nature portfolio

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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>.

Sta	atistic						
For	all statist	ical analyses, confirm that the following items are present in the figure legend, table legend, main text, or Methods section.					
n/a	Confirm						
	The	exact sample size (n) for each experimental group/condition, given as a discrete number and unit of measurement					
\boxtimes	A st	atement on whether measurements were taken from distinct samples or whether the same sample was measured repeatedly					
\boxtimes	11 1	statistical test(s) used AND whether they are one- or two-sided common tests should be described solely by name; describe more complex techniques in the Methods section.					
\boxtimes	A d	escription of all covariates tested					
\boxtimes	A d	escription of any assumptions or corrections, such as tests of normality and adjustment for multiple comparisons					
	A fu	Ill description of the statistical parameters including central tendency (e.g. means) or other basic estimates (e.g. regression coefficient) O variation (e.g. standard deviation) or associated estimates of uncertainty (e.g. confidence intervals)					
\boxtimes		null hypothesis testing, the test statistic (e.g. F , t , r) with confidence intervals, effect sizes, degrees of freedom and P value noted P values as exact values whenever suitable.					
\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	Esti	mates of effect sizes (e.g. Cohen's d , Pearson's r), indicating how they were calculated					
	1	Our web collection on <u>statistics for biologists</u> contains articles on many of the points above.					
So	ftware	e and code					
Poli	cy inform	ation about <u>availability of computer code</u>					
Data collection		tion n/a					
Data analysis		sis n/a					
		utilizing 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 crongly 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

Data associated with this study are presented in this paper.

Human resea	arcn par	ticipants		
Policy information a	about <u>studies</u>	s involving human research participants and Sex and Gender in Research.		
Reporting on sex and gender		n/a		
Population characteristics		n/a		
Recruitment		n/a		
Ethics oversight		n/a		
Note that full informat	tion on the ap	proval of the study protocol must also be provided in the manuscript.		
Field-spe	citic r	eporting		
Please select the on	ne below that	t is the best fit for your research. If you are not sure, read the appropriate sections before making your selection.		
Life sciences		Behavioural & social sciences		
For a reference copy of th	he document wi	th all sections, see <u>nature.com/documents/nr-reporting-summary-flat.pdf</u>		
Life scien	ices st	tudy design		
All studies must disc	close on thes	se points even when the disclosure is negative.		
Sample size	Sample size v	was chosen based on our previous experience and other publications.		
Data exclusions	No data exclu	uded.		
Replication	From multiple	e experiemnts.		
Randomization	Experiments	were randomized blindly.		
Blinding	Blinded durin	ng data collection.		
Reporting	σ for s	specific materials, systems and methods		
<u> </u>		rs about some types of materials, experimental systems and methods used in many studies. Here, indicate whether each material,		
*		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 & exp	perimental	systems Methods		
n/a Involved in the	e study	n/a Involved in the study		
Antibodies		ChIP-seq		
Eukaryotic cell lines		Flow cytometry		
Palaeontology and archaeology				
	d other organi	sms		
	Clinical data			
Dual use re	search of cond	cern Common Comm		
Antibodies				
Antibodies used		(Sigma-RAB0287-1KT), IFN-g (R & D System-DY485-05), IL-4, IL-10 (Invitrogen-BMS613, 88-7105-22) assay kit for ELISA, Granzyme igma-RAB0210-1KT) and mouse typer isotyping panel kit (BioRad-1722055)		
		d kit and followed the instruction.		

Eukar	votic (cell	lines
Lakai	y O Ci O (2011	

<u> </u>			_		
Policy information about <u>c</u> e	ell lines	and Sex and Gender in Research			
Cell line source(s)		HEK293T (Clontech)			
Authentication		Certified by supplier.			
Mycoplasma contaminat	ion	We check periodically using using commercial Lonza MycoAlert™ Mycoplasma Detection kit.			
Commonly misidentified lines (See <u>ICLAC</u> register)		This cell line is not listed in the Database of Cross-contaminated or Misidentified Cell Lines.			
Animals and othe	er res	earch organisms			
Policy information about <u>st</u> Research	tudies ir	nvolving animals; ARRIVE guidelines recommended for reporting animal research, and Sex and Gender in			
Laboratory animals	Balb/c	mice (Jackson lab) ~ 3months			
Wild animals	n/a				
Reporting on sex	Female	e-overy as a tissue of interest			
Field-collected samples	n/a				
Ethics oversight	IACUC,	Saint Louis University			
Flow Cytometry					
Plots					
Confirm that:					
The axis labels state t	he mar	ker and fluorochrome used (e.g. CD4-FITC).			
The axis scales are cle	early vis	ible. Include numbers along axes only for bottom left plot of group (a 'group' is an analysis of identical markers).			
All plots are contour	plots wi	th outliers or pseudocolor plots.			
A numerical value for	numbe	er of cells or percentage (with statistics) is provided.			
Methodology					
Sample preparation		Describe the sample preparation, detailing the biological source of the cells and any tissue processing steps used.			
Instrument		Identify the instrument used for data collection, specifying make and model number.			
Software		Describe the software used to collect and analyze the flow cytometry data. For custom code that has been deposited into a community repository, provide accession details.			
Cell population abundan	ce	Describe the abundance of the relevant cell populations within post-sort fractions, providing details on the purity of the samples and how it was determined.			
Gating strategy		Describe the gating strategy used for all relevant experiments, specifying the preliminary FSC/SSC gates of the starting cell population, indicating where boundaries between "positive" and "negative" staining cell populations are defined.			

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