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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 <u>Editorial Policies</u> and the <u>Editorial Policy Checklist</u>.

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For	all statistical an	alyses, confirm that the following items are present in the figure legend, table legend, main text, or Methods section.		
n/a	a Confirmed			
	The exact	sample size (n) for each experimental group/condition, given as a discrete number and unit of measurement		
	A stateme	nt 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.			
\boxtimes	A description of all covariates tested			
\boxtimes	A description of any assumptions or corrections, such as tests of normality and adjustment for multiple comparisons			
\boxtimes	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>			
\times	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	\square 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.			
Software and code				
Poli	cy information a	about <u>availability of computer code</u>		
Da	ata collection	Live-cell imaging was done using Micro-Manager.		
Da	ata analysis	For immunofluorescence and basic image analysis ImageJ 1.52p was used. Image segmentation was done using Ilastik. Boundary tracing and		

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:

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.

- Accession codes, unique identifiers, or web links for publicly available datasets

optical flow analysis was done using either Python or Matlab R2019.

- 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 reporting	
Please select the one	e below that is the best fit for your research. If you are not sure, read the appropriate sections before making your selection.	
Life sciences	e sciences Behavioural & social sciences Ecological, evolutionary & environmental sciences	
or a reference copy of the	e document with all sections, see <u>nature.com/documents/nr-reporting-summary-flat.pdf</u>	
_ife scien	ces study design	
All studies must disc	lose on these points even when the disclosure is negative.	
	The number of biological and technical replicates were chosen according to previously published experimental designs (Linnemann et al, 2015) and are specified for each experiment in the figure legends.	
Data exclusions	No data was excluded.	
Replication	All experiments were repeated for at least ten different organoids.	
	Donor samples were picked according to age and parity in order to provide a broad biological variation, as seen in the Supplementary Table. Organoids were chosen according to developmental stage, size and complexity to enhance comparability.	
Blinding	ng Blinding was not necessary as data interpretation is based on quantitative measurements with no data being excluded.	
Reporting	g for specific materials, systems and methods	
	n from authors about some types of materials, experimental systems and methods used in many studies. Here, indicate whether each material dis 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.	

Ma	terials & experimental systems	Methods	
n/a	Involved in the study	n/a Involved in the study	
\boxtimes	Antibodies	ChIP-seq	
\boxtimes	Eukaryotic cell lines	Flow cytometry	
\boxtimes	Palaeontology and archaeology	MRI-based neuroimaging	
\boxtimes	Animals and other organisms		
\boxtimes	Human research participants		
\boxtimes	Clinical data		
\boxtimes	Dual use research of concern		