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>.

Statistics

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		
	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	
×		Th e statistical test (s) u sed AND whether they are on e- 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)	
X		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 freed om and <i>P</i> value noted Give <i>P</i> values as exact values whenever suitable.	
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	
×		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 availability of computer code					
Datacollection	LCR instrument was used for of-body sensor data collection.				
Dataanalysis	Origin 2023 was used to analyze a data, p lot data and calculate the statistical parameters.				

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 man uscripts 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 thes tatement adheres to our policy

The data that support the finding of the study are available in this article and supplementary materials. All raw and analyzed datasets generated during the study are available from the corresponding authors on request.

Policy information about studies with human participants or human data. See also policy information about sex, gender (id entity/presentation), and sexual orientation and race, ethnicity and racism

Reporting on sex and gender	Findings and reports are not involved with sex and gender.	
Reporting on race, ethnicity, or other socially relevant group ings	Race, ethnicity, or other socially relevant groupings were not be categorized in this paper.	
Population characteristics	Healthy volunteers aged 20 to 30 years old were selected.	
Recruitment	The participants were recruited from the southern University of Science and Technology through advertisement by posted notices. They were no-self-selection biases or other biases. All participants gave written informed consent before participation in this study.	
Ethics oversight	Southern University of Science and Technology, SUSTech In stitutional Review Board.	

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

Field-specific reporting

Please select the one 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 nature.com/documents/nr-reporting-summary-flatpdf

Life sciences study design

All studies must disclose on these points even when the disclosure is negative.

Sample size	For on body evaluation of the wearable sensor, 2 subjects (1 female subject and 1 male subject) at the age range from 20 to 30 years old were involved.
Data exclusions	No data exclusions.
Replication	The experimental data are obtained from the results of at least three parallel experiments.
Randomization	The device was fabricated with same process and was tested in a participants under same conditions Randomization was therefore not relevant to the study.
Blinding	Not relevant, because a blinding process wound't influence the sampling results.

Reporting for specific materials, systems and methods

We require in formation 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.

Ma	tenals & experimental systems	Met
n/a	Involved in the study	n/a I
×	Antibodies	×
×	Eu karyotic cell lines	×
×	Palaeontology and archaeology	×
×	Animals and other organisms	
×	Clinical data	
×	Dual use research of concern	
×	Plants	

Methods

- /a Involved in the study
- X ChIP-s eq
- Flow cytometry
- 🗙 📄 MRI-based neuroimaging

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 tho segenerated by transgenic approaches, gene editing, chemical/radiation-based muta genesis 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
Authentication	applied. Describe any authentication procedures for each seed stock used or no velgenotype 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.