# nature research

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Last updated by author(s): 9/25/2020

## 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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For	all st	atistical analyses, confirm that the following items are present in the figure legend, table legend, main text, or Methods section.
n/a	Cor	nfirmed
	X	The exact sample size (n) for each experimental group/condition, given as a discrete number and unit of measurement
	×	A statement 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
	x	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>
X		For Bayesian analysis, information on the choice of priors and Markov chain Monte Carlo settings
X		For hierarchical and complex designs, identification of the appropriate level for tests and full reporting of outcomes
X		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 and code

Policy information about <u>availability of computer code</u>

Data collection

No software was used.

Transcriptomics and proteomics data analysis and integration was performed using R/Bioconductor. The metabolic models were developed and analyzed using COBRA toolbox in Matlab 2019a.

For manuscripts utilizing 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 reviewers. We strongly 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

The models generated in this study are available as SBML files in the Supplementary Dataset 1 and can also be accessed from the biomodels database under accession MODEL1909260003, MODEL1909260004, MODEL1909260005, MODEL1909260006. The accession numbers of all publicly available datasets used in this study are provided in relevant sections of the manuscript and supplementary data.

Field-spe	ecific reporting	
Please select the c	one below that is the best fit for you	r research. If you are not sure, read the appropriate sections before making your selection.
Life sciences	Behavioural & social	sciences Ecological, evolutionary & environmental sciences
For a reference copy of	the document with all sections, see <u>nature.co</u>	om/documents/nr-reporting-summary-flat.pdf
Life scier	nces study desig	n
	isclose on these points even when t	
Sample size	We have used 4 different human perivalidation.	pheral blood mononuclear cells (PBMC) and we used paired T-test one tailed for our statistical
Data exclusions	Data excluded from the study are the replicate	one who showed signs of bacteria or fungi contamination. Contamination biased measurement of
Replication	To confirm distribution within replica reduce error of handling samples dur	tes, each sample was analyzed in duplicate or triplicate to obtain the mean of every sample and to ing experimental manipulation.
Randomization	PBMC from different donors from age	e-matched cells have been used and cells were split into untreated and treated conditions.
Blinding	The cohort chosen are the age-match	ned for human samples. This will allow for a homogenous distribution of samples during treatment.
Reportin	ng for specific ma	aterials, systems and methods
		naterials, experimental systems and methods used in many studies. Here, indicate whether each material, not sure if a list item applies to your research, read the appropriate section before selecting a response.
Materials & ex	operimental systems	Methods
n/a Involved in t	he study	n/a Involved in the study
Antibodie	S	ChIP-seq

Materials & experimental systems	Methods
n/a Involved in the study	n/a Involved in the study
Antibodies	ChIP-seq
Eukaryotic cell lines	Flow cytometry
Palaeontology and archaeology	MRI-based neuroimaging
Animals and other organisms	·
Human research participants	
Clinical data	
Dual use research of concern	

#### **Antibodies**

Antibodies used

anti-CD3 (clone OKT3, Biolegend) and anti-CD28 (clone CD28.2, Biolegend).

Validation

Describe the validation of each primary antibody for the species and application, noting any validation statements on the manufacturer's website, relevant citations, antibody profiles in online databases, or data provided in the manuscript.