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.

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

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Coi	nfirmed
X	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
X	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.
×	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. <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>
	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 <u>statistics for biologists</u> contains articles on many of the points above.
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Software and code

Policy information about availability of computer code

Data collection Yokogawa CV8000 acquisition software

Data analysis CellProfiler v.4; Orange v.3; Python, PipelinePilot (Biovia) v.18, Cellpose v2.2, GraphPad Prism v.5

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

code and publicly available data is available via links in the manuscript. Raw data is available with the manuscript. Image data generated are available upon request

Research involving	ng human partici	pants, their data, or biological material	
Policy information about s and sexual orientation and		ipants or human data. See also policy information about sex, gender (identity/presentation), m.	
Reporting on sex and gend	and gender n/a		
Reporting on race, ethnicit other socially relevant grou			
Population characteristics	n/a		
Recruitment	n/a		
Ethics oversight	n/a		
Note that full information on	the approval of the study pro	otocol must also be provided in the manuscript.	
Field-specifi	c reporting		
Please select the one belo	w that is the best fit for yo	our research. If you are not sure, read the appropriate sections before making your selection.	
X Life sciences	Behavioural & socia	al sciences	
For a reference copy of the document	nent with all sections, see <u>nature</u>	.com/documents/nr-reporting-summary-flat.pdf	
Life science:	study desi	gn	
All studies must disclose o	n these points even when	the disclosure is negative.	
Sample size a mini	mum of 1000 cells/condition	was analyzed	
Data exclusions no dat	no data were excluded		
Replication we use	we used our published method for QC reproducibility of single cell data sets (Stossi et al., EHP 2022; PMID: 35167326)		
Randomization not re	not relevant		
Blinding not re	not relevant		
Reporting for	or specific m	naterials, systems and methods	
•	/ !	f materials, experimental systems and methods used in many studies. Here, indicate whether each material, re not sure if a list item applies to your research, read the appropriate section before selecting a response.	
Materials & experim	ental systems	Methods	
n/a Involved in the study	,	n/a Involved in the study	

IVIC	iteriais & experimental systems	IVIE	uious
n/a	Involved in the study	n/a	Involved in the study
×	Antibodies	×	ChIP-seq
	x Eukaryotic cell lines	×	Flow cytometry
×	Palaeontology and archaeology	×	MRI-based neuroimaging
×	Animals and other organisms		
×	Clinical data		
×	Dual use research of concern		
×	Plants		

Eukaryotic cell lines

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

Cell line source(s) all cell lines are commercially available and were obtained from ATCC or from BCM cell culture core

none of the cell lines was authenticated on-site but all were obtained from ATCC or BCM cell culture core (which performs in Authentication house validation)

Il cell lines were routinely tested for mycoplasma contamination using high magnification microscopy after DAPI staining. All Mycoplasma contamination

cell lines were negative for the duration of the experiments

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

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11	1/a

Plants

Seed stocks	n/a
Novel plant genotypes	n/a
Authentication	n/a