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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
	\boxtimes	The exact sample size (n) for each experimental group/condition, given as a discrete number and unit of measurement
	\boxtimes	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
X		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)
\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 <i>Give P values as exact values whenever suitable.</i>
\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
		Our web collection on <u>statistics for biologists</u> contains articles on many of the points above.

Software and code

Policy information about availability of computer code

Data collection

The cryo-EM data were collected using the Smart EPU Software for single particle analysis by Thermofisher Scientific. The bacterial growth curves were recordered using the Agilent BioTek Gen 5 software package.

The models were created using Coot v0.8.9.2 Data analysis

The models were analyzed and visualized using ChimeraX v1.4 and Chimera v1.14

The cryo-EM snapshots were processed using Relion v3.1 and CryoSPARC v4.3.0

The models were refined using Phenix v1.20.1-4487

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 $reviewers. \ We strongly \ encourage \ code \ deposition \ in \ a \ community \ repository \ (e.g. \ GitHub). \ See the \ Nature \ Portfolio \ \underline{guidelines \ for \ submitting \ code \ \& \ software} \ for \ further \ information.$

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

DATA AVAILABILITY

The Supplementary Data 1-9 presented in this manuscript are freely accessible and available through the public repository of data, FigShare, using the following link: https://figshare.com/s/374a95769c5f7e9cdc04. The PDB accession codes for Structures 1-6, along with the associated cryo-EM maps will be made available before the publication of this manuscript via the Protein Data Bank. Currently, these structural data are shared through a freely accessible link on Google Drive.

Research involving human participants, their data, or biological material

Policy information a and sexual orientation		with human participants or human data. See also policy information about sex, gender (identity/presentation), chnicity and racism.				
Reporting on sex a	on sex and gender We did not study any human populations in our study.					
Reporting on race, other socially relev groupings		We did not study any human populations in our study.				
Population charac	n characteristics We did not study any human populations in our study.					
Recruitment		We did not study any human populations in our study.				
Ethics oversight		We did not study any human populations in our study.				
Note that full informat	ion on the appro	oval of the study protocol must also be provided in the manuscript.				
Field-specific 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 Behavioural & social sciences Ecological, evolutionary & environmental sciences						
For a reference copy of th	e document with a	all sections, see <u>nature.com/documents/nr-reporting-summary-flat.pdf</u>				
Life scien	ces stu	ıdy design				
All studies must disclose on these points even when the disclosure is negative.						
	Each cryo-EM map presented in this study was calculated by using at least 10,287 individual ribosome snapshots to reconstruct the three-dimensional volume of the ribosome.					
	Some ribosome images were excluded from the analysis based on the signal corresponding to the ribosomal ligands in the masked regions as described in detail in the Materials and Methods and Supplementary Figure 2.					
	To measure the impact of deletions of the Balon-coding gene on P. arcticus growth, each measurement was made eight times using different cell samples for each measurement.					
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		ntal systems Me	ethods		
n/a	/a Involved in the study		Involved in the study		
\boxtimes	Antibodies	\boxtimes	ChIP-seq		
\boxtimes	Eukaryotic cell lines	\boxtimes	Flow cytometry		
Palaeontology and archaeology		rchaeology	MRI-based neuroimaging		
\boxtimes	Animals and other organisms				
\boxtimes	Clinical data				
\boxtimes	Dual use research of concern				
\boxtimes	X Plants				
Plants					
Seed stocks W		We did not study plants in our st	udy.		
No	vel plant genotypes	We did not study plants in our stu	udy.		

Authentication

We did not study plants in our study.