nature portfolio

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Reporting Summary

Provide your data availability statement here.

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

Statistics					
For all statistical ar	nalyses, confirm that the following items are present in the figure legend, table legend, main text, or Methods section.				
n/a Confirmed					
☐ ☐ The exact	act sample size (n) for each experimental group/condition, given as a discrete number and unit of measurement				
A stateme	A statement on whether measurements were taken from distinct samples or whether the same sample was measured repeatedly				
The statis Only comm	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 descrip	🔀 A description of all covariates tested				
A descrip	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 Give <i>P</i> values as exact values whenever suitable.					
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					
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 an	d code				
Policy information	about <u>availability of computer code</u>				
Data collection	N/A				
Data analysis	ata analysis N/A				
	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 Portfolio guidelines for submitting code & software for further information.				
Data					
All manuscripts m - Accession code - A description o	about <u>availability of data</u> nust include a <u>data availability statement</u> . This statement should provide the following information, where applicable: s, unique identifiers, or web links for publicly available datasets f any restrictions on data availability asets or third party data, please ensure that the statement adheres to our <u>policy</u>				

Human rese	arch part	icipants	
Policy information	about <u>studies</u>	involving human research participants and Sex and Gender in Research.	
Reporting on sex and gender		NOT APPLICABLE	
Population characteristics		NOT APPLICABLE	
Recruitment		NOT APPLICABLE	
Ethics oversight		NOT APPLICABLE	
Note that full information on the approval of the study protocol must also be provided in the manuscript.			
Field and	oific re	an orting	
Field-spe		·	
Life sciences		is the best fit for your research. If you are not sure, read the appropriate sections before making your selection. Behavioural & social sciences	
		all sections, see <u>nature.com/documents/nr-reporting-summary-flat.pdf</u>	
Life scier	nces st	udy design	
All studies must dis	close on these	e points even when the disclosure is negative.	
Sample size		composed of: control (n=9; standard chow), or chow with a 15% incorporation of pork mince (n=9), pork sausage (n=10), or ing frankfurter sausage (n=10).	
Data exclusions	Data were not	excluded.	
Replication		involved one replication but were longitudinal and where possible (e.g. blood/urine biomarkers, bodyweight etc.) repeated e taken at different time points.	
Randomization		re allocation to dietary groups at random. Tissue sections, blood, urine and feacal samples were analyses in a randomised order. counting, histology and biochemicals	
Blinding	Tumor counting and histology were conducted in a blinded manner and then unblinded for analysis. Metagenomic sequencing and mass spectrometry analysis took place in a blinded manner and were unblinded after quantitifications. Mouse groups and diets were not blinded.		
Reportin	σ for s	pecific materials, systems and methods	
We require information	on from authors	s 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	nerimental (systems Methods	
Materials & experimental systems n/a Involved in the study Methods n/a Involved in the study			
Antibodies ChIP-seq			
Eukaryotic		Flow cytometry	
Palaeontology and archaeology MRI-based neuroimaging			
Animals and other organisms Clinical data			
Dual use research of concern			
Antibodies			
Antibodies used	No individually purchased antibodies were used apart from a proprietary antibody within the malondialdehyde (MDA) ELISA kit supplied by Abcam, Cambridge, UK. The details of this antibody are not available.		

Validation

Not applicable.

Animals and other research organisms

Policy information about <u>studies involving animals</u>; <u>ARRIVE guidelines</u> recommended for reporting animal research, and <u>Sex and Gender in Research</u>

Laboratory animals	C57BL/6J-ApcMin
Wild animals	Not applicable.
Reporting on sex	All mice used were female to eliminate gender as a potential confounding factor.
Field-collected samples	Not applicable.
Ethics oversight	Studies in mice underwent Ethical Review and were performed in accordance with the UK Animals (Scientific Procedures) Act 1986.

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