nature portfolio

Corresponding author(s):	Julio Saez-Rodriguez, Bing Zhang
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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.

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	onfirmed
	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.
X	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>
x	For Bayesian analysis, information on the choice of priors and Markov chain Monte Carlo settings
x	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 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 Repository with o

Repository with data used in the presented analysis: https://zenodo.org/records/12566560 and repository with code used to obtain the data whenever applicable: https://github.com/saezlab/kinase_benchmark

Data analysis

Repository with code used for the presented analysis: https://github.com/saezlab/kinase_benchmark
Additionally the following packages were used for the analysis: OmniPathR (v3.11.1), decoupler v2.8.0), stats (v4.3.3), ssGSEA2 (v1.0.1), decoupler-py (v1.6.1)

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 benchmark data used in this study are available in the benchmarKIN package (https://github.com/saezlab/benchmarKIN) and have been deposited in a Zenodo

repository (https://zenodo.org/uploads/12566560). The kinase-substrate libraries used in this study are available: from the following sources: PhosphoSitePlus (https://www.phosphosite.org/staticDownloads#), PTMsigDB (https://proteomics.broadapps.org/ptmsigdb/), and iKiP-DB (https://pubs.acs.org/doi/suppl/10.1021/acs.jproteome.2c00198/suppl_file/pr2c00198_si_007.zip). NetworKIN can be accessed at http://netphorest.science/download/networkin_human_predictions_3.1.tsv.xz and is also included in the aforementioned Zenodo repository.

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Research involv	III BIII	uman p	ai ucipai	nus, unen	uata, or	DIDIORICAL	matenai

		vith human participants or human data. See also policy information about sex, gender (identity/presentation), thnicity and racism.			
Reporting on sex ar	nd gender	NA			
Reporting on race, other socially relev		NA			
Population characteristics NA		NA			
Recruitment		NA			
Ethics oversight NA					
Note that full informa	ation on the appr	oval of the study protocol must also be provided in the manuscript.			
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Field-spe					
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Life sciences		ehavioural & social sciences			
For a reference copy of t	the document with	all sections, see <u>nature.com/documents/nr-reporting-summary-flat.pdf</u>			
l ife scier	nces sti	udy design			
		points even when the disclosure is negative.			
Sample size	NA				
Data exclusions	Data from colon (COAD), ovarian (OV), and pancreatic (PDAC) cancer were excluded from our tumor-based benchmark approach. These cancer types exhibited lower median correlations (0.32, 0.26, and 0.31, respectively) compared to others. Since we previously observed that common targets of the same kinases were less well correlated in these three cancer types than in others, we excluded them to ensure more consistent correlation patterns across the remaining data.				
Replication	All attempts we	All attempts were successful			
Randomization	NA .				
Blinding	NA				
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•	<u> </u>	pecific 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.			
Materials & ex	<u> </u>	·			
		n/a Involved in the study X ChIP-seq			
Eukaryotic cell lines		Flow cytometry			
Palaeontology and archaeology MRI-based neuroimaging					
Animals an	nd other organism	is			
Clinical dat	ia .				
Dual use re	esearch of concer	n			
✗ ☐ Plants					

Plants

Seed stocks	NA
Novel plant genotypes	NA
Authentication	NA