nature portfolio

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Reporting Summary

- A description of any restrictions on data availability

The fMRI data is available via https://zenodo.org/deposit/4423384

- For clinical datasets or third party data, please ensure that the statement adheres to our policy

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>.

Statistics		
For 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 statis	tical test(s) used AND whether they are one- or two-sided non tests should be described solely by name; describe more complex techniques in the Methods section.	
A descript	ion of all covariates tested	
A descript	ion 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.		
For Bayes	ian analysis, information on the choice of priors and Markov chain Monte Carlo settings	
For hierar	chical 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.		
Software an	d code	
Policy information	about <u>availability of computer code</u>	
Data collection	The fMRI data is available via https://zenodo.org/deposit/4423384	
Data analysis	For the fMRI data analysis, SPM12 toolbox (Wellcome Department of Cognitive Neurology, Institute of Neurology, UCL, London, UK) running on matlab was used.	
	s 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 encourage code deposition in a community repository (e.g. GitHub). See the Nature Portfolio guidelines for submitting code & software for further information.	
Data		
All manuscripts m	about <u>availability of data</u> ust include a <u>data availability statement</u> . This statement should provide the following information, where applicable: s, unique identifiers, or web links for publicly available datasets	

Please select the one belo	by 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 docu	ment with all sections, see <u>nature.com/documents/nr-reporting-summary-flat.pdf</u>	
Robavioural	l & social sciences study design	
	,	
	on these points even when the disclosure is negative.	
Study description	The manuscript describes a new robotic procedure and method to study hallucinations in different populations.	
Research sample	The manuscript reuses the fMRI dataset from Bernasconi, Blondiaux et al., 2021 at https://zenodo.org/deposit/4423384	
Sampling strategy	Not applicable	
Data collection	No data were collected for this study. The manuscript reuses the fMRI dataset from Bernasconi, Blondiaux et al., 2021 at https://zenodo.org/deposit/4423384	
Timing	Not applicable	
Data exclusions	No data were excluded for the analysis	
Non-participation	No participants dropped out.	
Randomization	Participants were not allocated into experimental groups	
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Magnetic resonance imaging

Experimental design

Design type task functional MRI and block design

Design specifications	2 experimental blocks including two conditions (asynchronous and synchronous) of 30 sec with 20 sec rest (among those 0.5 sec and 1.5 sec are the "start" auditory cue and "stop" auditory cue). Each block lasts 12 min (8 repetition per conditions). The control tasks include one condition each (either robotic manipulation or robotic stimulation on the back) of 30 sec and also 20 sec of rest (including 0.5 sec and 1.5 sec are the "start" auditory cue and "stop" auditory cue). Those tasks lasted around 7 min (8 repetitions per condition).	
Behavioral performance measur	The movement parameters of the front robot along the X and Z axis are recorded at a frequency of 100 Hz. Questionnaire data is also assessed after the scanning session.	
Acquisition		
Imaging type(s)	functional and structural	
Field strength	ЗТ	
Sequence & imaging parameters	Users can adapt this part as the robotic procedure does not depend on the MR sequence used. The data used in the current manuscript (Bernasconi, Blondiaux et al., 2021) uses the following sequences: - For the functional images, we used a whole brain echo-planar imaging (EPI) with the following parameters: 43 continuous slices, FOV=230mm, TR=2.5s, TE=30ms, flip angle=90°, in-plane resolution=2.5x2.5mm2, slice thickness=2.5mm. We used a a 64-channel head-coil. The two experimental runs contained 320 volumes and the control conditions (in which there is only 8 repetitions per condition) included 160 volumes. - anatomical image was recorded using a T1-weighted MPRAGE sequence (TR=2.3s, TE=2.32 ms, Inversion time=900ms, flip angle=8°, 0.9mm isotropic voxels, 192 slices per slab and FOV=240mm).	
Area of acquisition	whole brain	
Diffusion MRI Used	Not used ■ Not used Not used	
Preprocessing		
Preprocessing software	Standard SPM12 preprocessing pipeline is applied to the images, which includes the following steps: first the functional images are corrected for slice timing and are spatially realigned. Then, the anatomical image is then co-registered with the mean functional image and segmented into grey matter, white matter and cerebro-spinal fluid (CSF) tissue. Finally, the anatomical and the functional images are normalized to the Montreal Neurological Institute (MNI) brain template. Functional images are then smoothed with a Gaussian kernel with full-width half-maximum of 6mm. The two experimental runs were filtered with a high-pass filter at 1/300 Hz to remove low frequency confounds, while the two control tasks were filtered with a high-pass filter at 1/100 Hz.	
Normalization	Normalised	
Normalization template	MNI template	
Noise and artifact removal	Head motion is assessed based on framewise displacement (FD) calculation.	
Volume censoring	lot applicable	
Statistical modeling & infere	ence	
Model type and settings	Mass univariate with fixed effects for the first and second level analysis	
Effect(s) tested	Effect of interest: asynchronous condition > synchronous condition For the control conditions: motor task > baseline and touch task > baseline Conjunction analysis: asynchronous > [motor + touch] ∩ synchronous > [motor + touch]	
Specify type of analysis: W	hole brain ROI-based Both	
Statistic type for inference (See <u>Eklund et al. 2016</u>)	Cluster-wise P = 0.001	
Correction	FWE-corrected threshold of P = 0.05	

Models & analysis

	Involved in the study
X	Functional and/or effective connectivity
X	Graph analysis
X	Multivariate modeling or predictive analysis