nature research

Corresponding author(s):	Elizabeth Hinde
Last updated by author(s):	Oct 1, 2020

Reporting Summary

Nature Research 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 Research policies, see our Editorial Policies and the Editorial Policy Checklist.

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

_			
C-	 Fic:	tico	•
_	 	111	

n/a	Confirmed
	$oxed{\boxtimes}$ The exact sample size (n) for each experimental group/condition, given as a discrete number and unit of measurement
	🔀 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.
\times	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	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.
So	ftware and code
Polic	cy information about <u>availability of computer code</u>
Da	ata collection Provide a description of all commercial, open source and custom code used to collect the data in this study, specifying the version used OR

state that no software was used.

Data analysis

Provide a description of all commercial, open source and custom code used to analyse the data in this study, specifying the version used OR state that no software was used.

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 Research 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 list of figures that have associated raw data
- A description of any restrictions on data availability

The datasets generated during and/or analysed during the current study are available from the corresponding author on reasonable request.

Field-spe	cific re	porting	
Please select the or	ne below that is	s the best fit for your research. If you are not sure, read the appropriate sections before making your selection.	
\times Life sciences	В	ehavioural & social sciences	
For a reference copy of t	he document with	all sections, see <u>nature.com/documents/nr-reporting-summary-flat.pdf</u>	
Life scier	ices stu	udy design	
		points even when the disclosure is negative.	
Sample size		wo region's of interest within N > 8 cells over a minimum of two biological experiments	
Data exclusions	We excluded m	icroscopy data where significant photobleaching or axial drift was observed during the acquisition	
Replication	We measured two region's of interest within N > 8 cells over a minimum of two biological experiments and confirmed results with relevant controls		
Randomization	NA		
Blinding	NA		
We require informatic system or method list Materials & exp n/a Involved in th	cell lines ogy and archaeol d other organism earch participant	n/a Involved in the study ChIP-seq Flow cytometry MRI-based neuroimaging State of the study MRI-based neuroimaging State of the study MRI-based neuroimaging	
Antibodies			
Antibodies used	γΗ2Α.>	γH2A.X (S139) (Cat. 9718S, Cell Signalling) and 53BP1 (Cat. 4937S, Cell Signalling)	
Validation	Was used to quantify live cell imaging data and the antibodies selected were highly cited.		
Eukaryotic c	ell lines		
Policy information			
Cell line source(s)			
Authentication DIvA cells (originally provided by Gaëlle Legube, LBCMCP, CNRS, Toulouse, France		DIvA cells (originally provided by Gaëlle Legube, LBCMCP, CNRS, Toulouse, France	

Confirm that the DIvA cell line tested negative for mycoplasma contamination

Mycoplasma contamination

Commonly misidentified lines (See <u>ICLAC</u> register)

NA