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Reporting Summary

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Sta	tistics					
For a	ll statistical analyse	es, confirm that the following items are present in the figure legend, table legend, main text, or Methods section.				
n/a	/a Confirmed					
	The exact sam	The exact sample 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.					
	A description of all covariates tested					
	\square 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.					
\boxtimes	For Bayesian a	analysis, information on the choice of priors and Markov chain Monte Carlo settings				
\boxtimes	For hierarchica	al and complex designs, identification of the appropriate level for tests and full reporting of outcomes				
	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.				
Sof	tware and c	ode				
Polic	y information abou	ut <u>availability of computer code</u>				
Da	ta collection	n.a.				
Data analysis n.a.		n.a.				
For ma	anuscripts utilizing custo	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.				
Dat	ta					
All r - -	, manuscripts must i Accession codes, uni A list of figures that l	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				
micro	microbiota data will be available at public database xxx					
Fie	eld-speci	fic reporting				
Pleas	se select the one be	elow that is the best fit for your research. If you are not sure, read the appropriate sections before making your selection.				
$\sum L$	ife sciences	Behavioural & social sciences Ecological, evolutionary & environmental sciences				

For a reference copy of the document with all sections, see <u>nature.com/documents/nr-reporting-summary-flat.pdf</u>

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All studies must discl	ose on these points even when the disclosure is negative.				
Sample size	mentioned in paper				
Data exclusions n	mantioned in paper				
Replication	not appilcable				
Randomization	mentioned in paper				
Blinding	nont applicable				
We require information	for specific materials, systems and methods from authors about some types of materials, experimental systems and methods used in many studies. Here, indicate whether each material, it 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 & expe	erimental systems Methods				
n/a Involved in the	ChIP-seq				
Eukaryotic ce					
	other organisms				
Human resea	arch participants				
Human reseai	rch participants				
Policy information ab	out studies involving human research participants				
Population characte	eristics patients with metabolic syndrome,				
Recruitment	recruited by newspaper advertisement				
Ethics oversight	approved by AMC ethics committee / IRB				
Note that full information	on on the approval of the study protocol must also be provided in the manuscript.				
Clinical data					
Policy information ab All manuscripts should c	oout <u>clinical studies</u> comply with the ICMJE <u>guidelines for publication of clinical research</u> and a completed <u>CONSORT checklist</u> must be included with all submissions.				
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Data collection	collected during trial (see manuscript)				
Outcomes	NTR 4913 www.trialregister.nl				