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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 Editorial Policies and the Editorial Policy Checklist.

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Fora	all statistical an	alyses, 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 stateme	ent on whether measurements were taken from distinct samples or whether the same sample was measured repeatedly		
	The statist	atistical test(s) used AND whether they are one- or two-sided mmon tests should be described solely by name; describe more complex techniques in the Methods section.		
	A descript	A description of all covariates tested		
	A descript	cription of any assumptions or corrections, such as tests of normality and adjustment for multiple comparisons		
	A full desc	escription of the statistical parameters including central tendency (e.g. means) or other basic estimates (e.g. regression coefficient) riation (e.g. standard deviation) or associated estimates of uncertainty (e.g. confidence intervals)		
	For null hy Give P value	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>P values as exact values whenever suitable</i> .		
	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 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 <u>availability of computer code</u>				
Da	ta collection	Animal trajectories are collected with home made labview program		
Da	ata analysis	Data are analysed using Python 3.8.5		

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

R version 4.1.0

- For clinical datasets or third party data, please ensure that the statement adheres to our $\underline{\text{policy}}$

Data Availability: Source data are provided with this paper. All data supporting the findings of this study are available from the corresponding author upon reasonable request.

Code Availability: Animal's trajectories are collected with homemade labview program (version 2014). Results were generated using code written in R (version 4.1.0) and Python (version 3.8.5). All codes used to run the analysis are available from the authors upon request. A sample code for the model and archetype is

publicly available at https://zenodo.org/record/5596424#.YXfBAy8ivgY				
Field-spe	cific reporting			
	ne below that is the best fit for your research. If you are not sure, read the appropriate sections before making your selection.			
X Life sciences	Behavioural & social sciences Ecological, evolutionary & environmental sciences			
For a reference copy of t	he document with all sections, see <u>nature.com/documents/nr-reporting-summary-flat.pdf</u>			
Lite scier	nces study design			
All studies must dis	close on these points even when the disclosure is negative.			
Sample size	No sample size calculation were performed. Suitable sample sizes were estimated based on previous experiments (Naudé et al Nat Neuro 2016) and are similar to those employed in the field.			
Data exclusions	the that did not learn the task correctly during the deterministic session were excluded (i.e. they did not self-stimulate above baseline spite increasing stimulation intensity, with a maximum of 200 uA, see Carlezon & Chartoff Nat. Protocols 2007). For the probabilistic sions, mice self-stimulating at least 35 times in a 5 minutes session were included for the behavioral analysis. We did not exclude animals ng statistical outliers.			
Replication	All experiments were replicated with sucess. All experiments are at least be replicated twice. The main experiments has been replicated more than 5 times.			
Randomization	The probability associated with each target in the probabilistic session was pseudorandomly assigned at the beginning of the probabilistic session and kept constant for each animal. We used C57b6 male mice. Mice that received saline or nicotine where chosen pseudorandomly. For optogenetic experiments, all mice were male DATiCre transgenic mice. Mice that received GFP, Jaws or Catch were chosen pseudorandomly. IFor each experiment the same number of variables were used and analysed.			
Blinding	v data collection is managed by the software (closed-loop reward delivery upon videotracking), without any investigators action. So ling is not an issue.			
Reportin	g for specific materials, systems and methods			
We require information	on from authors about some types of materials, experimental systems and methods used in many studies. Here, indicate whether each material, ed 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 & exp	perimental systems Methods			
n/a Involved in th	e study n/a Involved in the study			
Antibodies				
Eukaryotic				
Palaeontology and archaeology MRI-based neuroimaging				
Animals and other organisms				
Human research participants Clinical data				
	esearch of concern			
Antibodies				
Antibodies used	Mouse anti-TH (Sigma); Cy3 anti-mouse (Jackson ImmunoResearch); Chicken anti-YFP (Life technologies Molecular Probes) and AlexaFluor488 anti-chicken (Jackson ImmunoResearch)			
Validation	(Antibody) Ab1: Mouse anti-TH https://www.sigmaaldrich.com/FR/fr/product/sigma/t1299			
	(Antibody) Ab2: Cy3 anti-mouse No antibody was detected against non-immunoglobulin serum proteins. See details here: https://www.jacksonimmuno.com/catalog/products/715-165-150			
	(Antibody) Ab1: Chicken anti-YFP This Antibody was verified by Relative expression to ensure that the antibody binds to the antigen stated. View Details here: https://www.thermofisher.com/antibody/product/GFP-Antibody-Polyclonal/A-6455			

(Antibody) Ab2: AlexaFluor488 anti-chicken

No antibody was detected against non-immunoglobulin serum proteins. See details here: https://www.jacksonimmuno.com/catalog/products/711-225-152

Animals and other organisms

Policy information about studies involving animals; ARRIVE guidelines recommended for reporting animal research

Laboratory animals Male C57BL/6J mouse, and C57Bl/6Rj DATiCRE. 8-16 weeks old.

Wild animals no wild animals were used in the study

Field-collected samples | no field collected sample were used in the study

Ethics oversight Experiments were conducted at Sorbonne University, Paris, France, in accordance with the local

regulations for animal experiments as well as the recommendations for animal experiments issued by the

European Council (directives 219/1990 and 220/1990). The protocol was approved by the Committee on the Ethics of Animal

Experiments and the Ministry of Higher Education and Research (permit 01438.02)

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