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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 <u>Authors & Referees</u> and the <u>Editorial Policy Checklist</u>.

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For	all statistical analy	rses, confirm that the following items are present in the figure legend, table legend, main text, or Methods section.				
n/a	Confirmed					
	The exact sa	\sum 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				
\boxtimes	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					
	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)					
\boxtimes	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.					
\boxtimes	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	Estimates of	effect sizes (e.g. Cohen's d, Pearson's r), indicating how they were calculated				
	ı	Our web collection on <u>statistics for biologists</u> contains articles on many of the points above.				
So	ftware and	code				
Poli	cy information abo	out <u>availability of computer code</u>				
D	ata collection	The enzymatic activities and the protein concentrations were collected using the software SoftMax Pro 7.03. The signals of the Northern blots were quantified using the software ImageJ, version 152K.				
D	ata analysis	Data analysis was performed with Microsoft Excel 2013.				
		stom algorithms or software that are central to the research but not yet described in published literature, software must be made available to editors/reviewers.				

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 list of figures that have associated raw data
- A description of any restrictions on data availability

The authors declare that all data supporting the findings of this study are available within the paper and its supplementary files. The average raw data to the quantification of specific enzyme activities and transcript levels are shown in Supplementary Figures S3 – S6. The results fo the single measurements are summarized in the Source Data file. The results of the bioinformatics genome analyses are shown in Supplementary DataTable S1.

Field-specific reporting						
Please select the or	ne below that is the best fit for you	ur research. If you are not sure, read the appropriate sections before making your selection.				
\times Life sciences	Behavioural & social sciences Ecological, evolutionary & environmental sciences					
For a reference copy of t	For a reference copy of the document with all sections, see <u>nature.com/documents/nr-reporting-summary-flat.pdf</u>					
Life sciences study design						
All studies must dis	close on these points even when t	he disclosure is negative.				
Sample size	For all experimental analyses three biological replicates (independent cultures) were used. The specific enzyme activities (nkat/mg) of the reporter enzymes as well as the relative transcript levels were quantified, and average values and their standard deviations were calculated (see Supplementary Figures). The translational efficiencies were calculated as the quotients of specific activities and transcript levels (see Figures 4 and 5).					
Data exclusions	There have been no data exclusions.					
Replication	Three biological replicates were used for all experiments.					
Randomization	There has been no randomization.					
Blinding	There has been no binding.					
Reporting for specific materials, systems and methods						
	The state of the s	materials, experimental systems and methods used in many studies. Here, indicate whether each material, 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				
Antibodies		ChIP-seq				
Eukaryotic cell lines		Flow cytometry				
Palaeontology		MRI-based neuroimaging				

Human resea

Animals and other organisms

Human research participants