nature portfolio

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Last updated by author(s):	Nov 14, 2022

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>.

For all statistical analyses, confirm that the following items are present in the figure legend, table legend, main text, or Methods section.

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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 Give <i>P</i> values as exact values whenever suitable.				
\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				
	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	Software and code				
Poli	Policy information about <u>availability of computer code</u>				
Da	ata collection	Olympus Fluoview program			
Da	ata analysis	Matlab 2022, https://github.com/moustaam0/Algamal2022_analysis_w_OASIS, Graphpad Prism 9.0			
Forn	nanuscripts utilizing	g 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			

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:

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.

- 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

All data and code used to draw this article's conclusions are publicly available or presented in the paper and supplementary material. Time series data from all recordings stored as Matlab files are available at https://figshare.com/articles/dataset/Matlab_source_files_for_Algamal_et_al_2020/19740019. The code is available at https://github.com/moustaam0/Algamal2022_analysis_w_OASIS.

Human rese	arch part	icipants	
Policy information	about <u>studies</u> i	involving human research participants and Sex and Gender in Research.	
Reporting on sex	and gender	NA	
Population chara	acteristics	NA	
Recruitment		NA	
Ethics oversight		NA	
Note that full informa	ation on the app	roval of the study protocol must also be provided in the manuscript.	
E. 1.1			
Field-spe			
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	
For a reference copy of	the document with	all sections, see <u>nature.com/documents/nr-reporting-summary-flat.pdf</u>	
Life scier	nces st	udy design	
All studies must dis	sclose on these	points even when the disclosure is negative.	
Sample size	Sample sizes were decided based on on information from previous in vivo calcium imaging studies.		
Data exclusions	NA		
Replication		-SOM-Cre and 8 APP/PS1-SOM-Cre mice for SOM recordings, 5 WT-PV-Cre and 6 APP/PS1-PV-Cre mice for PV recordings, and 9 Cre and 4 WT-SOM-Cre mice) and 9 APP/PS1 (5 APP/PS1-PV-Cre and 4 APP/PS1-SOM-Cre mice) mice for excitatory neurons	
Randomization	No randomizat	cion.	
Blinding	Experimenters	were not blinded.	
We require informati	ion from authors	pecific materials, systems and methods about some types of materials, experimental systems and methods used in many studies. Here, indicate whether each material, o 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	nerimentals	systems Methods	
n/a Involved in th	•	n/a Involved in the study	
Antibodies	•	ChIP-seq	
Eukaryotic cell lines		Flow cytometry	
	logy and archaec		
	nd other organisr		
Clinical dat	-		
	esearch of conce	orn	
	escaron or conce	···	
Antibodies			
Antibodies used	anti-P	V (Sigma, P3088), anti-SOM (Millipore Sigma, MAB354), anti-GFP (CiteAb, A10262)	

Validation

As described on the company's website.

	ell lines	and Sex and Gender in Research
Cell line source(s)		NA
Authentication	NA	
Mycoplasma contaminat	nation NA	
Commonly misidentified (See <u>ICLAC</u> register)	lines	NA NA
alaeontology an	nd Arc	chaeology
Specimen provenance	NA	
Specimen deposition	NA	
Dating methods	NA	
Tick this box to confir	rm that t	the raw and calibrated dates are available in the paper or in Supplementary Information.
	NA	
		oval of the study protocol must also be provided in the manuscript.
nimals and other of the other o	er resetudies in	earch organisms avolving animals; ARRIVE guidelines recommended for reporting animal research, and Sex and Gender in L(cre)Zjh/J (Stock sed#013044, The Jackson Laboratory), B6;129P2-Pvalbtm1(cre)Arbr/J (Stock # 008069, The Jackson tory), and B6;C3-Tg(APPswe,PSEN1dE9)85Dbo/Mmjax (Stock #034829, The Jackson Laboratory). Males and Females were
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Clinical trial registration NA NA Study protocol NA Data collection NA Outcomes

Dual use research of concern

Policy information about <u>dual use research of concern</u>

Hazards

Could the accidental, deliberate or reckless misuse of agents or technologies generated in the work, or the application of information presented in the manuscript, pose a threat to:

No Yes Public health National security				
Crops and/or livestock				
Ecosystems				
Any other significant area				
Experiments of concer				
Does the work involve an	ny of the	ese experiments of concern:		
No Yes				
Demonstrate how	Demonstrate how to render a vaccine ineffective			
Confer resistance	to therap	peutically useful antibiotics or antiviral agents		
Enhance the virule	ence of a	pathogen or render a nonpathogen virulent		
Increase transmiss	sibility of	a pathogen		
Alter the host rang	ge of a pa	athogen		
Enable evasion of	diagnosti	ic/detection modalities		
Enable the weapon	nization (of a biological agent or toxin		
Any other potentia	ally harm	ful combination of experiments and agents		
ChIP-seq				
Data deposition				
	w and fir	nal processed data have been deposited in a public database such as <u>GEO</u> .		
Confirm that you have	e depos	ited or provided access to graph files (e.g. BED files) for the called peaks.		
Data access links May remain private before publi	ication.	NA		
Files in database submiss	sion	NA		
Genome browser session (e.g. <u>UCSC</u>)	Genome browser session (e.g. UCSC)			
Methodology				
Replicates	NA			
Sequencing depth	NA			
Antibodies	Antibodies NA			
Peak calling parameters NA				
Data quality	Data quality NA			
Software				
Flow Cytometry				
Plots				
Confirm that:				
	he mark	ker and fluorochrome used (e.g. CD4-FITC).		
		ible. Include numbers along axes only for bottom left plot of group (a 'group' is an analysis of identical markers).		
		th outliers or pseudocolor plots.		
A numerical value for	numbe	r of cells or percentage (with statistics) is provided.		

Methodology			
Sample preparation	NA		
Instrument	NA		
Software	NA		
Cell population abundance	NA		
Gating strategy	NA		
Tick this box to confirm that a	a figure exemplifying the gating strategy is provided in the Supplementary Information.		
Magnetic resonance ir	maging		
Experimental design			
Design type	NA		
Design specifications	NA		
Behavioral performance measure	es NA		
Acquisition			
Imaging type(s)	NA		
Field strength	NA		
Sequence & imaging parameters	NA NA		
Area of acquisition	NA		
Diffusion MRI Used	☐ Not used		
Preprocessing			
Preprocessing software	NA		
Normalization	NA		
Normalization template	NA		
Noise and artifact removal	NA NA		
Volume censoring	NA NA		
Statistical modeling & inference			
Model type and settings			
Effect(s) tested			
Specify type of analysis: Whole brain ROI-based Both			
Statistic type for inference (See <u>Eklund et al. 2016</u>)	Statistic type for inference NA		
Correction	NA		
Models & analysis			
n/a Involved in the study			
Functional and/or effective connectivity			
Graph analysis			
Multivariate modeling or predictive analysis			

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Functional and/or effective connectivity	NA
Graph analysis	NA
Multivariate modeling and predictive analysis	NA