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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 <u>Authors & Referees</u> and the <u>Editorial Policy Checklist</u>.

Statistics	
For all statistical	analyses, confirm that the following items are present in the figure legend, table legend, main text, or Methods section.
n/a Confirmed	
☐ ☐ The exa	act sample size $(n)$ for each experimental group/condition, given as a discrete number and unit of measurement
☐ X A state	ment on whether measurements were taken from distinct samples or whether the same sample was measured repeatedly
The sta	tistical test(s) used AND whether they are one- or two-sided nmon tests should be described solely by name; describe more complex techniques in the Methods section.
A descr	iption of all covariates tested
A descr	iption of any assumptions or corrections, such as tests of normality and adjustment for multiple comparisons
A full d	escription of the statistical parameters including central tendency (e.g. means) or other basic estimates (e.g. regression coefficient) riation (e.g. standard deviation) or associated estimates of uncertainty (e.g. confidence intervals)
For nul	hypothesis testing, the test statistic (e.g. $F$ , $t$ , $r$ ) with confidence intervals, effect sizes, degrees of freedom and $P$ value noted alues as exact values whenever suitable.
For Bay	resian analysis, information on the choice of priors and Markov chain Monte Carlo settings
For hie	rarchical and complex designs, identification of the appropriate level for tests and full reporting of outcomes
Estimat	tes 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 a	nd code
Policy information	on about <u>availability of computer code</u>
Data collection	Tissue samples were acquired during a warm autopsy, and frozen. Afterwards, DNA was extracted from the frozen samples and sequenced.
Data analysis	All analyses were performed using published open-source software (see Methods).
	ring custom algorithms or software that are central to the research but not yet described in published literature, software must be made available to editors/reviewers. ge code deposition in a community repository (e.g. GitHub). See the Nature Research guidelines for submitting code & software for further information.
Data	
All manuscripts - Accession co - A list of figur	on about <u>availability of data</u> must include a <u>data availability statement</u> . This statement should provide the following information, where applicable: des, unique identifiers, or web links for publicly available datasets es that have associated raw data of any restrictions on data availability
All raw data (BAM	files) were deposited in the SRA archive. All figures make use of these raw data.
Field-sp	ecific reporting
Please select the	one below that is the best fit for your research. If you are not sure, read the appropriate sections before making your selection.
✓ Life sciences	Behavioural & social sciences

## Life sciences study design

Life Scien	ices study design		
All studies must dis	close on these points even when the disclosure is negative.		
Sample size	No sample-size calculations were performed as this is a n=1 study. A total of 18 regional samples from a single colorectal cancer patient were analysed.		
Data exclusions	No samples were excluded from the analyses.		
Replication	not applicable.		
Randomization	not applicable.		
Blinding	not applicable.		
We require information	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 th  Antibodies			
Antibodies  Eukaryotic	ChIP-seq  Cell lines  Flow cytometry		
Palaeontolo			
	d other organisms		
	earch participants		
Clinical data			
<b>□</b>  □			
Antibodies			
Antibodies used	Anti-EpCAM (EBA1) antibody and DRAQ5 and 7ADD dyes		
Validation	Anti -EpCAM (EBA1) validation data link: http://www.bdbiosciences.com/ds/is/tds/23-5534.pdf DRAQ5 validation data link: https://www.thermofisher.com/order/catalog/product/62254 7AAD validation data link: https://www.thermofisher.com/order/catalog/product/00-6993-50		
Human resea	arch participants		
Policy information a	about <u>studies involving human research participants</u>		
Population charac	cteristics 1 colorectal cancer patient (age = 51)		
Recruitment	Written informed consent was provided by the patient's family.		
Ethics oversight	Sample collection was approved by a local ethics committee (CAEI Galicia 2014/015).		
Note that full informa	tion on the approval of the study protocol must also be provided in the manuscript.		
Flow Cutoma			
Flow Cytome	eury		
Plots			
Confirm that:	14 (22.5772)		
	s state the marker and fluorochrome used (e.g. CD4-FITC).		
	s are clearly visible. Include numbers along axes only for bottom left plot of group (a 'group' is an analysis of identical markers).		
	ontour plots with outliers or pseudocolor plots.		
A numerical v	alue for number of cells or percentage (with statistics) is provided.		

## Methodology

Sample preparation

Instrument

FACSAria III

Software

FACSDiva (BD Biosciences) and Flowlogic (Miltenyi Biotec) softwares were used to collect and analyze flow cytometry data.

Cell population abundance

EpCAM is expressed at a high level and frequency in colon cancer tissues (95%) and in most human adenocarcinomas.

The gating Strategy is shown in Supplementary Fig. 6

Tick this box to confirm that a figure exemplifying the gating strategy is provided in the Supplementary Information.