# nature portfolio

Corresponding author(s):	Peng Chen
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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	alyses, confirm that the following items are present in the figure legend, table legend, main text, or Methods section.
n/a	Confirmed	
	<b>x</b> 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
X		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 descript	tion of all covariates tested
×	A descript	tion of any assumptions or corrections, such as tests of normality and adjustment for multiple comparisons
	A full deso	cription of the statistical parameters including central tendency (e.g. means) or other basic estimates (e.g. regression coefficient) ation (e.g. standard deviation) or associated estimates of uncertainty (e.g. confidence intervals)
x		ypothesis testing, the test statistic (e.g. $F$ , $t$ , $r$ ) with confidence intervals, effect sizes, degrees of freedom and $P$ value noted ses as exact values whenever suitable.
×	For Bayes	ian analysis, information on the choice of priors and Markov chain Monte Carlo settings
×	For hierar	chical 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.
So <sup>.</sup>	ftware an	d code
Poli	cy information	about <u>availability of computer code</u>
Da	ata collection	Andor iO2 is used to collect data for live cell and in vitro imaging.

# 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.

The code is compatible with MATLAB R2019b Update 1 (9.7.0.1216025). A README file with detailed instructions is also provided in

Custom MATLAB codes are used for data analysis. MATLAB codes are included in the Supplementary Software 1.

#### Data

Data analysis

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

Supplementary Software 1.

 $\hbox{-} For clinical datasets or third party data, please ensure that the statement adheres to our \underline{policy}$ 

All data are available in the main text, Supplementary Information, and Source Data. Raw data supporting the findings of this study are available upon request due to the large volume.

## Research involving human participants, their data, or biological material

Policy information about st and sexual orientation and	rudies with <u>human participants or human data</u> . See also policy information about <u>sex, gender (identity/presentation),</u> <u>race, ethnicity and racism.</u>				
Reporting on sex and gende	r Not applicable.				
Reporting on race, ethnicity other socially relevant group					
Population characteristics	Not applicable.				
Recruitment	Not applicable.				
Ethics oversight	Not applicable.				
Note that full information on t	the approval of the study protocol must also be provided in the manuscript.				
E: 11 :C:					
Field-specific	c reporting				
Please select the one belov	w that is the best fit for your research. If you are not sure, read the appropriate sections before making your selection.				
<b>x</b> Life sciences	Behavioural & social sciences Ecological, evolutionary & environmental sciences				
For a reference copy of the docum	nent with all sections, see <a href="mailto:nature.com/documents/nr-reporting-summary-flat.pdf">nature.com/documents/nr-reporting-summary-flat.pdf</a>				
Life sciences	s study design				
All studies must disclose or	n these points even when the disclosure is negative.				
	Bootstrap analysis was done to determine statistical reliability of data. The data is provided in Supplementary Notes 3.3 (supplementary fig. 10). Sample sizes are either included in the figure or the figure caption.				
Data exclusions No data	a were excluded.				
Replication All atte	All attempts at replication were success.				
	E. coli cells were grouped together into 6-8 concentration groups and the divisions were chosen to ensure that each concentration groups in general have thousands of single-molecule tracking trajectories each. Detailed description is available in supplementary fig. 6.				
Blinding Blinding	Blinding is not relevant to our study.				
We require information from	er specific materials, systems and methods authors about some types of materials, experimental systems and methods used in many studies. Here, indicate whether each material, evant to your study. If you are not sure if a list item applies to your research, read the appropriate section before selecting a response.				
Materials & experime	ental systems Methods				
n/a Involved in the study	·				
Antibodies	ChIP-seq				
<b>x</b> Eukaryotic cell lines					
Palaeontology and					
X Animals and other of	organisms				
Clinical data  Dual use research of	of concern				
Plants					
1					
Antibodies					
Antibodies used	1) Rabbit-derived antiGFP primary antibody (1:10,000 dilution, Rockland Immunochemical, catalog number: 600-401-215) 2) Goat-derived Horseradish Peroxidase-conjugated Fab fragment anti-rabbit antibody (1:20,000 dilution, Rockland Immunochemical,				

catalog number:

611-103-122)

More details are described in Supplementary Note 2.1.

Validation

Validation statement on manufacturer's website: antibody has been tested by western blot and ELISA.

- 1) Rabbit-derived antiGFP primary antibody: https://www.rockland.com/categories/primary-antibodies/gfp-antibody-600-401-215/
- 2) Goat-derived Horseradish Peroxidase-conjugated Fab fragment anti-rabbit antibody: https://www.rockland.com/categories/secondary-antibodies/rabbit-igg-hl-secondary-antibody-peroxidase-conjugated-pre-adsorbed-611-103-122/? srsltid=AfmBOoqIR9rvM8IJZxeO0Jkt\_dxZSnKp81OLB1iuS38xC9YrWOY6\_6KQ

#### **Plants**

Seed stocks

Report on the source of all seed stocks or other plant material used. If applicable, state the seed stock centre and catalogue number. If plant specimens were collected from the field, describe the collection location, date and sampling procedures.

Novel plant genotypes

Describe the methods by which all novel plant genotypes were produced. This includes those generated by transgenic approaches, gene editing, chemical/radiation-based mutagenesis and hybridization. For transgenic lines, describe the transformation method, the number of independent lines analyzed and the generation upon which experiments were performed. For gene-edited lines, describe the editor used, the endogenous sequence targeted for editing, the targeting guide RNA sequence (if applicable) and how the editor was applied.

Authentication

was applied.
Describe any authentication procedures for each seed stock used or novel genotype generated. Describe any experiments used to assess the effect of a mutation and, where applicable, how potential secondary effects (e.g. second site T-DNA insertions, mosiacism, off-target gene editing) were examined.