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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 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.
X		A description of all covariates tested
\boxtimes		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

Automated Cryo-EM data collection was driven by the Leginon automation software package (v3.0)

Data analysis

Anisotropic magnification distortion was estimated by software mag_distortion_estimate (version 0.0.0). CTF estimation was performed using CTFFIND (version 4.1.8). Beam-induced motion was corrected by Unblur (version 1.0.0). Particle picking was performed using EMAN2 (version 2.2) e2helixboxer.py. Classification, helical reconstruction, and 3D refinement were performed in RELION (version 2.1.0). Refined maps were sharpened using phenix.auto_sharpen (version 1.13-2998). Atomic models were built in COOT (version 0.8.9.1). Model refinement was performed with phenix.real_space_refine (version 1.13-2998). Structures were presented using PyMOL (version 2.2.0). ThT curves were generated using GraphPad Prism (version 7.0c). Energetic calculations were performed using custom written software, and the software is available to anyone inquiring.

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 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

All structural data have been deposited into the Worldwide Protein Data Bank (wwPDB) and the Electron Microscopy Data Bank (EMDB) with the following accession codes: SegA-sym (PDB 6N37, EMD-9339), SegA-asym (PDB 6N3B, EMD-9350), SegA-slow (PDB 6N3A, EMID-9349), SegB A315E (PDB 6N3C, EMD-0334). All other data are available from the authors upon reasonable request.

Field-specific reporting						
Please select the one below that is the best fit for your research. If you are not sure, read the appropriate sections before making your selection.						
∑ Life sciences	Life sciences Behavioural & social sciences Ecological, evolutionary & environmental sciences					
For a reference copy of t	he document with all sections, see <u>nature</u> .	com/documents/nr-reporting-summary-flat.pdf				
Life sciences study design						
All studies must dis	close on these points even when	the disclosure is negative.				
Sample size	Sample sizes for ThT assays were 4 and are generally considered as sufficient for plate-reader based experiments.					
Data exclusions	No data were excluded from the results.					
Replication	All attempts at replication were successful.					
Randomization	Randomization is not relevant to our study. No research animal, human research participants or clinical studies are involved in our study.					
Blinding	Blinding was not relevant to our stu	dy. No subjective analysis were required.				
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 Meth		Methods				
n/a Involved in the study		n/a Involved in the study				
Antibodies		ChIP-seq				
Eukaryotic cell lines		Flow cytometry				
Palaeontol	ogy	MRI-based neuroimaging				

Animals and other organisms

Human research participants
Clinical data