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Corresponding author(s):	DBPR - COMMSMED-21-0167-T
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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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S †	· a:	tic	ŤΒ	\sim

For all statistical a	nalyses, confirm that the following items are present in the figure legend, table legend, main text, or Methods section.		
n/a Confirmed			
☐ ☐ The exact	t sample size (n) for each experimental group/condition, given as a discrete number and unit of measurement		
A statem	ent on whether measurements were taken from distinct samples or whether the same sample was measured repeatedly		
The statis	stical test(s) used AND whether they are one- or two-sided non tests should be described solely by name; describe more complex techniques in the Methods section.		
A descrip	tion of all covariates tested		
A descrip	tion of any assumptions or corrections, such as tests of normality and adjustment for multiple comparisons		
A full des	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 h	sypothesis testing, the test statistic (e.g. F , t , r) with confidence intervals, effect sizes, degrees of freedom and P value noted uses as exact values whenever suitable.		
For Bayes	sian analysis, information on the choice of priors and Markov chain Monte Carlo settings		
For hiera	rchical and complex designs, identification of the appropriate level for tests and full reporting of outcomes		
Estimate:	s 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 an	id code		
Policy information	about <u>availability of computer code</u>		
Data collection	Excel, GraphPad Prism		
Data analysis	Excel, GraphPad Prism		
	g 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 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 <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

Source data for the main figures are available as Supplementary Data 1. Additional data sets that support the findings of this study are available from the corresponding author upon reasonable request.

Field-spe	ecific re	eporting
Please select the o	ne below that i	is the best fit for your research. If you are not sure, read the appropriate sections before making your selection.
Life sciences	E	Behavioural & social sciences Ecological, evolutionary & environmental sciences
For a reference copy of t	the document with	all sections, see <u>nature.com/documents/nr-reporting-summary-flat.pdf</u>
Life scier	nces sti	udy design
All studies must dis	sclose on these	points even when the disclosure is negative.
Sample size	Retrospective a	analysis, no sample size calculation
Data exclusions	No data were e	excluded
Replication	Not applicable	in this retrospective analysis
Randomization	Retrospective a	analysis, no randomization
Blinding	Blinding was no	ot applicable in this retrospective analysis
Reportin	g for s	pecific materials, systems and methods
We require informati	on from authors	about some types of materials, experimental systems and methods used in many studies. Here, indicate whether each material, by 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		
n/a Involved in th		n/a Involved in the study
Antibodies	;	ChIP-seq
Eukaryotic		Flow cytometry
	logy and archaeo nd other organisr	
	search participan	
Clinical dat	ta	
Dual use re	esearch of conce	rn
Antibodies		
Antibodies used	NA	
Validation	NA	
Eukaryotic c	ell lines	
Policy information about <u>cell lines</u>		
Cell line source(s)	NA
Authentication		NA
Mycoplasma con	tamination	NA
Commonly misid		NA
(See <u>ICLAC</u> register	1	
Palaeontolo	gy and Ar	chaeology
Specimen prover	nance NA	

NA

Specimen deposition

Dating methods	NA	
Tick this box to confir	m that the raw and calibrated dates are available in the paper or in Supplementary Information.	
Ethics oversight	NA	
Note that full information on t	ne approval of the study protocol must also be provided in the manuscript.	
Animals and othe	r organisms	
Policy information about <u>st</u>	udies involving animals; ARRIVE guidelines recommended for reporting animal research	
Laboratory animals	NA	
Wild animals	NA	
Field-collected samples	NA	
Ethics oversight	NA	
Note that full information on t	ne approval of the study protocol must also be provided in the manuscript.	
Human research _l	participants	
Policy information about <u>st</u>	udies involving human research participants	
Population characteristic	S NA	
Recruitment	NA	
Ethics oversight	Approval for QoL assessment was obtained from the Ethics committee of the University of Dresden, Germany on June 12 2017 (EK 255062017), University of Zurich, Switzerland on April 20th 2015 (KEK-ZHNo: 2014-0631). The retrospective study on patients after pancreatectomy was approved by the TU Dresden Institutional Review Board (EK 310062019).	
Note that full information on t	ne approval of the study protocol must also be provided in the manuscript.	
	as approval of the staat, protects made also be promised in the manager pa	
Clinical data		
Policy information about <u>cl</u> All manuscripts should comply	nical studies with the ICMJE guidelines for publication of clinical research and a completed CONSORT checklist must be included with all submissions.	
Clinical trial registration	NA, retrospective analysis	
Study protocol	NA, retrospective analysis	
Data collection	Data were retrospectively collected from 2012 to 2018 in patients scheduled for pancreatic surgery at the Department of Visceral,-Thorax- and Vascular Surgery at the TU Dresden, Germany	
Outcomes	NA	
Dual use research	of concorn	
	ual use research of concern	
Hazards		
Could the accidental, deli in the manuscript, pose a	berate or reckless misuse of agents or technologies generated in the work, or the application of information presented threat to:	
No Yes		
Public health		
National security		
Crops and/or livestock		
Ecosystems		
Any other significant area		

Experiments of concer	n	
Does the work involve an	y of the	ese experiments of concern:
Confer resistance t Confer resistance t Enhance the virule Increase transmiss Alter the host rang Enable evasion of o	to theral nce of a libility of se of a p diagnost nization	
ChIP-seq		
Data deposition		
Confirm that both raw	v and fi	nal processed data have been deposited in a public database such as <u>GEO</u> .
Confirm that you have	e depos	sited or provided access to graph files (e.g. BED files) for the called peaks.
Data access links May remain private before public	cation.	NA
Files in database submiss	ion	NA
Genome browser session (e.g. <u>UCSC</u>)		NA
Methodology		
Replicates	NA	
Sequencing depth	NA	
Antibodies	NA	
Peak calling parameters	NA	
Data quality	NA NA	
Software	NA	
Flow Cytometry		
Plots		
Confirm that:		
The axis labels state the	he mar	ker and fluorochrome used (e.g. CD4-FITC).
		sible. Include numbers along axes only for bottom left plot of group (a 'group' is an analysis of identical markers).
		ith outliers or pseudocolor plots.
A numerical value for	numbe	er of cells or percentage (with statistics) is provided.
Methodology		
Sample preparation		(NA
Instrument	NA	
Software	NA	
Cell population abundance	ce	(NA

Gating strategy	NA
	a figure exemplifying the gating strategy is provided in the Supplementary Information.
Magnetic resonance in	maging
Experimental design	Tiaging
	NA .
Design type	NA
Design specifications	NA
Behavioral performance measur	es (NA
Acquisition	
Imaging type(s)	NA
Field strength	NA
Sequence & imaging parameters	5 NA
Area of acquisition	NA
Diffusion MRI Used	Not used ■ Not used
Preprocessing	
Preprocessing software	NA
Normalization	NA
Normalization template	NA
Noise and artifact removal	NA
Volume censoring	NA
Statistical modeling & infere	ence
Model type and settings	NA
Effect(s) tested	NA NA
Specify type of analysis: W	'hole brain ROI-based Both
Statistic type for inference (See <u>Eklund et al. 2016</u>)	NA
Correction	NA
Models & analysis	
n/a Involved in the study	
Functional and/or effective	e connectivity

n/a	Involved in the study
X	Functional and/or effective connectivity
X	Graph analysis
\boxtimes	Multivariate modeling or predictive analysis