# 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
	ightharpoonup 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 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>
	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
	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 availability of computer code

Data collection Origin 2022b, h5py 3.9.0

Data analysis Python 3.9.5, GPAW 21.6.0, Pytorch 2.1.1+cu118, Numpy 1.26.4, matplot 3.4.3

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 <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

The data (KS wavefunction and GW energies of 302 materials) generated in this study have been deposited in the Materials Data Facility repository under DOI https://doi.org/10.18126/rpa1-yp91. Source data are provided with this paper.

Research inv	olving ht	ıman participants, their data, or biological material	
		with <u>human participants or human data</u> . See also policy information about <u>sex, gender (identity/presentation),</u> ethnicity and racism.	
Reporting on sex ar	nd gender	not applicable	
Reporting on race, ethnicity, or other socially relevant groupings		not applicable	
Population characte	eristics	not applicable	
Recruitment		not applicable	
Ethics oversight not a		not applicable	
Note that full informa	ation on the app	roval of the study protocol must also be provided in the manuscript.	
Field-spe		eporting  is the best fit for your research. If you are not sure, read the appropriate sections before making your selection.	
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Life sciences  For a reference copy of t		Behavioural & social sciences	
_ife scier	nces st	udy design	
All studies must dis	close on these	e points even when the disclosure is negative.	
Sample size	Describe how sample size was determined, detailing any statistical methods used to predetermine sample size OR if no sample-size calculation was performed, describe how sample sizes were chosen and provide a rationale for why these sample sizes are sufficient.		
Data exclusions	Describe any data exclusions. If no data were excluded from the analyses, state so OR if data were excluded, describe the exclusions and the rationale behind them, indicating whether exclusion criteria were pre-established.		
Replication	Describe the measures taken to verify the reproducibility of the experimental findings. If all attempts at replication were successful, confirm this OR if there are any findings that were not replicated or cannot be reproduced, note this and describe why.		
Randomization	Describe how samples/organisms/participants were allocated into experimental groups. If allocation was not random, describe how covariates were controlled OR if this is not relevant to your study, explain why.		
Blinding	Describe whether the investigators were blinded to group allocation during data collection and/or analysis. If blinding was not possible, describe why OR explain why blinding was not relevant to your study.		
		social sciences study design	
All studies must dis	close on these	e points even when the disclosure is negative.	
,		y describe the study type including whether data are quantitative, qualitative, or mixed-methods (e.g. qualitative cross-sectional, titative experimental, mixed-methods case study).	
inform		the research sample (e.g. Harvard university undergraduates, villagers in rural India) and provide relevant demographic mation (e.g. age, sex) and indicate whether the sample is representative. Provide a rationale for the study sample chosen. For es involving existing datasets, please describe the dataset and source.	
Sampling strategy	prede ratior	tibe the sampling procedure (e.g. random, snowball, stratified, convenience). Describe the statistical methods that were used to etermine sample size OR if no sample-size calculation was performed, describe how sample sizes were chosen and provide a nale for why these sample sizes are sufficient. For qualitative data, please indicate whether data saturation was considered, and criteria were used to decide that no further sampling was needed.	
		de details about the data collection procedure, including the instruments or devices used to record the data (e.g. pen and paper, outer, eye tracker, video or audio equipment) whether anyone was present besides the participant(s) and the researcher, and	

Indicate the start and stop dates of data collection. If there is a gap between collection periods, state the dates for each sample cohort.

whether the researcher was blind to experimental condition and/or the study hypothesis during data collection.

Timing

Data exclusions If no data were excluded from the analyses, state so OR if data were excluded, provide the exact number of exclusions and the rationale behind them, indicating whether exclusion criteria were pre-established.

State how many participants dropped out/declined participation and the reason(s) given OR provide response rate OR state that no Non-participation participants dropped out/declined participation.

If participants were not allocated into experimental groups, state so OR describe how participants were allocated to groups, and if Randomization allocation was not random, describe how covariates were controlled.

# Ecological, evolutionary & environmental sciences study design

All studies must disclose on these points even when the disclosure is negative.

Briefly describe the study. For quantitative data include treatment factors and interactions, design structure (e.g. factorial, nested, Study description hierarchical), nature and number of experimental units and replicates. Describe the research sample (e.g. a group of tagged Passer domesticus, all Stenocereus thurberi within Organ Pipe Cactus National Research sample Monument), and provide a rationale for the sample choice. When relevant, describe the organism taxa, source, sex, age range and any manipulations. State what population the sample is meant to represent when applicable. For studies involving existing datasets, describe the data and its source. Sampling strategy Note the sampling procedure. Describe the statistical methods that were used to predetermine sample size OR if no sample-size calculation was performed, describe how sample sizes were chosen and provide a rationale for why these sample sizes are sufficient. Data collection Describe the data collection procedure, including who recorded the data and how. Timing and spatial scale Indicate the start and stop dates of data collection, noting the frequency and periodicity of sampling and providing a rationale for these choices. If there is a gap between collection periods, state the dates for each sample cohort. Specify the spatial scale from which the data are taken If no data were excluded from the analyses, state so OR if data were excluded, describe the exclusions and the rationale behind them, Data exclusions indicating whether exclusion criteria were pre-established. Describe the measures taken to verify the reproducibility of experimental findings. For each experiment, note whether any attempts to Reproducibility repeat the experiment failed OR state that all attempts to repeat the experiment were successful. Randomization Describe how samples/organisms/participants were allocated into groups. If allocation was not random, describe how covariates were controlled. If this is not relevant to your study, explain why. Blinding Describe the extent of blinding used during data acquisition and analysis. If blinding was not possible, describe why OR explain why blinding was not relevant to your study. Did the study involve field work?

## Field work, collection and transport

Field conditions Describe the study conditions for field work, providing relevant parameters (e.g. temperature, rainfall).

Location State the location of the sampling or experiment, providing relevant parameters (e.g. latitude and longitude, elevation, water depth).

Describe the efforts you have made to access habitats and to collect and import/export your samples in a responsible manner and in Access & import/export compliance with local, national and international laws, noting any permits that were obtained (give the name of the issuing authority,

the date of issue, and any identifying information).

Describe any disturbance caused by the study and how it was minimized. Disturbance

# 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	x	ChIP-seq	
×	Eukaryotic cell lines	x	Flow cytometry	
×	Palaeontology and archaeology	×	MRI-based neuroimaging	
×	Animals and other organisms			
×	Clinical data			
×	Dual use research of concern			
×	Plants			

### **Plants**

Seed stocks

Report on the source of all seed stocks or other plant material used. If applicable, state the seed stock centre and catalogue number. If plant specimens were collected from the field, describe the collection location, date and sampling procedures.

Novel plant genotypes

Describe the methods by which all novel plant genotypes were produced. This includes those generated by transgenic approaches, gene editing, chemical/radiation-based mutagenesis and hybridization. For transgenic lines, describe the transformation method, the number of independent lines analyzed and the generation upon which experiments were performed. For gene-edited lines, describe the editor used, the endogenous sequence targeted for editing, the targeting guide RNA sequence (if applicable) and how the editor was applied.

Authentication

Describe any authentication procedures for each seed stock used or novel genotype generated. Describe any experiments used to assess the effect of a mutation and, where applicable, how potential secondary effects (e.g. second site T-DNA insertions, mosiacism, off-target gene editing) were examined.