

Fig. S1. Plentiful high-purity Lin<sup>-</sup> Sca-1<sup>+</sup> SCs obtained from mice myocardium by enzymatic digestion and magnetic cell sorting. *A*: Lin<sup>-</sup> Sca-1<sup>-</sup> cells and Lin<sup>-</sup> Sca-1<sup>+</sup> SCs under optical microscope after magnetic cell sorting. *B*, *C*: Typical immunofluorescent stain images (*B*) and graphic representation (*C*) of flow cytometric data via magnetic cell sorting. *D*: The percentage of Lin<sup>-</sup> Sca-1<sup>-</sup> and Lin<sup>-</sup> Sca-1<sup>+</sup> cells in primary cell culture detected by flow cytometry. Mean  $\pm$  SEM,  $n = 8$  per group. \*\*\* $P < 0.001$  vs Lin<sup>-</sup> Sca-1<sup>-</sup> group.

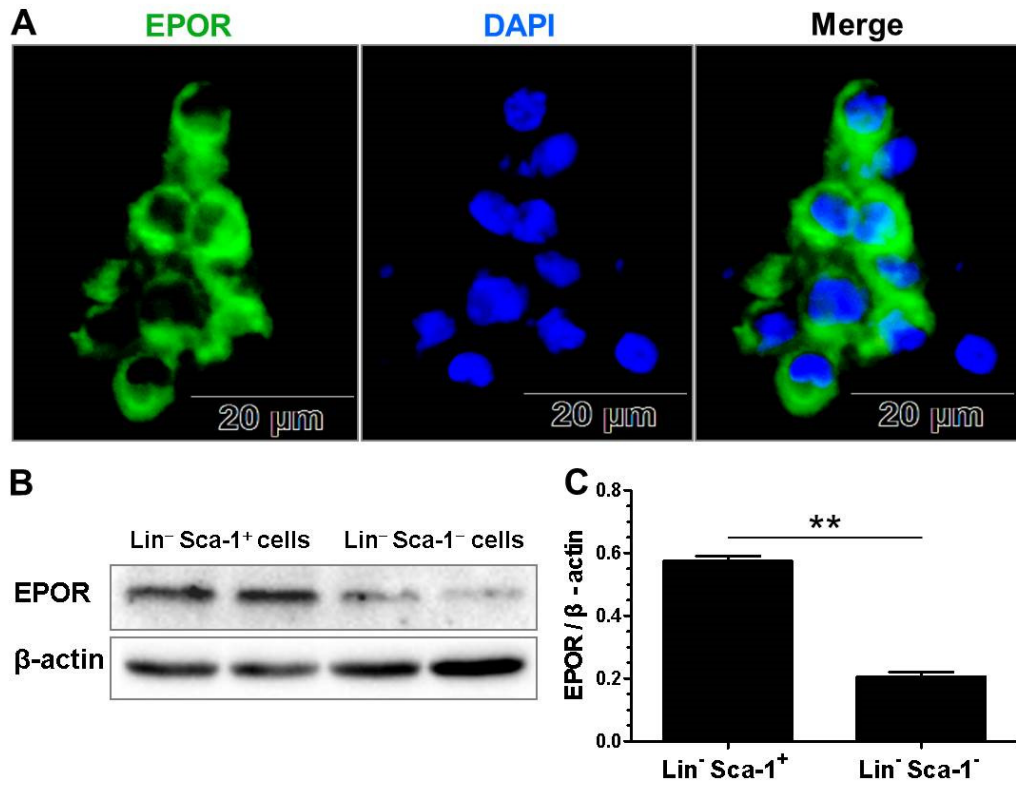


Fig. S2. Expression of EPO receptor (EPOR) in Lin<sup>-</sup> Sca-1<sup>-</sup> cells and Lin<sup>-</sup> Sca-1<sup>+</sup> SCs. *A*: Immunofluorescent staining of EPOR in Lin<sup>-</sup> Sca-1<sup>-</sup> cells and Lin<sup>-</sup> Sca-1<sup>+</sup> SCs. *B*, *C*: Western blot results of EPOR distribution on Lin<sup>-</sup> cells. Mean ± SEM, *n* = 3 per group. \*\**P* < 0.01.