytochemistry (I). Scale bar, 20 μM.

(J) X-gal staining reveals no β-galactosidase-positive cells in the liver of the E13.5 mouse embryo. Scale bar, 25 μM.
Figure 2. Tamoxifen-inducible Mds1CreERT2 allele construction and expression in adult hematopoiesis and fetal HSC but not EMP- or primitive-derived hematopoiesis

(A) A DNA cassette encoding the TAM-inducible Cre-ER fusion protein was introduced into the Mecom locus in embryonic stem (ES) cells via homologous recombination.

(B) ES cell DNA was screened by Southern blotting, cutting the genomic DNA with EcoRV and blotting with 5′ and 3′ probes situated outside of the recombination cassette. Clones B3 and B5 show the expected sizes of 15.1 Kb and 7.6 kb, with 5′ and 3′ probes, respectively. Mice were created from ES cell clone B5, which were backcrossed onto a C57Bl6 background and crossed with mice bearing the Rosa26 LSL-YFP allele.

(C–E) These mice were treated with TAM, and the peripheral blood was analyzed by flow cytometry at 30 days and bone marrow at days 4, 10, and 15 for YFP+ cells, which progressively increased. (D) Quantitation of percentage of bone-marrow YFP-positive cells following the first TAM treatment on the days indicated. Average ± SEM, n = 3. (E) Representative scatterplot (day 30) showing the appearance of YFP+ blood cells following TAM treatment.

(F and G) X-gal staining of fetal liver in E13.5 mouse embryos, after a single treatment with TAM at E7.5 (F) or E9.5 (G). Eosin counter-stain; bar denotes 10 μm.

(H) Flow cytometric analysis of E11.5 AGM, yolk sac, and blood for pre-HSC (kit+CD31+VECadherin+) versus EMP (kit+CD41+CD16/32+) as indicated. For each analysis, the percentage of total cells that were YFP+ is charted as average ± SEM, n = 5. See Figures S2A and S2B, for the gating strategy.

(I) Flow cytometric analysis of E11.5 blood and liver for lineage-positive cell subsets (EryP Ter119+ blood, EryD Ter119+FSChi to distinguish from circulating EryP in the liver) as
indicated. For each analysis, the percentage of total cells that were YFP+ is charted as average ± SEM, n = 5. See Figures S2D and S2E for the gating strategy.

(J) Flow cytometric analysis of E12.5 fetal liver and blood for cell subsets as indicated gated as in (I). For each analysis, the percentage of total cells that were YFP+ is charted as average ± SEM, n = 10.

(K) Analysis of mice competitively transplanted with 10^6 E12.5 liver cells from TAM-treated (at E9.5) Mds1CreERT2/WT, Rosa26LSL-YFP/LSL-YFP mice and 10^5 unlabeled adult bone-marrow cells. Flow cytometric analysis of circulating YFP+ and YFP-negative granulocyte/monocyte cells (CD11b+), B cells (CD19+), and T cells (CD3e+) at 12 weeks post-transplant. See Figure S2F for gating strategy. Average ± SEM, n = 3.
(A) Flow cytometric analysis of E14.5 liver cells. Levels of YFP positivity were normalized to the level found in LSK (Lin-Kit⁺Sca1⁺, average 23.3% YFP⁺) to control for excision rates. Averages ± SEM of 5 individual embryos are plotted for LK (Lin-Kit⁻Sca1⁻), erythroid (Ter119⁺FSC⁹⁺hi), granulocyte (Ly6G⁺), monocyte (Ly6C⁺), and B (CD19⁺) cells. See Figures S3A and S3B for gating strategy.

(B) Flow cytometric analysis of circulating blood cells and liver cells at E16.5. Averages ± SEM of 3 individuals are plotted. Gating is as in (A) with the addition of circulating definitive erythroid (EryD, Ter119⁺) cells gated as in Figure S3C. Levels of YFP positivity were normalized to the level found in LSK average 8.1% YFP⁺) to control for excision rates.

(C) Flow cytometric analysis of E16.5 progenitors in the liver. LSK was further refined into LT-HSC, ST-HSC, MPP2, MPP3, and MPP4 based on Flt3, CD150, and CD48 positivity (see Figures S3A and S3D). LK were further refined into lymphoid (CD127⁺), myeloid (CD16/32⁻), and erythroid/megakaryocyte (CD16/32⁻) progenitors. Values normalized to YFP levels in LSK (B). Averages ± SEM, n = 3.

(D) Analysis by flow cytometry of YFP⁺ F4/80⁺ macrophages from E14.5 and E16.5 brain and liver normalized to YFP positivity of LSK (panels A and B). Details of gating are found in Figure S3E. Average ± SEM, n = 3. Unpaired two tail Student’s t test was performed.
comparing YFP labeling in cell populations to LSK. *p < 0.05; **p < 0.01; ***p < 0.001; ****p < 0.0001.
Figure 4. TAM treatment at E9.5 of Mds1\textsuperscript{CreERT2} Rosa26\textsuperscript{LSL-YFP} efficiently labels HSCs

(A) Flow cytometric analysis of the proportion of YFP\textsuperscript{+} granulocyte/monocyte cells (GR-1\textsuperscript{+}), B cells (B220\textsuperscript{+}), and T cells (CD3\textsuperscript{ε}\textsuperscript{+}) in the circulation of 1-year-old Mds1\textsuperscript{CreERT2} Rosa26\textsuperscript{LSL-YFP} mice after TAM treatment at E8.5 (empty circles, n = 4) or at E9.5 (solid circles, n = 2).

(B) Flow cytometric analysis of bone marrow of the same mice as in (A) demonstrating a high percentage of YFP\textsuperscript{+} progenitor cells in LSK subsets, as well as of lineage-restricted lymphoid (CD127\textsuperscript{+}), myeloid (CD16/32\textsuperscript{+}), and erythroid/megakaryocyte (CD16/32\textsuperscript{-}) LK subsets. See Figure S4A for gating strategy. Total marrow CD45\textsuperscript{+} YFP\textsuperscript{+} cells shown in (C).

(C) Percentage of YFP\textsuperscript{+} F4/80\textsuperscript{+} tissue-resident macrophages in these mice. See Figure S4B for detailed gating strategy. In all panels, the average is plotted.
| Reagent or resource | Source | Identifier |
|---------------------|--------|------------|
| Hámster Anti-Mouse CD3ε PE-CF594 Clone 145–2C11 | BD | (BD Biosciences Cat# 562286, RRID:AB_11153307) |
| Anti-Mouse CD3ε PE Clone 145–2C11 | eBioscience | Cat# 17–0031-82 |
| Anti-Mouse CD3ε APC-AF780 Clone 17A2 | eBioscience | Cat# 47–0032-82 |
| Rat Anti Mouse -CD11b APC Clone M1/70 | BD | (BD Biosciences Cat# 553312, RRID:AB_398535) |
| Rat Anti Mouse -CD11b APC-EF780 Clone M1/70 | eBioscience | Cat# 27–0112-82 |
| Rat Anti Mouse CD11b PEcy7 Clone M1/70 | BD | (BD Biosciences Cat# 561098, RRID:AB_2033994) |
| Anti-Mouse CD16/32 PE Clone 93 | eBioscience | Cat# 12–0161-82 |
| Anti-Mouse CD16/32 PEcy7 Clone 93 | eBioscience | Cat# 25–0161-81 |
| Rat Anti-Mouse CD16/CD32 APC-Cy7 Clone 2.4G2 | BD | (BD Biosciences Cat# 560541, RRID:AB_1645229) |
| Rat Anti-mouse CD19 APCeFluor780 Clone ID3 | eBioscience | Cat# 47–0193-82 |
| Rat Anti-mouse CD19 PE Clone ID3 | eBioscience | Cat# 12–0193-82 |
| Rat Anti-mouse CD19 PerCP-Cy5.5 Clone ID3 | eBioscience | Cat #25–0193-82 |
| Rat Anti-Mouse CD34 AF700 Clone RAM34 | BD | (BD Biosciences Cat# 560518, RRID:AB_1727471) |
| Rat Anti-Mouse CD45 PE Clone 30-F11 | eBioscience | Cat #12–0451-82 |
| Rat Anti-Mouse CD45R/B220 PE-Cy7 Clone RA3–6B2 | BD | (BD Biosciences Cat# 552772, RRID:AB_394458) |
| Armenian Hamster anti-Mouse CD48 PE, Clone: HM48–1, | eBioscience | Cat# 12–0481-81 |
| Armenian Hamster anti-Mouse CD48 APC-eFluor780, Clone: HM48–1, | eBioscience | Cat# 50–112-4034 |
| Rat Anti-Mouse CD115 (CSF1R) PE clone AFS98 | eBioscience | Cat# 12–1152-82 |
| Rat Anti-Mouse CD117 (Kit) PE-cy5, Clone: 2B8 | eBioscience | Cat# 50–140-92 |
| Rat Anti-Mouse CD117 (Kit) PE Clone: 2B8 | eBioscience | Cat #12–1171-82 |
| Rat Anti-Mouse CD117 (Kit) PECF594 Clone: 2B8 | BD | (BD Biosciences Cat# 562417, RRID:AB_11154233) |
| Rat Anti-Mouse CD117 (Kit) PE-cy7, Clone: 2B8 | eBioscience | Cat #25–1171-82 |
| Rat Anti-Mouse CD127 BUV737 Clone SB/199 | BD | (BD Biosciences Cat# 612841, RRID:AB_2870163) |
| Rat Anti-Mouse CD127 APC clone A7R34 | eBioscience | Cat#17–1271-82 |
| Rat anti-Mouse CD135 (Flt3) PE, Clone: A2F10 | eBioscience | Cat# 50–106-20 |
| Rat anti-Mouse CD135 (Flt3) PE-CF594, Clone: A2F10 | BD | (BD Biosciences Cat# 562537, RRID:AB_2737639) |
| Rat anti-Mouse CD144 (VECadherin) EF660 Clone BV13 | eBioscience | Cat#50–106-20 |
| Rat anti-Mouse CD150 APC Clone: 9D1 | eBioscience | Cat# 50–150-87 |
| Rat Anti-mouse F4/80 APC Clone BM8 | eBioscience | Cat #17–4801-82 |
| Rat Anti-Mouse GR1 (Ly-6G and Ly6C) APC-Cy7 Clone RB6–8C5 | BD | (BD Biosciences Cat# 557661, RRID:AB_396775) |
| Rat anti-mouse Ly6C APC Clone HK1.4 | eBioscience | Cat #17–5932-82 |
| Rat anti-mouse Ly6C PEcy7 Clone HK1.4 | eBioscience | Cat #25–5932-82 |
| Reagent or resource          | Source         | Identifier                                                                 |
|-----------------------------|----------------|----------------------------------------------------------------------------|
| Rat antimouse Sca-1 (Ly-6A/E) PerCP-Cy5.5, Clone: D7 | eBioscience    | Cat# 50–158-66                                                            |
| Rat Anti-Mouse Ter119 Percp cy5.5 Clone TER-119   | BD             | (BD Biosciences Cat# 560512, RRID:AB_10561844)                            |
| Rat Anti-Mouse Ter119 APC-eFluor780 Clone TER-119 | eBioscience    | Cat #47–5921–80                                                            |
| Rabbit anti-Runx            | Abcam          | (Abcam Cat# ab92336, RRID:AB_2049267)                                      |
| Goat anti-rabbit IgG AF647   | ThermoFisher   | (Thermo Fisher Scientific Cat# A-21244, RRID:AB_253812)                    |
| ProLong Gold antifade reagent | ThermoFisher   | Cat # P36934                                                                |

**Bacterial and Virus Strains**

- bacterial line DY380: Neal Copeland (Liu et al., 2003) n/a

**Chemicals, Peptides, and Recombinant Proteins**

- PE-CF594 Streptavidin: BD Biosciences Cat# 562318
- Tamoxifen (CAS 10540–29–1): Santa Cruz Biotechnology Cat# sc-208414
- DAPI: ThermoFisher Cat# EN62248

**Critical Commercial Assays**

- SuperScript IV One-Step RT-PCR System: Thermofisher Cat# 12594025
- GoTaq: Promega Cat# M7422

**Experimental Models: Organisms/Strains**

- B6.129X1-Gt(Rosa)26Scnat1(1/EF5PR1)tm1(EYFP)Cos/J: Jackson Lab Stock No: 006148
- B6.129S4-Gt(Rosa)26Sortm1(FLP1)Dym/RainJ: Jackson Lab Stock No: 009086
- Mds1 CreEr: This paper n/a
- Mds1 LacZ: This paper n/a
- C57BL/6 wildtype Inbred mice: Charles River C57BL/6NCrl

**Oligonucleotides**

- M2 wtfw GGTGTCCAAAACGTGCAATGC: This paper n/a
- M2 fw ACTCACCCCTGAAGTTCAGG: This paper n/a
- M2 rv CGAGTTGCCCCACAGCG: This paper n/a
- YFP fw AAAGTCGCTCTGAGTTGTTAT: This paper n/a
- YFP rv AAGACCGCGAAGATTTTGTGC: This paper n/a

**Recombinant DNA**

- BAC RP24–120B18: BACPAC Resources Center (BPRC) RP24–120B18
- GS1650: This paper n/a
- GS1656: This paper n/a
- GS1690: This paper n/a

**Software and Algorithms**

- FlowJo 8.5.3 software (TreeStar): BD FJ8.5.3