nature portfolio

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Last updated by author(s):	Nov 16, 2022

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>.

For all statistical analyses, confirm that the following items are present in the figure legend, table legend, main text, or Methods section.

n/a	Confirmed						
	The exact	ct sample size (n) for each experimental group/condition, given as a discrete number and unit of measurement					
\times	A stateme	ent on whether measurements were taken from distinct samples or whether the same sample was measured repeatedly					
	The statis	stical test(s) used AND whether they are one- or two-sided mon tests should be described solely by name; describe more complex techniques in the Methods section.					
\boxtimes	A descript	otion of all covariates tested					
\boxtimes	A descript	otion of any assumptions or corrections, such as tests of normality and adjustment for multiple comparisons					
	A full desc	description of the statistical parameters including central tendency (e.g. means) or other basic estimates (e.g. regression coefficient) ariation (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						
	Estimates of effect sizes (e.g. Cohen's <i>d</i> , Pearson's <i>r</i>), 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							
Policy information about <u>availability of computer code</u>							
Da	ata collection	n/a					
Da	ata analysis n/a						
	For manuscripts 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 reviewers. We strongly 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 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

RNA-sequencing data are available through the Gene Expression Omnibus (GEO) repository under accession codes xxx and xxx.

Raw, uncropped images of western blots, southern blots of terminal restriction fragments, and RNA dot-blots are provided in Supplementary Figures. Source data for Figures 1-4 and Extended Data Figures 1-12 are provided.

Human rese	arch part	cicipants				
Policy information	about <u>studies</u>	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 informa	ation on the app	proval of the study protocol must also be provided in the manuscript.				
Field-spe	ecific re	eporting				
Please select the or	ne below that	is the best fit for your research. If you are not sure, read the appropriate sections before making your selection.				
X Life sciences		Behavioural & social sciences				
For a reference copy of t	he document wit	h all sections, see <u>nature.com/documents/nr-reporting-summary-flat.pdf</u>				
Life scier	ices st	udy design				
All studies must dis	sclose on thes	e points even when the disclosure is negative.				
Sample size	Sample size w	vas determined by significance.				
Data exclusions	No data were excluded.					
Replication	Data were rep	produced and number of repeats are indicated. Repetitions confirmed the initial results.				
Randomization	Randomizatio	n was not possible, since the experiments were completed by the same individuals.				
Blinding	Blinding was not possible, since the experiments were completed by the same individuals.					
We require informati	on from author	pecific materials, systems and methods s about some types of materials, experimental systems and methods used in many studies. Here, indicate whether each material, o 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	Materials & experimental systems Methods					
·		n/a Involved in the study ChIP-seq				
Antibodies Eukaryotic cell lines		Flow cytometry				
	Clinical data Dual use research of concern					
Dual use re	.scarcii Oi COIIC					
Antibodies						
Antibodies used	s used Antibodies are described in the methods section.					

Antibodies were validated by band size, and by si/sh suppression of targets.

Validation

Eukaryotic cell lines

Policy information about <u>cell lines and Sex and Gender in Research</u>

Cell line source(s)

Cell lines were purchased from Coriell cell repository.

Commercial c ell lines were authenticated by the provider (Coriell)

Mycoplasma contamination Our cells are tested for mycoplasma monthly. All cells in use were found negative consistently.

Commonly misidentified lines (See ICLAC register)