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## **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 Authors & Referees and the Editorial Policy Checklist.

Sta	atistics		
		ses, confirm that the following items are present in the figure legend, table legend, main text, or Methods section.	
n/a	Confirmed		
	The exact san	nple size (n) for each experimental group/condition, given as a discrete number and unit of measurement	
	A statement of	on whether measurements were taken from distinct samples or whether the same sample was measured repeatedly	
	The statistical Only common t	l test(s) used AND whether they are one- or two-sided rests should be described solely by name; describe more complex techniques in the Methods section.	
	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 descript AND variation	cion of the statistical parameters including central tendency (e.g. means) or other basic estimates (e.g. regression coefficient) in (e.g. standard deviation) or associated estimates of uncertainty (e.g. confidence intervals)	
	For null hypot	thesis testing, the test statistic (e.g. $F$ , $t$ , $r$ ) with confidence intervals, effect sizes, degrees of freedom and $P$ value noted sexact values whenever suitable.	
$\boxtimes$	For Bayesian	analysis, information on the choice of priors and Markov chain Monte Carlo settings	
$\times$	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		
	ı	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>	
D	ata collection	na	
D	ata analvsis	For gene-based and gene-set analyses we used MAGMA 1.06	

SNP heritability and genetic correlations were estimated using LD score regression (https://github.com/bulik/ldsc) and LD hub (http://ldsc.broadinstitute.org/).

eQTL for 44 GTEx tissues (https://gtexportal.org/)

PheWAS and drug target information (https://genetics.opentargets.org)

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/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 availability of data

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

Genotype and phenotype data from UKBB have already been processed and are available at https://www.ukbiobank.ac.uk/. The GWAS result from 23andMe are available by request from https://www.23andme.com/. Data from QSKIN are available on request from project leader Professor David Whiteman david.whiteman@qimrberghofer.edu.au.

Field-spe	ecific reporting			
Please select the o	ne 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 document with all sections, see <u>nature.com/documents/nr-reporting-summary-flat.pdf</u>			
Life scier	nces study design			
All studies must disclose on these points even when the disclosure is negative.				
Sample size	Rather than generate a sample size calculation we assembled the largest possible GWAS dataset to identify novel loci. This approach was successful as we identified numerous novel loci.			
Data exclusions	Samples excluded during GWAS quality control, including for ancestry, data quality and relatedness, are described in full in the methods section.			
Replication	tion Data was collected from multiple independent cohorts.			
Randomization	Samples were derived from existing cohorts and were not randomized.			
Blinding	GWAS cleaning and quality control was performed blind to the phenotypes.			
Reporting for specific materials, systems and methods				
We require information from authors about some types of materials, experimental systems and methods used in many studies. Here, indicate whether each material, system or method listed 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	Materials & experimental systems Methods			
n/a Involved in th	ne study n/a Involved in the study			
Antibodies	ChIP-seq			
Eukaryotic	cell lines			
Palaeontol	logy MRI-based neuroimaging			
Animals and other organisms				
Human res	Human research participants			
Clinical data				
Human research participants				

Policy information about  $\underline{\text{studies involving human research participants}}$ 

Population characteristics

Recruitment

Described in full in the methods section.

Ethics oversight

Described in full in the methods section.

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