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 Editorial Policies and the Editorial Policy Checklist.

Please do not complete any field with "not applicable" or n/a. Refer to the help text for what text to use if an item is not relevant to your study. For final submission: please carefully check your responses for accuracy; you will not be able to make changes later.

Statistics

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
	The exact sample size (n) for each experimental group/condition, given as a discrete number and unit of measurement
	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.
	A description of all covariates tested



- 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. *F*, *t*, *r*) with confidence intervals, effect sizes, degrees of freedom and *P* value noted Give *P* values as exact values whenever suitable.
 - For Bayesian analysis, information on the choice of priors and Markov chain Monte Carlo settings
 - For hierarchical and complex designs, identification of the appropriate level for tests and full reporting of outcomes

(IDA) description of any assumptions or corrections, such as tests of normality and adjustment for multiple comparisons

© Estimates of effect sizes (e.g. Cohen's d, Pearson's r), 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 availability of computer code

Data collection | Spectra data was collected using Bruker OPUS 8.8. Electrochemical data was collected using EC-Lab V11.43.

Data analysis The data was analyzed using Python 3.10 codes. The codes are provided here: https://zenodo.org/records/11122131 where requirements.txt

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 original spectrum data are provided in a zenodo record: https://zenodo.org/records/11122131 Source data of the figure are provided with this paper.

Research involving hu	man participants, their data, or biological material
Policy information about studies v sexual orientation and race, ethnic	with human participants or human data. See also policy information about sex, gender (identity/presentation), and city and racism.
Reporting on sex and gender	Not applicable
Reporting on race, ethnicity, or other socially relevant grouping	Not applicable s
Population characteristics	Not applicable
Recruitment	Not applicable
Ethics oversight	Not applicable
Note that full information on the appr	oval of the study protocol must also be provided in the manuscript.
Field-specific re	porting
Please select the one below that is	s the best fit for your research. If you are not sure, read the appropriate sections before making your selection.
OLife sciences OBe	ehavioural & social sciences
Life sciences stu	points even when the disclosure is negative.
Sample size	
Data exclusions	
Replication	
Randomization	
Blinding	
Behavioural & s	social sciences study design
All studies must disclose on these	points even when the disclosure is negative.
Study description	
Research sample	
Sampling strategy	
Data collection	
Timing	
Data exclusions	
Non-participation	
Randomization	
Ecological, evol	utionary & environmental sciences study design
All studies must disclose on these	points even when the disclosure is negative.
Study description	

Study description

Research sample

Sampling strategy

Data collection

Timing and spatial scale

Data exclusions

Reproducibility

Randomization

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Blinding	
Did the study involve field	d work? Oyes ONo
ield work, collect	tion and transport
Field conditions	
Location Access & import/export Disturbance	
Reporting fo	r specific materials, systems and methods
	uthors about some types of materials, experimental systems and methods used in many studies. Here, indicate whether each material, vant to your study. If you are not sure if a list item applies to your research, read the appropriate section before selecting a response.
Involved in the stu Antibodies Eukaryotic cell lines Palaeontology and arc Animals and other org Clinical data Dual use research of c	Involved in the study ChIP-seq Flow cytometry Chaeology MRI-based neuroimaging ganisms
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Commonly misidentified I (See ICLAC register)	ines
alaeontology and	d Archaeology
Specimen provenance Specimen deposition	
Dating methods Tick this box to confirm	n that the raw and calibrated dates are available in the paper or in Supplementary Information.
Ethics oversight ote that full information on the	he approval of the study protocol must also be provided in the manuscript.
animals and other	r research organisms

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

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Laboratory animals	
Wild animals	
Reporting on sex	
Field-collected samples	
Ethics oversight	
Note that full information on t	he approval of the study protocol must also be provided in the manuscript.
Clinical data	
Policy information about cl	inical studies
All manuscripts should comply	with the ICMJE guidelines for publication of clinical research and a completed CONSORT checklist must be included with all submissions.
Clinical trial registration	
Study protocol	
Data collection	
Outcomes	
Outcomes	
Dual use research	of concern
Policy information about du	ual use research of concern
Hazards	
Could the accidental, deli	berate or reckless misuse of agents or technologies generated in the work, or the application of information presented in
the manuscript, pose a th	nreat to:
No Yes	
OPublic health	
ONational security	
OCrops and/or livesto	ck
O Ecosystems	
OAny other significant	area
Experiments of concer	rn
Does the work involve an	y of these experiments of concern:
No Yes	y of these experiments of concern.
ODemonstrate how to	o render a vaccine ineffective
	therapeutically useful antibiotics or antiviral agents
© OEnhance the virulen	ce of a pathogen or render a nonpathogen virulent
OIncrease transmissib	ility of a pathogen
Alter the best range	of a pathogen

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lo	Yes
C	ODemonstrate how to render a vaccine ineffective
C	OConfer resistance to therapeutically useful antibiotics or antiviral agents
C	Enhance the virulence of a pathogen or render a nonpathogen virulent
C	OIncrease transmissibility of a pathogen
C	OAlter the host range of a pathogen
C	Enable evasion of diagnostic/detection modalities
C	Enable the weaponization of a biological agent or toxin
C	OAny other potentially harmful combination of experiments and agents
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Plants	
Seed stocks	
Novel plant genotypes	
Authentication	
ChIP-seq	
Data deposition	
	and final processed data have been deposited in a public database such as GEO.
Confirm that you have	deposited or provided access to graph files (e.g. BED files) for the called peaks.
Data access links May remain private before public	ation
Files in database submission	
Genome browser session (e.g. UCSC)	
Methodology Replicates	
Sequencing depth	
Antibodies	
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Peak calling parameters Data quality	
Software	
Flow Cytomotry	
Flow Cytometry	
Plots	
Confirm that:	
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	rly visible. Include numbers along axes only for bottom left plot of group (a 'group' is an analysis of identical markers).
	ots with outliers or pseudocolor plots.
■A numerical value for n	umber of cells or percentage (with statistics) is provided.
Methodology Sample preparation	
Instrument	
Software	
Cell population abundance	a
Gating strategy	
Tick this box to confirm	that a figure exemplifying the gating strategy is provided in the Supplementary Information.
Magnetic resonan	ice imaging
Experimental design Design type	

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Design specifications	
Behavioral performance measures	
Acquisition Imaging type(s)	
Field strength	
Sequence & imaging parameters	
Area of acquisition	
Diffusion MRI OUsed	ONot used
Preprocessing Preprocessing software	
Normalization	
Normalization template	
Noise and artifact removal	
Volume censoring	
Statistical modeling & inference Model type and settings Effect(s) tested Specify type of analysis: Whole	
Statistic type for inference	
(See Eklund et al. 2016)	
Correction	
Models & analysis a Involved in the study a Functional and/or effective conn Graph analysis Multivariate modeling or predict Functional and/or effective connecti Graph analysis	ive analysis
Multivariate modeling and predictive	e analysis