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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 our <u>Editorial Policies</u> and the <u>Editorial Policy Checklist</u>.

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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
	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.
	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 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>
	For Bayesian analysis, information on the choice of priors and Markov chain Monte Carlo settings
	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.
So	ftware and code
Poli	cy information about availability of computer code

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

Data collection

Data were collected by the Allen Institute Brain Observatory.

Data analysis

Data were analyzed using the Brain Observatory SDK, Sci-Kit Learn, and custom written Python code that is available at github.com/ achristensen56. Data were analyzed using python 3.7.9, allensdk 2.3.2, matplotlib 3.3.2, pandas 0.25.3, scikit-learn 0.23.2, scipy: 1.5.2, seaborn: 0.11.0

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 <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

Data is freely available via the Allen Institute Brain Observatory.

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 nr-reporting-summary-flat.pdf
Life scier	nces study design
All studies must dis	sclose on these points even when the disclosure is negative.
Sample size	No sample size calculation was done. We used all relevant data from the publicly available dataset we analyzed and effect sizes were estimated via bootstrapping.
Data exclusions	We excluded mice who did not have at least 50 trials of running and 50 trials of stationary data from our analysis. This was to ensure we had adequate sampling of both running and stationary periods across all stimuli presented. The numbers of excluded datasets are below:
	VISpm (9 / 21 experiments used) VISp (8 / 24 experiments used) VISal(8 / 20 experiments used) VISI (12 / 23 experiments used) VISrl (3 / 6 experiments used) VISam (7 /15 experiment used)
Replication	We did not independently replicate this study. The data were collected as a part of a large scale effort from the Allen Institute for Brain Science, and not by us. We do not have the ability to replicate the experimental paradigm.
Randomization	Data were randomly sub-sampled for experimental groups to create even numbers of data points in each group. We did not e.g. randomize subjects into experimental conditions because the experimental conditions were self-generated, to be concrete the mice chose when they run, not us. We balanced data for covariates like which stimuli class was presented by sub-sampling to ensure all classes were present in both running and stationary datasets, to control for the possibility mice chose to ran more during some stimuli than others.
	No blinding was performed. We did not perform the data collection, or any experiments where subjective results were reported.

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		
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