nature portfolio

Corresponding author(s):	Katharina Schlacher
Last updated by author(s):	Feb 6, 2023

Reporting Summary

Nature Portfolio wishes to improve the reproducibility of the work that we publish. This form provides structure for consistency and transparency in reporting. For further information on Nature Portfolio policies, see our <u>Editorial Policies</u> and the <u>Editorial Policy Checklist</u>.

_				
C-	-	Fic:	tica	•
_	_		111 >	

For	all statistical ar	nalyses, confirm that the following items are present in the figure legend, table legend, main text, or Methods section.		
n/a	Confirmed			
	The exact sample size (n) for each experimental group/condition, given as a discrete number and unit of measurement			
\boxtimes	A stateme	A statement on whether measurements were taken from distinct samples or whether the same sample was measured repeatedly		
	The statistical test(s) used AND whether they are one- or two-sided Only common tests should be described solely by name; describe more complex techniques in the Methods section.			
\boxtimes	A description of all covariates tested			
\boxtimes	A description of any assumptions or corrections, such as tests of normality and adjustment for multiple comparisons			
	A full description of the statistical parameters including central tendency (e.g. means) or other basic estimates (e.g. regression coefficient) AND variation (e.g. standard deviation) or associated estimates of uncertainty (e.g. confidence intervals)			
	For null hypothesis testing, the test statistic (e.g. <i>F</i> , <i>t</i> , <i>r</i>) with confidence intervals, effect sizes, degrees of freedom and <i>P</i> value noted <i>Give P values as exact values whenever suitable.</i>			
\boxtimes	For Bayesian analysis, information on the choice of priors and Markov chain Monte Carlo settings			
\boxtimes	For hierarchical and complex designs, identification of the appropriate level for tests and full reporting of outcomes			
\boxtimes	\square Estimates of effect sizes (e.g. Cohen's d , Pearson's r), indicating how they were calculated			
	•	Our web collection on <u>statistics for biologists</u> contains articles on many of the points above.		
Software and code				
Poli	cy information	about availability of computer code		
Da	ata collection	NA		
Da	ata analysis	described in the manuscript if applicable		
	,	g custom algorithms or software that are central to the research but not yet described in published literature, software must be made available to editors and encourage code deposition in a community repository (e.g. GitHub). See the Nature Portfolio guidelines for submitting code & software for further information.		

Data

Policy information about <u>availability of data</u>

All manuscripts must include a data availability statement. This statement should provide the following information, where applicable:

- Accession codes, unique identifiers, or web links for publicly available datasets
- A description of any restrictions on data availability
- For clinical datasets or third party data, please ensure that the statement adheres to our policy

All data pertaining to the results in the manuscript are available in the main text, the Supplementary materials, and Source Data file. The raw data in Supplementary Table 2 are protected and are not available due to data privacy laws. Material requests underlie compliance with institutional policies and can be made by contacting the corresponding authors.

Life sciences	ne below that i	s the best fit for your research. If you are not sure, read the appropriate sections before making your selection. Sehavioural & social sciences Ecological, evolutionary & environmental sciences all sections, see nature.com/documents/nr-reporting-summary-flat.pdf
Life scier	nces stu	udy design
All studies must dis	sclose on these	points even when the disclosure is negative.
Sample size	described in ma	anuscript, sample size was not predetermined, experiments performed in three biological replicas
Data exclusions	no data exclude	ed
Replication	sample sizes de	escribed in manuscript, minimally three biological replicas where possible
Randomization	described in th	e manuscript when applicable, samples were allocated at random
Blinding	described in th	e manuscript when applicable, samples were analyzed in a blinded fashion when possible
Reporting for specific materials, systems and methods We require information from authors about some types of materials, experimental systems and methods used in many studies. Here, indicate whether each material, system or method listed is relevant to your study. If you are not sure if a list item applies to your research, read the appropriate section before selecting a response. Materials & experimental systems Methods		
Antibodies used	listed	in the manuscript
Validation	validat	ted as shown in mauscript
Eukaryotic cell lines		
Policy information about cell lines		
Cell line source(s)	self (generated from experimental animals)
Authentication		genotyping
Mycoplasma con	tamination	tested
Commonly misidentified lines (See ICLAC register)		

Animals and other organisms

 $Policy\ information\ about\ \underline{studies\ involving\ animals};\ \underline{ARRIVE\ guidelines}\ recommended\ for\ reporting\ animal\ research$

Laboratory animals

genetic mouse model including strain (mixed) and ages described in detail in manuscript

Wild animals	no wild animals were used
Field-collected samples	no field collected samples were used in this study
Ethics oversight	MD Anderson IACUC committee for small animal work

Note that full information on the approval of the study protocol must also be provided in the manuscript.

Flow Cytometry

Plots

Confirm that:

- The axis labels state the marker and fluorochrome used (e.g. CD4-FITC).
- The axis scales are clearly visible. Include numbers along axes only for bottom left plot of group (a 'group' is an analysis of identical markers).
- All plots are contour plots with outliers or pseudocolor plots.
- A numerical value for number of cells or percentage (with statistics) is provided.

Methodology

Sample preparation	samples were prepared using mouse thymus and bone marrow cells
Instrument	LSRFortessa X-20 Analyzer was used
Software	FlowJo 10.4.2 (FlowJo, LLC).
Cell population abundance	no sorting occured
Gating strategy	Gating strategy was performed as described in Mayle, A., Luo, M., Jeong, M. & Goodell, M. A. Flow cytometry analysis of murine hematopoietic stem cells. Cytometry A 83, 27-37, doi:10.1002/cyto.a.22093 (2013).

X Tick this box to confirm that a figure exemplifying the gating strategy is provided in the Supplementary Information.