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Reporting Summary

x Life sciences

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Sta	atistics					
For	all statistical analys	es, confirm that the following items are present in the figure legend, table legend, main text, or Methods section.				
n/a	Confirmed					
	x The exact sam	1 ple size (n) for each experimental group/condition, given as a discrete number and unit of measurement				
	🗶 A statement o	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.					
x	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)					
×	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>					
×	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					
x	Estimates of e	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.				
So	ftware and o	code				
Poli	cy information abo	ut <u>availability of computer code</u>				
Data collection		Structural Data Collection: XDS (Mar 2019), DIALS 1.14.5, XIA2 0.5653. HPLC and Mass Spec Data Collaction: Chromeleon Chromatography Data System Software, HyStar v3.2.				
Da	ata analysis	Graphpad Prism 8, Bruker Compass Data Analysis 4.4, GlycoWorkBench, ClustalOmega, IMG database, CAZy, SeaView, Pfam, SMART, Pymol 1.8.x, Pointless 1.11.19, Aimless 0.7.4, MolRep 11.6.04, Phaser2.8.2, Phenix 1.18.3855, Coot 0.8.9.1, CCP4 suite 7.0, Privateer MKIII				
		om algorithms or software that are central to the research but not yet described in published literature, software must be made available to editors/reviewers. deposition in a community repository (e.g. GitHub). See the Nature Research guidelines for submitting code & software for further information.				
Da	ta					
All	manuscripts must - Accession codes, un - A list of figures that - A description of any	ut <u>availability of data</u> include a <u>data availability statement</u> . This statement should provide the following information, where applicable: ique identifiers, or web links for publicly available datasets have associated raw data restrictions on data availability				
The	data that support the	e findings in this paper are available upon request from the corresponding authors.				
Fi	eld-speci	fic 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.

___ Ecological, evolutionary & environmental sciences

Life sciences study design

Sample size	No sample size calculation was performed. Non-kinetic enzyme assays were repeated at least once for each substrate tested with different enzyme preparations. Bacterial growths are averages of triplicate cultures and were repeated at least once.
Data exclusions	No data were excluded.
Replication	Bacterial growth experiments on different glycans were carried out in triplicate and the experiments repeated at least once. The non-kinetic assays on recombinant enzymes were reproduced multiple times with different enzyme preparations on each of the different substrates tested. Where possible, positive and negative controls for enzyme assays were used throughout to be sure of the accuracy of results and experimental conditions. LC-MS/MS experiments were repeated at least once independently and the labelled glycans from a single enzyme assay often analysed twice to ensure the data was reproducible and in some cases optimise detection. In all cases replicates were consistent and the data was analysed independently by two people (Dr Lucy Crouch and Paulina Urbanowicz).
Randomization	N/A
Blinding	N/A
Reportin	g for specific materials, systems and methods
Ve require informat	ion from authors about some types of materials, experimental systems and methods used in many studies. Here, indicate whether each materia ted 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.
Materials & ex	perimental systems Methods

Involved in the study

Flow cytometry

MRI-based neuroimaging

ChIP-seq

X

×

Human	research	partici	pants

Animals and other organisms

Human research participants

Policy information about <u>studies involving human research participants</u>

Population characteristics

Involved in the study

Eukaryotic cell lines

Palaeontology

Clinical data

Antibodies

X

X

X

N/A

Recruitment

For both the IBD and NEC samples, the patients were not chosen, but were just the next ones that required surgery after we had all the ethical approvals in place. Only a small number of patients were included as this was only a proof-of-principle study.

Ethics oversight Newcas

Newcastle and North Tyneside Research Ethics Committee 1 (REC:17/NE/0361) and SERVIS study (approvals 10/H0908/39 and 15-NE-0334)

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