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 <u>Editorial Policies</u> and the <u>Editorial Policy Checklist</u>.

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For	all statistical analyses, confirm that the following items are present in the figure legend, table legend, main text, or Methods section.
n/a	Confirmed
\times	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
	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 <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
X	For hierarchical and complex designs, identification of the appropriate level for tests and full reporting of outcomes
	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 <u>availability of computer code</u>

Data collection

Bruker Topspin 4.1;MEME Suite version 5.5.2; Amber ff99sb; Amber ff14sb; GROMACS; AMOEBA; PROPKA3; Tinker HP

Data analysis

Data Availability Statement

Software and code: CcpNMR Analysis 2.5.2 for NMR data analysis; CNS for structure calculation; Aria 2.0 for structure refinement; Multicoil2 for prediction of coiled-coil structure; Fiji using the Coloc2 plug-in; R (cran.r-project.org) Prism 6.0 for statistical analysis; Simulations were analysed using tcl code (available under https://doi.org/10.5281/zenodo.14163696) and the VMD software.

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 guidelines for submitting code & software for further information.

Data

Data exclusions

Randomization

Replication

Blinding

no data exclusion

no randomization

no blinding

All attempts of replication were successful.

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

Chemical shifts have been deposited under the following accession codes StREM13_160-198 BMRB ID: 52390; StREM13_171-198 BMRB ID: 52391; StREM13_150-198 BMRB ID: 52393; AtREM11_156-175 BMRB ID: 52402; AtREM12_193-212 BMRB ID: 52403; AtREM13_171-190 BMRB ID: 52404; MtREM22_189-208 BMRB ID: 52405; AtREM41_277-296 BMRB ID: 52406; AtREM51_536-555 BMRB ID: 52407; AtREM62_490-509 BMRB ID: 52408; AtREM63_423-442 BMRB ID: 52409; AtREM64_408-427 BMRB ID: 52410; AtREM65_328-347 BMRB ID: 52411. Structural ensembles have been deposited for StREM1.3171-198, StREM160-198 and StREM150-198 under the following identifiers PDB ID 9F1E, PDB ID 9F1E and PDB ID 9F1G, respectively. Primary sequences of remorins used, detailed NMR restraints have been deposited in supplementary data 1. Data used to create the Figures have been deposited in the Supplementary data 4. MD simulation data have been deposited in Zenodo under the following access: https://doi.org/10.5281/zenodo.14163696

Human resea	h participants			
Policy information a	t studies involving human research participants and Sex and Gender in Research.			
Reporting on sex a	gender NA	NA		
Population charac	stics NA			
Recruitment	NA			
Ethics oversight	NA	NA		
Note that full informat	on the approval of the study protocol must also be provided in the manuscript.			
Field-spe	fic reporting			
Please select the on	elow that is the best fit for your research. If you are not sure, read the appropriate sections before making your selectio	า.		
☐ Behavioural & social sciences ☐ Ecological, evolutionary & environmental sciences				
For a reference copy of the document with all sections, see nature.com/documents/nr-reporting-summary-flat.pdf				
Life sciences study design				
All studies must disc	on these points even when the disclosure is negative.			
Sample size	Nicotiana Benthamiana leaves were aagroinfiltrated to transform cells with fluroescent tagged proteins, At least 23 cells were analyzed for each condition over the course of 3-4 experiments.			

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		
Antibodies	ChIP-seq		
Eukaryotic cell lines	Flow cytometry		
Palaeontology and archaeology	MRI-based neuroimaging		
Animals and other organisms			
Clinical data			
Dual use research of concern			
Dual use research of concern			
Policy information about <u>dual use research of concer</u>	<u>n</u>		
Hazards			
Could the accidental, deliberate or reckless misuse in the manuscript, pose a threat to:	of agents or technologies generated in the work, or the application of information presented		
No Yes			
Public health			
National security			
Crops and/or livestock			
Ecosystems			
Any other significant area			
Experiments of concern			
Does the work involve any of these experiments of	concern:		
No Yes			
Demonstrate how to render a vaccine ineffective			
-1-			
Enhance the virulence of a pathogen or render a nonpathogen virulent			
Increase transmissibility of a pathogen			
Alter the host range of a pathogen			
Enable evasion of diagnostic/detection modalities			
Enable the weaponization of a biological agent or toxin			
Any other potentially harmful combination of e	xperiments and agents		