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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St	at	isti	CS

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
	\boxtimes	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
\boxtimes		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
X		A description of any assumptions or corrections, such as tests of normality and adjustment for multiple comparisons
	\boxtimes	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)
\boxtimes		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
	\boxtimes	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

The 20 combinatorial mutagenesis datasets were downloaded from the publications referenced in Table 1. Some were edited as described in Methods to suit the purpose of our analysis.

Data analysis

All scripts used for analysis are available on our GitHub repository: http://github.com/whatdoidohaha/RFA

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

All sequence-function data were gathered from published studies (Table 1) and are available on GitHub (https://github.com/whatdoidohaha/RFA) and Zenodo (https://doi.org/10.5281/zenodo.8307147).

Research inv	volving hu	man participants, their data, or biological material	
		vith human participants or human data. See also policy information about sex, gender (identity/presentation), thnicity and racism.	
Reporting on sex	and gender	Our study does not involve human subjects.	
Reporting on rac other socially rela groupings		Our study does not involve human subjects.	
Population chara	ecteristics	Our study does not involve human subjects.	
Recruitment		Our study does not involve human subjects.	
Ethics oversight		Our study does not involve human subjects.	
Note that full informa	ation on the appro	oval of the study protocol must also be provided in the manuscript.	
Field-spe	ecific re	porting	
Please select the o	ne below that is	s 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	
For a reference copy of	the document with a	all sections, see <u>nature.com/documents/nr-reporting-summary-flat.pdf</u>	
Life scier	nces stu	udy design	
All studies must dis	sclose on these	points even when the disclosure is negative.	
Sample size	We searched all combinatorial mutagenesis datasets that meet the precision and coverage standard: the squared correlation among replicate measurements must be greater than 0.9, and these measurements must be available for at least 40% of possible genotypes. Among the many available small datasets with fewer than 100 possible genotypes, we focused on three datasets for which high-order epistasis has been reported. Otherwise, all datasets meeting the above standards were included for analysis. This resulted in total 20 datasets encompassing structurally and functionally diverse proteins.		
Data exclusions	For 2 of the 20	datasets, a subset of genotypes with large measurement noise were removed as described in Methods.	
Replication	Each combinatorial mutagenesis dataset derives from two or more replicate measurements. To minimize the impact of measurement noise, we only analyzed datasets for which the squared correlation among replicate measurements is greater than 0.9.		
Randomization	Randomization is not relevant to our study design.		
Blinding	Blinding is not relevant to our study design.		
We require informati	perimental sone study	Decific 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. Methods	
Animals ar	logy and archaeol nd other organism ta esearch of concer	is a second of the second of t	

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.

Describe any authentication procedures for each seed stock used or novel genotype generated. Describe any experiments used to

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.