nature portfolio

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Reporting Summary

Nature Portfolio 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 Portfolio policies, see our <u>Editorial Policies</u> and the <u>Editorial Policy Checklist</u>.

		ics

For	all statistical analyses, confirm that the following items are present in the figure legend, table legend, main text, or Methods section.
n/a	Confirmed
	\square 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)
\boxtimes	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>
\boxtimes	For Bayesian analysis, information on the choice of priors and Markov chain Monte Carlo settings
\boxtimes	For hierarchical and complex designs, identification of the appropriate level for tests and full reporting of outcomes
\boxtimes	Estimates of effect sizes (e.g. Cohen's <i>d</i> , Pearson's <i>r</i>), 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 availability of computer code

Data collection

ECG sensor equipped with an integrated ADS1292R signal acquisition converter, STM32F103C8T6 microcontroller, and Bluetooth transmission device; Neurosky TGAM EEG03 for EEG signal recording; ZTEMG-1100 PCB for EMG signal recording

Data analysis

No software was used

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 Portfolio 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 description of any restrictions on data availability
- For clinical datasets or third party data, please ensure that the statement adheres to our policy

The authors declare that the data supporting the findings of this study are available within the main article and the Supplementary Information. Source data are provided with this paper.

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Policy information a and sexual orientation		vith <u>human participants or human data</u> . See also policy information about <u>sex, gender (identity/presentation),</u> thnicity and racism.				
Reporting on sex and gender		In this work, human volunteers participated in physiological signal collection experiments after providing informed consent. The tests were taken on participates' skin. Gender was not taken into account as a variable.				
Reporting on race, ethnicity, or other socially relevant groupings		There is no socially constructed or socially relevant categorization variable in our manuscript.				
Population charac	teristics	There is no population characteristic relevant data.				
Recruitment		The participants in this work are the authors of this article.				
Ethics oversight		(Identify the organization(s) that approved the study protocol.				
Note that full informat	ion on the appro	oval of the study protocol must also be provided in the manuscript.				
Field-spe	cific re	porting				
Please select the on-	e below that is	s the best fit for your research. If you are not sure, read the appropriate sections before making your selection.				
Life sciences	В	ehavioural & social sciences				
For a reference copy of th	ne document with a	all sections, see <u>nature.com/documents/nr-reporting-summary-flat.pdf</u>				
Life scien	ces stu	udy design				
All studies must disc	close on these	points even when the disclosure is negative.				
Sample size	N/A					
Data exclusions	There is no data exclusion.					
Replication	We confirm the	experimental findings are reproducible.				
Randomization N/A						
Blinding	N/A					
		Decific materials, systems and methods about some types of materials, experimental systems and methods used in many studies. Here, indicate whether each material,				
		your study. If you are not sure if a list item applies to your research, read the appropriate section before selecting a response.				
Materials & exp	erimental s	ystems Methods				
						
Antibodies	шь	ChiP-seq				
	Eukaryotic cell lines Control C					
Clinical data	ı					
Dual use research of concern						
Plants						

Plants

Seed stocks

Report on the source of all seed stocks or other plant material used. If applicable, state the seed stock centre and catalogue number. If plant specimens were collected from the field, describe the collection location, date and sampling procedures.

Novel plant genotypes

Describe the methods by which all novel plant genotypes were produced. This includes those generated by transgenic approaches, gene editing, chemical/radiation-based mutagenesis and hybridization. For transgenic lines, describe the transformation method, the number of independent lines analyzed and the generation upon which experiments were performed. For gene-edited lines, describe the editor used, the endogenous sequence targeted for editing, the targeting guide RNA sequence (if applicable) and how the editor was applied.

Describe any authentication procedures for each seed stock used or novel genotype generated. Describe any experiments used to

Authentication

Describe any authentication procedures for each seed stock used or novel genotype generated. Describe any experiments used to assess the effect of a mutation and, where applicable, how potential secondary effects (e.g. second site T-DNA insertions, mosiacism, off-target gene editing) were examined.