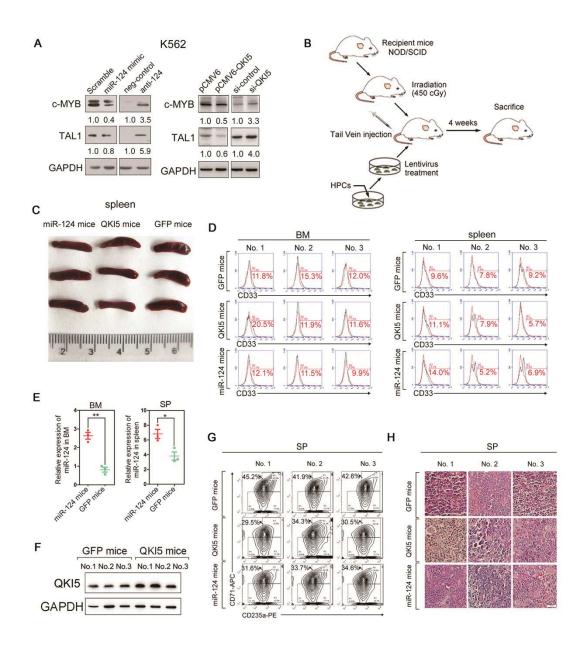
## Supplementary information, Figure S8



**Figure S8** QKI5 and miR-124 are negative regulators for *in vivo* erythroid differentiation. (**A**) Immunoblot of endogenous levels of c-MYB and TAL1 in K562 cells transfected with miR-124 mimic or Scramble control, and miR-124 inhibitors (anti-124) or negative control (neg-control), and pCMV6-QKI5 or pCMV6, and si-QKI5 or si-control. (**B**) A schematic representation of the establishment of the

human HPC-transplanted mouse model. (C) Pictures of mice spleen. (D) Monitoring of CD33<sup>+</sup> populations in BM and SP from mice transplanted with lenti-QKI5 transduced HPCs (QKI5 mice), GFP mice and miR-124 mice. (E) Q-PCR of miR-124 in BM and SP from miR-124 and GFP mice. (F) Immunoblot of QKI5 protein levels in BM of transplanted mice. (G) Monitoring of CD235a/CD71-stained fraction in SP from mice transplanted with lenti-QKI5 transduced HPCs (QKI5 mice), GFP mice and miR-124 mice. (H) Hematoxylin and eosin-stained sections of spleens harvested from mice at 4 weeks post-transplantation. A 400× magnification of a representative field is shown. Scale bar, 20  $\mu$ m.