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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>.

When statistical analyses are reported, confirm that the following items are present in the relevant location (e.g. figure legend, table legend, main

## Statistical parameters

text, or Methods section).				
n/a	Confirmed			
	$\boxtimes$	The $\underline{\text{exact sample size}}(n)$ for each experimental group/condition, given as a discrete number and unit of measurement		
	$\boxtimes$	An indication of whether measurements were taken from distinct samples or whether the same sample was measured repeatedly		
	$\boxtimes$	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.		
$\boxtimes$		A description of all covariates tested		
$\boxtimes$		A description of any assumptions or corrections, such as tests of normality and adjustment for multiple comparisons		
	$\boxtimes$	A full description of the statistics including <u>central tendency</u> (e.g. means) or other basic estimates (e.g. regression coefficient) AND <u>variation</u> (e.g. standard deviation) or associated <u>estimates of uncertainty</u> (e.g. confidence intervals)		
	$\boxtimes$	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 <i>Give P values as exact values whenever suitable.</i>		
$\times$		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		
$\boxtimes$		Estimates of effect sizes (e.g. Cohen's d, Pearson's r), indicating how they were calculated		
	$\boxtimes$	Clearly defined error bars State explicitly what error bars represent (e.g. SD, SE, CI)		

Our web collection on statistics for biologists may be useful.

## Software and code

Policy information about <u>availability of computer code</u>

Data collection BIAcore T200 Control Software v3.0

Data analysis

BIAcore T200 Evaluation Software, The custom code used for multivalent binding analysis are described in the supplement and the exact code is available upon request to the authors as per the code availability statement. Microsoft Excel 2016.

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 upon request. 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

Policy information about availability of data

All manuscripts must include a data availability statement. This statement should provide the following information, where applicable:

- Accession codes, unique identifiers, or web links for publicly available datasets
- A list of figures that have associated raw data
- A description of any restrictions on data availability

All associated raw data is provided in the supplementary information.

Field-specific reporting				
Please select the b	est fit for your research. If you are not sure, read the appropriate sections before making your selection.			
\times Life sciences	Behavioural & social sciences Ecological, evolutionary & environmental sciences			
For a reference copy of	the document with all sections, see <u>nature.com/authors/policies/ReportingSummary-flat.pdf</u>			
Life scier	nces study design			
All studies must dis	sclose on these points even when the disclosure is negative.			
Sample size	The experiments were done in repeats as stated. Usually 3 repeats as is common in similar experiments. These sample sizes were chosen without sample-size calculations. The variance then measured indicated that these sample sizes were sufficient in all cases.			
Data exclusions	No data were excluded from the analysis.			
Replication	Replication Replication was done in almost all experiments. In the cases where it was excluded (some low affinity monovalent binding experiments large quantities of antibodies required made those experiments technically difficult. All attempts at replication were successful.			
Randomization	Covariants were not deemed to be important for this study due to the well defined experimental setup, and therefore sample groups and randomization were not used.			
Blinding	Blinding of investigators was not relevant to our study as automatic fitting was made by computer software.			
Reportin	g for specific materials, systems and methods			
Materials & experimental systems Methods				
n/a Involved in th	ne study n/a Involved in the study			
Unique biological materials ChIP-seq				
Antibodies Flow cytometry				
Eukaryotic cell lines  MRI-based neuroimaging  Palaeontology				
Animals and other organisms				
Human research participants				
Unique biolo	ogical materials			
Policy information about <u>availability of materials</u>				
Obtaining unique materials All unique material can be obtained from the authors upon request.				
Antibodies				

Antibodies used

Dilutions and usages described in the methods. Anti-DIG Monoclonal Rabbit Antibody: Life Technologies; Clone 9H27L19; LOT 1668052A. Catalog number 700772. All human Abs were produced and purified in house according to the Methods and the references 43-47.

Validation

The commercially obtained Abs were validated by Life Technologies by Western blot analysis of recombinant Digoxigenin conjugated to BSA. The in-hose produced Abs were validated earlier by western blot binding to NP and NIP in refs. 43-47.