SCA-1 Expression Level Identifies Quiescent Hematopoietic Stem and Progenitor Cells

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SUMMARY

Blood cell generation depends on continuous cellular output by the sequential hierarchy of hematopoietic stem cell (HSC) and progenitor populations that all contain quiescent and actively cycling cells. Hematopoietic stem and progenitor cells (HSPCs) express the surface molecule Stem cell antigen 1 (SCA-1/LY6A). Using histone 2B-red fluorescent fusion protein label retention and cell-cycle reporter mice, we demonstrate that high SCA-1 expression (SCA-1 hi) identifies not only quiescent HSCs but quiescent cells on all hierarchical levels within the lineage ~/C0 SCA-1+KIT+ (LSK) population. Each transplanted SCA-1 hi HSPC population also displayed self-renewal potential superior to that of the respective SCA-1 lo population. SCA-1 expression is inducible by type I interferon (IFN). We show, however, that quiescence and high self-renewal capacity of cells with brighter SCA-1 expression at steady state were independent of type I IFN signaling. We conclude that SCA-1 expression levels can be used to prospectively isolate functionally heterogeneous HSPC subpopulations.

INTRODUCTION

The multi-potent and self-renewing hematopoietic stem cells (HSCs) at the top of the hierarchy give rise to hematopoietic progenitor cell (HPC) populations with gradually narrowing differentiation and self-renewal potential (Eaves, 2015). Murine HSCs and uncommitted progenitors express the glycosyl phosphatidylinositol-anchored cell surface protein Stem cell antigen 1 (SCA-1/LY6A) (Holmes and Stanford, 2007) and reside within the lineage ~/C0 SCA-1+KIT+ (LSK) population (Okada et al., 1992; Purton and Scadden, 2007; Spangrude et al., 1988) of the bone marrow (BM).

Numerous studies have demonstrated heterogeneity within each of the hematopoietic stem and progenitor cell (HSPC) populations with regard to cell-cycle activity (Wilson et al., 2008), lineage specification (Dykstra et al., 2007; Müller-Sieburg et al., 2002), and repopulation activity (Foudi et al., 2008; Qiu et al., 2014). Multiple cell-surface markers allowing for prospective isolation of the quiescent subset of HSCs (Balazs et al., 2006; Grinenko et al., 2014; Kiel et al., 2005; Osawa et al., 1996; Shin et al., 2014; Sudo et al., 2012) have been proposed. However, a marker uniformly associated with quiescence among the LSK population has been lacking so far. Two recent studies (Sawen et al., 2016; Wilson et al., 2015) showed increased quiescence and repopulation activity of immunophenotypic HSCs with high expression of SCA-1 (SCA-1 hi).

While the function of the SCA-1 molecule remains enigmatic (Holmes and Stanford, 2007), it is well established that SCA-1 is strongly upregulated by type I (Essers et al., 2009; Pietras et al., 2014) and type II (de Bruin et al., 2007) interferon (IFN). SCA-1 seems to regulate cellular responses to type I IFN as loss of SCA-1 abrogates type I IFN-induced proliferation of HSCs (Essers et al., 2009; Walter et al., 2015).

In the present work, we show that type I IFN-independent high expression of SCA-1 identifies quiescent cells with elevated repopulation activity not only in the HSC compartment but also in LSK progenitor populations.

RESULTS

High SCA-1 Expression Identifies Quiescent Cells Not Only in HSCs but Also in Progenitor Populations

To identify the most quiescent HSPCs, we induced expression of histone 2B-red fluorescent fusion protein (H2B-RFP) by doxycycline (DOX) treatment of R26RTA/Col1A1H2B-RFP mice (Egli et al., 2007) for 8 weeks and analyzed the BM after different chase periods for retention of H2B-RFP by flow cytometry (Figures S1A and S1B). In accordance with previous studies (Foudi et al., 2008; Qiu et al., 2014; Sawen et al., 2016; Wilson et al., 2008), we observed rapid dilution of H2B-RFP from HPCs (HPC-1, LSK CD48 hiCD150−; HPC-2, LSK CD48 hiCD150+) (Oguro et al., 2013) (see Figure S1C for gating), while a proportion of the HSC (LSK CD48−/CD150+), multipotent progenitor (MPP) populations (LSK CD48−/CD150−) (Kiel et al., 2005) retained the label for up to 19 weeks (see Figure S1D). We compared the expression of surface antigens between quiescent H2B-RFP+ cells and their respective parental populations. Most
Figure 1. Quiescent Hematopoietic Stem and Progenitor Cells Express High Levels of SCA-1

(A–C) R26rtTA/Col1A1H2B-RFP mice were DOX-pulsed, chased for either 4, 6, 13, or 19 weeks, and HSPC populations were analyzed by flow cytometry for retention of H2B-RFP (see also Figures S1A–S1D). (A) Representative histograms of BM HSCs (LSK CD48<sup>hi</sup>CD150<sup>+</sup>, upper row) or MPPs (LSK CD48<sup>hi</sup>CD150<sup>lo</sup>, middle row) from mice (n = 9) chased for 19 weeks are shown. H2B-RFP<sup>+</sup> label retaining cells (black lines) were overlaid on to the total parental population (solid gray histograms). Antigen median fluorescence intensities (MFI, lower row, normalized to the mean MFI of HSCs) of the total or H2B-RFP<sup>+</sup> populations were determined and compared. (B) Representative histograms for SCA-1 expression of BM HSCs (left column) and MPPs (middle column) isolated from R26rtTA/Col1A1H2B-RFP mice chased for either 4 (upper row, n = 2), 6 (middle row, n = 4), or 13 weeks (lower row, n = 8) are shown and H2B-RFP<sup>+</sup> (black lines) and H2B-RFP<sup>+</sup> cells (solid gray histograms) were overlaid. SCA-1 MFIs (right column, normalized to total HSC population) of H2B-RFP<sup>+</sup> and H2B-RFP<sup>+</sup> HSPCs were compared. (C) Representative SCA-1 expression histograms of BM HPC-1 (LSK CD48<sup>hi</sup>CD150<sup>+</sup>, left column) or HPC-2 cells (LSK (legend continued on next page)
prominently, we observed significantly higher expression of SCA-1 on label-retaining HSCs and MPPs after 4, 6, 13, and 19 weeks of chase (Figures 1A and 1B). Moreover, elevated SCA-1 expression was also detectable in both H2B-RFP+ HPC populations after 4 and 6 weeks of chase (Figure 1C). In addition, we found a significant down-regulation of endothelial cell-specific adhesion molecule (ESAM) (Yokota et al., 2008), CD117 (Grinenko et al., 2014; Shin et al., 2014), CD34, and CD48 (Qiu et al., 2014) on 19-week label-retaining HSCs and MPPs, while CD150 (Beerman et al., 2010; Kiel et al., 2005; Morita et al., 2010) was slightly, but significantly, upregulated on H2B-RFP+ HSCs (Figure 1A).

Next, we analyzed the BM of Ki67-RFPki/wt cell-cycle reporter mice (Basak et al., 2014), in which a Ki67-RFP fusion protein faithfully reports quiescent (Ki67-RFP−) and cycling (Ki67-RFP+) HSPCs (Figure S1E). RFP expression did not alter hematopoiesis in this model as judged by HSPC compartment size and competitive transplantation (Figures S1F and S1G). Cells with bright SCA-1 expression were significantly enriched among quiescent Ki67-RFP− HPC-1, MPP, and HSC populations, while cycling Ki67-RFP+ HSPCs expressed lower SCA-1 levels (Figure 1D).

To identify alternative markers of HSPC quiescence, we correlated expression of CD201 (EPCR) and CD27 (Balazs et al., 2006; Vazquez et al., 2015; Wiesmann et al., 2000) to SCA-1 and Ki67-RFP expression (Figures S1H and S1I). We found CD201 expression level to be extremely useful for prospective enrichment of Ki67-RFP− HSPCs, and expression of CD201 and SCA-1 showed a strong positive correlation. In contrast, CD27 expression appeared to be independent of cell-cycle activity and SCA-1. We did not observe any link between ESAM and Ki67-RFP expression (Figures S1H and S1I), while ESAM expression was heterogeneous among HPC-1 and MPP and without correlation to Ki67-RFP expression.

SCA-1hi HSPCs Have High Repopulation Activity upon Transplantation

To investigate whether repopulation activity of donor HSPCs correlates with SCA-1 expression, we fractionated HSCs, MPPs, and HPCs-1 into either SCA-1hi or SCA-1lo populations (Figure S2A) and competitively transplanted these cells into lethally irradiated congenic recipients (Figure 2A). We observed a range of SCA-1 fluorescence intensity of approximately two decades among BM LSK cells and arbitrarily divided these SCA-1+ cells into two populations, in which the SCA-1lo fraction comprised events from the lower decade, while the SCA-1hi fraction consisted of events from the higher decade. The sorting gates were placed to discriminate SCA-1-negative outliers as well as avoiding overlap between SCA-1lo and SCA-1hi populations after sorting (Figures S2A and S2B).

SCA-1hi donor HSCs displayed durable multi-lineage repopulation of recipient peripheral blood (PB) and BM (Figure 2B, columns I and II; Figures S2B–S2D), while SCA-1lo HSCs either contributed to a much lower extent in primary (Figure 2B) and secondary recipients (Figure S2E) or were devoid of long-term and serial repopulation activity (Figure S2D). In contrast to HSCs, SCA-1hi MPPs did not harbor stable multi-lineage repopulation potential as evidenced either by a decline in PB neutrophil and BM chimerism at 16 weeks after transplantation (Figure 2C, columns I and II) or by secondary transplantation (Figure S2E). However, the intermediate-term repopulation of all leukocyte lineages by transplanted SCA-1hi MPPs was significantly enhanced compared with SCA-1lo donor MPPs. Likewise, SCA-1hi donor HPC-1 cells exhibited significantly elevated B-cell chimerism 3 weeks after transplantation in contrast to the corresponding SCA-1lo population (Figure 2D, column I).

The comparison of SCA-1 expression among donor-derived (CD45.2+) BM cells isolated from primary recipients of SCA-1hi or SCA-1lo donor cells (B6 HSCs or MPPs) revealed that SCA-1lo donor cells or their progeny can reacquire high expression of SCA-1 (Figures 2B and 2C, columns III and IV; Figure S2F).

Higher SCA-1 Expression and Increased Repopulation Activity of Quiescent HSCs Does Not Depend on Type I IFN Signaling

SCA-1 expression is strongly upregulated by type I IFN (Essers et al., 2009), and Ifnar1−/− mice, which lack type I IFN signaling (Kamphuis et al., 2006), showed on average...
decreased steady-state SCA-1 expression, albeit similar maximum levels (Figure S2G). The frequency of LSK cells was reduced in Ifnar1<sup>del/del</sup> mice, but the frequency as well as cell-cycle activity of HSCs was unchanged in comparison with control mice. However, there was still heterogeneity of SCA-1 expression levels within the various LSK HSPC populations of Ifnar1<sup>del/del</sup> mice. To investigate whether the superior repopulation activity of SCA-1<sup>hi</sup> HSPCs depends on type I IFN, we competitively transplanted either SCA-1<sup>hi</sup> or SCA-1<sup>lo</sup> Ifnar1<sup>del/del</sup> HSCs (Figure S2A). The PB repopulation activity of SCA-1<sup>hi</sup> Ifnar1<sup>del/del</sup> donor HSCs was significantly elevated compared with their SCA-1<sup>lo</sup> counterparts (Figure 2E, column I). However, the chimerism level of SCA-1<sup>lo</sup> Ifnar1<sup>del/del</sup> recipients started to rise at the end of primary transplantation (Figure 2E, columns I and II) and upon secondary transplantation (Figure S2E, columns I and II). SCA-1<sup>lo</sup> Ifnar1<sup>del/del</sup> derived LSK cells of recipients showed reacquisition of bright SCA-1 expression (Figures 2E and S2E, columns III and IV). At the end of secondary transplantation, PB and BM chimerism as well as SCA-1 expression of both donor cell entities were indistinguishable (Figure S2E, columns I-IV). However, as

**Figure 2. Type I IFN-Independent SCA-1 Expression Predicts Repopulation Activity**

(A) HSPCs (either HSCs, MPPs, or HPCs-1 isolated from B6 mice or Ifnar1<sup>del/del</sup> HSCs, all CD45.2<sup>+</sup>) were fractionated according to SCA-1 expression (see Figure S2A) and 100 cells were transplanted together with 5 x 10<sup>5</sup> B6.CD45.1 competitor WBMCs into lethally irradiated B6.CD45.1/2 recipients. (B–E) Each row shows analysis of primary recipient mice transplanted with either B6 HSCs, B6 MPPs, B6 HPCs-1, or Ifnar1<sup>del/del</sup> HSCs (n = 3–4/group). (Column I) PB chimerism of recipient mice was monitored longitudinally (red circles, neutrophils; blue triangles, B cells; green boxes, T cells. Continuous line, filled symbols: SCA-1<sup>hi</sup> donor cells; dotted line, open symbols: SCA-1<sup>lo</sup> donor cells. Means ± SD are shown, significance calculated with repeated-measures two-way ANOVA with Bonferroni error correction). (Column II) BM LSK donor chimerism (individual recipients and means are shown, unpaired Student’s t test, BM chimerism of B6 HPC-1 recipients was not analyzed due to complete lack of PB neutrophil reconstitution). (Column III) Representative contour plots of donor-derived (CD45.2<sup>+</sup>) LSK cells are depicted. Frequency (mean ± SD) of the CD45.2<sup>+</sup> LSK population among total WBMCs is shown. (Column IV) SCA-1 MFI of CD45.2<sup>+</sup> LSK cells was calculated (individual recipients and means are shown, mice showing <0.001% of CD45.2<sup>+</sup> LSK cells among total WBMCs were excluded from analysis, unpaired Student’s t test). *p = 0.01–0.05, **p = 0.001–0.01, ***p < 0.001; ns, not significant.

(F) Representative contour plots of cell-cycle analysis of either SCA-1<sup>hi</sup> or SCA-1<sup>lo</sup> BM LSK cells isolated from Ifnar1<sup>del/del</sup> mice (n = 5). Right data plot shows frequencies of LSK cells in G<sub>0</sub>, G<sub>1</sub>, or S/G<sub>2</sub>/M phase (mean ± SD, paired Student’s t test with Bonferroni-Holm error correction). **p = 0.001–0.01. See also Figure S2.
recipients of SCA-1\textsuperscript{hi} \textit{Ifnar1}\textsuperscript{del/del} donor HSCs displayed significantly higher PB chimerism throughout primary transplantation and in the initial phase of secondary transplantation, and in the conclusion that constitutive type I IFN signaling does not account for the superior repopulation potential of HSPCs with high SCA-1 expression. Moreover, cell-cycle analysis of LSK cells from \textit{Ifnar1}\textsuperscript{del/del} mice corroborated our finding that the increased quiescence of SCA-1\textsuperscript{hi} HSPCs was type I IFN independent (Figure 2F).

**DISCUSSION**

We showed that high SCA-1 expression allows for prospective isolation of quiescent and potent hematopoietic progenitor cells. As previously reported, transplanted LSK CD48\textsuperscript{lo}/CD150\textsuperscript{+} MPPs did not exhibit long-term reconstitution, but showed robust intermediate-term contribution to all blood cell lineages (Oguro et al., 2013). We showed that the vast majority of this repopulation potential was confined to those cells which expressed the highest levels of SCA-1 within the MPP population. These cells were rarely dividing in situ, as evidenced by label retention characteristics similar to those of SCA-1\textsuperscript{hi} HSCs as previously reported (Foudi et al., 2008; Säwe´n et al., 2016). Likewise, the B-lymphoid potential of the HPC-1 population was strongly enhanced among SCA-1\textsuperscript{hi} cells. In addition, we confirmed the recent finding (Säwe´n et al., 2016; Wilson et al., 2015) that the majority of functional HSCs resided within the SCA-1\textsuperscript{hi} LSK CD48\textsuperscript{lo}/CD150\textsuperscript{+} population, while HSCs seemed to be rare or weak among those cells expressing lower amounts of SCA-1. However, SCA-1\textsuperscript{lo} donor cells could give rise to progeny with a SCA-1\textsuperscript{hi} phenotype, demonstrating plasticity of SCA-1 expression levels.

The BM lin\textsuperscript{−}CD201\textsuperscript{−}CD27\textsuperscript{+} population significantly overlaps with LSK cells in B6 mice and consists of early HSPCs. Lin\textsuperscript{−}CD201\textsuperscript{−}CD27\textsuperscript{+} cells have been proposed for alternative HSPC identification in mouse strains (e.g., BALB/c or NOD strains) or under stress conditions, which lack a distinct LSK immunophenotype (Vazquez et al., 2015). Our result that CD201 expression level correlates with quiescence of HSPCs similar to SCA-1 facilitates identification and isolation of quiescent subsets among lin\textsuperscript{−}CD201\textsuperscript{−}CD27\textsuperscript{+} HSPCs.

In contrast, ESAM, which was proposed to label HSCs under stress conditions in a cell-cycle-specific manner (Sudo et al., 2012), did not seem to be useful for prospective enrichment of quiescent HSPCs at steady state, as Ki67-RFP and ESAM expression showed poor correlation.

We demonstrate that the properties of SCA-1\textsuperscript{hi} cell populations were not a result of constitutive type I IFN signaling. The latter was surprising, as steady-state type I IFN partially accounted for SCA-1 expression (Figure S2G) and “tonic” (basal or constitutive) type I IFN signaling has been implied in HSC maintenance (Gough et al., 2012). However, our transplantation data revealed that SCA-1\textsuperscript{hi} as well as SCA-1\textsuperscript{lo} \textit{Ifnar1}\textsuperscript{del/del} HSCs outgrew wild-type (wt) competitor cells in secondary recipients, which may be explained either by a general competitive advantage of \textit{Ifnar1}\textsuperscript{del/del} HSCs over wt competitor HSCs or, due to lower overall SCA-1 expression in \textit{Ifnar1}\textsuperscript{del/del} mice, potent HSCs still reside in the SCA-1\textsuperscript{lo} sorted population. The latter is very likely, as our SCA-1 sorting strategy for \textit{Ifnar1}\textsuperscript{del/del} donor cells separated only the brightest fraction of HSCs into the SCA-1\textsuperscript{hi} subset, while SCA-1\textsuperscript{lo} cells contained the majority of the cell population. Nevertheless, purified SCA-1\textsuperscript{hi} \textit{Ifnar1}\textsuperscript{del/del} donor cells repopulated primary recipients faster and more robustly. Moreover, cell-cycle analysis of \textit{Ifnar1}\textsuperscript{del/del} HSCs revealed that the increased quiescence of the SCA-1\textsuperscript{hi} population was type I IFN independent.

Hematopoietic contribution, differentiation pattern, and the relationship between the HSPC subpopulations and diverse SCA-1 expression have not been addressed in situ so far. We speculate that discrepancies between recent studies reporting high (Sawai et al., 2016) or low (Busch et al., 2015; Schoedel et al., 2016; Sun et al., 2014) contribution of HSCs to steady-state hematopoiesis might reflect lineage tracing of exclusive HSC subpopulations that differ by SCA-1 expression and cell-cycle activity.

SCA-1 is already one of the most widely used markers for prospective isolation of murine HSPCs, and our finding that quiescent subpopulations are identified by differential SCA-1 expression easily allows for refined purification and analysis strategies.

**EXPERIMENTAL PROCEDURES**

**Mice**

C57Bl/6 wt, B6.CD45.1, B6CD45.1/2, \textit{Ifnar1}\textsuperscript{del/del} (Kamphuis et al., 2006), R26\textsuperscript{ERT2}/Col1A1\textsuperscript{H2B-RFP} (Egli et al., 2007) and \textit{Kit}\textsuperscript{lo}RFP\textsuperscript{hi/wt} (Basak et al., 2014) mice were housed at the Experimental Center, TU Dresden. R26\textsuperscript{ERT2}/Col1A1\textsuperscript{H2B-RFP} mice were induced by drinking water containing 1 mg/mL doxycycline (Applichem) and 1% sucrose ad libitum for 8 weeks.

All animal experiments were carried out in accordance with institutional guidelines and were approved by Landesdirektion Dresden Ref. No. 24-9168.11-1/2012-39.

**Cell Preparation**

Whole bone marrow cells (WBMCs) were isolated by crushing long bones with mortar and pestle using PBS/2% fetal calf serum (FCS)/2 mM EDTA and filtered through a 100-μm mesh. After erythrocyte lysis in NH₄Cl buffer, cells were filtered through a 40-μm mesh. Hematopoietic lineage+ cells were removed with the lineage cell depletion kit (Miltenyi Biotec).

PB was drawn by retrobulbar puncture.
Flow Cytometry
Cells were stained for 30 min with fluorochrome-labeled antibodies (see Table S1) diluted in PBS/2% FCS/2 mM EDTA, washed twice, and analyzed or sorted on BD FACSAria or Miltenyi MACSquant flow cytometers. Data were analyzed using FlowJo 9.9.4 (Treestar). Negative and positive populations were identified by fluorescence-cytometers. Data were analyzed using FlowJo 9.9.4 (Treestar). Negative and positive populations were identified by fluorescence-cytometers. Data were analyzed using FlowJo 9.9.4 (Treestar). Negative and positive populations were identified by fluorescence-cytometers. Data were analyzed using FlowJo 9.9.4 (Treestar). Negative and positive populations were identified by fluorescence-cytometers. Data were analyzed using FlowJo 9.9.4 (Treestar). Negative and positive populations were identified by fluorescence-cytometers. Data were analyzed using FlowJo 9.9.4 (Treestar). Negative and positive populations were identified by fluorescence-cytometers. Data were analyzed using FlowJo 9.9.4 (Treestar). Negative and positive populations were identified by fluorescence-cytometers. Data were analyzed using FlowJo 9.9.4 (Treestar). Negative and positive populations were identified by fluorescence-cytometers. Data were analyzed using FlowJo 9.9.4 (Treestar). Negative and positive populations were identified by fluorescence-cytometers. Data were analyzed using FlowJo 9.9.4 (Treestar). Negative and positive populations were identified by fluorescence-cytometers. Data were analyzed using FlowJo 9.9.4 (Treestar). Negative and positive populations were identified by fluorescence-cytometers. Data were analyzed using FlowJo 9.9.4 (Treestar). Negative and positive populations were identified by fluorescence-cytometers. Data were analyzed using FlowJo 9.9.4 (Treestar). Negative and positive populations were identified by fluorescence-cytometers. Data were analyzed using FlowJo 9.9.4 (Treestar). Negative and positive populations were identified by fluorescence-cytometers. Data were analyzed using FlowJo 9.9.4 (Treestar). Negative and positive populations were identified by fluorescence-cytometers. Data were analyzed using FlowJo 9.9.4 (Treestar). Negative and positive populations were identified by fluorescence-cytometers. Data were analyzed using FlowJo 9.9.4 (Treestar). Negative and positive populations were identified by fluorescence-cytometers. Data were analyzed using FlowJo 9.9.4 (Treestar). Negative and positive populations were identified by fluorescence-cytometers. Data were analyzed using FlowJo 9.9.4 (Treestar). Negative and positive populations were identified by fluorescence-cytometers. Data were analyzed using FlowJo 9.9.4 (Treestar). Negative and positive populations were identified by fluorescence-cytometers. Data were analyzed using FlowJo 9.9.4 (Treestar).
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