nature research

Corresponding author(s):	FAN XIA
Last updated by author(s):	Dec 21, 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>.

_				
C-	-	Fic:	tica	•
_	_		111	•

For all statistic	al analyses, confirm that the following items are present in the figure legend, table legend, main text, or Methods section.
n/a Confirme	d
☐ X The €	xact sample size (n) for each experimental group/condition, given as a discrete number and unit of measurement
A sta	rement on whether measurements were taken from distinct samples or whether the same sample was measured repeatedly
I V I I I	tatistical test(s) used AND whether they are one- or two-sided ommon tests should be described solely by name; describe more complex techniques in the Methods section.
∑ A des	cription of all covariates tested
⊠ A des	cription of any assumptions or corrections, such as tests of normality and adjustment for multiple comparisons
	description of the statistical parameters including central tendency (e.g. means) or other basic estimates (e.g. regression coefficient) variation (e.g. standard deviation) or associated estimates of uncertainty (e.g. confidence intervals)
	ull 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 values as exact values whenever suitable.
For B	ayesian analysis, information on the choice of priors and Markov chain Monte Carlo settings
For h	erarchical and complex designs, identification of the appropriate level for tests and full reporting of outcomes
Estim	ates of effect sizes (e.g. Cohen's d , Pearson's r), indicating how they were calculated
ı	Our web collection on <u>statistics for biologists</u> contains articles on many of the points above.
Software	and code
Policy informa	tion about <u>availability of computer code</u>
Data collecti	on This is not relevant to our study.
Data analysi	This is not relevant to our study.
	illzing 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 ingly 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 <u>data availability statement</u>. 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

All data of our paper are available.

Field-spe	cific reporting
Please select the or	e 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 Ecological, evolutionary & environmental sciences
For a reference copy of the	ne document with all sections, see <u>nature.com/documents/nr-reporting-summary-flat.pdf</u>
Life scien	ces study design
All studies must disc	close on these points even when the disclosure is negative.
Sample size	NA
Data exclusions	NA
Replication	90% of the attempts are successful, the unsuccessful 10% are due to the quality of nanochannel fabrication and modification.
Randomization	The samples and participants in the work are randomly allocated.
Blinding	The investigators were blinded to group allocation during data collection.
We require informatic system or method list. Materials & exp. n/a Involved in the Antibodies Eukaryotic of Palaeontolo Animals and Human reso	ChIP-seq cell lines Flow cytometry gy and archaeology MRI-based neuroimaging d other organisms earch participants
•	bout <u>clinical studies</u>
·	comply with the ICMJE guidelines for publication of clinical research and a completed CONSORT checklist must be included with all submissions.
Clinical trial regist	Reference cohort (Xu et al. 2016, doi:10.1038/am.2015.138); urine specimens of 40 individual research participants: 11 female, 19 male; 10 normal individuals' urine samples, 15 diabetes patients before insulin injection or other drug therapy and 15 diabetes patients after insulin injection or other drug therapy.
Study protocol	Reference cohort (Xu et al. 2016, doi:10.1038/am.2015.138); urine specimens of 40 individual research participants: 11 female, 19 male; 10 normal individuals' urine samples, 15 diabetes patients before insulin injection or other drug therapy and 15 diabetes patients after insulin injection or other drug therapy.
Data collection	Reference cohort (Xu et al. 2016, doi:10.1038/am.2015.138); urine specimens of 40 individual research participants: 11 female, 19 male; 10 normal individuals' urine samples, 15 diabetes patients before insulin injection or other drug therapy and 15 diabetes

Reference cohort (Xu et al. 2016, doi:10.1038/am.2015.138); urine specimens of 40 individual research participants: 11 female, 19

male; 10 normal individuals' urine samples, 15 diabetes patients before insulin injection or other drug therapy and 15 diabetes

patients after insulin injection or other drug therapy.

patients after insulin injection or other drug therapy.

Outcomes