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 st | atistical analyses, confirm that the following items are present in the figure legend, table legend, main text, or Methods section. |
|-----|--------|--|
| n/a | Cor | 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 |
| × | | 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 |
| 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 coefficient) AND variation (e.g. standard deviation) or associated estimates of uncertainty (e.g. confidence intervals) |
| x | | 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 statistics for biologists contains articles on many of the points above

Software and code

Policy information about availability of computer code

Data collection

Biolayer Interferometry (BLI) experiments: Raw data was preprocessed using the Octet Data Analysis Software version 11.1 (FortéBio Inc. San Jose, CA).

single molecule FRET (smFRET) experiments: Data were collected using Nikon NIS-Elements AR 4.4 (Nikon Instruments, Japan)

Data analysis

BLI: Analyses were done using Octet Data Analysis Software version 11.1 (FortéBio Inc. San Jose, CA) or custom software. Custom software for BLI data analysis is provided as Source Data.

smFRET data: Custom software in MATLAB (MATLAB 9.8) for trace extraction described in (Fei, J., Kosuri, P., MacDougall, D.D. & Gonzalez, R.L. Coupling of ribosomal L1 stalk and tRNA dynamics during translation elongation. Molecular Cell 30, 348-359 (2008)) and tMAVEN for data analyses (Verma, A.R., Ray, K.K., Bodick, M., Kinz-Thompson, C.D. & Gonzalez, R.L., Jr. Increasing the accuracy of single-molecule data analysis using tMAVEN. Biophys J (2024).)

Custom software for smFRET data analysis is provided as Source Data.

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 <u>guidelines for submitting code & software</u> 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 from the corresponding authors upon reasonable request. Source data for the figures and supplementary figures are provided as a Source Data file. Source data are provided in this paper.

Research involving human participants, their data, or biological material

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

| Reporting on sex and gender | N/A |
|--|-----|
| Reporting on race, ethnicity, or other socially relevant groupings | N/A |
| Population characteristics | N/A |
| Recruitment | N/A |
| Ethics oversight | N/A |

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

Field-specific reporting

x Life sciences

| Please select the one below tl | hat is the best fit for you | ir research. If you are not s | ure, read the appropriate s | sections before making your s | selection. |
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Ecological, evolutionary & environmental sciences

| Life sciences | Behavioural & social sciences | Ecological, evolutionar |
|-----------------------------------|---|-------------------------------|
| For a reference copy of the docum | nent with all sections, see nature.com/documents/ | nr-reporting-summary-flat.pdf |

Life sciences study design

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

| Sample size | Sample size is described in the manuscript. |
|-----------------|---|
| Data exclusions | Data were excluded as part of the analyses process. Data from old or degraded samples were excluded. During data collection, some data were excluded if they show inconsistencies, high noise, etc. due to technical issues either with the microscope or the sample. |
| Replication | The biological samples (i.e. the in vitro transcribed tRNAs and T-boxes) were generated at least twice, each considered as a biological replicate. For each biological replicate, at least three technical replicates were done. |
| Randomization | N/A |
| Blinding | N/A |

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.

| Materials & experimental systems | Methods |
|---------------------------------------|---------------------------|
| n/a Involved in the study | n/a Involved in the study |
| X Antibodies | X ChIP-seq |
| ▼ Eukaryotic cell lines | Flow cytometry |
| Palaeontology and archaeology | MRI-based neuroimaging |
| Animals and other organisms | · |
| X Clinical data | |
| ▼ Dual use research of concern | |
| x Plants | |
| · | |

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. 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 analized.

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.