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Reporting Summary

Nature Research 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 Research policies, see <u>Authors & Referees</u> and the <u>Editorial Policy Checklist</u>.

Statistics						
For all statistical analys	es, confirm that the following items are present in the figure legend, table legend, main text, or Methods section.					
n/a Confirmed	/a Confirmed					
☐ ☐ The exact sam	ple size (n) for each experimental group/condition, given as a discrete number and unit of measurement					
A statement of	n whether measurements were taken from distinct samples or whether the same sample was measured repeatedly					
The statistical Only common to	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 descript AND variation	ion of the statistical parameters including central tendency (e.g. means) or other basic estimates (e.g. regression coefficient) (e.g. standard deviation) or associated estimates of uncertainty (e.g. confidence intervals)					
X	hesis testing, the test statistic (e.g. F , t , r) with confidence intervals, effect sizes, degrees of freedom and P value noted exact values whenever suitable.					
For Bayesian a	analysis, information on the choice of priors and Markov chain Monte Carlo settings					
For hierarchic	al and complex designs, identification of the appropriate level for tests and full reporting of outcomes					
Estimates of e	ffect sizes (e.g. Cohen's d, Pearson's r), indicating how they were calculated					
1	Our web collection on <u>statistics for biologists</u> contains articles on many of the points above.					
Software and c	ode					
Policy information abou	ut <u>availability of computer code</u>					
Data collection	Stimuli were presented using E-Prime version 2.0					
Data analysis	Data were analyzed using AFNI, SPM8, and custom MATLAB code					
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/reviewers. We strongly encourage code deposition in a community repository (e.g. GitHub). See the Nature Research guidelines for submitting code & software for further information.						
Data						
- Accession codes, un - A list of figures that	ut <u>availability of data</u> nclude a <u>data availability statement</u> . This statement should provide the following information, where applicable: ique identifiers, or web links for publicly available datasets have associated raw data restrictions on data availability					
All data and code are available at https://osf.io/b7y9n.						
Field-speci	fic reporting					
Please select the one b	elow that is the best fit for your research. If you are not sure, read the appropriate sections before making your selection.					
☐ Behavioural & social sciences ☐ Ecological, evolutionary & environmental sciences						
For a reference copy of the do	ocument with all sections, see <u>nature.com/documents/nr-reporting-summary-flat.pdf</u>					

Life sciences study design

All studies must disclose on these points even when the disclosure is negative.				
Sample size	The targeted number of participants was based upon previous relevant literature. A sample size of 24 participants was targeted in each of the described studies as that number was within range of prior literature and allowed for perfect balancing of mappings described below.			
Data exclusions	2 participants with incomplete data sets were excluded for this report.			
Replication	Replication was the purpose of the study and all replication measures are described in detail in the manuscript.			
Randomization	Shape-to-rule, color-to-feature, and decision-to-finger mappings were randomly counter-balanced.			
Blinding	Blinding was not relevant given that all fMRI manipulations were repeated measures in a single experimental design.			

Reporting for specific materials, systems and methods

We require information from authors about some types of materials, experimental systems and methods used in many studies. Here, indicate whether each material, system or method listed 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 & experimental systems		Me	ethods
n/a	Involved in the study	n/a	Involved in the study
\times	Antibodies	\boxtimes	ChIP-seq
\boxtimes	Eukaryotic cell lines	\boxtimes	Flow cytometry
\boxtimes	Palaeontology		MRI-based neuroimaging
\boxtimes	Animals and other organisms		
	Human research participants		
\boxtimes	Clinical data		

Human research participants

Policy information about studies involving human research participants

Population characteristics Healthy young adults ranging in age from 18-28 were included. Participants were right-handed native English speakers with no history of neurological or psychiatric disorders. 28 females and 20 males were included.

> Recruitment was performed via online advertisements to the research subject volunteer pool listed in the University of California, Berkeley psychology website.

The protocol was approved by the Committee for Protection of Human Subjects at the University of California, Berkeley Ethics oversight

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

deviations.

Magnetic resonance imaging

Experimental design

Recruitment

Design type	Event-related
Design specifications	Half of the participants completed 1920 trials divided into 192 blocks with 24 blocks per cell of the 2 x 2 x 2 design. The other half of the participants completed 864 trials divided into 96 blocks with 12 blocks per cell of the 2 x 2 x 2 design.
Behavioral performance measures	Accuracy and reaction time were collected. Appropriate performance was verified by noting accuracies well above

Acquisition				
Imaging type(s)	functional			
Field strength	3 Tesla			
Sequence & imaging parameters	EPI, 22.0 cm^2 FoV, 64x64 matrix, 3.8 mm slice thickness, 35 slices, TR = 2000 ms, TE = 25 ms, flip angle = 70 degrees			
Area of acquisition	whole brain			
Diffusion MRI Used	Not used ■ Not used			
Preprocessing				
Preprocessing software	AFNI - 3dDespike, SPM8 - slice timing correction, realign and unwarp, segmented normalization, smoothing at 8 FWHM			
Normalization	non-linear using a T1-weighted MPRAGE			
Normalization template	(ICBM152			
Noise and artifact removal	For subjects with greater than 3mm total displacement or a framewise 0.5mm movement, motion regressors including linear, quadratic, differential, and quadratic differential terms were included as nuisance co-variates			
Volume censoring	Data were visually inspected for artifacts. No individual volumes were censored.			
Statistical modeling & inference				
Model type and settings	Mass univariate with AR1 correction for auto-correlation at the first level. Random effects t-tests at the second level.			
Effect(s) tested	Temporal control: (Dual+Delay)>(Restart+Control); Contextual control: (Dual+Restart)>(Delay+Control); Interaction: (Dual+Control)>(Delay+Restart); Spatial> (Delay+Restart); Spatial> (Delay+Control) (Delay+Restart); Spatial> (Delay+Control) (Delay+Restart); Spatial> (Delay+Control) (Del			
Specify type of analysis: Whole	brain ROI-based Both			
Statistic type for inference (See <u>Eklund et al. 2016</u>)	Cluster-wise at two levels: 0.05 height, 1019 voxel extent; 0.01 height, 300 voxel extent			
Correction	FWE p < 0.01 based upon 3dClustStim from AFNI			
Models & analysis				
n/a Involved in the study				
Functional and/or effective connectivity				
Graph analysis				
Multivariate modeling or predictive analysis				

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\boxtimes	Graph analysis				
\boxtimes	Multivariate modeling or predictive analys				