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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 Authors & Referees and the Editorial Policy Checklist.

For all statistical analyses, confirm that the following items are present in the figure legend, table legend, main text, or Methods section.

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n/a	Confirmed						
	The exact sam	The exact sample size (n) for each experimental group/condition, given as a discrete number and unit of measurement					
	A statement o	A statement on whether measurements were taken from distinct samples or whether the same sample was measured repeatedly					
	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	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						
\boxtimes	For hierarchical and complex designs, identification of the appropriate level for tests and full reporting of outcomes						
\square 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.							
So	ftware and c	ode					
Poli	cy information abou	ut <u>availability of computer code</u>					
D	ata collection	Leginon software was used for negative-stain microscopy data collection. ChromLab™ Software was used for collection of size-exclusion chromatography data. The Octet® Data Analysis HT software was used for collection of BLI data.					
D	ata analysis	Generally Graphpad Prism 7.0 was used for data and statistical analysis. Appion software was used for nsEM data processing. Relion 3.0 was used for 2D- and 3D-class average data collection. UCSF Chimera 1.13 nwas used for visualization, segmentation and figure preparation of 3D-refined maps. Glycopeptide fragmentation data was analyzed using Byos v3.9 (Protein Metrics Inc.).					
Forr	nanuscripts utilizing custo	om 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:

We strongly encourage code deposition in a community repository (e.g. GitHub). See the Nature Research guidelines for submitting code & software for further information.

- 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

Data availability

The data supporting the findings of the study are available from the corresponding authors upon reasonable request. 3D refined models of 16055 SOSIP complexed with polyclonal Fabs were submitted to EMDB. The list of EMDB IDs: 22714 (16055 SOSIP + r2463 Poly Fab); 22715 (16055 SOSIP + r2464 Poly Fab); 22716 (16055 SOSIP + r2465 Poly Fab); 22717 (16055 SOSIP + r2466 Poly Fab); 22718 (16055 SOSIP + r2467 Poly Fab); 22719 (16055 SOSIP + r2468 Poly Fab); 22720 (16055 SOSIP + r2469 Poly Fab), 22721 (16055 SOSIP + r2470 Poly Fab); 22722 (16055 SOSIP + r2471 Poly Fab); 22723 (16055 SOSIP + r2472 Poly Fab).

Field-spe	cific re	porting				
Please select the or	ne below that is	the best fit for your research. If you are not sure, read the appropriate sections before making your selection.				
✓ Life sciences	В	ehavioural & social sciences Ecological, evolutionary & environmental sciences				
For a reference copy of t	For a reference copy of the document with all sections, see nature.com/documents/nr-reporting-summary-flat.pdf					
Life scier	nces stu	ıdy design				
All studies must dis	isclose on these points even when the disclosure is negative.					
Sample size	No sample-size	sample-size calculations were performed.				
Data exclusions	No data was exc	No data was excluded from analyses.				
Replication	All data points v	All data points were repeated or performed in duplicate/triplicate.				
Randomization	Allocation was random.					
Blinding	Blinding was no	t relevant for our studies.				
We require informatic system or method list Materials & exp n/a Involved in th	on from authors a ced is relevant to perimental sy le study	n/a Involved in the study				
Antibodies						
Antibodies used		v-specific antibodies were obtained from either Michel Nussenzweig, James Robinson, Dennis Burton, Peter Kwong, Mark nnors, John Mascola or William Olson through the NIH AIDS Research and Reference Reagent Program.				
Validation	N.A.					
Eukaryotic c	ell lines					
Policy information a	about <u>cell lines</u>					
Cell line source(s))	HEK293F cells were obtained from Invitrogen, cat no. R79007. Ramos B cells were obtained from Drs. Li Wu and Vineet N. KewalRaman via the NIH AIDS Reagent Program, Division of AIDS, NIAID, NIH.				
Authentication		Cell lines were validated by vendors or collaborators.				
Mycoplasma contamination All cell lines were tested negative for mycoplasma contamination		All cell lines were tested negative for mycoplasma contamination				

Commonly misidentified lines (See <u>ICLAC</u> register)

None.

Animals and other organisms

Policy information about studies involving animals; ARRIVE guidelines recommended for reporting animal research

Laboratory animals New Zealand White Rabbits, 2.5-3kg, female.

Wild animals No wild animals were used.

Field-collected samples No field-collected samples were used.

Ethics oversight All procedures performed in rabbits were done by Covance and approved by Denver PA IACUC Committee, #0035-016

Note that full information on the approval of the study protocol must also be provided in the manuscript.