nature portfolio

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Last updated by author(s):	Jul 12, 2023

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
	×	The exact sample size (n) for each experimental group/condition, given as a discrete number and unit of measurement
	x	A statement on whether measurements were taken from distinct samples or whether the same sample was measured repeatedly
x		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
X		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)
x		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>
X		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
x		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

Leginon System version beta

Data analysis

MotionCor2 1.2.1, Relion 4.0-beta, CTFFIND v4.1.8, CryoSparc v3.3.1, Coot v0.9.1, Phenix dev-4694, GraphPad Prism 9, ImageJ 1.52a, and NIS-Elements Advance Research 5.2.6.

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

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

The cryo-EM reconstructions of the A61603-a1A-AR-Gq-scFv16 complex and the epinephrine-a1A-AR-Gq-scFv16 complex have been deposited in the Election Microscopy Data Bank (EMDB) under ID codes EMD-41267 and EMD-41268, respectively. The corresponding atomic models have been deposited in the Protein Data Bank (PDB) under ID codes 8THK and 8THL, respectively. Publicly available PDB entries used in this study are available under the accession codes 7B6W and 6WHA.

Human research participants						
Policy information	about <u>studies i</u>	nvolving human research participants and Sex and Gender in Research.				
Reporting on sex ar	nd gender	N/A				
Population characteristics		N/A				
Recruitment		N/A				
Ethics oversight		N/A				
Note that full informa	ation on the appr	roval of the study protocol must also be provided in the manuscript.				
Field-spe	ecific re	eporting				
		is the best fit for your research. If you are not sure, read the appropriate sections before making your selection.				
X Life sciences	□ B	Behavioural & social sciences				
For a reference copy of t	the document with	all sections, see nature.com/documents/nr-reporting-summary-flat.pdf				
Life scier	nces sti	udy design				
All studies must dis	sclose on these	points even when the disclosure is negative.				
Sample size	No Sample size calculations were performed. For cryo-EM samples, eight grids of each sample were pre-screened to identify the optimal grid for data collection. The number of grids screened were random and was not limited by any experimental parameter. The sample size was deemed sufficient as it allowed to determine the structures.					
Data exclusions	No data were e	excluded from the analyses.				
Replication		Ca2+ assays were repeated three times, and the data are represented as mean ± SD of the three independent experiments. All attempts olication were successful.				
Randomization	Division of part	ticles into random halves is automatically performed during 3D reconstruction by Relion 4.0-beta and CryoSparc v3.3.1.				
Blinding	Blinding is not a	inding is not applicable for this study, as group allocation is not used.				
	C					
Reportin	g for sp	pecific materials, systems and methods				
		about some types of materials, experimental systems and methods used in many studies. Here, indicate whether each material, your study. If you are not sure if a list item applies to your research, read the appropriate section before selecting a response.				
Materials & ex	perimental s	systems Methods				
n/a Involved in th	ne study	n/a Involved in the study				
X Antibodies	X Antibodies X ChIP-seq					
	Eukaryotic cell lines X D Flow cytometry					
	nd other organisn to	ns				
Clinical dat Dual use re	esearch of conce	rn				
E Dadi ase it	escurer or correct					
Eukaryotic cell lines						
Policy information about <u>cell lines and Sex and Gender in Research</u>						
Cell line source(s)	e source(s) Insect cell line Sf9 was obtained from Expression Systems. HEK293T cell lines were obtained from ATCC.					
Authentication Authentication was not performed for this study.		Authentication was not performed for this study.				

All cell lines were tested negative for mycoplasma contamination.

Mycoplasma contamination

No commonly misidentified cell lines were used.

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