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>.

Statistics

For	all st	atistical analyses, confirm that the following items are present in the figure legend, table legend, main text, or Methods section.
n/a	Cor	ifirmed
	×	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.
X		A description of all covariates tested
	×	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. F, t, r) with confidence intervals, effect sizes, degrees of freedom and P value noted Give P values as exact values whenever suitable.
X		For Bayesian analysis, information on the choice of priors and Markov chain Monte Carlo settings
×		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 d, Pearson's r), indicating how they were calculated
		Our web collection on statistics for biologists contains articles on many of the points above.

Software and code

Policy information about availability of computer code		
Data collection	Zen 2 pro, NMRPipe software	
Data analysis	Smoldyn-2.71, Fiji, MicrobeJ, POKY software suite	

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 <u>data availability statement</u>. 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

All data supporting the findings of this study are available within the paper and its Supplementary Information.

Research involving human participants, their data, or biological material

Policy information about studies with <u>human participants or human data</u>. See also policy information about <u>sex, gender (identity/presentation)</u>, <u>and sexual orientation</u> and <u>race, ethnicity and racism</u>.

Reporting on sex and gender	Not applicable
Reporting on race, ethnicity, or other socially relevant groupings	Not applicable
Population characteristics	Not applicable
Recruitment	Not applicable
Ethics oversight	Not applicable

Note that full information on the approval of the study protocol must also be provided in the manuscript.

Field-specific reporting

Please select the one below that is the best fit for your research. If you are not sure, read the appropriate sections before making your selection.

Behavioural & social sciences 📃 Ecological, evolutionary & environmental sciences

For a reference copy of the document with all sections, see <u>nature.com/documents/nr-reporting-summary-flat.pdf</u>

Life sciences study design

All studies must disclose on these points even when the disclosure is negative.

Sample size	Fig. 1b, n>300/time point for the bar chart.
	Fig. 1c, Samples from experiment shown in 1b.
	Fig. 1e, Representative images.
	Fig. 1f, Samples from experiment shown in 1e,f.
	Fig. 1g, Representative image.
	Fig. 1h, n>300/types of cell.
	Fig. 2b, Representative images; n=60 per cell types for the graphs.
	Fig. 2c, Samples from experiment shown in 2b.
	Fig. 2d, Representative images; n=10 per cell types for the graphs.
	Fig. 3a, Not applicable
	Fig. 3b, Representative images.
	Fig. 3c, Samples from experiment shown in 2d.
	Fig. 3d, n>20 per types of condensates.
	Fig. 4c, Representative images.
	Fig. 4d, n>300/types of cell.
	Fig. 5a, Representative images.
	Fig. 5b, Representative images; n=20 per cell types for the graphs.
	Fig. 5c, Representative gel images; n=3 for the graphs.
	Fig. 6a-c, Not applicable.
	Supplementary Fig. 1a, n=2
	Supplementary Fig. 1b, n=3
	Supplementary Fig. 1c, n=3
	Supplementary Fig. 2a, n=3
	Supplementary Fig. 2b, Representative images; n=60 per cell types for the graphs.
	Supplementary Fig. 2c, Representative images.
	Supplementary Fig. 2d, Representative images; n=10 per cell types for the graphs.
	Supplementary Fig. 3a-b, Not applicable.
	Supplementary Fig. 4 a-c, Representative images.
	Supplementary Fig. 5a, Representative images, n=20 per cell types for the graph.
	Supplementary Fig. 5b, Representative images, n=20 per cell types for the graph.
	Supplementary Fig. 6a, Representative gel images, n=3 for the graphs.
	Supplementary Fig. 7a, n=3.
	Supplementary Fig. 8a-c, Not applicable.
Data exclusions	4 c, Last two time points were not shown in the graphs as the signals were no higher than the background.
Replication	Fig. 1b, 5-10 images from a single representative experiment which was repeated two times and produced similar results;
	Fig. 1c, Experiment was repeated two times and produced similar results.

Blinding	Not applicable
Randomization	Not applicable
	with same parameters.
	Supplementary Fig. 8a-b, Representative data from individual runs. In this model, we observed no significant variation between runs initiated
	Supplementary Fig. 5a-b, Experiment was repeated three times and produced similar results.
	Supplementary Fig. 4a-c, Experiment was repeated three times and produced similar results.
	Supplementary Fig. 2d, Experiment was repeated three times and produced similar results.
	Supplementary Fig. 2a-b, Experiment was repeated three times and produced similar results.
	Supplementary Fig. 1a-c, Experiment was repeated three times and produced similar results.
	parameters.
	Fig. 6a-c, Representative data from individual runs. In this model, we observed no significant variation between runs initiated with same
	Fig. 5c, Experiment was repeated three times and produced similar results.
	Fig. 5b, Experiment was repeated three times and produced similar results.
	Fig. 5a, Experiment was repeated three times and produced similar results.
	Fig. 4c-d, Experiment was repeated three times and produced similar results.
	Fig. 3c, Experiment was repeated three times and produced similar results.
	Fig. 3b, Experiment was repeated three times and produced similar results.
	Fig. 2c, Experiment was repeated three times and produced similar results.
	Fig. 2b, Experiment was repeated three times and produced similar results.
	Fig. 1h, Experiment was repeated three times and produced similar results.
	Fig. 1g, Experiment was repeated three times and produced similar results.
	Fig. 1f, Experiment was repeated three times and produced similar results.
	Fig. 1e, Experiment was repeated three times and produced similar results.

Reporting for specific materials, systems and methods

We require information from authors about some types of materials, experimental systems and methods used in many studies. Here, indicate whether each material, system or method listed is relevant to your study. If you are not sure if a list item applies to your research, read the appropriate section before selecting a response.

MRI-based neuroimaging

Materials & experimental systems

M	et	ho	ds
	CC		00

n/a	Involved in the study	n/a	Involved in the study
	X Antibodies	×	ChIP-seq
×	Eukaryotic cell lines	×	Flow cytometry
×	Palaeontology and archaeology	×	MRI-based neuroim
×	Animals and other organisms		
×	Clinical data		
×	Dual use research of concern		
×	Plants		

Antibodies

	HA Epitope Tag, Invitrogen (2-2.2.140), Ref 26183, lot YB362804; GFP antibody, Gene tex GTX113617, Lot 42179. HRP conjugated secondary antibody Jackson ImmunoResearch.
Validation	Not applicable

Plants

Seed stocks	Not applicable
Novel plant genotypes	Not applicable
Authentication	Not applicable