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Reporting Summary

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Statistics			
For all statistical analys	es, confirm that the following items are present in the figure legend, table legend, main text, or Methods section.		
n/a Confirmed	Confirmed		
☐ ☐ The exact sam	ple size (n) for each experimental group/condition, given as a discrete number and unit of measurement		
A statement of	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.		
A description	A description of all covariates tested		
A description	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 Give <i>P</i> values as exact values whenever suitable.			
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			
\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			
Policy information abou	ut <u>availability of computer code</u>		
Data collection	We used the program GetData Graph Digitizer to extract the data from the Figures by Metzl-Raz (2017); see Methods.		
Data analysis	The saturation functions to describe the growth rate dependence of the active ribosome fractions in E. coli and yeast were fitted using the function nls() in gnu R; see Methods.		
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/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.			
Data			
Policy information about <u>availability of data</u> 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 list of figures that have associated raw data - A description of any restrictions on data availability			
The datasets used for Fig. 2 and Supplementary Figures 3 and 4 are available from the original sources (Refs.46–48, 50); see Data Availability statement.			
Field-specific reporting			
Please select the one b	elow that is the best fit for your research. If you are not sure, read the appropriate sections before making your selection.		
V Life sciences	Rehavioural & social sciences		

Life sciences study design

All studies must disclose on these points even when the disclosure is negative.				
Sample size	All experimental data was obtained from previous publications; thus, we had no control over sample sizes.			
Data exclusions	No data was excluded.			
Replication	All experimental data was obtained from previous publications. The smooth dependencies of the ribosomal protein fractions on growth rate support the reproducibility of this data. The error bars for the growth rate and dry mass density in Supplementary Figure 3 reflect the reproducibility reported by Cayley et al. (2000), Ref. 51.			
Randomization	There were no experimental groups in this study. Thus, randomization was not relevant.			
Blinding	There were no experimental groups in this study. Thus, blinding was not relevant.			

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	
\boxtimes	Antibodies	ChIP-seq	
\boxtimes	Eukaryotic cell lines	Flow cytometry	
\boxtimes	Palaeontology	MRI-based neuroimaging	
\boxtimes	Animals and other organisms		
\boxtimes	Human research participants		
\boxtimes	Clinical data		