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>.

Please do not complete any field with "not applicable" or n/a. Refer to the help text for what text to use if an item is not relevant to your study. For final submission: please carefully check your responses for accuracy; you will not be able to make changes later.

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For	all statistical an	alyses, confirm that the following items are present in the figure legend, table legend, main text, or Methods section.		
n/a	Confirmed			
	X The exact	sample size (n) for each experimental group/condition, given as a discrete number and unit of measurement		
	X A stateme	ent on whether measurements were taken from distinct samples or whether the same sample was measured repeatedly		
X		tical test(s) used AND whether they are one- or two-sided on tests should be described solely by name; describe more complex techniques in the Methods section.		
X	A descript	ion of all covariates tested		
X	X			
X	X 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)			
X	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>			
X	X For Bayesian analysis, information on the choice of priors and Markov chain Monte Carlo settings			
X	For hierarchical and complex designs, identification of the appropriate level for tests and full reporting of outcomes			
	X Estimates	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 an	d code		
Poli	cy information a	about availability of computer code		
D	ata collection	All code to collect data has been deposited in GitHub and linked in the Methods and Code Availability sections.		
D	ata analysis	All code to analyze data has been deposited in GitHub and linked in the Methods and Code Availability sections.		
	,	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		

Data

Policy information about availability of data

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

The data generated in this study have been deposited in the NCBI Sequence Read Archive; the accession code is in the Data Availability statement.

	ut studies with <u>human participants or human data</u> . See also policy information about <u>sex, gender (identity/presentation),</u> and <u>race, ethnicity and racism</u> .		
Reporting on sex and			
Reporting on race, et other socially relevar groupings	**		
Population character	istics N/A		
Recruitment	N/A		
Ethics oversight	N/A		
Note that full information	on the approval of the study protocol must also be provided in the manuscript.		
-	Behavioural & social sciences		
All studies must disclos	e on these points even when the disclosure is negative.		
Sample size Th	e number of sequencing reads in each sample was chosen to reach coverage of at least 1000x (or 100,000x for clustering).		
Data exclusions Se	Sequencing reads were excluded if they failed to align uniquely or pass the filters of SEISMIC-RNA (see Methods for details).		
Replication All	no-ASO experiments were performed at least twice to confirm reproducibility. ASO experiments were replicated if feasible.		
Randomization No.	randomization was performed.		
Blinding	Researchers were not blinded.		
Behaviour	al & social sciences study design		
All studies must disclos	e on these points even when the disclosure is negative.		
Study description			
	v samula		

Study description	
Research sample	
Sampling strategy	
Data collection	
Timing	
Data exclusions	
Non-participation	
Randomization	

Ecological, ev	olutionary & environmental sciences study design
All studies must disclose on	hese points even when the disclosure is negative.
Study description	
Research sample	
Sampling strategy	
Data collection	
Timing and spatial scale	
Data exclusions	
Reproducibility	
Randomization	
Blinding	
Did the study involve field Field work, collect	
Field conditions	
Location	
Access & import/export	
Disturbance	
Ne require information from au	specific materials, systems and methods thors about some types of materials, experimental systems and methods used in many studies. Here, indicate whether each material and 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 & experimer	·
n/a Involved in the study X Antibodies X Eukaryotic cell lines X Palaeontology and ar X Animals and other or X Clinical data X Dual use research of	ganisms

Antibodies

Antibodies used

Validation

Eukaryotic cell lines			
Policy information about <u>cell lines and Sex and Gender in Research</u>			
Cell line source(s)	ST Cells: ATCC CRL-1746		
Authentication	Not authenticated.		
Mycoplasma contamination	Not tested.		
Commonly misidentified lir (See <u>ICLAC</u> register)	Not in ICLAC register of cross-contaminated cell lines.		
Palaeontology and	Archaeology		
Specimen provenance			
Specimen deposition			
Dating methods			
Tick this box to confirm	that the raw and calibrated dates are available in the paper or in Supplementary Information.		
Ethics oversight			
Note that full information on the	e approval of the study protocol must also be provided in the manuscript.		
Animals and other	research organisms		
Policy information about <u>stud</u> <u>Research</u>	dies involving animals; ARRIVE guidelines recommended for reporting animal research, and <u>Sex and Gender in</u>		
Laboratory animals			
Wild animals			
Reporting on sex			
Field-collected samples			
Ethics oversight			
Note that full information on the	e approval of the study protocol must also be provided in the manuscript.		
Clinical data			
Policy information about <u>clin</u> All manuscripts should comply w	ical studies vith the ICMJE guidelines for publication of clinical research and a completed CONSORT checklist must be included with all submissions.		
Clinical trial registration			
Study protocol			
Data collection			
Outcomes			

Dual use research of concern

Policy information about $\underline{\text{dual use research of concern}}$

Hazards

Could the accidental, deliberate or reckless misuse of agents or technologies generated in the work, or the application of information presented in the manuscript, pose a threat to:

No Yes Public health National security Crops and/or liveste Ecosystems Any other significan			
Experiments of concern	n		
Does the work involve any	y of these experiments of concern:		
No Yes Demonstrate how to render a vaccine ineffective Confer resistance to therapeutically useful antibiotics or antiviral agents Enhance the virulence of a pathogen or render a nonpathogen virulent Increase transmissibility of a pathogen			
Alter the host range			
Enable evasion of d	iagnostic/detection modalities		
	ization of a biological agent or toxin		
Any other potential	lly harmful combination of experiments and agents		
Plants			
Seed stocks			
Novel plant genotypes			
Authentication			
ChIP-seq			
Data deposition			
	and final processed data have been deposited in a public database such as GEO.		
Confirm that you have	deposited or provided access to graph files (e.g. BED files) for the called peaks.		
Data access links May remain private before public	ation.		
Files in database submissi	on (
Genome browser session (e.g. <u>UCSC</u>)			
Methodology			
Replicates			
Sequencing depth			
Antibodies			
Peak calling parameters			
Data quality			

Software

Flow Cytometry	
The axis scales are clearly visil All plots are contour plots wit	er and fluorochrome used (e.g. CD4-FITC). ole. Include numbers along axes only for bottom left plot of group (a 'group' is an analysis of identical markers). h outliers or pseudocolor plots. of cells or percentage (with statistics) is provided.
Methodology	
Sample preparation	
Instrument	
Software	
Cell population abundance	
Gating strategy	
Tick this box to confirm that a	figure exemplifying the gating strategy is provided in the Supplementary Information.
NAti	
Magnetic resonance in	naging
Experimental design	
Design type	
Design specifications	
Behavioral performance measure	
Imaging type(s)	
Field strength	
Sequence & imaging parameters	
Area of acquisition	
Diffusion MRI Used	☐ Not used
Dransaccing	
Preprocessing Preprocessing software	
Normalization	
Normalization template	
Noise and artifact removal	
Volume censoring	
Statistical modeling & infere	nce
Model type and settings	
Effect(s) tested	

ROI-based

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Statistic type for inference	
(See Eklund et al. 2016)	
Correction	
Models & analysis	
n/a Involved in the study	
Functional and/or effective connect	zivity
Graph analysis	
Multivariate modeling or predictive	analysis
Functional and/or effective connectivity	
Graph analysis	

Multivariate modeling and predictive analysis