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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.) [α	1.51	

For	all statistical ar	nalyses, confirm that the following items are present in the figure legend, table legend, main text, or Methods section.		
n/a	Confirmed			
	The exact	The exact sample size (n) for each experimental group/condition, given as a discrete number and unit of measurement		
	A statem	A statement on whether measurements were taken from distinct samples or whether the same sample was measured repeatedly		
	The statis	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 des	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 Give <i>P</i> values as exact values whenever suitable.			
\boxtimes	For Bayes	sian 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	s 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 No software was used		No software was used		
Da	Data analysis SAS 9.4 and GraphPad Prism 9.4.1			
		g 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 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 <u>data availability statement</u>. 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

All data that support the findings of this study are available from the corresponding author upon reasonable request.

arch parti	zicipants			
Policy information about studies involving human research participants and Sex and Gender in Research.				
and gender	NA			
cteristics	NA			
	NA			
	NA			
ation on the appr	proval of the study protocol must also be provided in the manuscript.			
ecific re	eporting			
ne below that i	is the best fit for your research. If you are not sure, read the appropriate sections before making your selection.			
E	Behavioural & social sciences			
the document with	h all sections, see <u>nature.com/documents/nr-reporting-summary-flat.pdf</u>			
nces sti	udy design			
sclose on these	e points even when the disclosure is negative.			
	study had a predefined objective, and based on historical data obtained in-house, sample sizes were calculated by statisticians to detect etermined effect sizes with 80% power.			
No data were e	excluded			
	its included either control groups or control time-points validating the experiments. All experiments met the pre-set gatekeepers y of the experiment. Experiments were not replicated due to ethical reasons of animal experimentation			
allocation was	s random			
Operators were	re blinded to the objectives of the study			
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,				
	o 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				
ne study	n/a Involved in the study ☑ ChIP-seq			
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Animals and other organisms				
Clinical data Dual use research of concern				
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	about studies and gender cteristics ation on the app ecific re ne below that the document wit CCS St sclose on thes Each study ha predetermine No data were All experiment for the validit allocation was Operators we g for S on from author ted is relevant t perimental ne study cell lines ogy and archae and other organis			

Antibodies

Antibodies used

CV3-25 antibody (ImmunoPrecise)

Validation

CV3-25 binds the S2 subunit of the SARS-CoV-2 Spike protein [Jennewein, M. F. et al. Isolation and characterization of cross-neutralizing coronavirus antibodies from COVID-19+ subjects. Cell Rep. 36, 109353 (2021)]

Animals and other research organisms

Policy information about <u>studies involving animals</u>; <u>ARRIVE guidelines</u> recommended for reporting animal research, and <u>Sex and Gender in Research</u>

Laboratory animals	Female BALB/c mice aged 8-10 weeks and Female Syrian golden RjHan:AURA hamsters, aged 9-14 weeks
Wild animals	NA
Reporting on sex	Only female hamsters were used in these studies to reduce inter-individual variation, thereby limiting the number of animals required
Field-collected samples	NA
Ethics oversight	Animal experiments were approved by the Central Authority for Scientific Procedures on Animals (Centrale Commissie Dierproeven).

the institutional animal welfare body, and conducted in accordance with the European guidelines (EU directive on animal testing

2010/63/EU and ETS 123) and local Dutch legislation on animal experiments

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