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 Editorial Policies and the Editorial Policy Checklist.

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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			
	X A stateme	ent on whether measurements were taken from distinct samples or whether the same sample was measured repeatedly			
×		tical test(s) used AND whether they are one- or two-sided non tests should be described solely by name; describe more complex techniques in the Methods section.			
X	A description of all covariates tested				
X	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 coefficien 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>				
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 <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 an	d code			
Poli	cy information	about <u>availability of computer code</u>			
D	ata collection	SoftWorx v7.0.0 (GE) and LSM880 Zen v2.3 (Zeiss) were used to collect the data.			
D	ata analysis	SoftWorx v7.0.0 (GE), LSM880 Zen v2.3 (Zeiss) , and Microsoft Excel were used to amalyze the data.			
		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 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 supporting the findings of this study are available within the paper and its Supplementary information files.

Life sciences study design

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

N/A

LITE SCIET	1003 300	ady design			
All studies must dis	sclose on these	points even when the disclosure is negative.			
Sample size	Following timing of gene expression from transfected DNA, 594, 125, and 129 cells were monitored in the living state in Fig. 1d, Fig. 5b, and Fig. 5d, respectively.				
Data exclusions No data were excluded from analyses.					
Replication	CLEM (Fig. 3) and iCLEM (Fig. 4) analyses were repeated several times, and the data were obtained from at least three cells.				
Randomization	Randomization N/A				
Blinding N/A					
We require informatis system or method liss Materials & ex n/a Involved in the state of the system	ion from authors a sted is relevant to sperimental so he study	n/a Involved in the study ChIP-seq Flow cytometry MRI-based neuroimaging			
Clinical dat					
✗ ☐ Dual use re	esearch of concer	n			
Antibodies					
Antibodies used	All anti	ibodies used are described in Materials and Methods.			
Validation	All anti	ibodies used are described in Materials and Methods.			
<u>Eukaryotic c</u>	:ell lines				
Policy information	about <u>cell lines</u>				
Cell line source(s)		HeLa and U2OS cell lines were obtained from RIKEN cell bank and the Health Protection Agency, respectively.			
Authentication		Authenticated by RIKEN cell bank and the Health Protection Agency,			
Mycoplasma contamination		All cell lines were tested negative for mycoplasma contamination.			