# nature neuroscience

Corresponding Author:	Aaron D. Gitler	# Main Figures:	1
Manuscript Number:	NN-BC51241B	# Supplementary Figures:	4
Manuscript Type:	Brief Communication	# Supplementary Tables:	0
		# Supplementary Videos:	0

## Reporting Checklist for Nature Neuroscience

This checklist is used to ensure good reporting standards and to improve the reproducibility of published results. For more information, please read Reporting Life Sciences Research.

Please note that in the event of publication, it is mandatory that authors include all relevant methodological and statistical information in the manuscript.

### ▶ Statistics reporting, by figure

- Please specify the following information for each panel reporting quantitative data, and where each item is reported (section, e.g. Results, & paragraph number).
- Each figure legend should ideally contain an exact sample size (n) for each experimental group/condition, where n is an exact number and not a range, a clear definition of how n is defined (for example x cells from x slices from x animals from x litters, collected over x days), a description of the statistical test used, the results of the tests, any descriptive statistics and clearly defined error bars if applicable.
- · For any experiments using custom statistics, please indicate the test used and stats obtained for each experiment.
- Each figure legend should include a statement of how many times the experiment shown was replicated in the lab; the details of sample collection should be sufficiently clear so that the replicability of the experiment is obvious to the reader.
- For experiments reported in the text but not in the figures, please use the paragraph number instead of the figure number.

Note: Mean and standard deviation are not appropriate on small samples, and plotting independent data points is usually more informative. When technical replicates are reported, error and significance measures reflect the experimental variability and not the variability of the biological process; it is misleading not to state this clearly.

		TEST US	TEST USED n		DESCRIPTIVE STATS (AVERAGE, VARIANCE)		P VALUE		DEGREES OF FREEDOM & F/t/z/R/ETC VALUE			
	FIGURE NUMBER	WHICH TEST?	SECTION & PARAGRAPH #	EXACT VALUE	DEFINED?	SECTION & PARAGRAPH #	REPORTED?	SECTION & PARAGRAPH #	EXACT VALUE	SECTION & PARAGRAPH #	VALUE	SECTION & PARAGRAPH #
example	1a	one-way ANOVA	Fig. legend	9, 9, 10, 15	mice from at least 3 litters/group	Methods para 8	error bars are mean +/- SEM	Fig. legend	p = 0.044	Fig. legend	F(3, 36) = 2.97	Fig. legend
example	results, para 6	unpaired t- test	Results para 6	15	slices from 10 mice	Results para 6	error bars are mean +/- SEM	Results para 6	p = 0.0006	Results para 6	t(28) = 2.808	Results para 6
+	1h	unpaired Student's t- test	Fig. legend	6	6 separate wells	Methods para 6	error bars are mean +/- SEM	Fig. legend	p = 0.0057, p = 0.0024	Fig. legend		

		TEST US	SED	n		DESCRIPTIVE STATS (AVERAGE, VARIANCE)		P VALUE		DEGREES OF FREEDOM & F/t/z/R/ETC VALUE		
	FIGURE NUMBER	WHICH TEST?	SECTION & PARAGRAPH #	EXACT VALUE	DEFINED?	SECTION & PARAGRAPH#	REPORTED?	SECTION & PARAGRAPH #	EXACT VALUE	SECTION & PARAGRAPH#	VALUE	SECTION & PARAGRAPH #
+	1j	unpaired Student's t- test	Fig. legend	13 controls, 8 ALS	iN cells from patient (4 cells per patient)	Fig. legend	error bars are mean +/- SEM	Fig. legend	p = 0.0032	Fig. legend		

#### ▶ Representative figures

1.	Are any representative images shown (including Western blots and
	immunohistochemistry/staining) in the paper?

If so, what figure(s)?

2. For each representative image, is there a clear statement of how many times this experiment was successfully repeated and a discussion of any limitations in repeatability?

If so, where is this reported (section, paragraph #)?

Fig 1a, 1b, 1d, 1f, 1i

Yes

For 1a, 1b, 1d and 1f the statement is in Methods, paragraph 3 For 1i the statement in in Methods, paragrap 7

# ▶ Statistics and general methods

1. Is there a justification of the sample size?

If so, how was it justified?

Where (section, paragraph #)?

Even if no sample size calculation was performed, authors should report why the sample size is adequate to measure their effect size.

2. Are statistical tests justified as appropriate for every figure?

Where (section, paragraph #)?

- a. If there is a section summarizing the statistical methods in the methods, is the statistical test for each experiment clearly defined?
- b. Do the data meet the assumptions of the specific statistical test you chose (e.g. normality for a parametric test)?

Where is this described (section, paragraph #)?

No, but statistically significant results are presented for each experiment.

Yes

For Fig 1h the statistical test is defined in the figure legend. For Fig 1j the statistical test is defined in the figure legend.

Yes

	C.	Is there any estimate of variance within each group of $$ data?	Standard error bars are reported in Fig 1h, j
		Is the variance similar between groups that are being statistically compared?	
		Where is this described (section, paragraph #)?	
	d.	Are tests specified as one- or two-sided?	two-sided T-test for Fig 1h, j
	e.	Are there adjustments for multiple comparisons?	N/A
3.	Are crite	ria for excluding data points reported?	N/A
	Was this	criterion established prior to data collection?	
	Where is	this described (section, paragraph #)?	
4.	samples)	ne method of randomization used to assign subjects (or to the experimental groups and to collect and process data.	N/A
	If no rand	domization was used, state so.	
	Where d	oes this appear (section, paragraph #)?	
5.		ement of the extent to which investigator knew the group n during the experiment and in assessing outcome included?	Samples were blinded before image taking and analysis by an independent investigator. Only after quantification for each sample was done, the identities were reviled.
	If no blin	ding was done, state so.	was done, the identities were reviied.
	Where (s	section, paragraph #)?	Method section, paragraphs 6 and 7
6.		riments in live vertebrates, is a statement of compliance with uidelines/regulations included?	N/A
	Where (s	section, paragraph #)?	
7.		ecies of the animals used reported?	Method section, paragraphs 6
	where (s	section, paragraph #)?	
8.		ain of the animals (including background strains of KO/ic animals used) reported?	N/A
	Where (s	section, paragraph #)?	
9.	Is the sex	x of the animals/subjects used reported?	N/A
٥.		section, paragraph #)?	
	where (s	section, paragraph #):	
10.	Is the ag	e of the animals/subjects reported?	N/A
	Where (s	section, paragraph #)?	
11.	For anim	als housed in a vivarium, is the light/dark cycle reported?	N/A
	Where (s	section, paragraph #)?	

	For animals housed in a vivarium, is the housing group (i.e. number of animals per cage) reported?	N/A
	Where (section, paragraph #)?	
	For behavioral experiments, is the time of day reported (e.g. light or dark cycle)?	N/A
	Where (section, paragraph #)?	
	Is the previous history of the animals/subjects (e.g. prior drug administration, surgery, behavioral testing) reported?	N/A
	Where (section, paragraph #)?	
	a. If multiple behavioral tests were conducted in the same group of animals, is this reported?	N/A
	Where (section, paragraph #)?	
15.	If any animals/subjects were excluded from analysis, is this reported?	N/A
	Where (section, paragraph #)?	
	How were the criteria for exclusion defined?	N/A
	Where is this described (section, paragraph #)?	,
	<ul> <li>Specify reasons for any discrepancy between the number of animals at the beginning and end of the study.</li> </ul>	N/A
	Where is this described (section, paragraph #)?	
<b>.</b>	Doogonto	
<u> </u>	Reagents	
	Have antibodies been validated for use in the system under study (assay and species)?	Yes
	a. Is antibody catalog number given?	Method section, paragraphs 4, 6, 7
	Where does this appear (section, paragraph #)?	
	<ul> <li>b. Where were the validation data reported (citation, supplementary information, Antibodypedia)?</li> </ul>	Antibody catalog information, found with the catalog number reported in Methods, paragraph 4, 6, 7
	Where does this appear (section, paragraph #)?	
2	Call line identity	NO
2.	•	NO
	a. Are any cell lines used in this paper listed in the database of commonly misidentified cell lines maintained by <u>ICLAC</u> and <u>NCBI Biosample</u> ?	

Where (section, paragraph #)?

b.	If yes, include in the Methods section a scientific
	justification of their useindicate here in which section and
	paragraph the justification can be found.

N/A	

- c. For each cell line, include in the Methods section a statement that specifies:

  - have the cell lines been authenticated? If so, by which
  - have the cell lines been tested for mycoplasma contamination?

Where (section, paragraph #)?

- the source of the cell lines

N/A			

### Data deposition

Data deposition in a public repository is mandatory for:

- a. Protein, DNA and RNA sequences
- b. Macromolecular structures
- c. Crystallographic data for small molecules
- d. Microarray data

Deposition is strongly recommended for many other datasets for which structured public repositories exist; more details on our data policy are available here. We encourage the provision of other source data in supplementary information or in unstructured repositories such as Figshare and Dryad.

We encourage publication of Data Descriptors (see Scientific Data) to maximize data reuse.

1. Are accession codes for deposit dates provided?

Where (section, paragraph #)?



### ▶ Computer code/software

Any custom algorithm/software that is central to the methods must be supplied by the authors in a usable and readable form for readers at the time of publication. However, referees may ask for this information at any time during the review process.

1. Identify all custom software or scripts that were required to conduct the study and where in the procedures each was used.

. / ^			
I/A			

2. If computer code was used to generate results that are central to the paper's conclusions, include a statement in the Methods section under "Code availability" to indicate whether and how the code can be accessed. Include version information as necessary and any restrictions on availability.

I/A			

### Human subjects

1. Which IRB approved the protocol?

Where is this stated (section, paragraph #)?

N/A			

2.	Is demographic information on all subjects provided?	N/A		
	Where (section, paragraph #)?			
3.	Is the number of human subjects, their age and sex clearly defined?	N/A		
	Where (section, paragraph #)?			
4.	Are the inclusion and exclusion criteria (if any) clearly specified?	N/A		
	Where (section, paragraph #)?			
_		(		
5.	How well were the groups matched?	N/A		
	Where is this information described (section, paragraph #)?			
6	Is a statement included confirming that informed consent was	N/A		
0.	obtained from all subjects?	19/74		
	Where (section, paragraph #)?			
7.	For publication of patient photos, is a statement included confirming	N/A		
	that consent to publish was obtained?			
	Where (section, paragraph #)?			
<b>.</b> 4	EN ADL atuation			
▶ fMRI studies				
	iviki studies			
For	r papers reporting functional imaging (fMRI) results please ensure that the primation is clearly provided in the methods:	ese minimal reporting guidelines are met and that all this		
For	r papers reporting functional imaging (fMRI) results please ensure that th ormation is clearly provided in the methods:			
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For	r papers reporting functional imaging (fMRI) results please ensure that the properties of the provided in the methods:  Were any subjects scanned but then rejected for the analysis after the data was collected?  a. If yes, is the number rejected and reasons for rejection			
For	r papers reporting functional imaging (fMRI) results please ensure that the properties of the provided in the methods:  Were any subjects scanned but then rejected for the analysis after the data was collected?	N/A		
For	r papers reporting functional imaging (fMRI) results please ensure that the properties of the provided in the methods:  Were any subjects scanned but then rejected for the analysis after the data was collected?  a. If yes, is the number rejected and reasons for rejection	N/A		
For info	r papers reporting functional imaging (fMRI) results please ensure that the provided in the methods:  Were any subjects scanned but then rejected for the analysis after the data was collected?  a. If yes, is the number rejected and reasons for rejection described?  Where (section, paragraph #)?	N/A N/A		
For info	r papers reporting functional imaging (fMRI) results please ensure that the ormation is clearly provided in the methods:  Were any subjects scanned but then rejected for the analysis after the data was collected?  a. If yes, is the number rejected and reasons for rejection described?	N/A		
For info	r papers reporting functional imaging (fMRI) results please ensure that the provided in the methods:  Were any subjects scanned but then rejected for the analysis after the data was collected?  a. If yes, is the number rejected and reasons for rejection described?  Where (section, paragraph #)?	N/A N/A		
For info	r papers reporting functional imaging (fMRI) results please ensure that the provided in the methods:  Were any subjects scanned but then rejected for the analysis after the data was collected?  a. If yes, is the number rejected and reasons for rejection described?  Where (section, paragraph #)?  Is the number of blocks, trials or experimental units per session and/or subjects specified?	N/A N/A		
For info	r papers reporting functional imaging (fMRI) results please ensure that the provided in the methods:  Were any subjects scanned but then rejected for the analysis after the data was collected?  a. If yes, is the number rejected and reasons for rejection described?  Where (section, paragraph #)?  Is the number of blocks, trials or experimental units per session and/or subjects specified?	N/A N/A		
For info	r papers reporting functional imaging (fMRI) results please ensure that the provided in the methods:  Were any subjects scanned but then rejected for the analysis after the data was collected?  a. If yes, is the number rejected and reasons for rejection described?  Where (section, paragraph #)?  Is the number of blocks, trials or experimental units per session and/or subjects specified?  Where (section, paragraph #)?  Is the length of each trial and interval between trials specified?	N/A  N/A  N/A		
For info	r papers reporting functional imaging (fMRI) results please ensure that the provided in the methods:  Were any subjects scanned but then rejected for the analysis after the data was collected?  a. If yes, is the number rejected and reasons for rejection described?  Where (section, paragraph #)?  Is the number of blocks, trials or experimental units per session and/or subjects specified?  Where (section, paragraph #)?	N/A  N/A		
For info	r papers reporting functional imaging (fMRI) results please ensure that the provided in the methods:  Were any subjects scanned but then rejected for the analysis after the data was collected?  a. If yes, is the number rejected and reasons for rejection described?  Where (section, paragraph #)?  Is the number of blocks, trials or experimental units per session and/or subjects specified?  Where (section, paragraph #)?  Is the length of each trial and interval between trials specified?  Is a blocked, event-related, or mixed design being used? If applicable,	N/A  N/A  N/A		
For info 1. 2. 4.	r papers reporting functional imaging (fMRI) results please ensure that the ormation is clearly provided in the methods:  Were any subjects scanned but then rejected for the analysis after the data was collected?  a. If yes, is the number rejected and reasons for rejection described?  Where (section, paragraph #)?  Is the number of blocks, trials or experimental units per session and/or subjects specified?  Where (section, paragraph #)?  Is the length of each trial and interval between trials specified?  Is a blocked, event-related, or mixed design being used? If applicable, please specify the block length or how the event-related or mixed design was optimized.	N/A  N/A  N/A  N/A		
For info 1. 2. 4.	r papers reporting functional imaging (fMRI) results please ensure that the provided in the methods:  Were any subjects scanned but then rejected for the analysis after the data was collected?  a. If yes, is the number rejected and reasons for rejection described?  Where (section, paragraph #)?  Is the number of blocks, trials or experimental units per session and/or subjects specified?  Where (section, paragraph #)?  Is the length of each trial and interval between trials specified?  Is a blocked, event-related, or mixed design being used? If applicable, please specify the block length or how the event-related or mixed	N/A  N/A  N/A		

6.	How was behavioral performance measured?	N/A
7.	Is an ANOVA or factorial design being used?	N/A
0	For data acquisition, is a whole brain scan used?	N/A
Ο.	roi data acquisition, is a whole brain scan used:	N/A
	If not, state area of acquisition.	
	a. How was this region determined?	N/A
	a. How was this region determined:	N/A
9.	s the field strength (in Tesla) of the MRI system stated?	N/A
	a. Is the pulse sequence type (gradient/spin echo, EPI/spiral)	N/A
	stated?	
	b. Are the field-of-view, matrix size, slice thickness, and TE/TR/ flip angle clearly stated?	N/A
	nip angle clearly stated?	
10.	Are the software and specific parameters (model/functions,	N/A
	smoothing kernel size if applicable, etc.) used for data processing and	
	pre-processing clearly stated?	
11	Is the coordinate space for the anatomical/functional imaging data	N/A
11.	clearly defined as subject/native space or standardized stereotaxic	IV A
	space, e.g., original Talairach, MNI305, ICBM152, etc? Where (section,	
	paragraph #)?	
	F0	
12.	If there was data normalization/standardization to a specific space	N/A
	template, are the type of transformation (linear vs. nonlinear) used	
	and image types being transformed clearly described? Where (section,	
	paragraph #)?	
13.	How were anatomical locations determined, e.g., via an automated	N/A
	labeling algorithm (AAL), standardized coordinate database (Talairach	
	daemon), probabilistic atlases, etc.?	
1/	Were any additional regressors (behavioral covariates, motion etc)	N/A
14.	used?	IV A
	useu.	
15.	Is the contrast construction clearly defined?	N/A
16.	Is a mixed/random effects or fixed inference used?	N/A
	a. If fixed effects inference used, is this justified?	NI/A
	a. It fixed effects intereffice used, is this justified?	N/A
17.	Were repeated measures used (multiple measurements per subject)?	N/A
	a. If so, are the method to account for within subject	N/A
	correlation and the assumptions made about variance	
	clearly stated?	

18. If the threshold used for inference and visualization in figures varies, is this clearly stated?	N/A			
19. Are statistical inferences corrected for multiple comparisons?	N/A			
a. If not, is this labeled as uncorrected?	N/A			
20. Are the results based on an ROI (region of interest) analysis?	N/A			
a. If so, is the rationale clearly described?	N/A			
b. How were the ROI's defined (functional vs anatomical localization)?	N/A			
21. Is there correction for multiple comparisons within each voxel?	N/A			
22. For cluster-wise significance, is the cluster-defining threshold and the corrected significance level defined?	N/A			
▶ Additional comments				
Additional Comments	N/A			