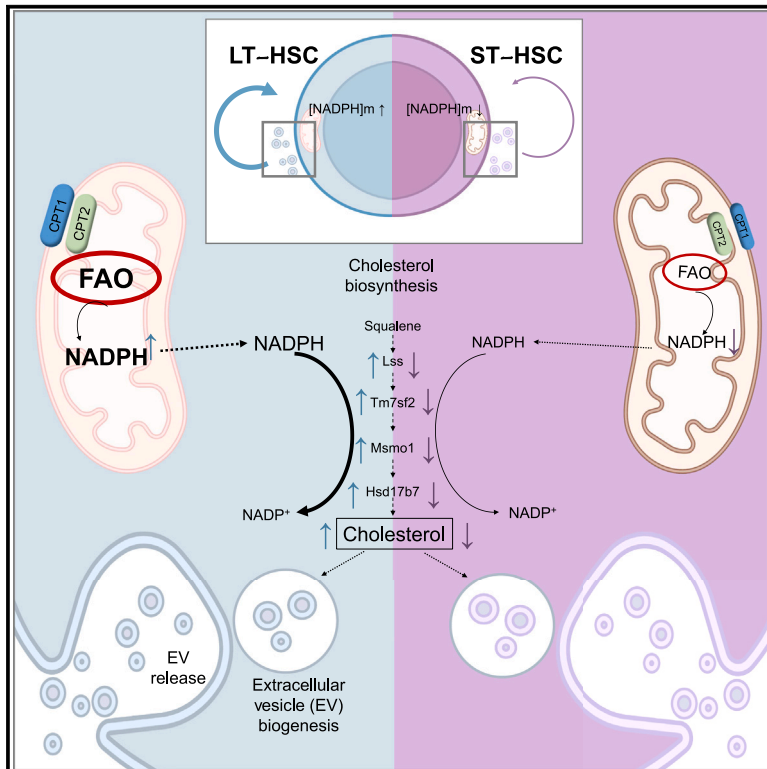


A mitochondrial NADPH-cholesterol axis regulates extracellular vesicle biogenesis to support hematopoietic stem cell fate

Graphical abstract



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In brief

Ito et al. investigate the mechanism by which mitochondrial metabolism controls hematopoietic stem cell (HSC) fate. They reveal that FAO (fatty acid oxidation)-generated NADPH fuels cholesterol biosynthesis in HSCs. A mitochondrial NADPH-cholesterol axis controls the biogenesis of extracellular vesicles that is required for proper HSC self-renewal and hematopoietic homeostasis.

Highlights

- HSCs have high mitochondrial NADPH that fuels cholesterol biosynthesis
- High NADPH and cholesterol synthesis in HSCs are dependent on FAO
- The NADPH-cholesterol axis supports proper HSC self-renewal and homeostasis
- HSC cholesterol levels impact autocrine EV signaling



Article

A mitochondrial NADPH-cholesterol axis regulates extracellular vesicle biogenesis to support hematopoietic stem cell fate

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SUMMARY

Mitochondrial fatty acid oxidation (FAO) is essential for hematopoietic stem cell (HSC) self-renewal; however, the mechanism by which mitochondrial metabolism controls HSC fate remains unknown. Here, we show that within the hematopoietic lineage, HSCs have the largest mitochondrial NADPH pools, which are required for proper HSC cell fate and homeostasis. Bioinformatic analysis of the HSC transcriptome, biochemical assays, and genetic inactivation of FAO all indicate that FAO-generated NADPH fuels cholesterol synthesis in HSCs. Interference with FAO disturbs the segregation of mitochondrial NADPH toward corresponding daughter cells upon single HSC division. Importantly, we have found that the FAO-NADPH-cholesterol axis drives extracellular vesicle (EV) biogenesis and release in HSCs, while inhibition of EV signaling impairs HSC self-renewal. These data reveal the existence of a mitochondrial NADPH-cholesterol axis for EV biogenesis that is required for hematopoietic homeostasis and highlight the non-stochastic nature of HSC fate determination.



INTRODUCTION

Cellular metabolism in hematopoietic stem cells (HSCs) has recently become the focus of intense research interest.¹ Mitochondria participate in multiple cellular functions, including regulation of cell metabolism, through the tricarboxylic acid (TCA) cycle and the respiratory chain.^{2,3} In this multifaceted biochemical process, the 4 respiratory complexes in the inner mitochondrial membrane transport electrons from nicotinamide adenine dinucleotide (NADH)/flavin adenine dinucleotide (FADH₂) to oxygen in an electron transport chain (ETC), which generates a proton gradient across the mitochondrial membrane.

Early evidence indicated that HSC mitochondria are functionally dispensable and activated only when commitment is demanded.^{4,5} However, deeper investigation into HSC function has revealed its dependence on mitochondrial fatty acid oxidation (FAO)—the catabolism of fatty acids in the mitochondrial matrix that feeds the TCA cycle^{6,7}—or the expression of subunits of respiratory complex III (*Uqcrcs1*) or complex II (*SdhD*).^{8,9} Recent research has also highlighted the necessity of using a dye-independent method to improve the accuracy of assessment of HSC mitochondria,^{10,11} and multiple studies have since confirmed that mitochondrial content and membrane potential in hematopoietic stem and progenitor cells (HSPCs) has been consistently underestimated in a manner that can correlate with assessments of stem cell function.^{10–13}

FAO also produces metabolites required to re-model the lipid landscape of HSCs. Some lipid species such as sphingolipids, glycolipids, and sterols are known to contribute to hematopoiesis.¹⁴ Genetic loss of cholesterol efflux transporter has been shown to regulate mobilization^{15,16} and expansion¹⁷ of the HSPC pool by enhancing the signaling for critical cytokines (e.g., interleukin-3[IL-3]/granulocyte-macrophage colony-stimulating factor [GM-CSF] receptor).¹⁸

These observations imply the possibility that mitochondria play previously unappreciated roles in HSC biology. How exactly mitochondrial respiration and FAO steer HSC fate decisions, however, remains unclear.

RESULTS

HSCs have high mitochondrial NADPH

FAO feeds mitochondrial metabolism, which reduces NADH, FADH₂, and nicotinamide adenine dinucleotide phosphate (NADPH) to drive redox reactions in the ETC and support multiple biosynthetic processes. We have observed that HSCs have limited expression of complex I while relying on complex II to support ETC activity,^{19,20} supporting the idea that HSCs accumulate high levels of reduced cofactors.

Taking advantage of these spectral properties of NAD(P)H, we observed that NAD(P)H autofluorescence appeared brighter in CD150⁺CD48⁺CD135⁺Lin⁺Sca-1⁺c-Kit⁺ cells (hereafter, HSCs) and displayed a dimming trend with hematopoietic differentiation (Figures 1A and S1A). Fully differentiated myeloid cells apparently deviate from this trend. As these populations dramatically vary in size²¹ (Figures S1B and S1C), we corrected NAD(P)H autofluorescence for forward scatter area (FSC-A, used as reporter of cell size, Figure S1B). The resulting readout of NAD(P)H density per cell (labeled as nNAD(P)H) confirms the dimming

trend of NAD(P)H levels with differentiation, which reaches lower levels in fully differentiated cells (Figure S1D).

Higher autofluorescence was observed in more primitive CD34[−] HSCs compared to CD34⁺ HSCs (Figures 1B and 1C), and endothelial protein C receptor (EPCR) positivity^{22–24} correlated with higher NAD(P)H levels in both HSCs and CD34[−] HSCs (Figures S1E).

Co-staining with the mitochondrial marker tetramethylrhodamine, methyl ester (TMRM) confirmed the mitochondrial localization of NAD(P)H (Figure 1C). To assess whether the high mitochondrial NAD(P)H levels in primitive HSCs were mainly due to NADH or NADPH pools, we leverage the differing sensitivities of the NAD(P)H pool to the mitochondrial uncoupler carbonyl cyanide-p-trifluoromethoxyphenylhydrazone (FCCP), the oxidizer H₂O₂, and the ETC inhibitor rotenone.²⁵ This analysis suggests that CD34[−] HSCs have a significant enrichment of mitochondrial NADPH compared to CD34⁺ HSCs (Figures S1J and S1L).

Fluorescence lifetime imaging (FLIM) has enabled the *in situ* determination of NAD(P)H lifetime pools in living cells, thanks to the different fluorescence time decay of NADPH and NADH.^{25–27} FLIM assay confirmed that CD34[−] HSC mitochondria display a higher NADPH/NADH ratio than the mitochondria of CD34⁺ HSCs (Figure S1M). Further, NADPH- and NADH-dependent bioluminescent enzymatic systems (Figures 1E and 1F) corroborated that cytosolic NADPH ([NADPH]_c) was comparable (Figure S1N), and no significant differences were observed in either mitochondrial or cytosolic NADH levels (Figures 1G and S1O).

Collectively, these data show that long-term HSCs (CD34[−] and/or EPCR⁺) have significantly higher levels of mitochondrial NADPH than CD34⁺ HSCs (and/or EPCR[−]).

NAD(P)H autofluorescence marks HSC capacity

To assess the association between NADPH and HSC functions, CD34[−] HSCs were divided into two fractions reflecting high and low levels of NAD(P)H (Figures 1H and S2A and STAR Methods). Approximately 60% of CD34[−] HSCs were included in the NAD(P)H^{hi} gate, and a downward trend in NAD(P)H levels correlated with hematopoietic differentiation (Figure S2B). The percentage of NAD(P)H^{hi} cells was significantly higher in EPCR⁺ HSCs or EPCR⁺CD34[−] HSCs compared to EPCR[−] (Figures S2C). In agreement, NAD(P)H^{hi} bone marrow mononuclear cells (BMMNCs) displayed higher percentages of cKit⁺Sca-1⁺, CD150⁺CD48[−], and EPCR⁺ cells compared to the NAD(P)H^{lo} counterpart (Figures S2D).

Both NAD(P)H^{hi} and NAD(P)H^{lo} CD34[−] HSCs are quiescent as the two populations displayed no significant differences in the timing of first division (Figure S2E) or positivity for the proliferation marker ki67 (Figure S2F) or in the G0 fraction determined by Pyronin Y (Figure S2G). Interestingly, active (Pyronin Y⁺) HSCs displayed a moderate reduction in NAD(P)H autofluorescence when compared to dormant (Pyronin Y[−]) HSCs (Figure S2H). When HSCs are forced to exit quiescence by *in vivo* stimulation with pl:pC, we observed a reduction in NAD(P)H autofluorescence (Figure S2I), possibly reflecting the stimulation in mitochondrial maturation previously reported.²⁸

Neither features of mitochondrial morphology such as volume, elongation, compactness, or ultrastructure (cristae content or

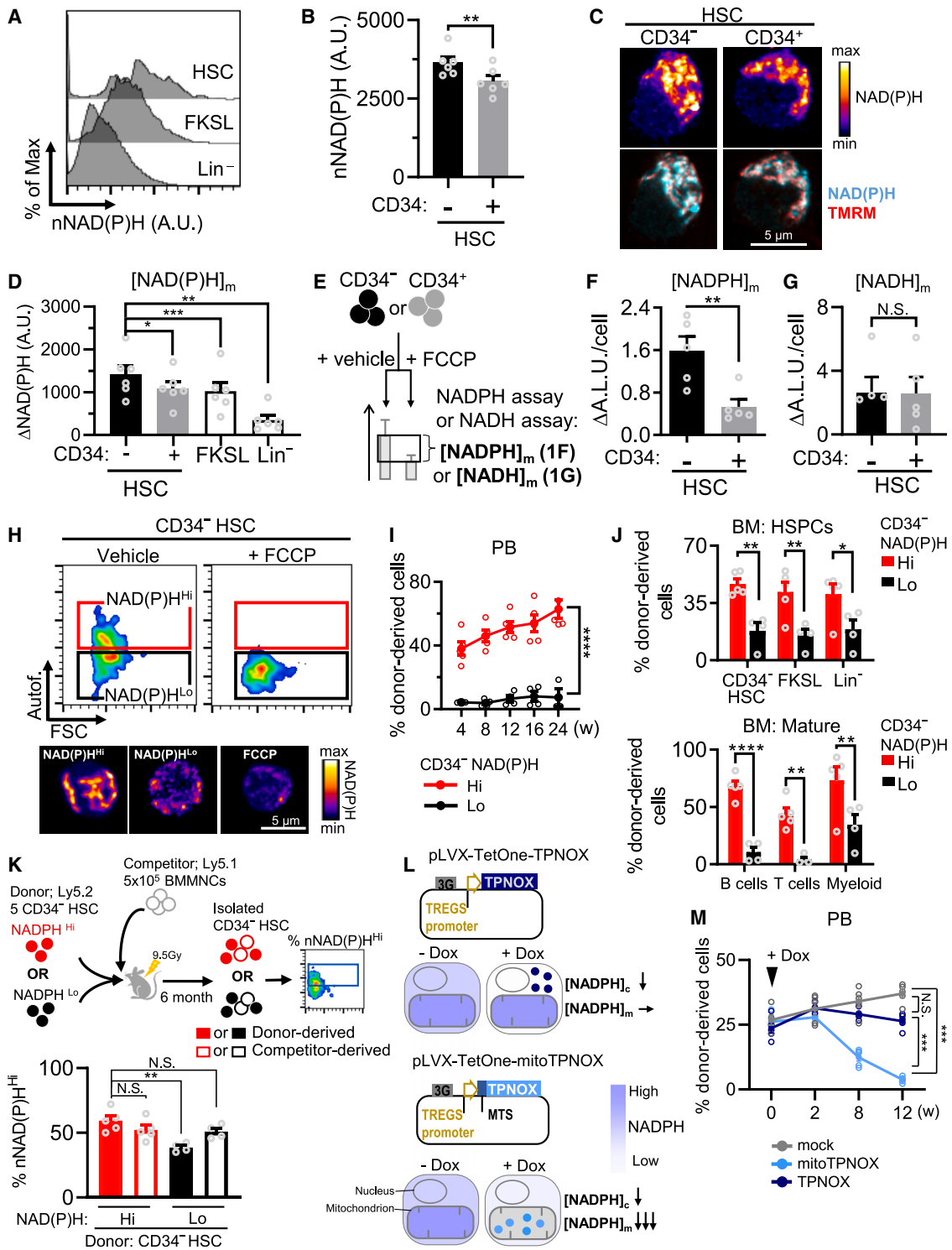


Figure 1. Mitochondrial NADPH levels mark stem cell capacity

(A) Flow cytometric histogram of the normalized NAD(P)H autofluorescence in HSCs, FKSL, and Lin⁻ cells (also see Figure S1A).

(B) Normalized NAD(P)H autofluorescence levels in CD34⁻ and CD34⁺ HSCs.

(C) Fluorescence microscopy images of NAD(P)H autofluorescence (top) and co-localization with the mitochondrial marker TMRM (bottom) obtained from CD34⁻ and CD34⁺ HSCs. Scale bar, 5 μm.

(D) Mitochondrial NAD(P)H determined as the difference of NAD(P)H autofluorescence after exposure to FCCP 1 μM in the HSPC populations investigated.

© Experimental design for the enzymatic determination of mitochondrial NADPH and NADH. Sorted CD34⁻ or CD34⁺ HSCs were exposed *in vitro* to FCCP 1 μM to dissipate mitochondrial NAD(P)H.

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morphology) nor reactive oxygen species (ROS) production were found to be altered in NAD(P)H^{Hi} versus NAD(P)H^{Lo} HSCs (Figures S2J–S2L), indicating that low levels of NAD(P)H are not dependent on the extra physiologic oxygen shock/stress induced in HSCs during isolation from bone marrow.²⁹

In vitro long-term culture-initiating cell (LTC-IC) capacity was enriched in the NAD(P)H^{Hi} fraction (Figure S2M). In competitive bone marrow transplantation (BMT) settings, NAD(P)H^{Hi} CD34⁺ HSCs exhibited higher reconstitution capacity compared to NAD(P)H^{Lo} CD34⁺ HSCs (Figure 1I). NAD(P)H^{Hi} cells gave rise to higher levels of chimerism in HSPCs and Lin[−] cells as well as mature cells without lineage bias (B, T, and myeloid cells) (Figure 1J). Taken together, these results suggest that high NAD(P)H levels as measured by autofluorescence can serve as a marker for HSC capacity. Further, CD34⁺ HSCs derived from NAD(P)H^{Hi} donors maintained a level of NAD(P)H autofluorescence comparable to that of competitor HSCs. In contrast, CD34⁺ HSCs derived from NAD(P)H^{Lo} donors had much lower NAD(P)H levels when compared to both NAD(P)H^{Hi}-derived HSCs or competitor HSCs (Figure 1K). This indicates that NAD(P)H^{Lo} HSCs are defective in their reconstitution potential as well in the capacity to generate their physiological NAD(P)H pool.

To confirm the compartmentalization of NADPH, untargeted triphosphopyridine nucleotide oxidase (TPNOX), which decreases cytosolic NADPH, or mitochondria-targeted TPNOX (mitoTPNOX), which specifically consumes intra-mitochondrial NADPH, was utilized to manipulate cellular NADPH levels, a system under the control of the doxycycline (Dox)-inducible promoter³⁰ (Figures 1L, S2N, and S2O). Both TPNOX and mitoTPNOX decreased colony-replating capacity after long-term culture (Figure S2N, left). Critically, mitoTPNOX significantly decreased LTC-IC capacity *in vitro* and mitochondrial NAD(P)H autofluorescence (Figures S2N, right and S2O) to a much greater extent than TPNOX, suggesting that mitochondrial NADPH is critical to regulate HSC function. We thus tested *in vivo* expression of mitoTPNOX in a transplantation setting, where it was induced with Dox 3 weeks after BMT. The mitoTPNOX, but not TPNOX, reduced hematopoietic contributions of donor cells from 4 weeks after induction, while no significant changes in chimerism were found 2 weeks after induction in either case (Figure 1M). These data further support the important roles played by mitochondrial NADPH in hematopoietic homeostasis.

Mitochondrial NADPH segregation controls HSC fate determination

We next measured the kinetics of mitochondrial NAD(P)H levels in HSCs during cell division *in vitro*. Single CD34⁺ HSCs were cultured and visually tracked until their first division. Corresponding daughter cells were stained with TMRM to perform 3D quantitation of NAD(P)H per single mitochondria (Figure 2A). We used a statistical approach to determine symmetric or asymmetric distribution of mitochondria between corresponding daughters (Figures 2A and S3A and STAR methods). With this strategy, 33% of divisions were labeled as asymmetric divisions, 36% as symmetric high, and the remaining 31% as symmetric low (Figure 2B), proportions similar to those obtained by paired daughter cell assays.^{6,7}

We speculated that mitochondria can be asymmetrically inherited by daughter cells during HSC mitosis. NAD(P)H^{Hi} CD34⁺ HSCs were isolated and subjected to cell-tracer staining followed by quantitative live imaging to track NAD(P)H inheritance during division. First division occurs at ~38 h from isolation (Figures S3B), consistent with cumulative division frequency analysis (Figures S2E). We observed that daughter cells can exhibit either the same NAD(P)H level as the mother or a significantly lower NAD(P)H level, but no daughter received a higher NAD(P)H level than its related mother cell (Figure 2C). We classified three distinct outputs of cell division (Figures 2D and S3C): a symmetric high where the two daughters have NAD(P)H comparable to the mother (41.5%); a symmetric low, with both daughters having lower NAD(P)H than the mother (28%); and an asymmetric output, where the daughters have different NAD(P)H levels (30.5%). The ratio of NAD(P)H intensity between two daughters confirmed the distribution pattern of NAD(P)H in those divisions' outputs determined as in Figure 2B (Figure S3D). A stochastic-segregation mitochondria during division should lead to cases of asymmetric segregation where one daughter cell has higher NAD(P)H than the mother cell. Critically, none of the daughter cells (236 cells) exhibited higher NAD(P)H than the mother at a level of statistical significance (Figures 2C and 2D), supporting the notion that asymmetry in NAD(P)H levels between corresponding daughter cells is the result of a regulated/non-stochastic process.

To further understand the significance of different levels of NADPH inherited by HSCs, we isolated daughter after first

(F and G) Enzymatic determination of levels of mitochondrial NADPH (F) and NADH (G) in CD34⁺ and CD34⁺ HSCs.

(H) Gating strategy to define NAD(P)H^{Hi} (red box) and NAD(P)H^{Lo} (black box) CD34⁺ HSCs. Images of NAD(P)H autofluorescence of NAD(P)H^{Hi} CD34⁺ HSCs, NAD(P)H^{Lo} CD34⁺ HSCs, or FCCP-exposed CD34⁺ HSCs are also shown (bottom). Scale bar, 5 μm.

(I) Hematopoietic contribution of donor cells in peripheral blood of recipient mice during competitive BMT. Five NAD(P)H^{Hi} or NAD(P)H^{Lo} CD34⁺ HSCs were isolated and then transplanted to the irradiated recipient mice with 4.0 × 10⁵ competitor bone marrow mononuclear cells (BMMNCs).

(J) Percentages of donor-derived cells in the indicated fractions of the bone marrow of the recipient mice 6 months after BMT.

(K) Schematic representation (top) and quantitation (bottom) of the percentage of nNAD(P)H^{Hi} cells in donor-derived (filled bars) and competitor-derived (open bars) CD34⁺ HSCs, 6 months after BMT.

(L) Schematic representation of the plasmid encoding for tetracycline (TET)-inducible expression of TPNOX or mitoTPNOX and their effect on cytoplasmic or mitochondrial NADPH after the induction with doxycycline (Dox).

(M) *In vivo* repopulation capacity of mitoTPNOX-infected HSPCs. 1,000 KSL (Ly5.2) infected with mitoTPNOX (light blue), TPNOX (dark blue), or empty vector (gray) was transplanted into the irradiated recipient mice (Ly5.1) with competitor cells (4.0 × 10⁵ BMMNCs). Percentage of donor-derived cells in the peripheral blood the indicated weeks after Dox induction (black arrow) were shown. HSCs, Lin[−]Sca-1⁺c-Kit⁺CD135⁺CD150⁺CD48[−]; KSL, c-Kit⁺Sca-1⁺Lin[−]; FKSL, CD135⁺c-Kit⁺Sca-1⁺Lin[−]; A.U., arbitrary unit; A.L.u., arbitrary luminescent units; MTS, mitochondrial targeting sequence; PB, peripheral blood; w, week. Bar graphs represent means ± SEM, circles represent each replicate. (B), (F), and (G): Student's paired t test. (I), (J), and (M): two-way ANOVA with Sidak's multiple comparison test. (K): one-way ANOVA with Dunnett's multiple comparison test. ****, p < 0.0001; ***, 0.0001 ≤ p < 0.001; **, 0.001 ≤ p < 0.01; *, 0.01 ≤ p < 0.05; N.S., not significant, p ≥ 0.05. See also Figures S1 and S2.

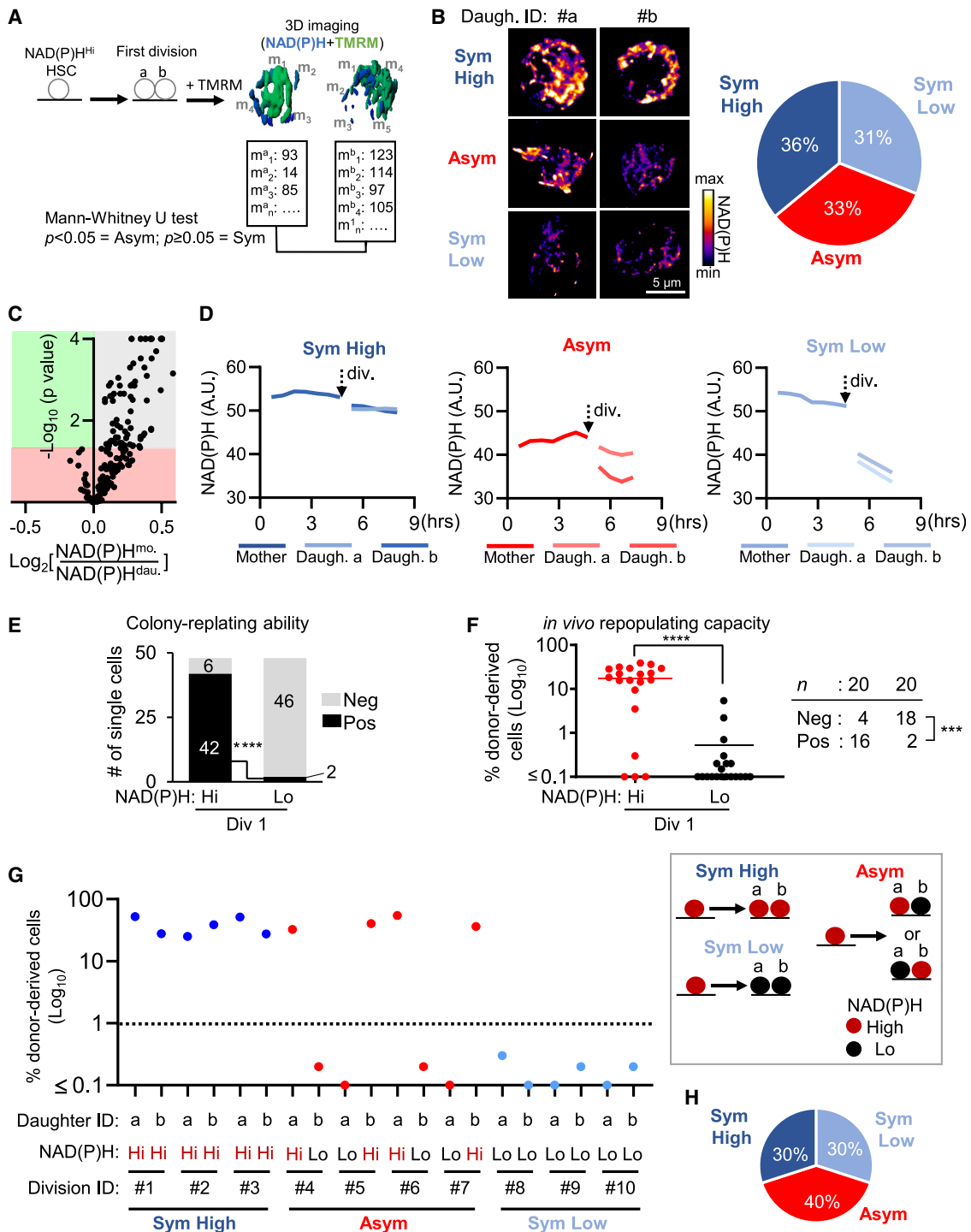


Figure 2. Mitochondrial NADPH levels mark HSC fate determination

(A) Experimental strategy for the determination of symmetric or asymmetric segregation of mitochondrial NAD(P)H after first division in the cultured CD34⁺ HSCs. Fluorescence microscopic analysis was performed to determine NAD(P)H autofluorescence intensity in the individual mitochondrion of each daughter cell. NAD(P)H intensities were subsequently compared between paired daughter cells by Mann-Whitney U test. (also see Figure S3A).

(B) Proportion of distribution of mitochondrial NAD(P)H after first division of the cultured CD34⁺ HSCs ($n = 36$, right). Images of NAD(P)H of the corresponding daughter cells are also shown (left). Scale bar, 5 μm .

(C and D) Determination of NAD(P)H segregation by live imaging.

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division that displayed NAD(P)H^{Hi} or NAD(P)H^{Lo} cells and performed functional assays. After first division (Div 1), the self-renewal capacity, measured by single-cell colony-replating ability after *in vitro* long-term culture (Figure 2E) and reconstitution capacity after single-cell transplantation (Figure 2F), was significantly enriched in NAD(P)H^{Hi} cells, supporting that NAD(P)H^{Hi} acts as a marker of HSC capacity during HSC division.

In vivo paired daughter cell (PDC) assay can determine the division patterns of HSCs retrospectively by assessing the reconstitution capacity of each daughter cell (Figure 2G).^{6,7,31} PDC assay revealed that daughter cells receiving symmetrically high NAD(P)H levels exhibited significant reconstitution potential, indicative of a symmetric division (30%). Division where NAD(P)H levels differ from between the two daughters also returned asymmetric reconstitution (40%). Ultimately, daughters that shared a low NAD(P)H level also shared a defective repopulation capacity (symmetric commitment, 30%). The percentage of the division patterns calculated by PDC assay overlapped with those obtained by live imaging of NAD(P)H (Figure 2H). Overall, these confirmed that NAD(P)H segregation upon Div 1 corresponds to the segregation of HSC function and their fate determination.

The intensity of TMRM was comparable in both NAD(P)H^{Hi} or NAD(P)H^{Lo} daughters, suggesting no major alterations in their respiration (Figure S3E) and corroborating the initial hypothesis that NAD(P)H levels in HSCs depend on FAO. Rate-limiting steps in FAO are represented by carnitine O-palmitoyltransferases 1 and 2 (Cpt1 and Cpt2).³² HSC daughters who retained NAD(P)H^{Hi} (Div 1) not only had higher expression of *Cpt1a* compared to NAD(P)H^{Lo} but also had no observed alterations of gene expression for *Ndufv1*, the subunit of respiratory complex I responsible for NADH binding (Figure S3F). Most strikingly, *Cpt1a* enabled us to clearly distinguish NAD(P)H^{Hi} cells from NAD(P)H^{Lo} cells at a single-cell level (Figure S3F). This observation indicates that during HSC division, daughter cells can differentially segregate their FAO capacity, which ultimately results in the asymmetry of NAD(P)H levels.

NADPH sustains cholesterol biosynthesis in HSCs

To identify biochemical pathways supported by NADPH in HSCs, we used 3 public datasets to investigate gene expression variation between mouse CD34⁻ and CD34⁺ HSCs^{33–35} (Figures S4A and S4B). We identified 77 NADPH-related pathways and compared their gene expression levels by Reactome

database (Figure 3A). Then we used the protein-metabolite Graphite networks³⁶ to retain NADPH metabolites in pathway-derived graphs. The resulting list of the NADPH-dependent pathways enriched in HSCs included the detoxification of ROS and linoleic acid metabolism, pathways already considered relevant for HSC maintenance (Figure 3B).

We selected the cholesterol biosynthesis pathway as that of greatest interest.^{15,18,37} Four genes involved in this pathway (*Lss*, *Tm7sf2*, *Msmo1*, and *Hsd17b7*) had significantly higher expression in CD34⁻ compared to CD34⁺ HSCs, with 3 of them catalyzing NADPH-dependent reactions (Figures 3B, 3C, and S4C). We confirmed the increased expression of *Tm7sf2*, *Msmo1*, and *Hsd17b7* in sorted CD34⁻ compared to CD34⁺ HSCs (Figure 3D).

To confirm the bioinformatic predictions, untargeted metabolomic analysis was performed. Metabolic pathway enrichment analysis highlighted the differential contribution of pathways linked to lipid synthesis (especially phospholipid and phosphatidylcholine biosynthesis), carbohydrate metabolism, and others (Figure S5A and Table S3). Steroid synthesis is also found among the enriched pathways, although it is not one of most prominent. This is most likely because we performed our untargeted metabolomics on the polar phase, while sterols are non-polar metabolites.

To overcome this limitation, we subjected the non-polar metabolite extracts of separated NAD(P)H^{Hi} and NAD(P)H^{Lo} CD34⁻ HSCs to a sterol analysis by gas chromatography-coupled mass spectrometry. We detected cholesterol and 12 other sterol species. We clustered 2 groups based on their specific sterol signature by partial least squares-discriminant analysis (PLS-DA) (Figure 4A, left). The levels of cholesterol were significantly lower in NAD(P)H^{Lo} compared to NAD(P)H^{Hi} CD34⁻ HSCs (Figure 4A, right; Table S4).

We used staining of fixed cells with filipin III (hereafter referred to as filipin) and flow cytometry to confirm cholesterol levels.³⁸ CD34⁻ HSCs showed higher intensity of filipin compared to CD34⁺ HSCs (Figure S5B). Further, fluorescence microscopy analysis of sorted HSCs indicated that NAD(P)H^{Hi} CD34⁻ HSCs have higher filipin intensity than NAD(P)H^{Lo} CD34⁻ HSCs (Figure 4B). Enzymatic determination of total intracellular cholesterol confirmed its enrichment in NAD(P)H^{Hi} CD34⁻ HSCs compared to NAD(P)H^{Lo} (Figure 4C). Consistently, long-term HSC purification based on EPCR positivity displayed the same trend (Figure S5C).

(C) Volcano plot relating the difference in NAD(P)H intensity between each daughter and its mother. (n = 3 independent experiments). Red box: daughters with not statistically different from the mother; gray box: daughter with NAD(P)H significantly lower than the mother; green box: daughter with NAD(P)H significantly higher than the mother.

(D) Timelapse traces of NAD(P)H intensity for each class of division. Cytokinesis is marked by a black arrow; the x axis represents hours of time-lapse imaging since the start of tracking of each mother cell.

(E and F) HSC capacity is enriched in NAD(P)H^{Hi} cells of Div 1 cells. 36 h after *in vitro* culture of CD34⁻ HSCs, single NAD(P)H^{Hi} or NAD(P)H^{Lo} Div 1 cells were isolated onto 96-well plate (also see Figure 2A) then followed by a long-term culture (E) or single-cell transplantation with 4.0×10^5 competitor BMMNCs (F). Colony replating capacity of these single cells (E, n = 48 cells) and the frequency of successful engraftment (>1% donor contribution in multi-lineage hematopoiesis) (F, n = 20) were determined.

(G and H) Percentages of hematopoietic contribution in peripheral blood of recipient mice transplanted with paired daughter cells. Single CD34⁻ HSC division was identified *in vitro* and then each daughter was imaged to determine NAD(P)H levels and individually transplanted with 2×10^5 competitor BMMNCs into irradiated recipient mice. Chimerism was evaluated at 4 months after BMT (10 divisions, for a total of 20 daughter cells analyzed). The division patterns of the investigated division based on donor contribution (H).

BMMNCs, bone marrow mononuclear cells; HSC, Lin⁻Sca-1⁺c-Kit⁺CD135⁻CD150⁺CD48⁻. (E): Student's paired t test. (F): nonparametric Kolmogorov-Smirnov test. ****, p < 0.0001; *, 0.01 ≤ p < 0.05; N.S., not significant, p ≥ 0.05. See also Figure S3.

Farnesyl pyrophosphate, an intermediate of the cholesterol pathway, can be a substrate for protein prenylation as well.³⁹ Results obtained from NAD(P)H^{Hi} and NAD(P)H^{Lo} CD34⁻ HSCs indicate that high NAD(P)H correlates with high levels of protein farnesylation (Figure S5D), confirming the higher flux through the mevalonate pathway. Finally, sorted NAD(P)H^{Hi} CD34⁻ HSCs exposed to FCCP or H₂O₂ display a cholesterol content comparable to the NAD(P)H^{Lo} counterparts (Figure S5E).

Next, we isolated HSPCs and then induced TPNOX and mitoTPNOX expression as previously described (Figures 1L, S2N, and S2O). After 2 weeks of culture, filipin staining indicates that mitoTPNOX, but not cytosolic TPNOX, significantly reduces cholesterol levels (Figure 4D).

Cholesterol biosynthesis requires cytosolic acetyl-CoA as a carbon source. We speculated that in HSCs, FAO supports cholesterol biosynthesis also by providing acetyl-CoA. Mitochondrial acetyl-CoA is transported to the cytosol in form of citrate by the mitochondrial citrate transport protein (CTP, coded by *Slc25a1*), which is then converted to acetyl CoA by ATP citrate lyase (ACLY).^{40,41} CD34⁻ HSCs displayed significantly higher expression of *Slc25a1* compared to CD34⁺ HSCs (Figure S5F), and NAD(P)H^{Hi} CD34⁻ HSCs, exposed *in vitro* to inhibitors of SLC25A1 or ACLY (BTC and BMS-303141, respectively), reduced cholesterol synthesis (Figure S5G). Similar to *Cpt1* levels, single-cell qPCR has confirmed that *Slc25a1* expression correlates with NAD(P)H levels in single HSC daughter cells (Figures S3F and S5H). These results suggest that NAD(P)H production (via FAO) and citrate shuttling undergo the same mechanism of segregation during HSC division.

Both the *in vitro* administration of BIBB-515 (inhibitor of 2,3-oxidosqualene cyclase [OSC], catalyst of a mandatory step of cholesterol biosynthesis)⁴² and the exposure to methyl- β -cyclodextrin (MBCD, an oligosaccharide able to extract cholesterol from the plasma membrane)⁴³ significantly reduced LTC-IC capacity of NAD(P)H^{Hi} but not NAD(P)H^{Lo} CD34⁻ HSCs (Figures S5I and S5J).

These data prompted us to test cholesterol inhibition on an *in vivo* setting. Statins are inhibitors of β -hydroxy β -methylglutaryl-CoA (HMG-CoA)-synthase widely tested in mice and humans. The *in vitro* treatment with lovastatin reduced filipin intensity in NAD(P)H^{Hi} CD34⁻ HSCs to levels comparable to those of NAD(P)H^{Hi} CD34⁻ HSCs exposed to MBCD or NAD(P)H^{Lo} CD34⁻ HSCs (Figure 4E). Low dosage of lovastatin reduced *in vitro* LTC-IC capacity of NAD(P)H^{Hi} CD34⁻ HSCs (Figure S5K) and significantly impaired the *in vivo* reconstitution capacity of NAD(P)H^{Hi} CD34⁻ HSCs in competitive reconstitution assays (Figures 4F and 4G). Further, HSC cholesterol deprivation via MBCD also significantly impaired the *in vivo* reconstitution capacity of NAD(P)H^{Hi} CD34⁻ HSCs (Figures 4F and 4G). The incubation with cholesterol-saturated MBCD (MBCD:Chol) increases cholesterol content in NAD(P)H^{Lo} CD34⁻ HSCs⁴⁴ (Figure 4H), leading to enhanced colony-replating capacity of CD34⁻ HSCs after long-term culture (Figure 4I).

The observations prompted us to test the impact of media composition on the NADPH-FAO-cholesterol axis. Reference media was supplemented with 1% BSA and 50 ng/mL stem cell factor (SCF)/thrombopoietin (TPO) (see STAR Methods and Figure S5L), and we tested the dilution or substitution of BSA as well as lowered the concentration of SCF/TPO to 1 ng/ml (Fig-

ure S5L). In sorted CD34⁻ Lin⁻Sca-1⁺c-Kit⁺ (KSL) cells, only low BSA concentration or its substitution with polyvinyl alcohol (PVA) potentiated the NADPH-cholesterol axis (Figures S5M and S5N). These observations are consistent with previous data showing that a low BSA concentration in the media supports HSC maintenance *in vitro*^{45,46} and stimulates the synthesis of cholesterol and fatty acids.⁴⁵

Altogether, these data indicate that *in vitro* manipulation of cholesterol levels directly impacts the function of HSCs.

FAO sustains NADPH and cholesterol synthesis

We hypothesized that most of the electrons required to reduce the NAD(P)H pool in NAD(P)H^{Hi} CD34⁻ HSCs are derived from FAO.⁴⁷ In agreement, NAD(P)H^{Hi} CD34⁻ HSCs exposed *in vitro* to the Cpt1 inhibitor etomoxir display reduced cholesterol levels (Figure 5A). We thus decided to genetically inactivate FAO. Cpt1 is reported to be sensitive to adenosine monophosphate-activated protein kinase (AMPK) activity,⁴⁸ a known regulator of HSCs,^{49–51} and pharmacological inhibition of Cpt1 can induce oligomerization of the pro-apoptotic members of the Bcl-2 family Bax and Bak.^{52,53} Because of these connections between Cpt1 and factors external to FAO that could bias the data interpretation, we decided to develop *Cpt2* conditional knockout mice (*Cpt2*^{fl/fl} *Vav-iCre*⁺ or *Cpt2* cKO). At steady state, *Cpt2*^{fl/fl} *Vav-iCre*⁺ mice displayed a reduced frequency of CD34⁻ HSCs (Figure S6A) and a significantly diminished proportion of CD34⁻ in the NADPH^{Hi} gate (Figure 5B), while no alterations of mitochondrial mass (Figure S6B) were detected. Reduced intensity of filipin staining was also found in CD34⁻ HSCs from *Cpt2*^{fl/fl} *Vav-iCre*⁺ mice (Figure 5C). Enrichment of cholesterol by MBCD:Chol restores the colony-forming capacity of *Cpt2*-deleted HSCs after long-term culture (Figures 5D and 5E). We then tested the self-renewal capacity of HSCs from *Cpt2* cKO mice. We first analyzed the cell fate impact of NAD(P)H per single mitochondria segregation on the first division of single NAD(P)H^{Hi} CD34⁻ HSCs from *Cpt2*^{fl/fl} *Vav-iCre*⁻ and *Vav-iCre*⁺ mice, as described in Figure 2A. The proportions of asymmetric, symmetric high, and symmetric low distribution in NAD(P)H^{Hi} CD34⁻ *Cpt2*^{fl/fl} *Vav-iCre*⁻ were comparable to that observed in control mice (Figures 2B and 5F). Interestingly, no symmetric high division was observed for NAD(P)H^{Hi} CD34⁻ HSCs from *Cpt2*^{fl/fl} *Vav-iCre*⁺ (0 out of 23), while the proportion of symmetric low rose from 36% (8 out of 22) in the *Cpt2*^{fl/fl} *Vav-iCre*⁻ to 70% (16 out of 23) upon deletion of *Cpt2*. Consistently, competitive reconstitution and serial transplantation assays confirmed reconstitution impairment in recipients of HSCs from *Cpt2* cKO mice (Figures S5G–S5I and S6C).

Cholesterol synthesis regulated by PPAR-FAO contributed to HSC capacity

Peroxisome proliferator-activated receptor- δ (PPAR δ)-specific agonist GW501516, which enhances FAO, was utilized to assess the impact of targeting the FAO-NADPH-cholesterol axis on HSC capacity. Exposure to GW501516 can potentially increase cholesterol levels in NAD(P)H^{Lo} CD34⁻ HSCs (Figure S6D). We then assessed cKO mice for Ppar gamma coactivator 1- α (*Ppargc1a*) in hematopoietic tissues. The *Ppargc1a* gene (or *Pgc1a*) is a co-activator stimulating PPAR factors to induce mitochondrial biogenesis and favor multiple catabolic

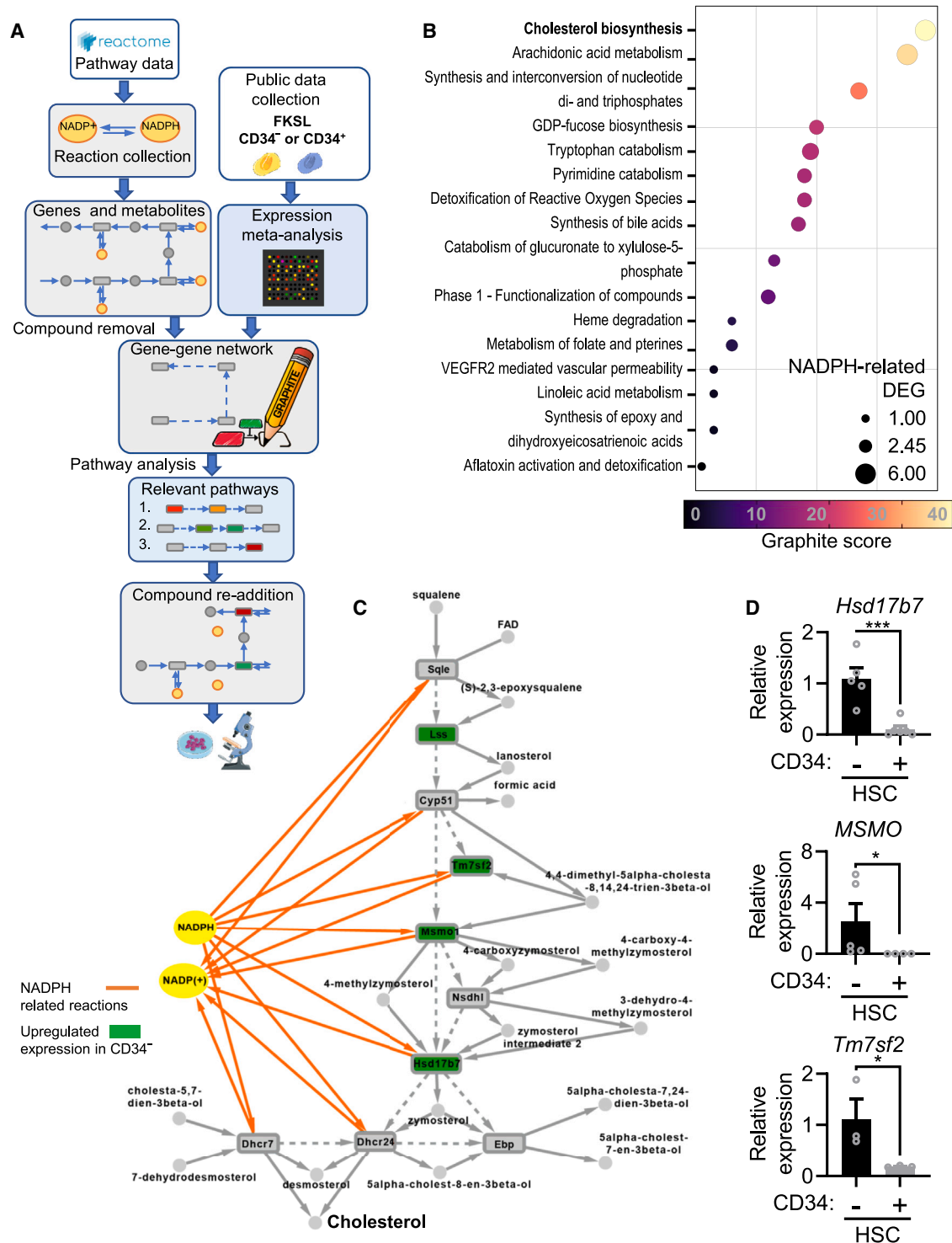


Figure 3. Identification of NADPH-dependent pathways in HSCs

(A) Flow diagram of the pathway analysis used to identify NADPH-dependent reactions differentially regulated in CD34⁺ compared to CD34⁻ HSCs.

(B) NADPH-dependent pathway enriched in the CD34⁺ HSCs and ranked after their Graphite score. Dot size represents the number of NADPH-related differentially expressed genes (DEGs) identified.

(C) Schematic representation of the cholesterol biosynthesis pathway derived from the pathway analysis. Green nodes are differentially expressed genes (all upregulated in CD34⁺ compared to CD34⁻ HSCs). Orange arrows highlight NADPH-dependent reactions.

(D) Levels of *Hsd17b7*, *MSMO*, and *Tm7sf2* in sorted CD34⁻ and CD34⁺ HSCs quantified by qPCR relative to *ActB*. HSC, Lin⁻Sca-1⁺c-Kit⁺CD135⁻CD150⁺CD48⁻; DEG, differentially expressed genes. Bar graphs represent means \pm SEM, circles represent each replicate. (D): unpaired Student's t test.

***, 0.0001 \leq p < 0.001; *, 0.01 \leq p < 0.05. See also Figure S4.

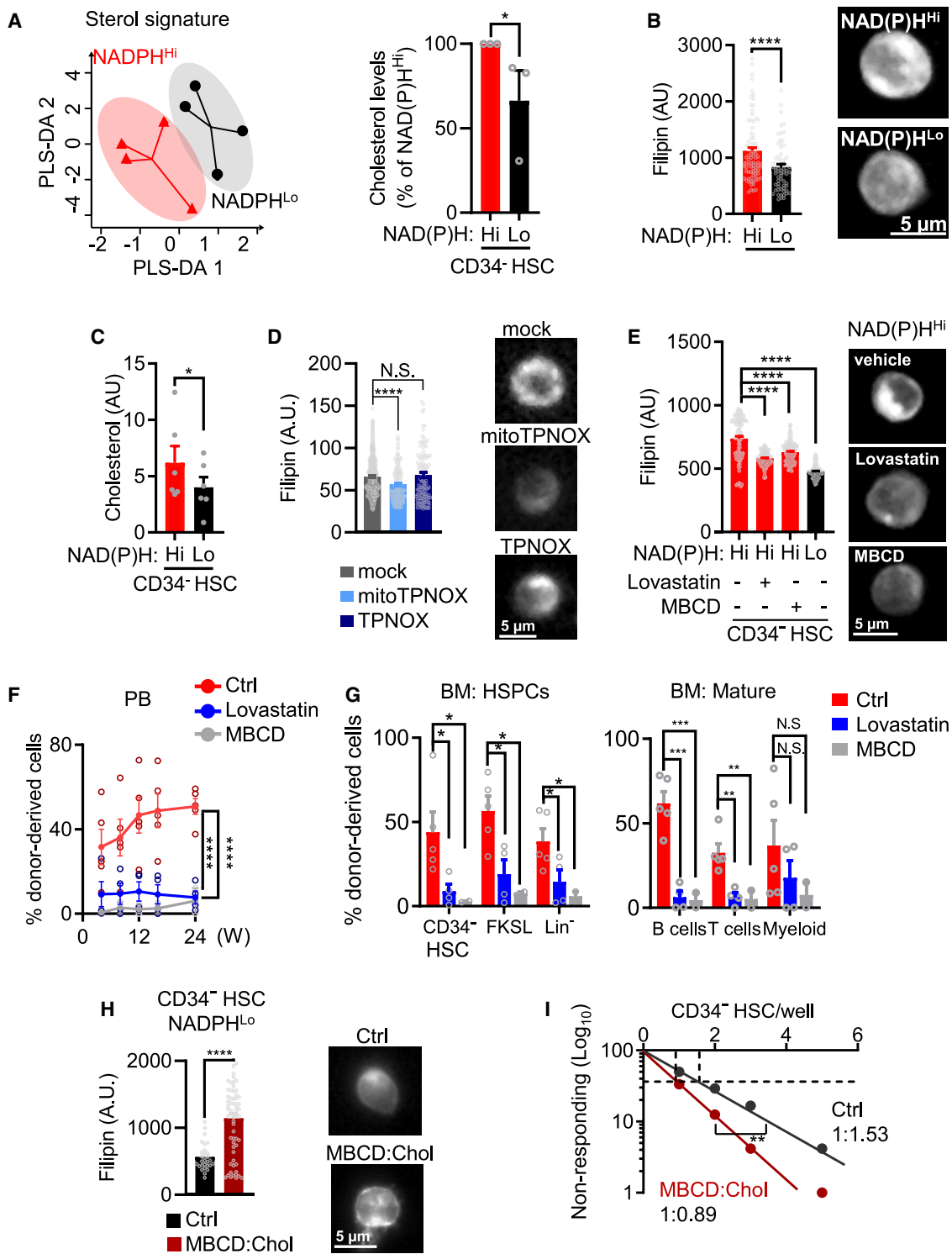


Figure 4. NADPH sustains cholesterol biosynthesis in HSC

(A) Partial least squares-discriminant analysis (PLS-DA) of sterol signature (left) and relative quantitation of cholesterol (right) identified by mass spectrometry in NAD(P)H^{Hi} or NAD(P)H^{Lo} CD34⁺ HSCs (n = 4). Ellipses display confidence level at 95%.

(B) Quantitation (left) and staining (right) of the cholesterol sensitive dye filipin in NAD(P)H^{Hi} and NAD(P)H^{Lo} CD34⁺ HSCs (n = NAD(P)H^{Hi}: 85 cells, NADPH^{Lo}: 67 cells).

(C) Enzymatic detection of total intracellular cholesterol in sorted NAD(P)H^{Hi} or NAD(P)H^{Lo} CD34⁺ HSCs.

reactions.^{54,55} *Pgc1a* deletion induces significant reduction of FAO rates, which can be rescued by administration of GW501516.⁵⁶ Although *Ppargc1a^{fl/fl} Vav-iCre⁺* mice showed no alterations in bone marrow composition or mitochondria numbers (Figures S6E and S6F), CD34⁺ HSCs from *Ppargc1a^{fl/fl} Vav-iCre⁺* mice showed lower NAD(P)H autofluorescence and cholesterol levels than littermate controls (Figures S6G and S6H). *In vitro* GW501516 exposure restored the cholesterol content of *Pgc1a*-depleted HSCs, an effect overcome by forcing NADPH consumption via co-exposure to both FCCP or H₂O₂ (Figures S6H). In a competitive BMT setting (Figure S6I), *Pgc1a*-deleted HSCs showed impaired reconstitution capacity, which was partially but significantly rescued by GW501516 (Figure S6I).

NADPH supports the biogenesis of EVs in HSCs

Our results indicate that proper levels of NADPH support the synthesis of cholesterol, suggesting a role for NADPH in the biogenesis of the plasma membrane or other components of the endomembrane system. Because NAD(P)H^{hi} and NAD(P)H^{lo} CD34⁺ HSCs are both quiescent (Figures S2E–S2G), indicating no demand for membrane biogenesis to support cell division, we hypothesized that higher cholesterol synthesis would be required for the biogenesis of extracellular vesicles (EVs).⁵⁷

Thus, we cultured HSCs in fetal bovine serum (FBS)-free media to allow the enrichment in the media of any spontaneously released EVs. Nanoparticle tracking analysis (NTA) identifies a significant amount of EVs released by HSCs, with an average diameter of ~95 nm (Figure 6A) confirmed by a tunable resistive pulse-sensing (TRPS) system (median diameter of ~112 nm) (Figure S7A). Finally, we used transmission electron microscopy to confirm that EVs from HSCs exhibit a cup-like morphology typical of exosomes (Figure S7B).

Next, we tested whether HSCs could produce transferable EVs using PKH26, a lipid-specific fluorescent marker, which incorporates with double-layered membranes and is retained by EVs after their release (Figure 6B, top). As a negative control, the same number of compensation beads (with no EV release) were processed in parallel (Figure 6B, bottom). HSCs exposed to PKH26-stained EVs only displayed a marked PKH26 signal at the cell edge and in the cytosol (Figure 6C).

NTA further showed that other bone marrow populations, including Nestin⁺ mesenchymal stem cells (MSCs), could also produce EVs (Figure S7C).

To determine the impact of defects in the NADPH-cholesterol axis on EV biogenesis, CD34⁺ HSCs were isolated from *Cpt2^{fl/fl} Vav-iCre⁻* or *Vav-iCre⁺* mice and cultured for 48 h. The conditional deletion of *Cpt2* significantly affected the number of EVs retrieved, but this was partially recovered by exposure of HSCs to MBCD:Chol (Figure 6D). The diameter of these EVs was not affected (Figure S7D).

Tetraspanins are a family of transmembrane proteins found in EVs that participate in their biogenesis and cargo selection.⁵⁸ Cholesterol metabolism supports the cycling of tetraspanins (including CD9 and CD63) within the endomembrane system, and inhibition of cholesterol biosynthesis impairs tetraspanin loading into EVs.^{59–62} Thus, we investigated the expression of CD63 and CD9 on EVs derived from *Cpt2*-deleted HSCs using a flow cytometry and immunomagnetic beads-based methodology. We confirmed that genetic inactivation of FAO resulted in the drastic reduction of CD63 and CD9 in EVs from HSCs (Figures S7E and S7F), and these defects can be reverted by the replenishment of intracellular cholesterol (Figures S7E and S7F).

Magnetic enrichment of EVs cannot resolve marker expression for single EVs but rather provides its expression for the whole EV population. To test the content of EV markers at single EV levels, we used enhanced super-resolution radial fluctuations (eSRRF) microscopy that has been demonstrated to resolve structure close to 100 nm.⁶³ We used commercially available fluorescent beads, which confirmed that eSRRF imaging is in a resolution domain compatible with EV investigation (Figures S7G and S7H). Next, HSC-derived EVs were immunostained for CD63 and imaged using eSRRF microscopy (Figure S7I).

This single-EV technique suggests that conditional *Cpt2* deletion affects CD63 levels in EVs, which were partially recovered by the replenishment of intracellular cholesterol (Figure 6E). Finally, we use a recently developed high-resolution single-particle platform, NanoFCM NanoAnalyzer (nFCM), to investigate the expression of CD63 on HSC-derived EVs.⁶⁴ The nFCM analysis confirmed that CD63 loading into EVs is dependent on the FAO-cholesterol axis and that external supplementation of cholesterol can partially recover CD63 loading in FAO-deficient HSCs (Figure 6F).

Because HSCs actively produce EVs and their biology depends on FAO and cholesterol, we investigated the impact of EVs on HSC function. We measured NAD(P)H levels and mitochondrial mass of CD34⁺ HSCs cultured in the presence of

(D) Quantitation (left) and images (right) of filipin staining in the cultured KSL expressing TPNOX, mitoTPNOX, or empty vector (mock) after 2 weeks of culture with doxycycline (mock, n = 551 cells; TPNOX, n = 119 cells; mitoTPNOX, n = 140 cells).

(E) Quantitation (left) and images of filipin (right) in sorted NAD(P)H^{hi} and NAD(P)H^{lo} CD34⁺ HSCs then cultured for 24 h with 37.5 nM lovastatin, an inhibitor of cholesterol synthesis, or for 2 h after cholesterol sequestration by 1.5 mM methyl- β -cyclodextrin (NAD(P)H^{hi}, n = 141 cells; NAD(P)H^{hi} + Lovastatin, n = 137 cells; NAD(P)H^{hi} + MBCD, n = 102 cells; NAD(P)H^{lo}, n = 107 cells).

(F and G) Analysis of BMT of five NAD(P)H^{hi} CD34⁺ HSCs exposed to 37.5 nM lovastatin (for 24 h *in vitro*), 1.5 mM MBCD (2 h *in vitro*) or Vehicle (Ctrl), with 4.0 \times 10⁵ competitor BMMNCs. (F) Percentages of donor-derived cells in the peripheral blood of recipient mice. (G) Percentages of donor-derived cells in HSPCs (left) or mature cells (right) from the bone marrow of recipient mice at 6 months after BMT.

(H) Quantitation (left) and images (right) of filipin staining in NAD(P)H^{lo} CD34⁺ HSCs exposed to cholesterol-saturated MBCD. n \geq 12 replicates, 3 independent experiments. (I) LTC-IC capacity of CD34⁺ HSCs exposed to cholesterol-saturated MBCD (Ctrl, n = 29, MBCD:Chol, n = 64). PLS-DA, Partial least squares-discriminant analysis; Lov, Lovastatin; HSC, Lin⁻Sca-1⁺c-Kit⁺CD135⁻CD150⁺CD48⁻; MBCD, methyl- β -cyclodextrin. Scale bars, 5 μ m. Bar graphs represent means \pm SEM, circles represent each replicate. (A): nonparametric Mann-Whitney U test. (B) and (H): unpaired Student's t test. (C) and (I): Student's paired t test.

(D) and (E): one-way ANOVA with Dunnett's multiple comparison test. (F) and (G): two-way ANOVA with Sidak's multiple comparison test. ****, p < 0.0001; ***, 0.0001 \leq p < 0.001; **, 0.001 \leq p < 0.01; *, 0.01 \leq p < 0.05; N.S., not significant, p > 0.05. See also Figure S5.

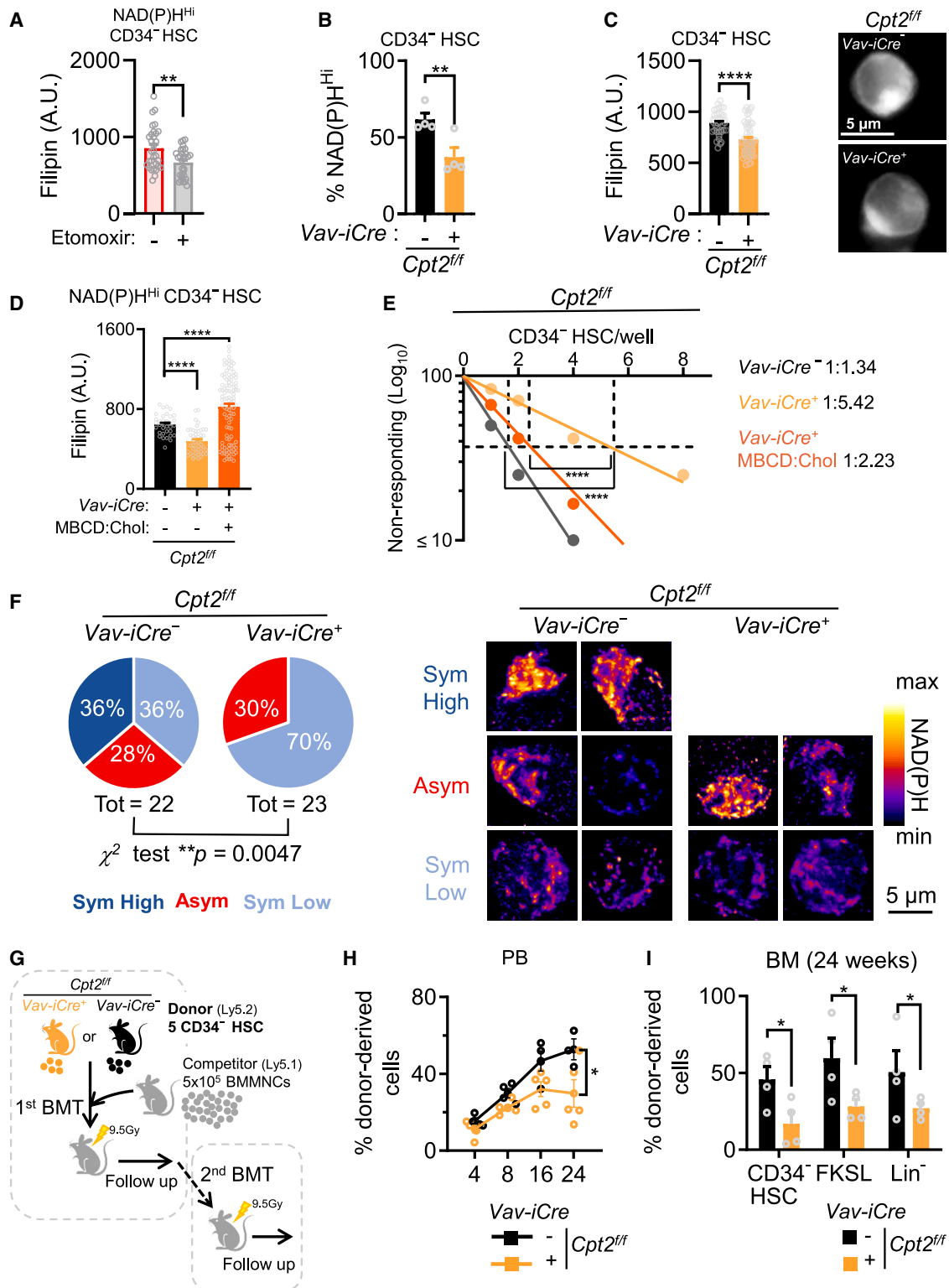


Figure 5. FAO sustains NADPH and cholesterol axis

(A) Quantitation of filipin staining in NAD(P)^H^{Hi} CD34⁻ HSCs exposed to etomoxir for 24 h *in vitro* (vehicle, n = 30 cells; etomoxir, n = 31 cells).

(B) Proportion of NAD(P)^H^{Hi} cells in CD34⁻ HSCs from Cpt2^{fl/fl} Vav-iCre⁻ and Vav-iCre⁺ mice. NAD(P)^H^{Hi} gate was defined as in Figure 1H.

(C) Quantitation (left) and images (right) of filipin staining in CD34⁻ HSCs from Cpt2^{fl/fl} Vav-iCre⁻ and Vav-iCre⁺ mice (Vav-iCre⁻, n = 40 cells; Vav-iCre⁺, n = 56).

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HSC-derived EVs or compensation bead-derived negative controls. EV exposure elevates NAD(P)H levels in acceptor HSCs while lowering their mitochondrial mass (Figures S7J and S7K), two features of mitochondrial physiology that distinguish HSCs from committed progenitors, as indicated by our observation (Figures 1A–1J) and multiple reports.^{7,10,65} In addition, the exposure of HSCs to HSC-derived EVs significantly increased the expression of SCF. This effect was not observed when EVs were obtained from committed progenitor (FKSL or Lin⁻) cells or MSCs (Figures S7L), although HSCs do appear to actively take up EVs from these other origins (Figures S7L and S7M).

We then test whether EV supplementation regulated HSC function by measuring the reconstitution potential *in vivo*. CD34⁻ HSCs were isolated and cultured for 48 h with HSC-derived EVs or vehicle and then subjected to competitive BMT (Figure 6G, left). Strikingly, HSCs supplemented with EVs displayed a higher capacity to reconstitute the hematopoietic system without lineage bias compared to HSCs receiving vehicle in both peripheral blood (Figure 6G) and bone marrow (Figure 6H). These results confirm that HSCs can stimulate their self-renewal potential in an autocrine manner through EV-mediated signaling.

To confirm this new paradigm, we inhibited EV formation *in vitro* by treating HSPCs with GW4869 (inhibitor of neutral sphingomyelinase, widely used to block EV generation)⁶⁶ and then measuring their colony-forming capacity upon serial replating. While vehicle-treated HSPCs were able to produce colonies after three re-platings, cells treated with GW4869 only formed colonies in the first two re-platings, indicating loss of self-renewal (Figure S7N).

Rab27a regulates EV secretion by promoting multivesicular endosome docking at the plasma membrane.⁶⁷ To confirm *in vivo* the inhibition of EV formation on HSC function, we used a Dox-inducible short hairpin RNA (shRNA) for Rab27a, able to reduce the production of EVs *in vitro* (Figure S7O). Sorted FKSL cells were infected *in vitro* and selected with G418 (1 μg/mL) before undergoing competitive BMT (Figure 6I, left). Dox induction (performed 2 weeks after BMT) resulted in a dramatic loss of reconstitution potential without lineage bias in both peripheral blood (Figure 6I) and bone marrow (Figure 6J) 2 weeks after Dox administration and remained stable up to 18 weeks. These findings prompted us to investigate the impact of Rab27a downregulation on HSPCs. We replicated the competitive BMT with infected FKSL cells (as in Figure 6I), focusing the follow-up during the first

3 weeks of Dox induction. In peripheral blood, the effect was visible a week after and became progressively more obvious in the following weeks (Figure S7P). After 3 weeks of Dox induction, mice were euthanized, and defective reconstitution was observed in all the populations tested (Figure S7Q). These results imply that downregulation of Rab27a had a specific impact not only on HSCs but also potentially on the function of progenitor cells.

HSCs reside in a special microenvironment in the bone marrow called the HSC niche (or the niche) that maintains hematopoietic homeostasis through interactions between various niche cells/factors and HSCs.⁶⁸ It has been proposed that EVs participate in this complex microenvironment.^{69–71} Although our data propose that HSC-derived EVs stimulate HSC function in an autocrine manner, we hypothesized that this process could also involve the participation of supporting stromal cells.

Thus, we isolated EVs from HSCs, FKSL cells, and Lin⁻ cells stained with PKH26, as described in Figure 6B. Fluorescently labeled EVs were next administered to bone marrow stromal cells *in vitro*, and the uptake of EVs was evaluated by fluorescence microscopy. Stromal cells preferentially take up EVs released by HSCs compared to those obtained from committed progenitors (Figure S7R). Interestingly, EVs derived from HSCs, but not from committed bone marrow progenitors, were able to potently stimulate SCF expression in stromal cells (Figure S7S). This suggests that EV-mediated signaling instructs stromal cells to provide a microenvironment favorable for HSC maintenance.

To finally corroborate this model, we measured the colony-forming ability of CD34⁻ HSCs after 6 weeks of co-culture with stromal cells in the presence of GW4869. We observed that GW4869 significantly impairs HSC maintenance *in vitro*, confirming the importance of EV biogenesis in hematopoietic homeostasis (Figure S7T).

Overall, these data show that an NADPH-cholesterol axis controls the biogenesis of EVs in HSCs that participate in HSC maintenance through both autocrine- and paracrine-signaling mechanisms.

DISCUSSION

In the current study, we propose that mitochondria can directly support cholesterol synthesis under specific non-respiratory conditions in HSCs. Cholesterol is not synthesized in mitochondria, but we expect that the elevation of TCA cycle metabolites

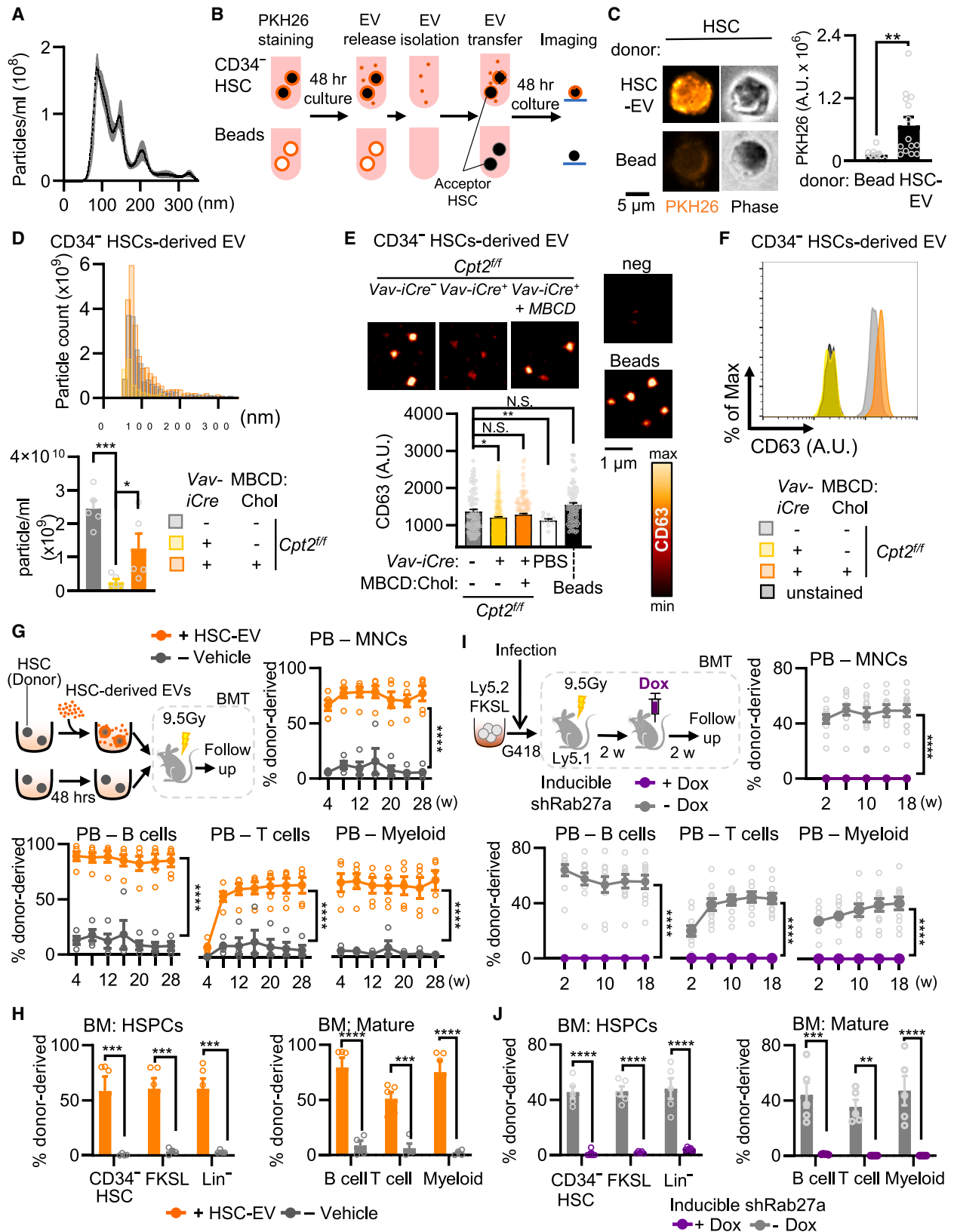
(D and E) Quantification of filipin staining in NAD(P)H^{hi} CD34⁻ HSCs (D; *Cpt2^{fl/fl} Vav-iCre^{-/-}*, n = 34, *Cpt2^{fl/fl} Vav-iCre^{+/+}*, n = 47, *Cpt2^{fl/fl} Vav-iCre^{+/+}* MBDC:Chol, n = 99) and LTC-IC capacity of CD34⁻ HSCs from the indicated genotyped mice (E, n ≥ 12 replicates, 3 independent experiments) exposed to cholesterol-saturated MBDC.

(F) Proportion of symmetric high (Sym High), asymmetric (Asym), or symmetric low (Sym low) distribution of mitochondrial NAD(P)H after first division of the cultured CD34⁻ HSCs from *Cpt2^{fl/fl} Vav-iCre^{-/-}* and *Cpt2^{fl/fl} Vav-iCre^{+/+}* mice (left). Asymmetric distribution of NAD(P)H was determined as in Figure 2G (*Vav-iCre^{-/-}*, n = 22; *Vav-iCre^{+/+}*, n = 23). Images of TMRM and NAD(P)H of pair daughter cells are also shown (right).

(G) Experimental design for serial transplantation with *Cpt2^{fl/fl} Vav-iCre^{-/-}* and *Vav-iCre^{+/+}* CD34⁻ HSCs. Five CD34⁻ HSCs from *Cpt2^{fl/fl} Vav-iCre^{-/-}* or *Vav-iCre^{+/+}* were transplanted into lethally irradiated Ly5.1 recipient mice together with 5x10⁵ competitor BMMNCs (1st BMT). After 6 months of follow up, BMMNC were isolated from each recipient mouse with positive reconstitution and transplanted into secondary recipient mice (1 x 10⁶ BMMNCs/secondary recipient mouse). The hematopoiesis of the recipient mice was followed for 16 weeks.

(H and F) Percentages of donor-derived cells in peripheral blood (H) or bone marrow HSPCs (F) of recipient mice at the indicated weeks after 1st BMT (also see Figure S6C).

BMT, bone marrow transplantation. Scale bar, 5 μm. Bar graphs represent means ± SEM, circles represent each replicate. (A) and (E): Student's paired t test. (B) and (C): unpaired Student's t test. (D): one-way ANOVA with Dunnet's multiple comparison test. (F), χ² test. (H) and (I): two-way ANOVA with Sidak's multiple comparison test. ****, p < 0.0001; ***, 0.0001 ≤ p < 0.001; **, 0.001 ≤ p < 0.01; *, 0.01 ≤ p < 0.05; N.S., not significant, p > 0.05. See also Figure S6.



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and mitochondrial NADH (driven by the high rate of FAO in HSCs) promote the elevation of NADPH through mitochondrial shuttle systems and the redundant reactions that keep NADH and NADPH in equilibrium.⁷² Mitochondrial metabolism in HSCs may generate an outward flux of reduced nicotinamide cofactors and metabolites (e.g., citrate), which would then become available for the synthesis of cholesterol in the cytosol and the endoplasmic reticulum (ER) membrane. Exchange of NADPH between the mitochondria and ER could be favored by their sites of close association, which have been reported to be enriched in cholesterol.⁷³ Also, genetic lowering of these contacts has been shown to affect HSC self-renewal capacity.⁷⁴

A recent study described the impact of citrate conversion to acetyl-CoA in HSCs after 5-fluorouracil (5-FU) induced myeloablation in mouse.⁷⁵ In this report, in the first days after 5-FU administration, HSCs undergo expansion, and in a subsequent phase, they change fate toward differentiation. In this setting, citrate-derived acetyl-CoA supports both self-renewal and differentiation by regulating the histone acetylation landscape and gene transcription.⁷⁵ Our study offers a deeper insight into this mechanism, reporting that in HSCs, citrate and acetyl-CoA derived from FAO are required for cholesterol synthesis (Figure S5G), which is instrumental for self-renewal. This also implies that regulation of histone acetylation might be relevant for HSC differentiation more than self-renewal. In support, our data show that inhibition of the mevalonate pathway by lovastatin or BIBB-515 (expected to boost acetyl-CoA levels) causes HSC exhaustion, strongly deteriorating long-term reconstitution capacity rather than supporting self-renewal capacity (Figures 4F, 4G, S5I, and S5J).

Previous reports have indicated that cholesterol efflux regulates HSC maintenance and that elevation of cholesterol plasma levels or inhibition of its efflux induces proliferation and mobilization of HSCs and progenitor cells via increased expression of the IL-3/GM-CSF receptor¹⁸ or the production of

G-CSF.⁷⁶ Despite these reports, cholesterol levels in purified HSCs and their functional role had yet to be determined. Here, we define that HSCs exhibit high cholesterol synthesis, which favors their self-renewal through a cell-autonomous function. Our results suggest, however, that this is (at least in part) mediated by an NADPH-cholesterol axis supporting the biogenesis of EVs, most likely exosomes.

EVs regulate cell-to-cell communication,⁷⁷ and multiple evidences indicate that some EVs can promote HSC maintenance by several mechanisms.^{78,79} We discovered that HSCs actively release EVs and that EV biogenesis supports the proper maintenance of HSC capacity in concert with the NADPH-cholesterol axis. Most interestingly, we observed that the NADPH-cholesterol axis is required for the proper loading of CD9 and CD63 into HSC-derived EVs. This speculation is confirmed by an early study reporting asymmetric partitioning of the tetraspanins CD53 and CD63 in human cord blood-derived HSCs upon cell division *in vitro*⁸⁰ and is further supported by a recent study that demonstrated that murine HSCs have heterogeneous levels of CD63. The long-term *in vivo* repopulation capacities are enriched in CD63-high HSCs when compared to their CD63-low counterparts.⁸¹

Together with our *in vitro* findings, this evidence suggests that EVs released from HSCs support self-renewal via an autocrine-signaling mechanism. In addition, we observed that other bone marrow populations can produce EVs at levels comparable to HSCs. These include progenitor cells (FKSL), more differentiated precursor cells (Lin⁻), and MSCs. We showed that while HSCs can take up EVs released from all these cells, only HSC-derived EVs can significantly stimulate the expression of SCF in HSCs. Critically, we found that bone marrow stromal cells also preferentially take up EVs isolated from HSCs compared to the ones isolated from more differentiated cells, again inducing a potent expression of SCF. This observation opens the possibility

Figure 6. NADPH supports the biogenesis of extracellular vesicles (EVs) in HSC

(A) Distribution of the diameter of EVs derived from HSCs determined by nanoparticle tracking analysis (NTA).
 (B) Experimental strategy of EV transfer between donor and acceptor HSCs. Donor CD34⁺ HSCs (black) or compensation beads (white) were sorted into FBS-free media, and stained with the lipid marker, PKH26 (orange). EVs were isolated and supplemented to freshly isolated acceptor HSCs, then EV-related PKH26 signal (orange) was determined by fluorescence microscopy.
 (C) Images (left) and quantitation (right) of PKH26 staining in acceptor CD34⁺ HSCs exposed to HSC-derived EVs (HSC-EV) or negative control (bead) (HSC-EV, n = 14 cells; bead, n = 11 cells).
 (D) Particle size distributions of EVs from CD34⁺ HSCs, measured by TRPS system. CD34⁺ HSCs isolated from *Cpt2^{fl/fl} Vav-iCre^{-/-}* and *Vav-iCre^{+/+}* mice were cultured *in vitro* for 48 h. A fraction of cells from *Cpt2^{fl/fl} Vav-iCre^{+/+}* mice were exposed to cholesterol-enriched methyl- β -cyclodextrin (MBCD:Chol). EVs were isolated and analyzed by TRPS (n = 5 independent experiments), frequency distribution of the diameter (top) and average concentration (bottom) are shown.
 (E) The expression of EV marker CD63 at single EV level was determined by enhanced super-resolution radial fluctuation (eSRRF). Images of CD63 in CD34⁺ HSCs-derived EVs (top) and quantitation of CD63 (bottom) in each (*Cpt2^{fl/fl} Vav-iCre^{-/-}*, n = 87; *Cpt2^{fl/fl} Vav-iCre^{+/+}*, n = 332; *Cpt2^{fl/fl} Vav-iCre^{+/+}* + MBCD:Chol, n = 138; PBS, n = 10; beads, n = 83). See also Figures S7G–S7I.
 (F) Nanometric flow cytometry analysis of HSC-derived EVs immunostained for CD63 and investigated by NanoFCM NanoAnalyzer.
 (G) Experimental design (left) and percentages of donor-derived cells (right) in various lineages of peripheral blood of recipient mice transplanted with HSCs supplemented with HSC-derived EVs. Per each recipient mouse, 25 CD34⁺ HSCs were sorted and cultured for 48 h. During this period, HSCs were supplemented with HSC-derived EVs (+HSC-EV; 1.5×10^6 EV per single HSC). EV-free PBS was used as a vehicle. After 48 h of culture, the whole culture was transplanted into lethally irradiated Ly5.1 recipient mice together with 4×10^5 competitor BMMNCs.
 (H) Percentages of donor-derived cells in the indicated fractions of the bone marrow of the recipient mice 28 weeks after BMT.
 (I) Experimental design (left) and percentages of donor-derived cells (right) in various lineages of peripheral blood of recipient mice after Rab27a interfering. After infection and selection with G418, 2,000 live FKSL cells were transplanted into lethally irradiated Ly5.1 recipient mice together with 1×10^6 competitor BMMNCs. Doxycycline was supplemented in water starting at week 2 after BMT. The x axis represents weeks after doxycycline administration.
 (J) Percentages of donor-derived cells in the indicated fractions of the bone marrow of the recipient mice 20 weeks after BMT.
 EV, extracellular vesicles; HSC-EV, HSC-derived extracellular vesicles; FKSL, CD135⁺c-Kit⁺Sca-1⁺Lin⁻. Scale bar, 5 μ m. Bar graphs represent means \pm S.E.M, circles represent each replicate. (C): Unpaired Student's t test. (D) and (E): one-way ANOVA with Dunnett's multiple comparison test. (G), (H), (I), and (J): two-way ANOVA with Sidak's multiple comparison test. ****, p < 0.0001; ***, 0.0001 \leq p < 0.001; **, 0.001 \leq p < 0.01; *, 0.01 \leq p < 0.05; N.S., not significant, p > 0.05. See also Figure S7.

that HSC-derived EVs might regulate HSC self-renewal also by activating paracrine signaling within the complex bone marrow environment where HSCs reside.^{68,82,83} EV-mediated communications among bone marrow cells could lead distinct outcomes influenced by the administration of GW4869. For instance, the study by Hurwitz et al.⁸⁴ focused primarily on HSPCs and short-term exposure to GW4869 and reported that short-term *ex vivo* exposure of HSPCs to GW4869 had an impact on multipotent progenitor (MPP) expansion and potentiated *in vivo* reconstitution (although this was biased toward myeloid lineages).⁸⁴ Our study likewise demonstrated that *ex vivo* exposure of HSPCs to GW4869 increased the number of colonies after initial replating. However, we found that prolonged exposure to GW4869 affected HSC activity, and the GW4869-treated colonies exhausted more rapidly than those produced by vehicle-treated cells, resulting in reduced numbers of colonies formed in subsequent replatings. We speculate that the discrepancies between the experimental designs and the observed results may highlight the existence of EV-mediated communications impacting MPP and possibly HSC function.

In summary, we propose that mitochondria maintain a pool of reduced NADPH in HSCs, which sustains cholesterol biosynthesis and EV biogenesis for proper HSC fate decision and homeostasis. Further studies will evaluate the potential application of this concept to HSC-related diseases and the improvement of HSC-transplantation techniques.

Limitations of the study

The pharmacological or genetic inhibition of EV biogenesis (by means of GW4869 or Rab27a downregulation, respectively) demonstrated the importance of proper EV biogenesis in the maintenance of HSCs and its potential impacts on the function of progenitor cells. We therefore cannot fully exclude the possibility that GW4869- or Rab27a-mediated EVs from HSCs may affect not only HSCs but also the function of progenitor cells or differentiated blood cells. Our experiments likewise cannot exclude that GW4869 exposure or knockdown of Rab27a affects the EV supply from committed hematopoietic cells. Rab27a and the target of GW4869 (neutral sphingomyelinase) are widely recognized as tools for blocking EV biogenesis^{67,85–88}; however, Rab27a is also a regulator of the late endocytic pathway, while neutral sphingomyelinase is important in lipidome regulation. Our data cannot fully exclude a cell-intrinsic role for Rab27a or neutral sphingomyelinase in HSC function. However, we reached the same conclusions from multiple experimental settings/conditions that produced results supporting a positive correlation between EV levels and HSC function. Importantly, the effects of GW4869 and EV supplementation are mediated by mechanisms that are distinct from those of Rab27a inhibition. We expect that future mechanistic exploration of cargo selection in HSCs and progenitor cells will yield new insights at the single-cell level into the specific communications (e.g., autocrine, and paracrine) involved in HSC retention and/or self-renewal capacity within the physiological microenvironment.

STAR★METHODS

Detailed methods are provided in the online version of this paper and include the following:

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SUPPLEMENTAL INFORMATION

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AUTHOR CONTRIBUTIONS

M.B., C.M., N.v.G., Ky. I., I.Z., Y.Z., L.F., and Y.J. contributed to the performance of the experiments and/or analysis of the data. E.C., G.S., and P.M. developed the bioinformatic algorithm and performed bioinformatics assays. N.v.G. performed metabolomics assays. T.N., F.M.L., T.F., C.P.L., B.Z., P.P., I.G., C.R., and D.T.S. provided advice and interpreted data. Ke. I. conceived and directed the study. M.B., C.M., N.v.G., and Ke. I. wrote the manuscript. All the authors contributed to the design of experiments, discussed the results, and commented on the manuscript.

DECLARATION OF INTERESTS

T.F. is a founder and equity holder in Genierian Pharmaceuticals and Coloma Therapeutics. D.T.S. is a director and equity holder of Agios Pharmaceuticals, Magenta Therapeutics, Editas Medicines, Clear Creek Bio, and LifeVaultBio; he is a founder of Fate Therapeutics and Magenta Therapeutics and a consultant to FOG Pharma, VcanBio, and Flagship Pioneering.

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STAR★METHODS

KEY RESOURCES TABLE

REAGENT or RESOURCE	SOURCE	IDENTIFIER
Antibodies		
CD11b-biotin	BD Biosciences	Cat# 553782 ; RRID: AB_394773
CD19-biotin	BD Biosciences	Cat# 553784; RRID: AB_395048
CD45R/B220-biotin	BD Biosciences	Cat# 553086; RRID: AB_394615
CD4-biotin	BD Biosciences	Cat# 553045; RRID: AB_394581
NK-1.1-biotin	BD Biosciences	Cat# 553163; RRID: AB_394675
TER-119-biotin	BD Biosciences	Cat# 553672; RRID: AB_394985
CD117 (c-kit)-APC/CY7	Biolegend	Cat# 105826; RRID: AB_1626278
CD117 (c-kit)-PE	Biolegend	Cat# 105808; RRID: AB_313217
CD150 (SLAM)-PerCp/Cy5.5	Biolegend	Cat# 115922; RRID: AB_2303663
CD48-Pacific Blue	Biolegend	Cat# 103418; RRID:AB_756140
Ly6G/Ly6C (Gr-1)-biotin	Biolegend	Cat# 108404; RRID: AB_313369
CD127-biotin	eBioscience	Cat# 13-1271-85; RRID: AB_466588
CD135 (Flt3)-biotin	eBioscience	Cat# 13-1351-82; RRID: AB_466599
CD34-APC	eBioscience	Cat# 50-0341-82; RRID: AB_10596826
CD34-FITC	eBioscience	Cat# 11-0341-85; RRID: AB_465022
CD3e-biotin	eBioscience	Cat# 13-0031-85; RRID: AB_466320
CD48-APC	eBioscience	Cat# 17-0481-82; RRID: AB_469408
CD63-APC	Life Technologies	Cat# A15712; RRID: AB_2534492
CD8a-biotin	eBioscience	Cat# 13-0081-85; RRID: AB_466347
CD201 (EPCR)-APC	eBioscience	Cat# 17-2012-82; RRID: AB_10717805
Ly-6A/E (Sca-1)-PE/Cy7	eBioscience	Cat# 25-5981-81; RRID: AB_469668
Mouse IgM-biotin	eBioscience	Cat# 13-5790-85; RRID: AB_466676
Streptavidin-APC	eBioscience	Cat# 17-4317-82
Streptavidin-Pacific Blue	eBioscience	Cat# 48-4317-82
CD45-biotin	BD Biosciences	Cat# 553077; RRID: AB_394608
Streptavidin-PE	Biolegend	Cat# 405203
CD31-APC	Biolegend	Cat# 102409; RRID: AB_312905
CD45.1-PE	Biolegend	Cat# 110707; RRID: AB_313497
CD45.2-PE-Cy7	Biolegend	Cat# 109830; RRID: AB_1186098
CD3e-APC	Biolegend	Cat# 100312; RRID: AB_312677
B220-PacificBlue	Biolegend	Cat# 103227; RRID: AB_492876
CD11b-FITC	Ebioscience	Cat# 53-0112-82; RRID: AB_469901
Gr-1-FITC	Biolegend	Cat# 108405; RRID: AB_313371
CD63 magnetic beads	Thermo Fisher Scientific	Cat# 10606D
PE mouse anti-human CD63	eBioscience	Cat# 12-0639-42; RRID: AB_2572565
PerCP/Cyanine5.5 anti-human CD9	BioLegend	Cat# 312109; RRID: AB_2728251
Anti-CD63	Abcam	Cat# ab134045; RRID: AB_2800495
Anti-Farnesyl	Sigma-Aldrich	Cat# 341286
Anti- Ki-67 (D3B5)	Cell Signaling	Cat# 9129; RRID: AB_2687446
Goat anti-Rabbit IgG-Alexa Fluor 488	Thermo Fisher Scientific	Cat# A32731; RRID: AB_2633280
Goat anti-Rabbit IgG-Alexa Fluor 594	Thermo Fisher Scientific	Cat# A11037; RRID: AB_2534095
Chemicals, peptides, and recombinant proteins		
Animal-Free Recombinant Murine TPO	PeptoTech	Cat# AF-315-14
BIBB 515	Cayman Chemical	CAS: 10010517

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REAGENT or RESOURCE	SOURCE	IDENTIFIER
Bovine Serum Albumin	Sigma-Aldrich	Cat# A4503
BMS-303141	Merk	Cat# SML0784; CAS: 943962-47-8
BTC	Merk	Cat# 115983; CAS: 2672-58-4
Etomoxir	Merk	Cat# E1905; CAS: 828934-41-4
FCCP (carbonylcyanide <i>p</i> -trifluoromethoxy-phenylhydrazone)	Millipore-Sigma	Cat# C2920; CAS: 370-86-5
Fetal bovine serum exosome-depleted	Capricorn scientific	Cat# FBS-ED-12F
GW4869	Merk	Cat# D1692; CAS: 6823-69-4
GW-501516	Santa Cruz Biotechnology	Cat# sc-202642< CAS: 317318-70-0
Hydrogen peroxide (H ₂ O ₂)	Millipore-Sigma	Cat# H1009; CAS: 7722-84-1
ITSX	Thermo Fisher Scientific	Cat# 51500056
Lovastatin	United State Pharmacopoea	Cat# 1370600 CAS: 73573-88-3
MethoCult GF M3434	STEMCELL Technologies	Cat# 03434
Methyl-beta-cyclodextrin (MBCD)	Millipore-Sigma	Cat# 332615; CAS: 128446-36-6
Rotenone	Cayman Chemical	Cat# 13995 CAS: 83-79-4
Poly(vinyl alcohol)	Sigma-Aldrich	Cat# 341584; CAS: 9002-89-5
Recombinant Murine SCF	PeproTech	Cat# 250-03
StemSpan™ SFEM	STEMCELL Technologies	Cat# 09600
Tetramethylrhodamine, Methyl Ester, Perchlorate (TMRM)	Thermo Fisher Scientific	Cat# T668
Verapamil	Millipore-Sigma	Cat# V4629; CAS: 152-11-4
Pyronin Y	Merk	Cat# 213519; CAS: 92-32-0
PEG 8000	Merk	Cat# 89510; CAS: 25322-68-3
PKH26	Sigma-Aldrich	Cat# MINI26-1KT
Critical commercial assays		
RNAeasy Mini kit	QIAGEN	Cat# 74104
Cholesterol Assay Kit (Cell based)	Abcam	Cat# ab133116
Total Cholesterol Assay Kits	Cell Biolabs	Cat# STA-384
NADP/NADPH-Glo	Promega	Cat# G9081
NAD/NADH-Glo	Promega	Cat# G9071
SPHERO™ Particle Size Standard Kit #PPS-6K	Spherotech	Cat# PPS-6K
Experimental models: Cell lines		
OP-9	ATCC	ATCC: CRL-2749
Deposited data		
Mendeley data	Source data	https://data.mendeley.com/preview/3g6snt5yzt? a=67b3935a-4810-4a22-af31-7336351b6dcf
MetaboLights	Raw metabolomic data	MTBLS6401
Experimental models: Organisms/strains		
B6.SJL-Ptprca Pepcb/BoyJ	The Jackson lab	JAX: 002014
B6N.Cg-Commd10Tg(Vav1-iCre)A2Kio/J	The Jackson lab	JAX: 008610
B6N.129(FVB)-Ppargc1atm2.1Brsp/J	The Jackson lab	JAX: 009666
C57BL/6	The Jackson lab	JAX: 000664
C57BL/6 Cpt2f/f	T. Finkel Lab, University of Pittsburgh Nomura et al. ⁸⁹	n.a.
Nes-Gfp transgenic mice	S. Frenette Lab, Albert Einstein College of Medicine Mendez-Ferrer et al. ⁹⁰	n.a.
Recombinant DNA		
pLKO-Tet-On-shRab27a	Yang et al. ⁹¹	Addgene 120930
pUC57-TPNOX	Cracan et al. ³⁰	Addgene 87853

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REAGENT or RESOURCE	SOURCE	IDENTIFIER
pUC57-mitoTPNOX	Cracan et al. ³⁰	Addgene 87854
Oligonucleotides		
<i>Scf</i> F- CCTCTCGCAATGCAAAGAAGG	This paper	N/A
<i>Scf</i> R- GAGGTCGGTAGCATATTTGGAAG	This paper	N/A
<i>Msmo1</i> F- AAACAAAAGTGTGGCGTGTC	This paper	N/A
<i>Msmo1</i> R- AAGCATTCTTAAAGGGCTCCTG	This paper	N/A
<i>Tm7sf2</i> F- GTCGCGGCTTTACTGATCCT	This paper	N/A
<i>Tm7sf2</i> R- CAGGCAGATAGGCCGGTAG	This paper	N/A
<i>Hsd17b7</i> F- CCTCTCGCAATGCAAAGAAGG	This paper	N/A
<i>Hsd17b7</i> R- GAGGTCGGTAGCATATTTGGAAG	This paper	N/A
<i>ActB</i> F- GGCTGTATTCCCCTCCATCG	This paper	N/A
<i>ActB</i> R- CCAGTTGGTAACAATGCCATGT	This paper	N/A
Software and algorithms		
GraphPad Prism	GraphPad	https://www.graphpad.com/
Fiji - ImageJ	Schindelin et al. ⁹²	Fiji (imagej.net)
NanoJ-eSRRF	Gustafsson et al. ⁹³	https://github.com/HenriquesLab/NanoJ-eSRRF
CellProfiler	McQuin et al. ⁹⁴	https://cellprofiler.org/
Graphite	Sales et al. ³⁶	Graphite Web (unipd.it)
FlowJo	BD	https://www.flowjo.com/
L-CALC	STEMCELL Technologies	https://www.stemcell.com/l-calc-software.html
Olympus - scanR	EVIDENT scientific	https://lifescience.evidentscientific.com.cn/en/video/scanr-product-overview/
Compound Discoverer 3.0	ThermoFisher Scientific	https://thermo.flexnetoperations.com/control/thmo/login
MetaboAnalyst	Xia et al. ⁹⁵	https://www.metaboanalyst.ca/
Izon Control Suite Software (V3.4)	Izon Science	https://support.izon.com/qnano-control-suite-software

RESOURCE AVAILABILITY

Lead contact

Further information and requests for resources and reagents (including mouse strains, cell lines and plasmids) should be directed to and will be fulfilled by the lead contact, Keisuke Ito (keisuke.ito@einsteinmed.edu).

Materials availability

This study did not generate new unique reagents.

Data and code availability

Source data have been deposited to Mendeley Data and are available at <https://data.mendeley.com/preview/3g6snt5yzzr?af31-7336351b6dcf>. The raw metabolomics data are deposited to MetaboLights (reference number MTBLS6401). Any additional information required to reanalyze the data reported in this paper is available from the **lead contact** upon request.

EXPERIMENTAL MODELS AND STUDY PARTICIPANT DETAILS

C57BL/6 mice (B6-CD45.2, or Ly5.2), B6N.Cg-Commd10Tg(Vav1-iCre)A2Kio/J (*Vav-iCre*), B6N.129(FVB)-Ppargc1atm2.1Brsp/J (*Ppargc1af/f*) and B6.SJL-Ptprca Pepcb/BoyJ (B6-CD45.1, or Ly5.1) were purchased from The Jackson Laboratory. C57BL/6 *Cpt2ff* were kindly gifted by Dr. Finkel.⁸⁹ *Nes-Gfp* transgenic mice were kindly gifted by Dr. Frenette.⁹⁰ GW-501516 was administered at 5 mg per kg body weight per day by oral gavage.⁷ All animal experiments were approved by the Institutional Animal Care and Use Committee of the Albert Einstein College of Medicine. All mice were 2–4 month old at the time of their utilization. Both male and female mice were used.

METHOD DETAILS

Flow cytometry and cell sorting

Bone marrow mononuclear cells (BMMNCs) were isolated by the flushing technique. Briefly, bone marrow from the femora and tibia was flushed in FACS buffer (PBS +2% FBS). After red blood cells lysis with ACK Lysing Buffer (Gibco), the bone marrow was filtered using 75mm cell-strainer cap (Falcon). After isolation, BMMNCs were incubated with the following mix of monoclonal antibodies against lineage: Ly6G/Ly6C (Gr-1)-biotin (Cat# 108404) from Biolegend; CD11b-biotin (Cat# BDB553309), CD19-biotin (Cat# BDB553784), CD45R/B220-biotin (Cat# BDB553086), CD4-biotin (Cat# BDB553782), NK-1.1-biotin (Cat# BDB553163) and TER-119-biotin (Cat# BDB553672) from Fisher Scientific; CD135 (Flt3)-biotin (Cat# 13-1351-82) from eBioscience; CD127-biotin (Cat# 13-1271-85), CD3e-biotin (Cat# 13-0031-85), CD8a-biotin (Cat# 13-0081-85), Mouse IgM-biotin (Cat# 13-5790-85) from Life Technologies, and then resuspended in 2% FBS-PBS for 30 min in ice. Lineage antibody staining was followed by incubation with antibody mix for the following HSPC markers: CD117 (c-kit)-APC/CY7 (Cat# 105826) or CD117 (c-kit)-PE (Cat# 105808), CD150 (SLAM)-PerCp/Cy5.5 (Cat# 115922), CD48-Pacific Blue (Cat# 103418) from Biolegend; CD201(EPCR)-APC (17-2012-82), CD34-APC (Cat# 50-0341-82) or CD34-FITC (Cat# 11-0341-85), CD48-APC (Cat# 17-0481-82), Ly-6A/E (Sca-1)-PE/Cy7 (Cat# 25-5981-81), Streptavidin-APC (Cat# 17-4317-82) or Streptavidin-Pacific Blue (Cat# 48-4317-82) from eBioscience; resuspended in 2% FBS-PBS for 30 min in ice. All antibodies were used at a 1:100 dilution.

Samples were acquired on an LSR II flow cytometer (Becton Dickinson) then data was analyzed using FlowJo 10 (Becton Dickinson). For Pyronin Y, freshly isolated BM stained for surface markers were stained with Pyronin Y 1 μ M (Sigma-Aldrich) at 37°C for 20 min and analyzed by flow cytometry. For cell sorting, BMMNCs stained with antibody mix were sorted directly into StemSpan SFEM (STEMCELL Technologies, Cat# 09600). Stem and progenitors cell fractions were isolated from lineage-depleted samples, as necessary. Cell sorting of CD34⁻ HSC was performed on a BD FACSAria II (Becton Dickinson). To sort NAD(P)H^{Hi} versus NAD(P)H^{Lo} CD34⁻ HSC were sorted using MoFlo XDP (Beckman Coulter). Briefly, BMMNCs stained for surface markers were subdivided into two aliquots then incubated for 5 min at 37°C in 2% FBS-PBS in presence or absence of FCCP 1 μ M. To ensure the consistency of our strategy, we confirmed that this gating process identified <2% of FCCP-exposed CD34⁻ HSCs as NAD(P)H^{Hi} (Figures 1H and S2A), while ~60% of untreated CD34⁻ HSCs were included in the NAD(P)H^{Hi} gate.

For MSC sorting, BM primary cells were isolated from femurs and tibia of Nes-*Gfp* transgenic mice. BM were crushed and digested with 2 mg/mL Collagenase type IV (Gibco, 17104-019) and 4 mg/mL Dispase (Gibco, 17105-041) in HBSS (Gibco) for 30 min at 37°C, following erythrocyte lysis. For flow cytometry sorting, cells were stained and enriched by immunomagnetic depletion using anti-biotin magnetic beads (Miltenyi Biotec) binding CD45-biotin (BD Biosciences, 553077) and Ter119-biotin (BD Biosciences, 553672), according to the manufacturer's recommendations. Cells were stained by Streptavidin-PE (Biolegend, 405203), CD31-APC (Biolegend, 102409), and sorted on a FACSAria (BD) to >95% purity using Nes-*Gfp* as MSC marker.⁹⁰

NAD(P)H analysis by flow cytometry

BMMNCs were isolated and stained for surface markers as described above. After staining, samples were subdivided in 3–5 fractions, each of them was incubated for 5 min at 37°C in 2% FBS-PBS in presence of FCCP 1 μ M, or H₂O₂ 100 μ M or rotenone 100 nM. After incubation, samples were recorded at LSRII, NAD(P)H autofluorescence was detected in the Indo-Violet channel (excitation laser at 350 nm, band-pass emission 450/50 nm). When indicated, samples exposed with FCCP or H₂O₂ were eventually incubated with H₂O₂ 100 μ M or rotenone 100 nM (respectively) at 37°C for additional 5 min then acquired at LSRII. To compensate for artifact related to cell size, autofluorescence intensity was normalized on FSC-A values, then scaled by multiplying for 10⁴. This index was utilized as a reporter of NAD(P)H concentration (Figure S1D). The use of size calibration beads (SPHERO Particle Size Standard Kit #PPS-6K, Spherotech, Lake Forest, IL, USA) confirms that the FSC-A can be used as a readout of cell size.

HSCP culture conditions

Sorted HSC or KSL were cultured in StemSpan SFEM (STEMCELL Technologies, Cat# 09600) supplemented with SCF (PeproTech, Cat# 250-03) and TPO (PeproTech, Cat# AF-315-14) at 50 ng/mL unless otherwise described. According to the experimental need, the media could be supplemented with one of the following chemicals: Etomoxir 10 μ M (Merk, #Cat E1905), Lovastatin 37.5 nM (United State Pharmacopoea, #Cat 1370600), GW4869 10 μ M (Merk, #Cat D1692), GW501516 0.1 μ M (Santa Cruz Biotechnology, #Cat sc-202642), BIBB 515 12.5 μ M (Cayman Chemical, #Cat 10010517), BTC 5 mM (Merk, #Cat 115983), BMS-303141 1 mM (Merk, #Cat SML0784), MBCD 1.5mM (Sigma- Aldrich, #Cat 332615). For media testing (Figure S5L–S5N) sorted CD34⁻ KSL were cultured with a handmade media adapted from.⁴⁶ Briefly the media was composed of F12, 1% ITSX, 1% (Thermo Fisher Scientific, Cat# 51500056), 10 mM HEPES, 1% P/S/G. This base media was further supplemented with bovine serum albumin (1% or 0.1%; Sigma-Aldrich, Cat# A4503), or poly(vinyl alcohol) 0.1% (Sigma-Aldrich, Cat# 341584), or exosome free-FBS 1% (Capricorn scientific, Cat# FBS-ED-12F), and SCF and TPO (50 ng/mL or 1 ng/mL).

Imaging of mitochondrial markers

Sorted cells were centrifuged at 400 g for 3 min at 4°C, resuspended in 30 μ L of StemSpan SFEM, then seeded on a Lab-Tek II Chamber Slide (Thermo Fisher Scientific, Cat # 154534PK) coated with Retronectin (Takahara Bio, Cat# T100B), stained with TMRM 2 nM, and incubated for 60 min to let cells adhere.

Each cell was acquired on a Delta Vision Core (GE Healthcare Life Sciences) epifluorescence microscope equipped with a 60X oil immersion lens (NA 1.4). The nicotinamide moiety of NADH absorbs light at approximately 340 nm, and emits fluorescence at 460 nm.⁹⁶ The intensity of this fluorescence reflects NADH and/or NADPH (cumulatively referred to as NAD(P)H hereafter) and it was collected using excitation filter 377/50nm and emission filter 447/60 nm. For NAD(P)H analysis in Figure S7J, single planes were acquired. After flat field correction and background subtraction, the average intensity of NAD(P)H for each cell was collected. For 3D imaging in Figures 1C, 1H, 2A, 2B, 5F, S2J, S6B, and S6E z stack were acquired with a z spacing of 0.3 μm and pixel size of 107nm. After acquisition, stacks were reconstructed by digital deconvolution using the Richardson-Lucy algorithm implemented in the DeconvolutionLab2 plugin available for Fiji.⁹² PSFs were manually measured by imaging PS-Speck Microscope Point Source Kit (Thermo Fisher Scientific, Cat# P7220).

After deconvolution, TMRM and Autofluorescence signals were segmented to identify mitochondria using a custom macro for Fiji. Briefly, to compensate for variation in expression levels, the histogram was stretched setting a maximal value for the top 0.1% of the original values. TMRM signal was emphasized by a 3D Top Hat filter (kernel $8 \times 8 \times 2$ pixel), then thresholded using Yen algorithm. Segmentation objects were identified and measured for fluorescence intensities and shape using the 3D suite.⁹⁷ Representative image renderings were obtained by Fiji.

3D imaging of mitochondrial NAD(P)H symmetry between daughter cells

Single CD34⁻ HSC were sorted and cultured as above mentioned. At 24 h after sorting, cells were visually inspected on an hourly basis for the occurrence of first division. Daughter cells were then moved to a RetroNectin-coated 35 mm glass bottom dish (Thermo Fisher Scientific, Cat# 150680), resuspended in 50 μL of StemSpan SFEM media supplemented with TMRM 2 nM and verapamil 50 μM . After 60 min of incubation 3D images were acquired, processed, and quantified as previously described. Average autofluorescence intensities for each mitochondrial particle were grouped for each daughter and compared with the values obtained from the paired daughter by Mann-Whitney U test. This offered a statistical comparison of mitochondrial autofluorescence distribution between paired daughters. Those pairs whose test provided a p value <0.05 were considered the result of an asymmetric segregation. The remaining were labeled symmetric. Those symmetric paired daughters whose median NAD(P)H intensity was above the median value of the whole dataset were considered symmetric high, on opposite, those symmetric paired daughters whose median NAD(P)H intensity was below the median value of the whole dataset were considered symmetric low (Figure S3A). The proportion of division between the genetic background described in Figure 5F was evaluated for statistical significance using χ^2 test.

Live NAD(P)H imaging during HSC first division

For NAD(P)H imaging in Figures 2C and 2D, and 3B-3D CD34⁻ HSC were sorted, and culture as described above. At 24 h from isolation, cells were imaged on a Delta Vision Core using a 40x/1.35 oil immersion lens (UAPON40XO340-2, Olympus) under controlled environmental conditions (temperature at 37°C and CO₂ at 5%). Images were acquired every ~ 40 min to minimize toxicity, for approximately 20 h. For every time point brightfield images were acquired to correct focus drifts. Images were corrected for shading and background, then cell divisions were manually tracked. NAD(P)H images were thresholded to collect only mitochondrial signal then average intensity was collected per each cell at each time point. Image processing and quantitation was obtained via FIJI software.⁹² The NAD(P)H intensity of each tracked daughter and its related mother was compared by 1-way ANOVA. Divisions where only one daughter was significantly different from the mother was classified as asymmetric. Those divisions with both daughters significantly different from mother were classified as symmetric low, and the remaining division were classified as symmetric high.

Enzymatic NADH and NADPH assay

4000-5000 CD34⁻ and CD34⁺ HSC were sorted as aforementioned. Samples were centrifuged at 400g for 3 min, deprived of the supernatant, then washed once in ice-cold PBS. After washing samples were split into two parts, and one exposed to FCCP 1 μM , NADPH and NADH were analyzed using the NADP/NADPH-Glo or NAD/NADH-Glo enzymatical kit (Promega, Cat# G9081, and Cat# G9071) as described by the manufacturer. Final luminometric readings were normalized on the number of cells sorted for each sample.

FLIM analysis

Images were acquired using a Leica TCS SP8 confocal microscope equipped with a tunable (680–1300 nm) femtosecond laser (InSight Deep See; Spectra-Physics) and an oil-immersion 40 \times objective (NA 1.3). TPEF images of the TMRM signal (1024 \times 1024 pixels, 290.6 \times 290.6 μm) from the fractions of interest were acquired at 720 nm excitation and 624 \pm 20 nm emission ranges. Location co-registered NAD(P)H lifetime data were then acquired at 720 nm excitation and 460 \pm 25 nm emission ranges with 2 min integration time (512 \times 512 pixels) using a Picoquant PicoHarp 300 time-correlated single photon counter and SymPhoTime analysis software. A combination of two-dimensional (2D) discrete Fourier transform and power spectral density (PSD) methods were applied on binarized images to segment out the whole cell regions.²⁷ Roundness ($4\pi \times \text{area}/\text{perimeter}^2$) of each segmented object was calculated and only the ones that above 0.6 were considered as cells to avoid taking debris or cell fragments into account. Afterward, for each cell, a three-level global Otsu's threshold was applied on the TMRM intensity images, and the pixels above the second threshold level were considered as mitochondria regions. Binary masks for whole cell region and mitochondria region were then saved for the following lifetime analyses. FLIMJ (ImageJ plugin) and MATLAB were used for lifetime analysis. A kernel size of four and an MLE (maximum likelihood estimation) noise model were used for pixel-wised 2-component lifetime curve fittings in the FLIMJ. The

resultant longer lifetime τ bound maps were calculated and exported. The pixel-wise NADPH/NADH ratio values were calculated based on method previously published.²⁵ For each cell, the averaged NADPH/NADH ratio values from all pixels in the mitochondria and whole cell regions were calculated in MATLAB based on the corresponding segmented binary masks.

Transmission electron microscopy

For mitochondrial analysis, 5000 sorted stem and progenitor cells were fixed in 2.5% glutaraldehyde, 2% paraformaldehyde in sodium cacodylate buffer 0.1 M, pH 7.4 for 3 h at 4°C. Samples were post-fixed in 1.0% aqueous osmium tetroxide (pH 7.4) followed by 1% uranyl acetate, dehydrated in a graded series of ethanol, and embedded in LX112 resin (LADD Research Industries). Ultrathin (80nm) sections were cut on a Leica UC7, contrasted with uranyl acetate followed by lead citrate, and viewed on a JEOL 1200EX transmission electron microscope (Jeol Ltd.) at 80 kV. Images were manually segmented by the TrakEM2 plugin available on Fiji.⁹⁸

For isolated EVs, EVs were deposited on carbonated grids, fixed in 2% PFA and stained with a contrasting solution of methyl-cellulose – as reported by Corona M.L. and colleagues.⁹⁹ Samples were then imaged with a TEM Zeiss EM 910 microscope (Zeiss).

Competitive reconstitution assay

Sorted HSC fractions from Ly5.2 donor were transplanted into lethally irradiated Ly5.1 congenic mice in competition with BMMNC. Reconstitution of donor (Ly5.2) cells and repopulation of donor myeloid and lymphoid cells were monitored by staining blood cells with antibodies against CD45.1-PE (Biolegend, Cat# 110707), CD45.2-PE-Cy7 (Biolegend, Cat# 109830), CD3e-APC (for T cell, Biolegend, Cat# 100312), B220-PacificBlue (Biolegend, Cat# 560472), CD11b-FITC and Gr-1-FITC (for myeloid cells, Ebioscience, Cat# 53-0112-82, and Biolegend, Cat# 108405 respectively). Analysis of chimerism in HSPC and fully differentiated hematopoietic cells in the bone marrow was obtained at 24 weeks after BMT. Isolation, staining and analysis of BMMNC at this stage was obtained as described in the ‘flow cytometry and cell sorting’ section.

To assess the roles of mitochondrial NADPH in a transplantation setting (Figure 1M), KSL (Ly5.2) cells were infected with mitoTPNOX, TPNOX or vector control. After puromycin selection, the infected 1000 KSL cells were transplanted into the irradiated recipient mice (Ly5.2) with 4×10^5 competitor BMMNCs (Ly5.1). Then after 3 weeks after transplantation, mice were kept with Doxycycline (1% in 5% sucrose water) to induce transgene expression. In some experiments, donor contribution in the immature hematopoietic lineages was also evaluated. For the serial transplantation assay, donor-derived cells were collected from recipient mice 6 months after BMT and transplanted into the recipient mice.^{6,7}

To assess EV functionality *in vivo* (Figure 6G), 125 sorted CD34⁺ HSC (Ly5.2) were cultured *in vitro* in StemSpan SFEM media supplemented with SCF and TPO (both at 50 ng/mL). Immediately after sorting, cells were supplemented with a bolus of HSC-derived EV (+HSC-EV). After 24 h of culture a second bolus of HSC-derived EV was administered to the same cultured HSC. Each bolus contained 0.4×10^9 EV resuspended in 10 μ L of filtered PBS, so to have approximately 1.5×10^6 EV per HSC. In parallel, a second group of HSCs were treated with the same volume of filtered PBS free from EV (vehicle). After 48 h from sorting the culture of donor cells was split into 5 parts, and each of them (estimated 45 donor cells) transplanted into one lethally irradiated Ly5.1 recipient mouse together with 4×10^5 competitor BMMNCs (Ly5.1).

To confirm *in vivo* the effect of the inhibition of EV formation on HSC function (Figure 6I), we used a Doxycycline inducible shRNA for Rab27a. Sorted FKSL (Ly5.2) cells were cultured in StemSpan SFEM media supplemented with SCF and TPO (both at 50 ng/mL) and infected with shRab27a inducible lentivirus (diluted 1:500). After 48 h of G418 (1 mg/ml) selection, 2000 live FKSL/mouse were transplanted in lethally irradiated recipient mice in competition with 1×10^6 BMMNCs (Ly5.1). After 2 weeks from BMT, mice were kept with 1% Doxycycline to induce transgene expression.

Lentiviral particle production and infection

pUC57-TPNOX and pUC57-mitoTPNOX were gifted from Vamsi Mootha (Addgene plasmid # 87853; <http://n2t.net/addgene:87853>; RRID:Addgene_87853 and plasmid # 87854; <http://n2t.net/addgene:87854>; RRID:Addgene_87854).³⁰ For expression in mammalian cells synthetic TPNOX and mitoTPNOX genes were subcloned into a pLVX-TetOne vector (Clontech, Cat# 631844) using BamHI and AgeI restriction sites. pLKO-Tet-On-shRab27a was a gift from Mien-Chie Hung (Addgene plasmid # 120930; <http://n2t.net/addgene:120930>; RRID:Addgene_120930).¹⁰⁰

For particles production, 5×10^6 HEK293T cells were seeded in a 10 cm dish 24 h before transfection and cultured in DMEM medium. Cells were then transfected with 3 μ g of VSVG, 5 μ g of pMDL, 2.5 μ g of pRev, and 10 μ g of Transfer Plasmid with 30 μ L of lipofectamine 2000 (Thermo Fisher Scientific, Cat# 11668019). At 24 h after transfection media was refreshed and supplemented with sodium butyrate 1mM. At 24 h after media change, the lentivirus-enriched media was collected and filtered through a 0.45 μ M filter and stored at -80°C . the procedure as repeated twice.

Collected media was ultracentrifuged at 90000 x g at 4°C in an SW40 Ti rotor (Beckman Coulter, Cat# 331301) for 90 min. After spinning, the supernatant was discarded, and the pellet gently resuspended in 300 μ L of PBS for 16 h at 4°C upon gentle shaking. Lentiviral particle batches were all stored at -80°C .

Optimal infection was experimentally determined for each lentiviral batch. 1000 FKSL cells were sorted into StemSpan SFEM and infected with 50; 25; 12.5; or 6.12 μ L of lentiviral preparation. At 24 h after infection cell toxicity was determined by trypan blue incorporation. The maximum amount of lentiviral preparation with minimal toxicity was chosen for subsequent experiments. Infected cells used then selected with puromycin (1 μ g/mL) or G418 (1 mg/mL) for 48h before BMT.

Colony-forming assays

Frequencies of LTC-ICs were calculated using a limiting-dilution method (maximum likelihood from the proportion of negative wells measured for each input dilution of cells assessed). In each experiment, at least 12 (LTC-IC) replicates of each dilution were performed, and a split-plot analysis of variance was used to investigate the difference between the two conditions. The analysis was performed using L-CALC software (STEMCELL Technologies, <https://www.stemcell.com/l-calc-software.html>). 12.5 μM BIBB 515 and 1.5mM MBCD (Sigma- Aldrich, 332615, 2 h exposure/week) was added to the media where indicated.

The colony formation assay was performed using MethoCult GF M3434 (STEMCELL Technologies, Cat# 03434) according to the manufacturer's instructions. Briefly, Lin-Sca-1+c-Kit+ cells (300 cells/plate, triplicate culture) were sorted and seeded into methylcellulose-based medium. Colonies were counted under microscope after 7 days incubation at 37°C, 5% CO₂. For replating assay with infected cells, sorted KSL were seeded in StemSpan SFEM to run infection and selection as described above before seeding into MethoCult GF M3434. Where indicated liquid or semisolid media were supplemented with reagents to induce Tet-On system (Dox 300 ng/mL) or manipulating cholesterol or exosome biogenesis.

Paired daughter cell assays

Single CD34⁺ HSC were sorted and cultured to track first division as described above. Daughter cells derived from a single parent cell were separated and were subjected to the competitive transplantation assay.^{6,7,31} After initial division, daughter cells from the same parent cell were separated, and NAD(P)H intensity was determined by fluorescence microscopy. After checking NAD(P)H intensity, a single daughter cell (Ly5.2) was transplanted into lethally irradiated recipient mice (Ly5.1) in a competitive assay. Donor chimerism was analyzed periodically. Long-term repopulation was achieved when more than 1% of the contribution in the peripheral blood of recipient mice was observed at the indicated time after transplantation.

Identification of NADPH dependent pathways

Initially, all mouse pathways containing NADPH (CHEBI: 16474; 57783; 77177; 77312) were collected from the Reactome database.³³ Topological pathway analysis was used to analyze the 77 collected NADPH pathways and compare the gene expression in ST and LT-HSCs. For this purpose, 3 public expression datasets were collected and re-analyzed (GSE94663,³⁴ GSE77098,³⁵ GSE15907⁹¹). Datasets were chosen by similarity of sample preparation. Only genes common to all 3 datasets were considered, with a resulting matrix of 17592 genes of which 3737 displayed significant variations. Gene expression was corrected to avoid batch effects using the ComBat method.¹⁰¹

Normalized expression matrix was used to screen the NADPH pathways using Clipper methods, (details in^{102,103}) and protein-metabolite Graphite networks³⁶ modified to retain NADPH metabolites in pathway-derived graphs. The results of this analysis were (i) a new pathway (represented in Figures 3C and S4C), obtained by merging all the chains of reactions involving NADPH that are differentially regulated between the two class of samples (CD34⁺ and CD34⁻), and (ii) a rank of the studied chains of reactions utilizing their clipper max score (Figure 3B). Differentially expressed genes were identified with the Samr package in R software once the data were normalized and mapped on the network with different node colors.

Metabolomic analysis

10000 to 20000 NAD(P)H^{Hi} and NAD(P)H^{Lo} CD34⁻ HSC were sorted directly into 500 μL 80% ice-cold methanol supplemented with 13C-labeled cholesterol (400nM 23,24,25,26,27-13C5-cholesterol, Cambridge Isotope Laboratories, Cat# CLM-9587). Polar and non-polar metabolites were extracted using methanol-chloroform phase separation.

For untargeted metabolomics analysis, polar phases were dried under nitrogen gas, resuspended in 120 μL 70% acetonitrile in water containing 2.5 mM of an internal standard (13C-, 15N-labeled amino acid mix; Cambridge Isotope Laboratories, Cat# CLM-9587), transferred to glass vial with microinserts, re-dried and finally resuspended in 25 μL 70% acetonitrile in water. Samples were run on a ThermoFisher Q-Exactive Orbitrap mass spectrometer equipped with Zic-pHILIC column (150 \times 2.1 mm, 5 μm ; Merck). A volume of 5 μL was injected and metabolites were monitored in full-scan, polarity-switching, mode (0–45 min, resolution 70,000, AGC target 3 \times 10⁶, m/z range 66.7–1000). Mobile phase A for chromatography consisted of 20 mM ammonium carbonate, 0.1% ammonium hydroxide, in water and mobile phase B of 97% acetonitrile in water. A pooled sample was created from all samples and used for MS/MS runs. Metabolite measurements were normalized to the internal 13C/15N-labelled amino acid standard. Data were processed with Compound Discoverer 3.0 (ThermoFisher Scientific), among all detected compounds (2965), 100 were identified based on their mass spectra using the mzCloud database. Another 643 could be putatively identified using their molecular weight by searching the Human Metabolome database and Kyoto Encyclopedia of Genes and Genomes (KEGG) Compound database (Table S1). The metabolomics data were then further analyzed using the MetaboAnalyst software. Statistically, 32 compounds differed significantly between the groups (t-test, $p < 0.01$; Table S2).

For sterol analysis, the non-polar phases were dried under nitrogen gas and resuspended in 500 μL transesterification solution (methanol:acetyl chloride in a 9:1 ratio). 300 μL hexane was added, samples were vortexed and incubated at 95°C for 10 min. Samples were then supplemented with 1 mL water and 500 μL hexane, vortexed for 1 min and centrifuged at 14,000 \times g for 15 min. The upper phase was transferred in a glass vial, dried under nitrogen gas, resuspended in 500 μL of -20°C chloroform, vortexed for 1 min and centrifuged. The bottom phase was then transferred to a new glass vial, and samples were extracted in chloroform a second time and re-dried. Samples were finally resuspended in 25 μL hexane to which 25 μL derivatization agent (N-Methyl-N-(trimethylsilyl) trifluoroacetamide +1% trimethylchlorosilane) was added, vortexed for 1 min and incubated at 80°C for 40 min. Samples were

transferred to glass inserts and analyzed on a Q-Exactive mass spectrometer (Thermo Fisher Scientific) equipped with an Agilent DB-5MS 30m, 0.25mm, 0.25 μ m column. Injection parameters: automatic injection, 1 μ L, 300°C, splitless for 1.5 min before purge at 50 mL/min. Gas chromatography parameters: oven at 60°C for 5 min, then to 300 °C at 16 °C/min, final temperature for 30 min; column in constant flow mode at 1 mL/min, all transfer lines at 350°C. Mass spectrometry parameters: EI + ion source at 350°C, full MS, 60,000 resolution, AGC target 1e6, autoIT, scan range 66.7–1000 m/z, 7.5 min solvent delay. Data were analyzed in Compound Discoverer (version 3.0) and normalized to the ¹³C-labeled cholesterol internal standard. Of the sterols identified, only those specific to animal cells and detectable in all replicates were kept for analysis.

Filipin staining

Sorted NAD(P)H^{Hi} and NAD(P)H^{Lo} CD34⁻ cells were centrifuged at 400 g for 3 min at 4°C, resuspended in 30 μ L of StemSpan SFEM then seeded on Lab-Tek II Chamber Slide (Thermo Fisher Scientific, Cat# 154526) coated with Fibronectin (Sigma-Aldrich, Cat# F1141) and incubated for 30 min to let cells adhere. After incubation cells were fixed and stained with filipin according to the Cholesterol Assay Kit (Cell based) (Abcam, Cat# ab133116) protocol. Stained cells were mounted in freshly prepared mounting media and imaged using a ZEISS AXIO examiner D1 microscope (Zeiss) equipped with a 40 \times oil Plan-APOCHROMAT objective and a Coolsnap HQ2 camera (Photometrics). Fluorescence detection was performed using a conventional DAPI filter set as recommended by the manufacturer (excitation filter 377/50nm and emission filter 447/60 nm). Images were processed and analyzed by CellProfiler (<https://cellprofiler.org/>)⁹⁴ using a customized pipeline. Briefly, images were subtracted from the background, segmented via the Otsu algorithm, then average intensity was collected for each cell per field of view and used as readout of cholesterol content.

Enzymatic cholesterol analysis

2000–3000 sorted NAD(P)H^{Hi} and NAD(P)H^{Lo} CD34⁻ cells were centrifuged at 400 G for 3 min at 4°C, then washed one time in ice-cold PBS. Subsequently, cholesterol was extracted and detected using the Total Cholesterol Assay Kits (Cell Biolabs, Cat# STA-384) according to manufacturer instructions. Final fluorescence reading was normalized based on the number of cells sorted for each sample.

Cholesterol enrichment

To force elevation of cholesterol in HSC plasma membrane, a cholesterol saturated MBCD (MBCD:chol) was prepared as following: MBCD 5 mM was solved in StemSpan SFEM media then incubated with cholesterol at a molar ratio 10:1 for 24 h at 37°C. Sorted HSCs were incubated with MBCD:chol for 60 min at 37°C then washed twice with StemSpan SFEM.

RT-qPCR

RNA from HSCs sorted from two-month-old mouse bone marrow were isolated using Qiagen RNeasy Kit and subjected to cDNA synthesis using Superscript III First-Strand synthesis system (Invitrogen, Cat# 18080051) or QuantiNova Reverse Transcription Kit (Qiagen, Cat# 205413). Real time quantitative PCR for *Slc25a1*, *Cpt1a*, *Ndufv1*, and *ActB* was performed using TaqMan Assay Mix (Applied Biosystems, Cat # 4304437) in QuantStudio 6 Flex Real-Time PCR with the dedicated Taqman probes (*Slc25a1*: Mm00467666_m1; *Cpt1a*: Mm01231183_m1; *Ndufv1*: Mm00504941_m1; *ActB*: Mm02619580_g1). Real time quantitative PCR for *Hsd17b7*, *Msmo1*, *Tm7sf2*, *Scf* and *ActB* was performed using Quantinova SYBR Green PCR Kit (Qiagen, Cat# 208056) in a Rotor-Gene Q (Qiagen, Cat# 9001862) using the following oligonucleotides.

Gene	Sequences
<i>Scf</i>	F- CCTCTCGCAATGCAAAGAAGG R- GAGGTCGGTAGCATATTTGGAAG
<i>Msmo1</i>	F- AAACAAAAGTGTGGCGTGTTT R- AAGCATTCTTAAGGGCTCCTG
<i>Tm7sf2</i>	F- GTCGCGGCTTACTGATCCT R- CAGGCAGATAGGCCGGTAG
<i>Hsd17b7</i>	F- CCTCTCGCAATGCAAAGAAGG R- GAGGTCGGTAGCATATTTGGAAG
<i>ActB</i>	F- GGCTGTATCCCCTCCATCG R- CCAGTTGGTAACAATGCCATGT

The PCR profile included a 10 min, 95°C hot-start to activate the Taq polymerase, followed by 40 cycles of a two-step program: 15 s at 95°C (denaturation) and 60 s at 60°C (annealing and extension). Relative gene expression level was analyzed by comparative Ct method and was normalized to *ActB*, as mentioned in the figure legends.

EVs isolation

3000 HSC or FKSL or Lin⁻ Nestin⁺ mesenchymal stem cells were sorted and cultured in StemSpan SFEM (STEMCELL Technologies) supplemented with 50 ng/mL stem cell factor (PeproTech, Cat# 250-03) and 50 ng/mL thrombopoietin (PeproTech, Cat# AF-315-14)

48 h. EVs were isolated by a modified PEG based method. EV-enriched media were exposed to 8% PEG 8000 (Merk, Cat# 89510) at 4°C for 24 h, then EVs were centrifuged at 3214 g for 1 h at 4°C. Supernatant was discarded and EV pellet resuspended in 30 μ L of particle-free PBS.¹⁰⁴

Tunable resistive pulse sensing (TRPS)

EVs concentration and size distribution were assessed using a qNano instrument (Izon Science) equipped with an NP150 nanopore (70–420 nm). 35 μ L of each sample were loaded in the upper fluid cell, and voltage, pressure and stretch of the system were adjusted accordingly. TRPS (Tunable resistive pulse sensing) measurements as well as data analysis were performed with the Izon Control Suite Software (V3.4).

Nanoparticle tracking analysis

Nanoparticle tracking analysis (NTA) uses laser light scattering and Brownian motion to determine EVs size and concentration. This allows determining the size of particles in a suspension that flows through a focused laser beam. The light scattered by individual particles is recorded over time to visualize and track their movement. The mean square displacement for each particle along its track allows to calculate individual particle sizes using the Stokes-Einstein equation.¹⁰⁵ Measurement of particle size and particle size distribution was performed with Nanosight NS300 (Malvern) instrument equipped with a 488 nm laser. All samples were diluted in filtered PBS to a final volume of 1 mL. Ideal measurement concentrations were found by pre-testing the ideal particle per frame value (20–100 particles/frame). For each measurement, four to five 1-min videos were captured under temperature 25°C and syringe pump speed 30. Data are represented as averaged finite track length adjustment (FTLA) concentration/size.

EV immunomagnetic isolation and flow cytometry

35 μ L of EVs suspension was incubated with CD63 magnetic beads (Thermo Fisher Scientific, Cat# 10606D). Isolated CD63-positive EVs were then labeled with an anti-CD63 monoclonal antibody (PE mouse anti-human CD63; eBioscience, Cat# 12-0639-42) or anti-CD9 monoclonal antibody (PerCP/Cyanine5.5 anti-human CD9, BioLegend, Cat# 312109). Negative control was performed by staining PBS (vehicle) instead of EVs. Flow cytometric experiments were conducted with Attune NxT Acoustic Focusing Cytometer (Life Technologies), and data were analyzed using FlowJo 10 (Becton Dickinson, Mountain View, CA, USA).

EV transfer

Sorted CD34⁺ HSC were stained with 2 μ M PKH26 (Sigma-Aldrich, Cat# MINI26-1KT) for 2 min at room temperature, then washed three times with StemSpan SFEM. PKH26-stained cells were plated in a 96 well plate and cultured in StemSpan SFEM medium as described above. After 48 h, the media was collected and EVs isolated as described above. Isolated EVs were administered to freshly isolated CD34⁺ HSC every 24 h for 48 h.

Super-resolution radial fluctuation imaging of EVs markers

Isolated EVs were seeded on a coverslip cleaned by acid wash and coated with poly-lysine for 60 min at 37°C to allow adhesion. EVs were then fixed with filtered 2% PFA for 10 min then blocked with filtered 5% BSA for 60 min at room temperature. Anti-CD63 (Abcam, Cat# ab134045) antibody was diluted 1:100 in filtered 5% BSA and incubated overnight at 4°C. After primary antibody hybridization, samples were washed three times with filtered PBS, then incubated with secondary antibodies goat anti-rabbit conjugated with Alexa 488 (Thermo Fisher Scientific, Cat# A32731). A coverslip containing fluorescent beads (TetraSpeck, Thermo Fisher Scientific, Cat# T7279) or PBS only were used as a positive and negative references for EVs detection.

Imaging was performed using a Nikon ECLIPSE Ti inverted widefield microscope equipped with 60 \times objective (CFI Plan Apo Lambda 60x/1.4 oil) and an additional 1.5 \times magnification was used to collect fluorescence onto an EMCCD camera (iXonEM+885, Andor), yielding a pixel size of 88 nm. Excitation was provided by a Lambda LS Stand-Alone 175W Xenon Arc Lamp filtered through a FITC excitation filter (FF01-494/20, Semrock) and a neutral density filter (ND8, Nikon). For each field of view was acquired a 200 frames stack with exposure of 164 ms.

Images were processed through the eSRRF plugin on ImageJ (<https://github.com/HenriquesLab/NanoJ-eSRRF>).⁹³ Before processing the whole dataset, parameter sweep function was used then radius value was set at 4, sensitivity was set to 4. The processed eSRRF images were then analyzed using Olympus scanR. Briefly, spots were identified in all images using the edge detection function. For each spot, area, circularity and CD63 total intensity were collected. The gating strategy was also applied for analysis using Olympus scanR. Representative gating for Figure S7I was obtained using FlowJo 10 (Becton Dickinson).

nFCM analysis

For each staining reaction, 2 \times 10⁸–2 \times 10⁹ of purified EVs were incubated with CD63-APC (a15712, life technologies) primary antibodies for 1 h at 37°C under shaking, protected from light. Unbound antibodies were removed by repeated washing (6–8 times) with 500 μ L PBS prior to analysis. Samples were next acquired using the NanoAnalyzer (nanoFCM Inc.).

Before each experiment, the NanoAnalyzer was aligned using polystyrene QC beads (nanoFCM Inc.). Size and concentration standard nanospheres (nanoFCM Inc.) were read directly after to calibrate the instrument for EV analysis. Once nFCM was aligned and calibrated, EV samples were diluted in filtered PBS (blank) to the optimal range for measurement (10⁸ particles/mL). Samples and blanks (200–800 events) were measured for 1 min, applying a laser power of 15 mW as excitation source, constant pressure of

1 kPa, and at an event rate between 2500 and 12,000 events/min (as recommended by manufacturers). SSC was set as the trigger channel, each particle that generated a signal above the SSC threshold was acquired as an event. For each event that also generated a signal above thresholds set in the fluorescent channels, the fluorescence intensity was registered. Empty staining reactions (without EVs) were performed as a control for all fluorescent reagents and measured under the same conditions as complete reactions (with EVs). Data analysis was performed using FlowJo 10 (Becton Dickinson).

Immunostaining

Sorted cells were resuspended in 30 μ L of StemSpan SFEM supplemented with 50 ng/mL SCF and 50 ng/mL thrombopoietin (PeproTech) and then seeded on Lab-Tek II Chamber Slide (Thermo Fisher Scientific, Cat# 154526) coated with Fibronectin (Sigma-Aldrich, Cat# F1141). Samples were then immunostained as described above. Anti-Farnesyl, rabbit polyclonal antibody (Sigma-Aldrich, Cat# 341286, dilution 1:100) or Anti-Ki-67 (D3B5) rabbit monoclonal antibody were used for detection of protein farnesylation or ki67 respectively. After primary antibody hybridization, samples were washed three times with filtered PBS, then incubated with secondary antibodies goat anti-rabbit conjugated with Alexa Fluor 488 (Thermo Fisher Scientific, Cat# A32731) or anti-rabbit conjugated with Alexa Fluor 594 (Thermo Fisher Scientific, Cat# A11037). Images were acquired using a ZEISS AXIO examiner D1 microscope (Zeiss) equipped with a 40 \times oil Plan-APOCHROMAT objective and a Coolsnap HQ2 camera (Photometrics). Next were processed and analyzed by CellProfiler (<https://cellprofiler.org/>)⁹⁴ using a customized pipeline. Briefly, images were subtracted from the background, segmented via the Otsu algorithm, then average intensity was then collected for each cell per field of view and used as readout of protein farnesylation.

QUANTIFICATION AND STATISTICAL ANALYSIS

For each dataset, the normality assumption was verified with the D'Agostino-Pearson Omnibus Test. Sample sizes and reproducibility for each figure are denoted in the figure legends. For mouse experiments, no statistical method was used to predetermine sample size. Experiments were not randomized, and the investigators were not blinded to allocation during experiments and outcome assessment. All data were collected and analyzed by Prism 7 (GraphPad). Samples that satisfied the normality assumption were evaluated through T-test, one-way or two-way ANOVA, appropriate multiple comparison correction is indicated in the figure legend for each data. When was not satisfied they were tested by Wilcoxon signed rank test or Kruskal-Wallis test. The used test is indicated in the figure legend for each data. All graphs show mean values. Error bars indicate \pm SEM, unless otherwise indicated.