# nature neuroscience

Corresponding Author:	Daniel O'Connor	# Main Figures:	4
Manuscript Number:	NN-A56441-T	# Supplementary Figures:	9
Manuscript Type:	Article	# 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	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
+												

ı		TEST US	SED		n		DESCRIPTIVE S' (AVERAGE, VARIA		P VALU	JE	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#
+	2a	two-tailed paired t-test	Fig. legend, Results para 4	1370	Number of responsive neurons from S1 sessions	Fig. legend	Error bars are mean +/- SEM across 5 mice, Hit-Miss: mean +/- 95% CI FA-CR: mean +/- 95% CI	Result s para 4, Meth ods, statist ics	p=1.17e-57 for Hit vs Miss p=0.130 for FA vs CR	Fig. legend	t(1369)=16.79 for Hit vs Miss t(1369)=1.512 for FA vs CR	Fig. legend
+	2b	two-tailed paired t-test	Fig. legend, Results para 4	607	Number of responsive neurons from S2 sessions	Fig. legend	Error bars are mean +/- SEM across 5 mice, Hit-Miss: mean +/- 95% CI FA-CR: mean +/- 95% CI	Result s para 4, Meth ods, statist ics	p=2.22e-70 for Hit vs Miss p=5.33e-5 for FA vs CR	Fig. legend	t(606)=20.32 for Hit vs Miss t(606)=4.07 for FA vs CR	Fig. legend
+	2c	two-tailed Wilcoxon signed-rank test	Fig. legend	1370, 607	Number of responsive neurons from S1 and S2 sessions	Fig. legend	Errorbars are mean +/- SEM across 5 mice	Meth ods, statist ics	p=1.73e-110 for hit vs miss in S1, p=3.76e-79 for hit vs miss in S2	Fig. legend	S1: z=22.33 S2:z=18.84	Fig. legend
+	2c	two-tailed Wilcoxon rank sum test	Fig. legend	1370, 607	Number of responsive neurons from S1 and S2 sessions	Fig. legend	Errorbars are mean +/- SEM across 5 mice	Result s para 4, Meth ods, statist ics	p = 8.38e-14 for normalized miss in S1 vs normalized miss in S2	Fig. legend	z = 7.46	Fig. legend
+	2d left	two-tailed K-S test	Fig. legend	2490, 1471	Number of neurons from S1 and S2 sessions	Fig. legend	All data points are plotted in fig., median in legend	Result s para 5, Fig. legend	p=0.0013	Fig. legend	K-S stat = 0.063	Fig. legend
+	2d midd le	two-tailed K-S test	Fig. legend	2490, 1471	Number of neurons from S1 and S2 sessions	Fig. legend	All data points are plotted in fig., median in legend	Result s para 5, Fig. legend	p=0.056	Fig. legend	K-S stat = 0.044	Fig. legend
+	2d right	two-tailed K-S test	Fig. legend	2490, 1471	Number of neurons from S1 and S2 sessions	Fig. legend	All data points are plotted in fig., median in legend	Result s para 5, Fig. legend	p=1.96e-7	Fig. legend	K-S stat = 0.093	Fig. legend
+	2g	bootstrap test	Fig. legend ,Meth ods	26898 (SP), 12387 (DP), 431962 (All)	Number of pairwise distances for SP, DP and all neurons	Fig. legend	All data points are plotted in fig., medians in text.	Result s para 5, Fig. legend	p<5e-5 (20,000 bootstrap iterations)	Fig. legend	NA	NA
+	2i	ANCOVA	Fig. legend	5 / 78	Number of mice/ number of binned values	Fig. legend	Errorbars are mean +/- SEM across mice	Fig. legend	p=1.711e-10 (test of zero slope) p=0.062 (difference between SP vs DP)	Fig. legend	F(1,74)=54.84 (slope) F(1,74)=3.58 (difference between SP and DP slopes)	Fig. legend
+	2j	ANCOVA	Fig. legend	5/58	Number of mice/ number of binned values	Fig. legend	Errorbars are mean +/- SEM across mice	Fig. legend	p=0.08 (test of zero slope) p=0.34 (difference between SP vs DP)	Fig. legend	F(1,54)=3.11 (slope) F(1,54)=0.92 (difference between SP and DP slopes)	Fig. legend

+	2k	one-tailed Wilcoxon rank sum test	Fig. legend	7 (S1), 9 (S2)	Number of sessions	Fig. legend	Errorbars are mean +/- SEM across mice	Fig. legend	p=0.045	Fig. legend	ranksum = 76	Fig. legend
+	21	one-tailed Wilcoxon rank sum test	Fig. legend	7 (S1), 9 (S2)	Number of sessions	Fig. legend	Errorbars are mean +/- SEM across mice	Fig. legend	p=0.021	Fig. legend	ranksum=40	Fig. legend
+	2m	two-tailed permutation test	Fig. legend	1370 (S1), 607 (S2)	Number of responsive neurons	Fig. legend	All data points are plotted in fig., mean +/- SEM across mice in text	Fig. legen d, Result s para 9	p<5e-5 (20,000 bootstrap iterations)	Fig. legend	NA	NA
+	2m	two-tailed permutation test	Fig. legend	536 (S1), 287 (S2)	Number of significant SP neurons	Fig. legend	All data points are plotted in fig.	Fig. legend	p<5e-5 (20,000 bootstrap iterations)	Fig. legend	NA	NA
+	2m	two-tailed permutation test	Fig. legend	338 (S1), 259 (S2)	Number of significant DP neurons	Fig. legend	All data points are plotted in fig.	Fig. legend	p<5e-5 (20,000 bootstrap iterations)	Fig. legend	NA	NA
+	3f	two-tailed Wilcoxon signed-rank test	Fig. legend	88 (s2p), 648 (unlabell ed)	Number of responsive neurons from 3 mice	Fig. legend	Errorbars are mean +/- SEM across mice	Result s para 12	s2p: p=7.43e-11 unlabeled: p=2.11e-43	Fig. legend,R esults para 12	s2p: z=6.511, unlabeled: z=13.81	Fig. legend
+	3f	two-tailed permutation test	Fig. legend	88 (s2p), 648 (unlabele d)	Number of responsive neurons from 3 mice	Fig. legend	Errorbars are mean +/- SEM across mice	Fig. legend	p=0.031	Fig. legend	NA	NA
+	3g	two-tailed permutation test	Fig. legend	133 (s2p), 1134 (unlabele d)	Number of neurons from 3 mice	Fig. legend	All data points are plotted in fig. Medians in legend	Fig. legend	p<5e-5 (20,000 bootstrap iterations)	Fig. legend	NA	NA
+	3h	two-tailed permutation test	Fig. legend	133 (s2p), 1134 (unlabele d)	Number of neurons from 3 mice	Fig. legend	All data points are plotted in fig. Medians in legend	Fig. legend	p<5e-5 (20,000 bootstrap iterations)	Fig. legend	NA	NA
+	3i	one-tailed Wilcoxon signed-rank test	Fig. legend	8	Number of sessions	Fig. legend	Errorbars are mean +/- SEM across mice	Fig. legend	p=0.012	Fig. legend	signedrank=34	Fig. legend
+	3j	one-tailed Wilcoxon signed-rank test	Fig. legend	8	Number of sessions	Fig. legend	Errorbars are mean +/- SEM across mice	Fig. legend	p=0.055	Fig. legend	signedrank=30	Fig. legend
+	3k	two-tailed permutation test	Fig. legend	88 (s2p), 648 (unlabele d)	Number of responsive neurons from 3 mice	Fig. legend	Errorbars are mean +/- SEM across sessions	Fig. legen d,Res ults para 13	p<5e-5 (20,000 bootstrap iterations)	Fig. legend	NA	NA
+	4e	two-tailed paired t-test	Fig. legend ,Result s para 15	160	Number of axons from 4 mice	Fig. legend	Errorbars are mean +/- SEM across mice, text gives Hit-Miss mean +/- 95% CI	Fig. legen d,Res ults para 15	p=0.0014	Fig. legend,R esults para 15	t(159) = 3.249	Fig. legend
+	4f	two-tailed paired t-test	Fig. legend ,Result s para 15	440	Number of axons from 4 mice	Fig. legend	Errorbars are mean +/- SEM across mice, text gives Hit-Miss mean +/- 95% CI	Fig. legen d,Rsul ts para 15	p = 0.0019	Fig. legend,R esults para 15	t(439) = 3.127	Fig. legend

+	4g	two-tailed sign test	Fig. legend	160(S1 axons), 440 (S2 axons)	Number of axons from 4 mice each	Fig. legend	Errorbars are mean +/- SEM across mice	Fig. legend	p=0.0034 (S1 axons), p=1.15e-18 (S2 axons)	Fig. legend	sign=99 (S1 axons), sign=313 (S2 axons)	Fig. legend
+	4g	two-tailed Wilcoxon rank sum test	Fig. legend	160(S1 axons), 440 (S2 axons)	Number of axons from 4 mice each	Fig. legend	Errorbars are mean +/- SEM across mice	Fig. legend	p=1.33e-4	Fig. legend	z=3.821	Fig. legend
+	4h	two-tailed Wilcoxon rank sum test	Fig. legend	7(S1 axons), 10(S2 axons)	Number of sessions from 4 mice each	Fig. legend	Errorbars are mean +/- SEM across mice	Fig. legend	p=0.044	Fig. legend	z=2.05	Fig. legend
+	4i	two-tailed Wilcoxon rank sum test	Fig. legend	7(S1 axons), 10(S2 axons)	Number of sessions from 4 mice each	Fig. legend	Errorbars are mean +/- SEM across mice	Fig. legend	p=0.109	Fig. legend	z=-1.47	Fig. legend
+	4m	two-tailed paired t-test	Fig. legend	12(S1), 15(S2), 12(Cntl)	Number of sessions from S1, S2 and control experiments	Fig. legend	Errorbars are mean +/- SEM across 4 mice	Fig. legend	p=9.22e-7 (S1) p=2.00e-5 (S2) p=0.023 (Cntl)	Fig. legend	t(11)=9.781 (S1) t(14)=6.286 (S2) t(11)=2.651 (Control)	Fig. legend
+	4m	two-tailed Wilcoxon rank sum test	Fig. legend	12(S1), 15(S2), 12(Cntl)	Number of sessions from S1, S2 and control experiments	Fig. legend	Errorbars are mean +/- SEM across 4 mice	Fig. legend	p=0.421 (S1 vs S2) p=0.001 (S2 vs Control)	Fig. legend	z=-0.805 (S1 vs S2) z=-3.270 (S2 vs Control)	Fig. legend
+	S1c	two-tailed K-S test	Fig. legend	338(S1), 502(S2)	Number of first-lick times from S1 and S2 sessions	Fig. legend	All data points are plotted in fig.	Fig. legend	p=0.436	results para 2, fig. legend	K-S stat=0.0597	Fig. legend
+	S3e left	two-tailed paired t-test	Fig. legend	226	Number of S1 neurons from 4 mice	Fig. legend	Errorbars are mean +/- SEM across 4 mice	Fig. legend	p=1.53e-21	Fig. legend	t(225)=10.60	Fig. legend
+	S3e right	two-tailed paired t-test	Fig. legend	214	Number of S2 neurons from 4 mice	Fig. legend	Errorbars are mean +/- SEM across 4 mice	Fig. legend	p=3.50e-13	Fig. legend	t(213)=7.76	Fig. legend
+	S3f	two-tailed Wilcoxon rank sum test	Fig. legend	226(S1), 214(S2)	Number of S1 and S2 neurons from 4 mice	Fig. legend	Errorbars are mean +/- SEM across 4 mice	Fig. legend	p=0.006	Fig. legend	z=2.721	Fig. legend
+	S3h	two-tailed permutation test	results para 11, fig. legend	4910	Number of mean noise correlations between S1 and S2 neurons	Fig. legend	Errorbars are mean +/- SEM across sessions	Fig. legend	p<2e-5 (20,000 bootstrap iterations)	Fig. legend	NA	NA
+ -	S3i left S3i midd le S3i right	two-tailed permutation tests	Fig. legend	226(S1), 214(S2)	Number of S1 and S2 neurons from 4 mice	Fig. legend	All data points are plotted in fig. median and 3rd quartile in legend.	Fig. legend	p=0.003 p=0.392 p=0.023	Fig. legend	NA	NA
+	S3j	one-tailed Wilcoxon sign rank test	Fig. legend	9	Number of sessions	Fig. legend	Error bars are mean +/- SEM across 3 mice	Fig. legend	p=0.037 (S1&S2 combined vs S1) p=0.005 (S1&S2 combined vs S2)	Fig. legend	signrank=7 signrank=2	Fig. legend

+	S3k	one-tailed Wilcoxon sign rank test	Fig. legend	9	Number of sessions	Fig. legend	Error bars are mean +/- SEM across 3 mice	Fig. legend	p=0.014 (S1 vs S2) p=0.037 (S1&S2 combined vs S1) p=0.010 (S1&S2 combined vs S2)	Fig. legend	signrank=41 signrank=7 signrank=3	Fig. legend
+	S9a row 1	two-tailed K-S test	Fig. legend	2490(S1), 1471(S2)	Number of neurons from S1 and S2 sessions	Fig. legend	All data points are plotted in fig.	NA	p=2.20e-4 p=0.060 p=0.002	Fig. legend	K-S stat=0.070 K-S stat=0.044 K-S stat=0.060	Fig. legend
+	S9a row 2	two-tailed K-S test	Fig. legend	2490(S1), 1471(S2)	Number of neurons from S1 and S2 sessions	Fig. legend	All data points are plotted in fig.	NA	p=5.14e-7 p=0.060 p=0.002	Fig. legend	K-S stat=0.090 K-S stat=0.044 K-S stat=0.060	Fig. legend
+	S9a row 3	two-tailed K-S test	Fig. legend	2490(S1), 1471(S2)	Number of neurons from S1 and S2 sessions	Fig. legend	All data points are plotted in fig.	NA	p=9.76e-6 p=0.041 p=7.15e-5	Fig. legend	K-S stat=0.081 K-S stat=0.046 K-S stat=0.074	Fig. legend
+	S9a row 4	two-tailed K-S test	Fig. legend	2490(S1), 1471(S2)	Number of neurons from S1 and S2 sessions	Fig. legend	All data points are plotted in fig.	NA	p=8.17e-8 p=0.051 p=1.76e-4	Fig. legend	K-S stat=0.096 K-S stat=0.045 K-S stat=0.071	Fig. legend
+	3d	two-tailed paired t-test	Fig. legend	88	Number of responsive S2p neurons from 3 mice	Fig. legend	Error bars are mean +/- SEM across mice	Fig. legend	p=9.24e-7	Fig. legend	t(87)=5.284	Fig. legend
+	3e	two-tailed paired t-test	Fig. legend	648	Number of responsive unlabeled neurons from 3 mice	Fig. legend	Error bars are mean +/- SEM across mice	Fig. legend	p=2.43e-35	Fig. legend	t(647)=13.187	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. 1e-g; Fig. 3b,c; Fig. 4b,d; Supp. Fig. 3a-d; Supp. Fig. 4c; Supp. Fig. 8a,b.

Images are example traces or schematics for experiments whose N values are reported in the legends and methods.

#### ▶ 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 #)? No statistical methods were used to predetermine sample sizes. Sample sizes are similar to what have been reported in the field. This is described in the Methods section titled "Statistics".

Statistical tests are justified as appropriate in multiple subsections of the Methods.

	a.	If there is a section summarizing the statistical methods in the methods, is the statistical test for each experiment clearly defined?	Yes, in the Methods section titled "Statistics".
	b.	Do the data meet the assumptions of the specific statistical test you chose (e.g. normality for a parametric test)?	Yes, this is described in the Methods section titled "Statistics".
		Where is this described (section, paragraph #)?	
	C.	Is there any estimate of variance within each group of data?  Is the variance similar between groups that are being statistically compared?	Yes, all data are reported with standard error of the mean or median with interquartile range. The variance is similar between groups. Dispersion is reported directly with the data. This is described in "data analysis" subsections of the Methods.
		Where is this described (section, paragraph #)?	described in data analysis subsections of the Methods.
	d.	Are tests specified as one- or two-sided?	Yes. In Methods and throughout.
	e.	Are there adjustments for multiple comparisons?	No.
3.	bar grap bar grap plots (wi	note transparency, <i>Nature Neuroscience</i> has stopped allowing hs to report statistics in the papers it publishes. If you have hs in your paper, please make sure to switch them to dotith central and dispersion statistics displayed) or to box-and-plots to show data distributions.	We now show dot-plots instead (superimposed on bar graphs).
4.	Was this	eria for excluding data points reported? s criterion established prior to data collection? s this described (section, paragraph #)?	Yes. This is described in multiple subsections titled "data analysis" under Methods.
5.	samples If no ran	he method of randomization used to assign subjects (or ) to the experimental groups and to collect and process data.  domization was used, state so.  loes this appear (section, paragraph #)?	Assignment was not randomized, as described in Methods section titled "Statistics".
6.	allocation	ement of the extent to which investigator knew the group on during the experiment and in assessing outcome included? Inding was done, state so.  Section, paragraph #)?	No blinding was done, as described in Methods section titled "Statistics".
7.		eriments in live vertebrates, is a statement of compliance with uidelines/regulations included?	Yes, reported in first line of Methods section.
	Where (	section, paragraph #)?	

8. Is the species of the animals used reported?

Where (section, paragraph #)?

Yes, in Methods section titled "Mice".

9.	Is the strain of the animals (including background strains of KO/transgenic animals used) reported?	Yes, in Methods section titled "Mice" and in Supp. Fig. 6.
	Where (section, paragraph #)?	
10.	Is the sex of the animals/subjects used reported?	Yes, in Methods section titled "Mice" and in Supp. Fig. 6.
	Where (section, paragraph #)?	
11.	Is the age of the animals/subjects reported?	Yes, in Methods section titled "Mice".
	Where (section, paragraph #)?	
12.	For animals housed in a vivarium, is the light/dark cycle reported?	Yes, in Methods section titled "Mice".
	Where (section, paragraph #)?	
13.	For animals housed in a vivarium, is the housing group (i.e. number of animals per cage) reported?	Yes, in Methods section titled "Mice".
	Where (section, paragraph #)?	
14.	For behavioral experiments, is the time of day reported (e.g. light or dark cycle)?	Yes, in Methods section titled "Mice".
	Where (section, paragraph #)?	
15.	Is the previous history of the animals/subjects (e.g. prior drug administration, surgery, behavioral testing) reported?	Mice were only used for the experiments reported here, as described in the Methods.
	Where (section, paragraph #)?	
	a. If multiple behavioral tests were conducted in the same group of animals, is this reported?	Behavioral testing is described fully in the Methods section titled "Behavioral task."
	Where (section, paragraph #)?	
16.	If any animals/subjects were excluded from analysis, is this reported?	No animals were excluded.
	Where (section, paragraph #)?	
	a. 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 #)?	

# ▶ Reagents

1.		ibodies been validated for use in the system under study d species)?	N/A. No antibodies were used.
	a.	Is antibody catalog number given?	N/A
		Where does this appear (section, paragraph #)?	
	b.	Where were the validation data reported (citation, supplementary information, Antibodypedia)?	N/A
		Where does this appear (section, paragraph #)?	
2.	Cell line i	dentity	N/A. No cell lines were used.
	a.	Are any cell lines used in this paper listed in the database of	
		commonly misidentified cell lines maintained by <u>ICLAC</u> and	
		NCBI Biosample?	
		Where (section, paragraph #)?	
	b.	If yes, include in the Methods section a scientific	N/A
		justification of their useindicate here in which section and paragraph the justification can be found.	
		paragraph the justification can be found.	
	C.	For each cell line, include in the Methods section a	N/A
		statement that specifies:	
		- the source of the cell lines	
		- have the cell lines been authenticated? If so, by which	
		method?	
		- have the cell lines been tested for mycoplasma	
		contamination?	
	Wł	nere (section, paragraph #)?	

#### ▶ Data deposition

Provide a Data availability statement in the Methods section under "Data availability", which should include, where applicable:

- Accession codes for deposited data
- Other unique identifiers (such as DOIs and hyperlinks for any other datasets)
- At a minimum, a statement confirming that all relevant data are available from the authors
- Formal citations of datasets that are assigned DOIs
- A statement regarding data available in the manuscript as source data
- A statement regarding data available with restrictions

See our data availability and data citations policy page for more information.

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.

Where is the Data Availability statement provided (section, paragraph #)?

Data availability is described in Methods section titled "Data availability".

#### ▶ 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.

N/A

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

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.

N/A			

### ▶ Human subjects

1.	Which IRB approved the protocol?	N/A
	Where is this stated (section, paragraph #)?	
2		
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 #)?	
c	le a statement included confirming that informed concept was	N/A
О.	Is a statement included confirming that informed consent was obtained from all subjects?	N/A
	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 #)?	
•	fMRI studies	
	TWIN Studies	
	papers reporting functional imaging (fMRI) results please ensure that th	ese minimal reporting guidelines are met and that all this
inf	ormation is clearly provided in the methods:	
1	Were any subjects scanned but then rejected for the analysis after the	N/A
1.	data was collected?	IVA
	a. If yes, is the number rejected and reasons for rejection	N/A
	dagaribad?	
	described?	
	described?  Where (section, paragraph #)?	
2.	Where (section, paragraph #)?	N/A
2.		N/A
2.	Where (section, paragraph #)?  Is the number of blocks, trials or experimental units per session and/	N/A
	Where (section, paragraph #)?  Is the number of blocks, trials or experimental units per session and/ or subjects specified?  Where (section, paragraph #)?	
	Where (section, paragraph #)?  Is the number of blocks, trials or experimental units per session and/ or subjects specified?	N/A N/A
3.	Where (section, paragraph #)?  Is the number of blocks, trials or experimental units per session and/ or subjects specified?  Where (section, paragraph #)?	

5.	Is the task design clearly described?	N/A
	Where (section, paragraph #)?	
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
		IVA
	If not, state area of acquisition.	
	a. How was this region determined?	N/A
9. I	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?	14/7
	b. Are the field-of-view, matrix size, slice thickness, and TE/TR/	N/A
	flip 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	N/A
	space, e.g., original Talairach, MNI305, ICBM152, etc? Where (section, paragraph #)?	
	ραι αβι αφι ι π):	
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 labeling algorithm (AAL), standardized coordinate database (Talairach	N/A
	daemon), probabilistic atlases, etc.?	
14.	Were any additional regressors (behavioral covariates, motion etc) used?	N/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?	N/A
	a. II fixed effects inference used, is this justified?	IN/M
17.	Were repeated measures used (multiple measurements per subject)?	N/A

a. If so, are the method to account for within subject correlation and the assumptions made about variance clearly stated?	N/A
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	