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Last updated by author(s):	Aug 16, 2022

Reporting Summary

Nature Research 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 Research policies, see our Editorial Policies and the Editorial Policy Checklist.

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For a	all statistical ana	alyses, confirm that the following items are present in the figure legend, table legend, main text, or Methods section.
n/a	Confirmed	
\boxtimes	The exact s	sample size (n) for each experimental group/condition, given as a discrete number and unit of measurement
\boxtimes	A statemer	nt on whether measurements were taken from distinct samples or whether the same sample was measured repeatedly
\boxtimes	1 1	ical test(s) used AND whether they are one- or two-sided on tests should be described solely by name; describe more complex techniques in the Methods section.
\boxtimes	A descripti	on of all covariates tested
	A descripti	on of any assumptions or corrections, such as tests of normality and adjustment for multiple comparisons
	A full descr AND variat	ription of the statistical parameters including central tendency (e.g. means) or other basic estimates (e.g. regression coefficient) ion (e.g. standard deviation) or associated estimates of uncertainty (e.g. confidence intervals)
\boxtimes	1 1 '	pothesis testing, the test statistic (e.g. F , t , r) with confidence intervals, effect sizes, degrees of freedom and P value noted as as exact values whenever suitable.
\boxtimes	For Bayesia	an analysis, information on the choice of priors and Markov chain Monte Carlo settings
\boxtimes	For hierard	chical 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.
Sof	tware and	d code
Polic	y information a	bout <u>availability of computer code</u>
Da	ta collection	All the codes and data is publicly available and can be downloaded via a web browser.
Da	ta analysis	The TIAToolbox software library presented in the paper was used for analysis.
	,	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 ncourage code deposition in a community repository (e.g. GitHub). See the Nature Research guidelines for submitting code & software for further information.

Data

Policy information about <u>availability of data</u>

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 list of figures that have associated raw data
- A description of any restrictions on data availability

All datasets analysed during the production of TIAToolbox, except for one private oral dysplasia cohort dataset for HoVer-Net+, are publicly available. They can be accessed for research and non-commercial use at the following web addresses:

- The Cancer Genome Atlas (TCGA): https://www.cancer.gov/tcga
- $\bullet \ PanNuke: https://warwick.ac.uk/fac/cross_fac/tia/data/pannuke\\$
- PatchCamelyon (PCam): https://github.com/basveeling/pcam
- Kather 100k: https://zenodo.org/record/1214456
- Kumar (MoNuSeg Subset): https://monuseg.grand-challenge.org/
- MoNuSAC: https://monusac-2020.grand-challenge.org/

' ''	rwick.ac.uk/fac/cross_fac/tia/data/hovernet/ lasia cohort dataset is not available because we do not currently have ethical approval to share this dataset but the trained model is already
Field-spe	cific reporting
•	be below that is the best fit for your research. If you are not sure, read the appropriate sections before making your selection. Behavioural & social sciences Ecological, evolutionary & environmental sciences
	ne document with all sections, see nature.com/documents/nr-reporting-summary-flat.pdf
Life scier	ices study design
	close on these points even when the disclosure is negative.
Sample size	Publicly available or previously published datasets were used. Hence the sample size was dependent on the data availability.
Data exclusions	Publicly available or previously published datasets were used.
Replication	Models were trained on multiple datasets. Previously published results were also replicated.
Randomization	Original randomized splits were used for publicly available datasets were used where possible. When performing training with the Kather 100k dataset, we used an 80/20 split on the non-normalized data variant due to the availability of only stain-normalized test data. We used non-normalized data to improve generality and computational efficiency of the algorithm. If original data splits were not available, a randomized split was used.
Blinding	Publicly available or previously published datasets were used.
We require information system or method list	g for specific materials, systems and methods on from authors about some types of materials, experimental systems and methods used in many studies. Here, indicate whether each material, ed 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.
	perimental systems Methods n/a Involved in the study
n/a Involved in th	e study n/a Involved in the study ChIP-seq
Eukaryotic	
Palaeontolo	pgy and archaeology MRI-based neuroimaging
Animals an	d other organisms
	earch participants
Clinical dat	
Dual use re	search of concern
Human rese	arch participants
Policy information a	about <u>studies involving human research participants</u>
Population chara	Publicly available or previously published datasets were used.
Recruitment	Publicly available or previously published datasets were used. No recruitment was required.
Ethics oversight	N/A. Publicly available or previously published datasets were used.
Note that full informa	tion on the approval of the study protocol must also be provided in the manuscript.
Clinical data	
	about <u>clinical studies</u> I comply with the ICMJE <u>guidelines for publication of clinical research</u> and a completed <u>CONSORT checklist</u> must be included with all submissions.

Clinical trial registration N/A

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Study protocol

Data collection

N/A

Outcomes

N/A

N/A