# nature research

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## **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 <u>Editorial Policies</u> and the <u>Editorial Policy Checklist</u>.

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FOL	all statistical analyses, confirm that the following items are present in the figure regend, table regend, main text, or Methods Section.
n/a	Confirmed
	$\mathbf{x}$ 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)
x	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
	For hierarchical and complex designs, identification of the appropriate level for tests and full reporting of outcomes
	$\blacksquare$ 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 R version 3.4.4 (R Development Core Team, 2019)

Data analysis

minfi Bioconductor package version 1.24.0, limma package version 3.34.5, Rtsne version 0.13, R package randomForest version 4.6-12, R package glmnet version 2.0-18, R-package conumee, R-package RF\_Purify, also see https://github.com/mwsill/mnp\_training

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

Methylation data required for building the sarcoma classifier (reference set) were deposited at the public repository Gene Expression Omnibus (GEO; http://www.ncbi.nlm.nih.gov/geo) under the accession number (GSE140686). Supplementary Table 1 indicates the IDAT file names for each case. The remaining data are available within the Article, Supplementary Information or available from the authors upon request.

Field-sp	ecific reporting				
Please select the	one below that is the best fit for y	our research. If you are not sure, read the appropriate sections before making your selection.			
X Life sciences	Behavioural & soc	ial sciences Ecological, evolutionary & environmental sciences			
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Life scie	nces study desi	ign			
	lisclose on these points even whe	<u> </u>			
Sample size	Classifier methylation classes need at least 7 cases (pre-established, see the brain tumour classifier "Capper et al., Nature 2018)				
Data exclusions	No data sets were excluded.				
Replication	The separation of samples into the defined DNA methylation classes was reliably reproduced by iterative random downsampling of the reference cohort.				
Randomization	The methylation classifier reference cohort was compiled to recapitulate the entities established in the WHO classification of soft tissue and bone tumours. No randomization was performed.				
Blinding	Blinding was not relevant to this study, because the classifier prediction result is an objective measure, which was compared with the institutional diagnosis.				
We require informa	tion from authors about some types	naterials, systems and methods of materials, experimental systems and methods used in many studies. Here, indicate whether each material, 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			
X Antibodies		ChIP-seq			
	ic cell lines	Flow cytometry			
Palaeontology and archaeology		MRI-based neuroimaging			
<b>≭</b> Human n	X   Human research participants				

### Human research participants

Dual use research of concern

Clinical data

Policy information about studies involving human research participants

Population characteristics

Reference cohort: 1077 cases, 502 females + 575 males
Validation cohort: 428 cases, 195 females + 233 males

Recruitment

Reference cohort: national and international sarcoma (reference) centres
Validation cohort: MASTER, INFORM, MNP2.0, PTT2.0 trials

Ethics oversight

This study has been performed in accordance with the Declaration of Helsinki.

Note that full information on the approval of the study protocol must also be provided in the manuscript.