

Supplementary Materials for
**Deep brain stimulation of the thalamus restores signatures of consciousness
in a nonhuman primate model**

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This PDF file includes:

Supplementary Text
Figs. S1 to S12
Tables S1 to S7

Supplementary Text

-Extended Materials and Methods

Deep brain stimulation (DBS) methodology

Surgery using neuro-navigation

We implanted a four-lead (0, 1, 2, 3) MRI-compatible clinical DBS electrode (Medtronic 3389, USA) in two macaques (monkey N and T) with the ultimate goal of performing simultaneous DBS-fMRI acquisitions. Electrode leads were 1.5 mm long and spaced by 0.5 mm. The external diameter was 1.27 mm. We targeted the right centro-median thalamus (CM) and performed aseptic stereotaxic surgery under general anesthesia using a neuro-navigation system (BrainSight, Rogue, Canada) guided by anatomical 3T MRI (Prisma Fit, Siemens, Germany) (MPRAGE, T1 weighted, repetition time TR = 2200ms; inversion time TI=900ms, 0.80mm isotropic voxel size, sagittal orientation, mono-channel 1Tx-1Rx circular surface coil of 12.5 cm diameter). The head of the monkey was placed in a stereotaxic frame and maintained with ears and ocular bars. All devices were built in plastic and MR compatible materials. Gadolinium fiducials were placed on the frame, as well as on the macaque skull and the temporary headpost that held additional fiducials. These landmarks, recognized by the neuro-navigation system, were put all around the skull in a non-coplanar manner. Pre-operative MRI images aimed at defining the target and trajectory. We assessed the target location and trajectory according to the MNI macaque brain coordinates (x, y, z) and Paxinos Atlas reference space (L, B, S). We confirmed the contact spot following three other approaches: i) anterior-posterior commissure (AC-PC) system; ii) distance to different anatomical area landmarks such as the right caudate nucleus and iii) Saleem Atlas. Lead placement trajectory was simulated with the neuro-navigation module prior to surgery.

We drilled craniotomy, positioned a plastic cannula to guide the DBS electrode and fixed an anchoring device system (Stim-lock, Medtronic, USA) covering the craniotomy. This element aimed at stabilizing and blocking the lead extremity to avoid lead migration. Per-operative MRI were acquired to control convergence between the theoretical and the practical implantation spot (effective target reached during surgery versus desired planned location). The extracranial part of the lead was

protected with a plastic MR compatible chamber that was home-made by 3D-printing and was fixed to the skull with screws and dental acrylic.

Verification of the DBS settings and underlying behavioral responses

To ensure efficiency across experiments, impedances between leads and through electrode to an external reference were first measured outside the MRI environment with the DBS programmer device provided by the manufacturer (8840 N'Vision, Activa Clinician DBS Programmer, Medtronic, USA). Ultimately, we used an oscilloscope (Wave Runner 44XI, LeCroy, USA) to check the electrical current delivered to each lead at the beginning and at the end of each experimental session. The stimulation also generated an artifact on the EEG signal, which provided a final benchmark during the fMRI acquisitions.

Behavioral responses (see behavioral assessment section) to electrical thalamic stimulation were assessed in each animal outside the scanner at least 20 days after the DBS implantation. We empirically explored different voltage amplitudes and pulse widths while keeping a monopolar stimulation at a frequency of 130Hz, applied successively to each of the four DBS contacts. The DBS lead targeted the centro-median thalamus. On the target contact (centered in CM), we determined the voltage level for high central thalamic (CT) DBS as the voltage just above the threshold at which a significant behavioral response was observed. Low CT-DBS corresponded to a lower current delivery below the voltage level that led to an arousal pattern. For comparison and reproducibility purposes, we kept the exact same DBS settings for the control stimulation site (ventro-lateral thalamus (VL) DBS).

fMRI statistical analysis

Block-design fMRI analysis

We generated plots by extracting the activations responses to high CT-DBS with the hemodynamic function in frontal (area 6V; 9/46; 8A), parietal (ventral intraparietal VIP, parietal area PFG), cingulate (anterior ACC: posterior PCC) and temporal cortex (temporo-parieto-occipital area TPO). Activity profiles were plotted as percentage of signal changes across time.

Resting-state fMRI analysis

For the static resting state, we calculated for each experimental condition and sessions the average of positive and negative Z-values and performed a Student t-test with the null hypothesis of zero correlation to test for statistical significance of connectivity between the different experimental conditions.

We computed the static functional correlations by estimating for each experimental condition (noted e) for the awake state, anesthesia, low CT-DBS, high CT-DBS, low VL-DBS and high VL-DBS and acquired run (noted r) the covariance matrix $C_{e,r}$. This value was obtained by extracting and averaging across all runs r the time series of all voxels included in each selected anatomical ROI. We referred as static functional correlation or stationary functional connectivity the entry matrix $C_{e,r}(i,j)$ where each cell represented the mean strength of the functional correlation between the i - j pair. For each covariance matrix, a Fisher transformation was applied to calculate the Z-score. The Z-score matrices were averaged across runs to obtain one matrix per experimental condition. To assess statistical significance, Student t-test at the threshold p value < 0.0001 and a false discovery rate correction were applied on the correlations of all pairs of brain regions.

Dynamic resting state fMRI analysis

Covariance values between all ROIs were included ($[82 \times (82-1)]/2 = 3,321$ features per matrix). $Z_{c,s,w}$ matrices were subsampled along the time dimension (w) before clustering. The resulting centroids or median clusters (BS_n with $n=1-7$; each BS_n is sized 82×82) were then used to initialize a clustering of all data, obtaining a matrix of brain states $B_{c,s,w}$, which, for a given arousal condition c and session s , is a vector of length 464, valued 1–7, because each matrix in $Z_{c,s,w}$ is assigned a BS_n .

The similarity score was computed from the correlation coefficient between the vectorized structural matrix and each vectorized brain state from the clustering analysis. All brain states were ordered in ascending order of similarity to the structure using the similarity score. To quantify the relation between the probability of occurrence of a BS and the similarity score, for each arousal condition, a regression analysis was done, to quantify the beta value (β), the R^2 and a P value. The differences in BS composition across arousal states was evaluated through a fixed-effects ANOVA, with mean rank similarity, that is, the result of averaging each BS time series, valued from 1 to 7, as a dependent

variable and the vigilance condition as the in-dependent variable. A fixed-effects ANOVA was run to quantify the effect of sedation on the probability of brain state 7. For this, we followed the same procedure, but the mean rank similarity was calculated considering only BS 7 (window w valued at 1; any other state window w valued at 0).

To explore specifically the fluctuations of intervoxel correlation within nodes of the “macaque global neuronal workspace” and sensori-motor areas (anterior cingulate cortex, ACC; dorsolateral prefrontal cortex, PFCdl; frontal eye fields, FEF; dorsolateral premotor cortex, PMCdl; primary somatosensory cortex, S1; primary motor cortex, M1; intraparietal cortex, Pcip; primary auditory cortex, A1; inferior temporal cortex, TCi; visual area 1, V1 and posterior cingulate cortex, PCC), we extracted the values from the whole brain matrices and applied a one-way analysis of variance (ANOVA). FC across the brain states were highlighted by displaying Z-score in inter-region matrices.

Event-related task fMRI analysis

We generated plots by obtaining the β -weight of SPM regressions of individual macaque data with the hemodynamic functions of the appropriate stimulus categories and then plotted the mean and SE of these β -weights. These values estimate, in percentages of the whole-brain fMRI signal, the size of the fMRI activation relative to the implicit rest baseline that divides trials.

The first level analyses consisted in the convolution of the stimulus categories with the MION canonical hemodynamic response function (HRF) and its time derivative. We also added motion regressors and heart rate as variables of non-interest to the event-related regressors. Activation time series of all the fMRI voxels were computed for each fMRI run and signal change expressed in T-score maps for the different stimulus categories relative to rest periods. Global standard trials that immediately followed a global deviant trial were excluded.

-Extended Results

Thalamic DBS effects on resting-state networks in anesthetized macaques

Static Functional Correlations (Figure 4, S6)

To test for statistical significance of connectivity between brain regions in different experimental

conditions, Student t-tests were performed with the null hypothesis of zero correlation. We calculated for each experimental condition c and sessions the average of positive and negative Z values of $Z_{c,s}$. The average positive Z-value was $0.43 \pm 2.9e-4$ in the awake state, $0.22 \pm 2.6e-4$ under anesthesia, $0.27 \pm 2.5e-4$ during low CT-DBS, $0.39 \pm 2.8e-4$ during high CT-DBS, $0.28 \pm 3.1e-4$ during low VL-DBS and $0.19 \pm 2.4e-4$ during high VL-DBS (*Figure 4A*). Positive Z-values were significantly different under anesthesia and high CT-DBS ($p < 0.001$, FDR corrected (*Figure 4A*)). The average negative Z-value was $0.24 \pm 4.2e-4$ in the awake state, $0.09 \pm 1.7e-4$ under anesthesia, $0.12 \pm 2.7e-4$ during low CT-DBS, $0.11 \pm 3.2e-4$ during high CT-DBS, $0.08 \pm 2.7e-4$ during low VL-DBS and $0.08 \pm 2.2e-4$ during high VL-DBS (*Figure 4A*). Negative Z-values were significantly different under anesthesia and high CT-DBS ($p < 0.001$, FDR corrected) (*Figure 4A*). In the awake state, the frontal cortex (areas 9/46, 8A, 6V and M1), parietal cortex (parietal area PFG and ventral intraparietal area), anterior and posterior cingulate cortices, temporal cortex (area A1) and occipital cortex (area V1) were strongly correlated to each other (*left column, Figure 4B*).

Dynamic Functional Correlations (Figure 5, S7-S8, Table S4)

We applied k-means to the whole acquired dataset (including all experimental conditions) to cluster brain states (*Figure S7*). We also applied k-means to two data subsets, subset CT and subset VL, to specifically characterize the effects of CT-DBS and VL-DBS respectively. Subset CT included data from awake, anesthesia and anesthesia + high CT-DBS conditions (*Figure 5A-C*). Subset VL included data from awake, anesthesia and anesthesia + high VL-DBS conditions (*Figure 5D-E*).

-Clustering subset CT (data from awake, anesthesia and anesthesia + high CT-DBS conditions) (Figure 5A-C)

In the awake state, all 7 brain states were represented with a similar probability of occurrence ($\beta=0.45$; $R^2=0.22$; $p=0.28$). During anesthesia, state 7 (with the highest function-structure similarity) was dominant and state 1 (with the lowest function-structure similarity) never occurred ($\beta=1.91$; $R^2=0.67$; $p=0.02$). Under high CT-DBS, consistent with partial recovery of consciousness, the probability of occurrence of state 7 decreased in favor of all the other brain states, especially state 2 and 3 ($\beta=0.64$; $R^2=0.28$; $p=0.22$). We computed the slope of the linear relation between structural and functional

correlations for each recording session and compared the slope distributions in the awake, anesthesia and DBS conditions. Awake and high CT-DBS slopes were significantly lower than the anesthesia slopes, indicating that a greater diversity of states were explored in the wake state (awake versus anesthesia: t-test, $p=6e-5$ and $BF_{10}=338$, high CT-DBS versus anesthesia: t-test, $p=0.001$ and $BF_{10}=23$). Importantly no differences were observed between awake and high CT-DBS slopes (t-test, $p=0.42$, $BF_{01}=3.28$) (*Figure 5A-B, Table S6*). In the awake state, the mean rank of brain states was 4 (4.38 ± 1.28), for anesthesia, the mean rank was 6 (5.70 ± 1.40) and during high CT-DBS, the mean rank was 5 (4.55 ± 1.17). This brain state distribution was significantly different (ANOVA; $F(2;120)=12.52$; $p=1.15e-7$). Also, the frequency of brain state 7 was moderate in the awake experiments (probability=0.24), high during anesthesia (probability=0.58) and low again during high CT-DBS (probability=0.26; $p<0.0001$) (*Figure 5B*). The probability of brain state 7 was higher in anesthesia compared to the awake state (t-test, $p=1e-6$, $BF_{10}=9063$) and to high CT-DBS (t-test, $p=1e-5$, $BF_{10}=1017$) which did not differ significantly (t-test, $p=0.74$, $BF_{01}=4.18$). The mean similarity with the anatomical connectivity was also significantly different with 0.24 (± 0.06) for the awake state, 0.31 (± 0.07) for anesthesia and 0.25 (± 0.06) for high CT-DBS (ANOVA; $F(2,120)=14.75$; $p=1.87e-6$). Anatomically, the functional brain states 1, 2 and 3, that were most characteristic of the awake, presented strong correlations within the “macaque GNW” prefrontal (dorsolateral prefrontal cortex, PFCdl; dorsolateral premotor cortex, PMCdl), parietal (intraparietal cortex, PCip) and cingulate nodes (anterior cingulate cortex, ACC; posterior cingulate cortex, PCCr), whereas state 7 displayed low or null Z-score values across the same entire cortical network (*Figure 5C*). During high CT-DBS, the average duration of brain state 7 decreased compared to anesthesia (high CT-DBS versus anesthesia, $p=2.48e-3$; bootstrap analysis) and was similar to the awake state (*Figure S8*).

- Clustering subset VL (data from awake, anesthesia and anesthesia + high VL-DBS conditions) (*Figure 5D-E*)

In the awake state, all seven brain states were present ($\beta=0.16$; $R^2=0.02$; $p=0.74$). Under anesthesia, brain state 7 was dominant ($\beta=1.21$; $R^2=0.33$; $p=0.18$), as under high VL-DBS ($\beta=1.69$; $R^2=0.43$; $p=0.11$) (*Figure 5F-G*). We also computed the slope corresponding to each recording session and

compared the slope distributions in the awake, anesthesia and high VL-DBS conditions. Awake slopes were significantly lower compared to anesthesia and high VL-DBS slopes (awake versus anesthesia: t-test, $p=0.0007$ and $BF_{10}=39$, awake versus high CT-DBS: t-test, $p=1e-8$ and $BF_{10}=636824$). The slopes under anesthesia were smaller than the high VL-DBS slopes (t-test, $p=0.01$, $BF_{10}=3.81$).

For the awake state, the mean rank of brain states was 4 (4.41 ± 0.93), under anesthesia, the mean rank was 6 (5.81 ± 1.39) and under high VL-DBS, the mean rank was 6 (6.41 ± 0.38). The brain state distribution was significantly different (ANOVA; $F(2;102)=32.03$; $p=1.61e-11$). Brain state 7 was balanced in the awake state (awake brain state 7 probability=0.19), dominant under anesthesia (anesthesia brain state 7 probability=0.60) and high VL-DBS (high VL-DBS brain state 7 probability=0.72; $p<0.0001$) (*Figure 5F*). The probability of state 7 was smaller in the awake state compared to anesthesia (t-test, $p=10e-9$, $BF_{10}=2e7$) and compared to high VL-DBS (t-test, $p=10e-17$, $BF_{10}=1e15$). However, we found no evidence for a difference nor a similarity between anesthesia and high VL-DBS (t-test, $p=0.15$, $BF_{10}=0.64$, $BF_{01}=1.54$).

Brain state 1 highlighted strong correlations to all the tested cortical areas. Brain state 7 presented weak Z-score values with prefrontal (PFCdl; PMCdl), parietal (PCip) and cingulate cortex (ACC; PCC) (*Figure 5C*).

With high VL-DBS, the average duration of brain state 7 increased compared to the awake state (high VL-DBS v/s awake, $p<0.0001$, bootstrap analysis) and was similar to the anesthesia state (*Figure S8*).

-Clustering the whole dataset (data from awake, anesthesia, low CT-DBS, high CT-DBS, low VL-DBS and high VL-DBS) (Figure S7)

The occurrence of brain states in the awake condition was equiprobable ($\beta=0.05$; $R^2=0.003$; $p=0.91$). Under anesthesia, this probability was shaped by the brain state 7 ($\beta=1.87$; $R^2=0.59$; $p=0.04$). For the DBS sessions, brain state probability of occurrence was partly dominated by brain state 7 in the low CT-DBS condition ($\beta=1.09$; $R^2=0.49$; $p=0.08$), balanced under high CT-DBS ($\beta=0.28$; $R^2=0.059$; $p=0.6308$), partly dominated by the brain state 7 during the low VL-DBS experiments ($\beta=1.52$; $R^2=0.59$; $p=0.04$) and dominated by brain state 7 in the high VL-DBS condition ($\beta=2.57$; $R^2=0.73$; $p=0.01$) (*Figure S7*).

The mean rank was 4 in the awake state (3.81 ± 1.05), 5 under anesthesia (5.37 ± 1.34), 5 in the low CT-DBS condition (4.81 ± 0.93), 4 in the high CT-DBS condition (4.14 ± 1.03), 5 in the low VL-DBS condition (5.35 ± 0.86) and 6 in the high VL-DBS condition (6.01 ± 0.53). The brain state distribution was significantly different between structural and functional correlations (ANOVA; $F(5;193)=19.65$; $p=8.31e-16$) (*Figure S7*). The probability of occurrence of brain state 7 was low in the awake state (0.20), and high under anesthesia (0.54). Crucially, even though anesthesia continued, low CT-DBS and high CT-DBS reduced this probability down to an aware level (respectively 0.37 and 0.23). The probability of state 7 also decreased with low VL-DBS (0.38) but returned to high (0.63, $p<0.001$) under high VL-DBS. The mean similarity with the anatomical connectivity was also significantly different with 0.26 (± 0.04) for the awake state, 0.32 (± 0.06) for anesthesia, 0.29 (± 0.04) for low CT-DBS, 0.27 (± 0.04) for high CT-DBS, 0.31 (± 0.04) for low VL-DBS and 0.35 (± 0.03) and for high VL-DBS (ANOVA; $F(5,193)=17.72$; $p=1.94e-14$). The average duration of brain state 7 significantly decreased with high CT-DBS compared to the anesthesia state ($p=1.61e-9$, bootstrap analysis) and was similar to the awake state. Low CT-DBS and low VL-DBS decreased the duration the brain state 7 compared to anesthesia (low CT-DBS v/s anesthesia, $p=6.72e-8$; low VL-DBS v/s anesthesia, $p=1.88e-7$, bootstrap analysis). Under high VL-DBS, duration of the brain state 7 was similar to the anesthesia state (high VL-DBS v/s anesthesia, not significant) (*Figure S7*).

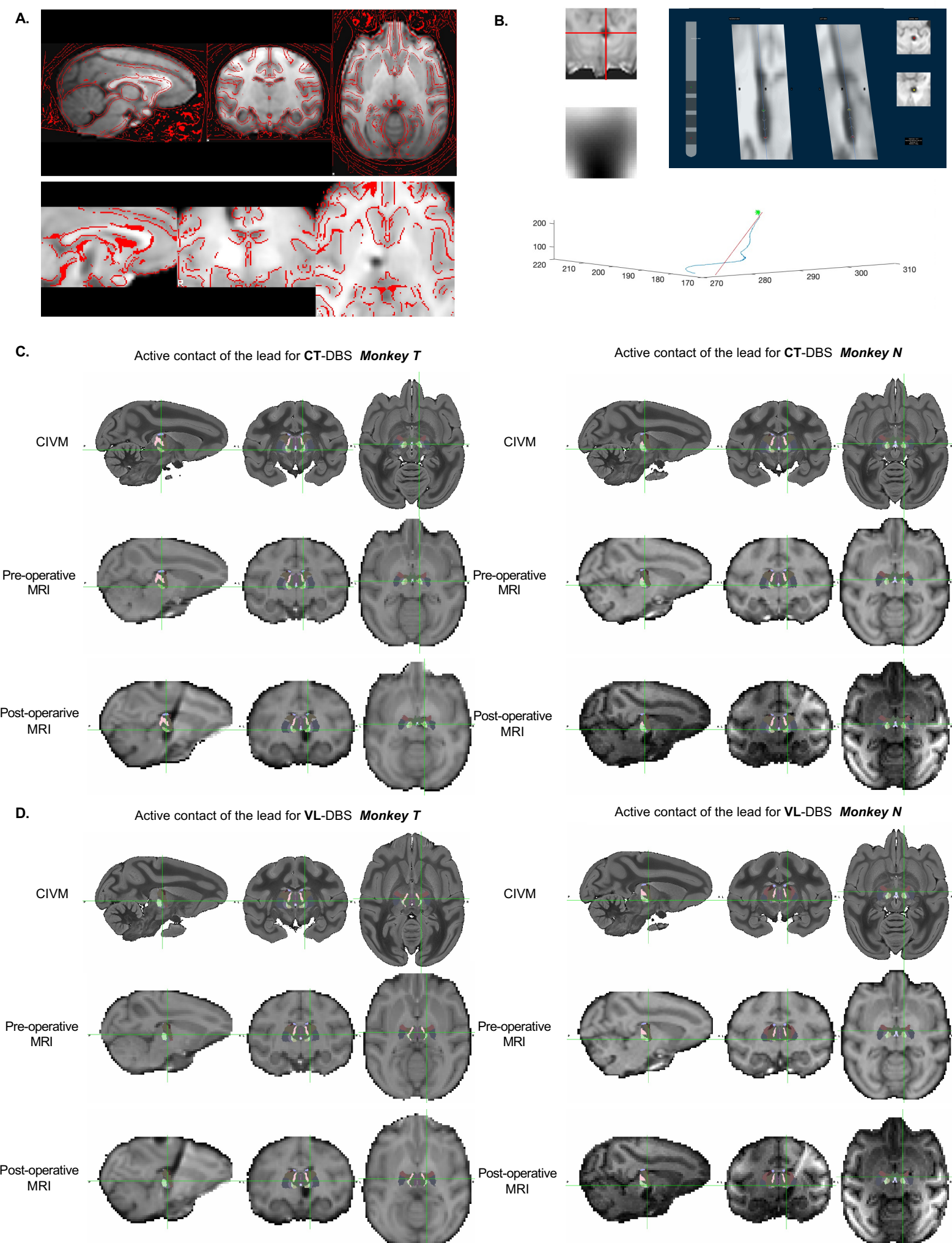
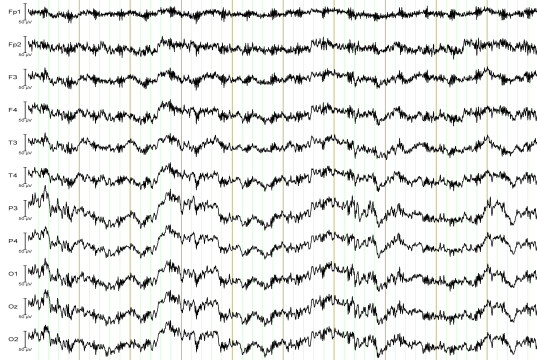


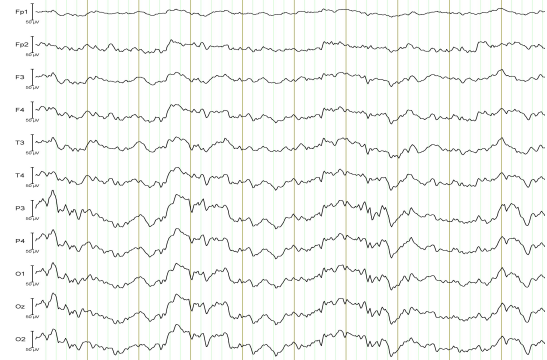
Figure S1: Localization of the DBS electrode contacts using the Lead-DBS macaque toolbox(71). (A) Coregistration of the pre and post-operative MRI anatomical images (upper panel) and between MRI post-operative and MNI macaque brain atlas(73) (lower panel) for monkey T. (B) Pre-reconstruction of the electrode lead trajectory using the entry point on the anatomical MRI image and manual correction of electrode localization adjusting the most inferior (contact 0) and most superior (contact 3) DBS contacts according to the electrode artifact in two dimensional planes presented orthogonally. (C) Location of the centro-median (CM) DBS contact and (D) ventral-lateral thalamus (VL) DBS contact in monkey T (left column) and monkey N (right column) on the sagittal, coronal and axial plan. The target is displayed in the CIVM MRI atlas(76) (upper panel), pre-operative structural MRI (middle panel) and post-operative structural MRI (lower panel) warped in the MNI macaque space(73).

EEG - MR B₀ field

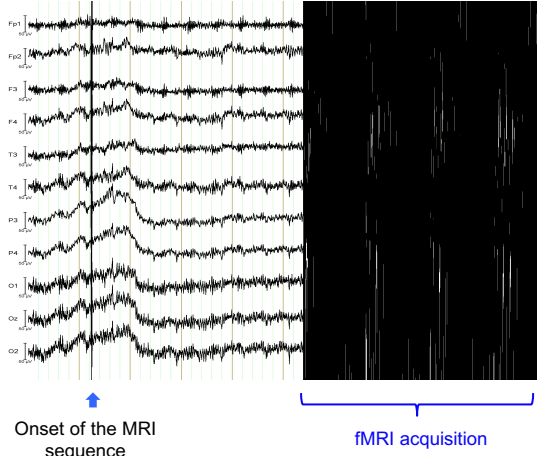


EEG-MRI artifact correction

EEG - MR B₀ field after artifact correction

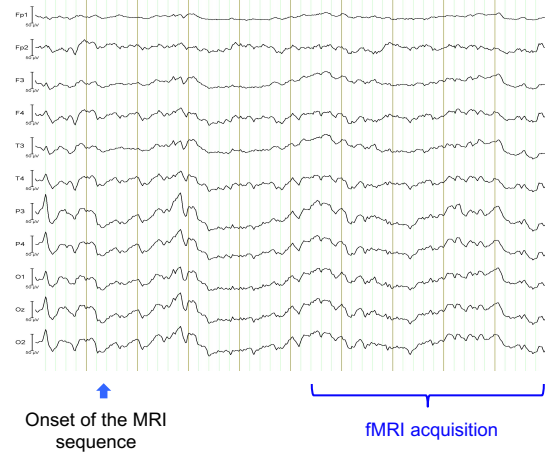


EEG - MR Gradient (fMRI)

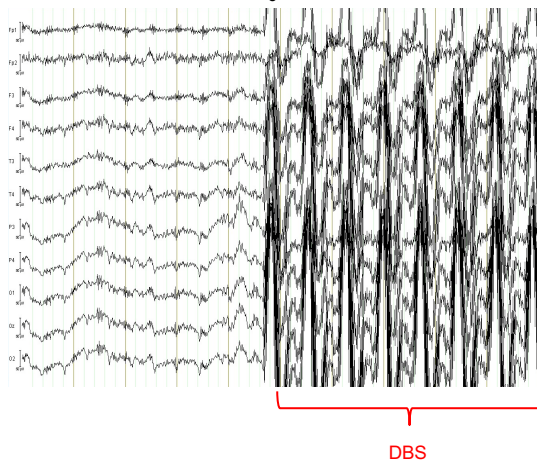


EEG-fMRI artifact correction

EEG - MR Gradient (fMRI) after artifact correction



EEG - MR B₀ field + DBS

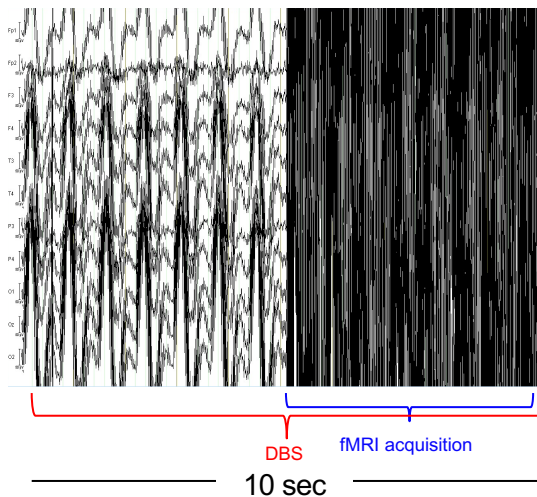


EEG-MRI-DBS artifact correction

EEG - MR B₀ field + DBS after artifact correction



EEG - MR Gradient (fMRI) + DBS



EEG-fMRI-DBS artifact correction

EEG - MR Gradient (fMRI) + DBS after artifact correction

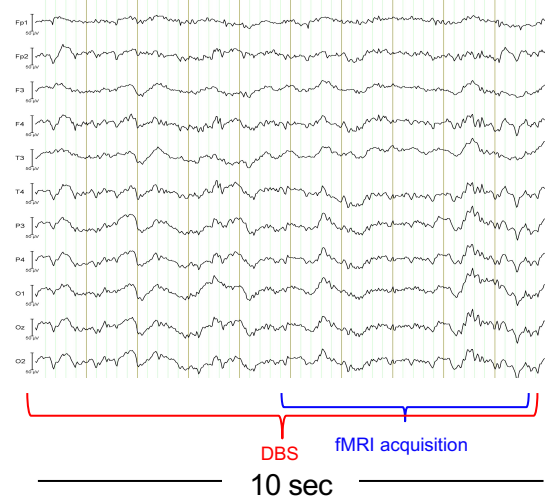


Figure S2: Suppression of EEG artifacts related to MR B₀ field, MR Gradients during fMRI acquisition and DBS. Examples of EEG recordings in anesthetized macaques inside a 3T MRI scanner without and with DBS of central thalamus (CT) thalamus at 3V.

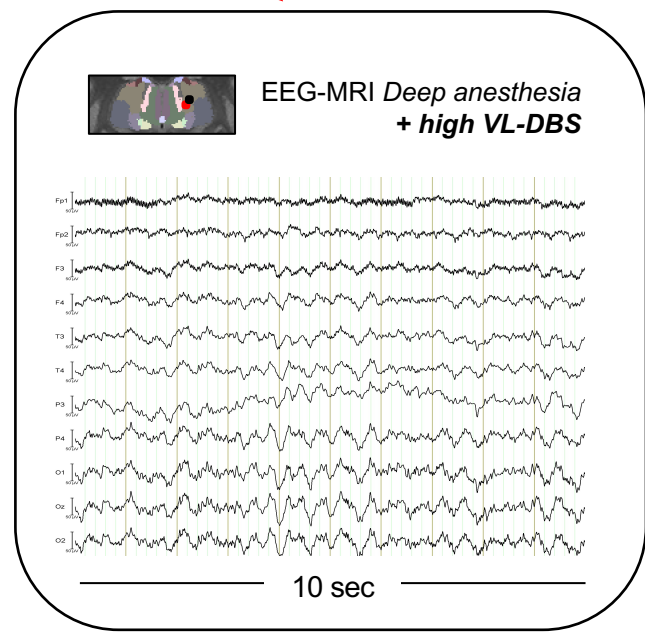
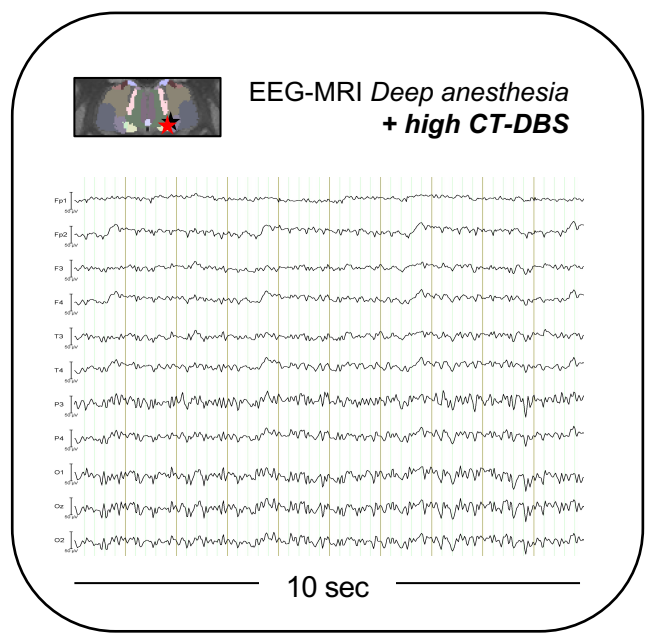
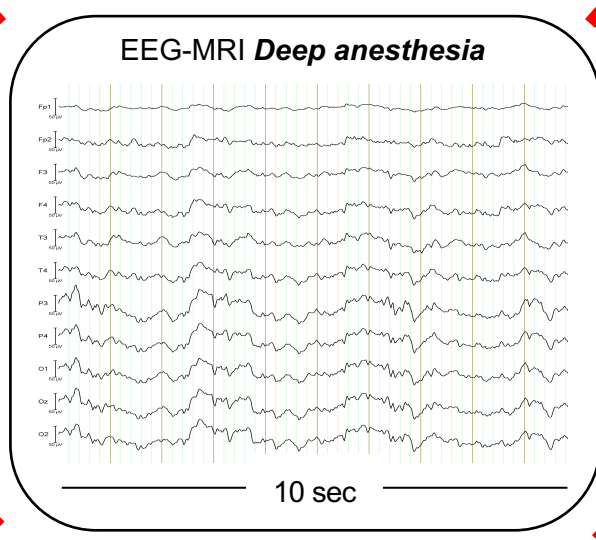
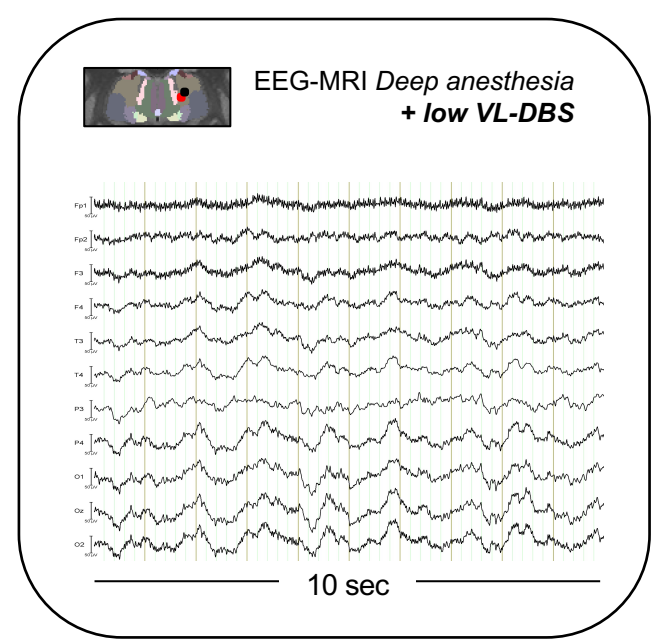
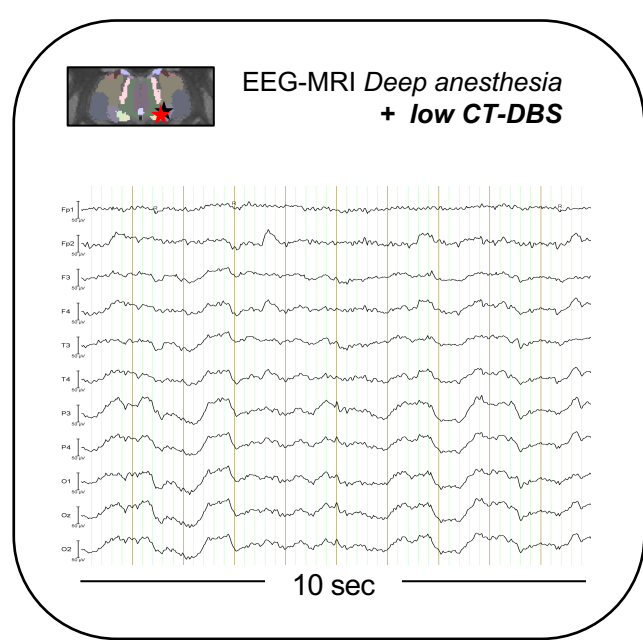


Figure S3: Effects of thalamic DBS on EEG. Examples of EEG recordings in anesthetized macaques inside a 3T MRI scanner with DBS of central thalamus (CT) or ventral-lateral thalamus (VL) at low or high voltages. The DBS-induced changes in cortical activity depends on the anatomical site of the active DBS lead contact and the intensity of the electrical stimulation.

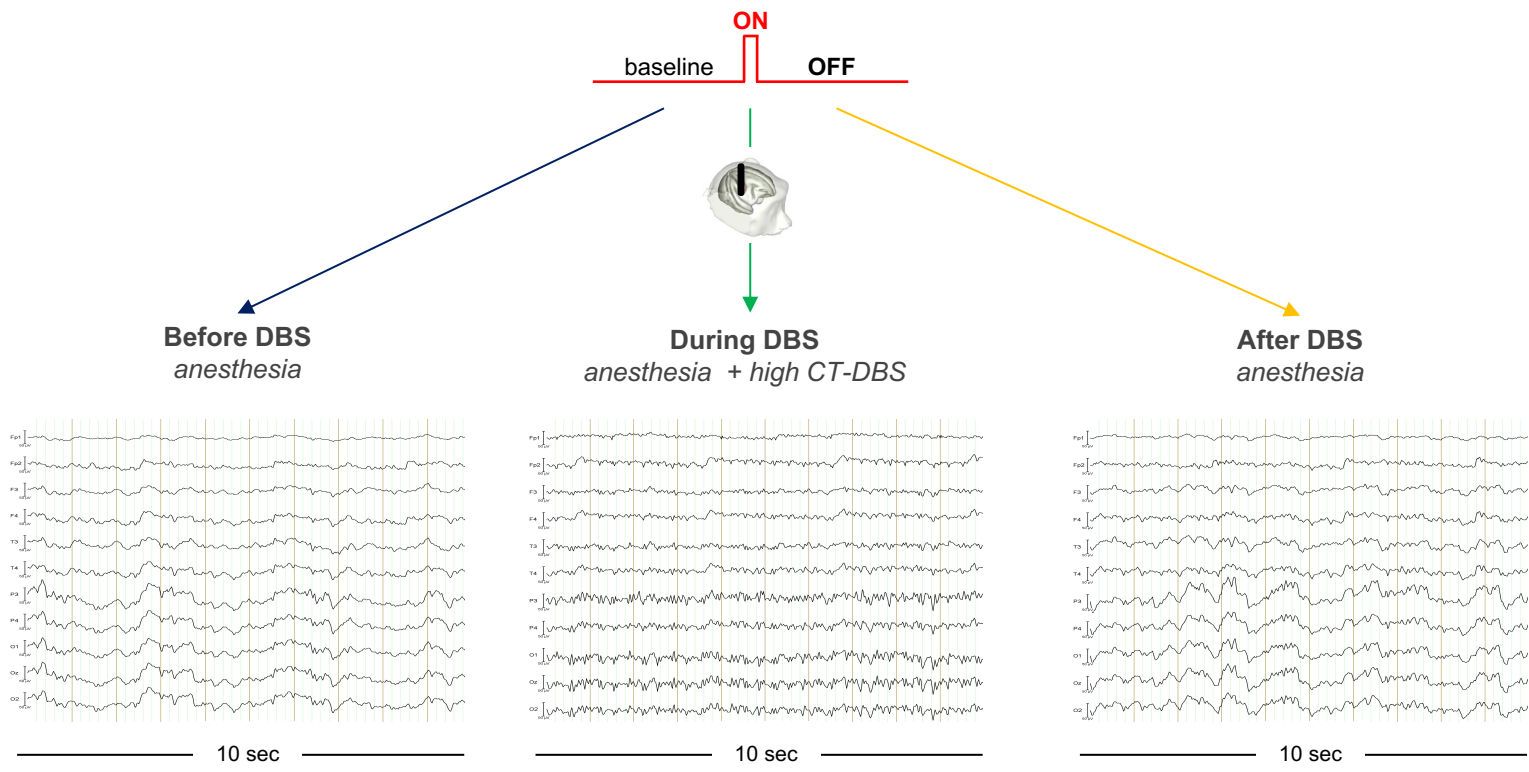


Figure S4: Cortical activity dynamic during and immediately after DBS

Examples of EEG recordings in anesthetized macaques inside a 3T MRI scanner before, during and after DBS of central thalamus (CT) at high voltage.

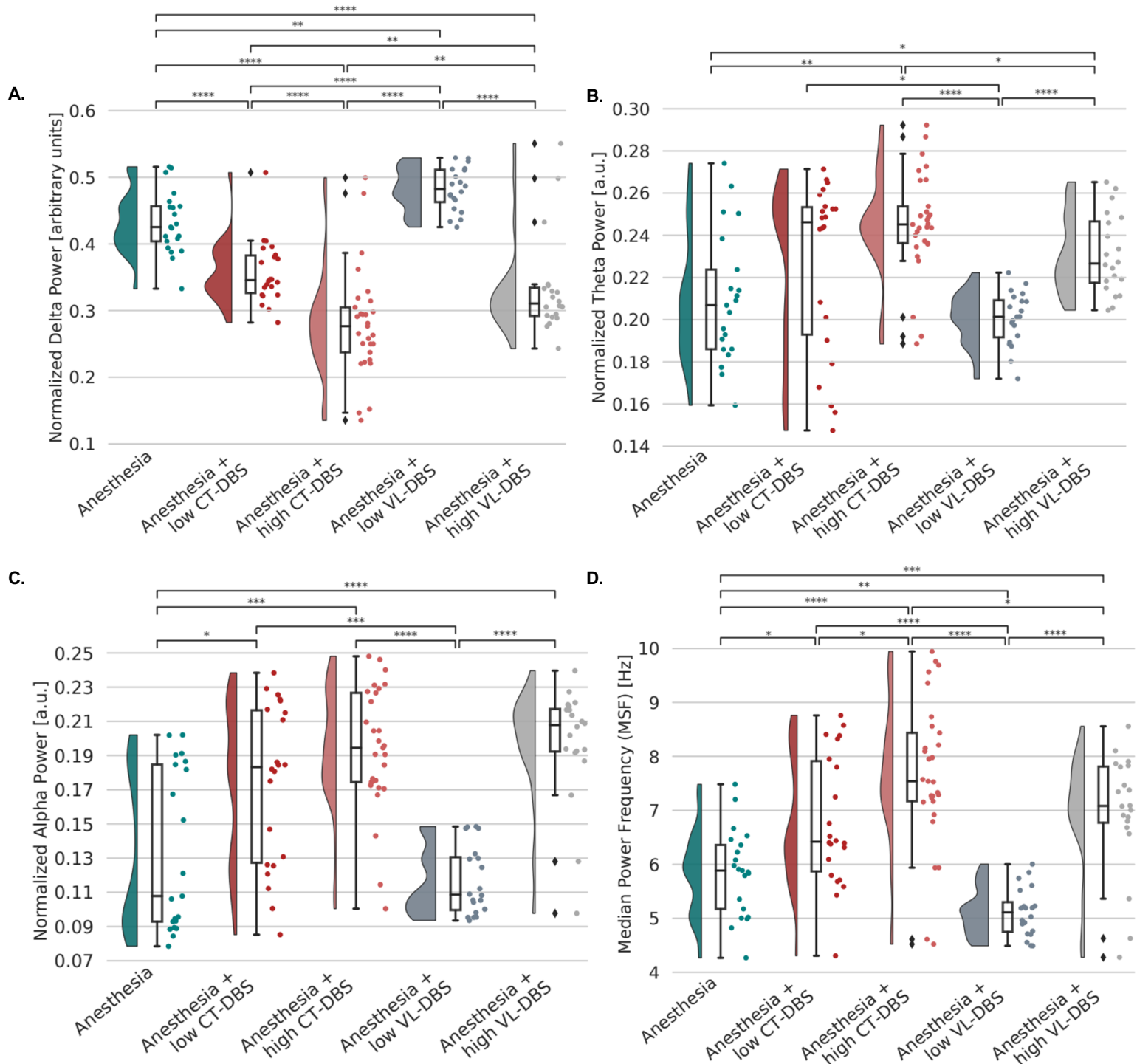


Figure S5: Modulation of normalized spectral power and median power frequency in the DBS conditions compared to anesthesia.

Distributions of the average values of normalized spectral power of (A) delta (1-4 Hz), (B) theta (4-8 Hz) and (C) alpha (8-13 Hz) oscillatory bands and (D) median power frequency (MSF) calculated on the epochs of the four stimulation conditions and under anesthesia. MSF is the frequency that divides the power spectrum in two equal areas. The figures consist of a distribution - smoothed version of a histogram, a box plot and a representation of the data points. Each dot in the figure represents the average value of a given marker across epochs during one recording session. a.u., arbitrary units. The significance lines represent FDR corrected Mann-Whitney U two-sided tests (see Methods). p-value annotation legend: ns: $5.00e-02 < p \leq 1.00e+00$, *: $1.00e-02 < p \leq 5.00e-02$, **: $1.00e-03 < p \leq 1.00e-02$, ***: $1.00e-04 < p \leq 1.00e-03$, ****: $p \leq 1.00e-04$. P-values are FDR corrected.

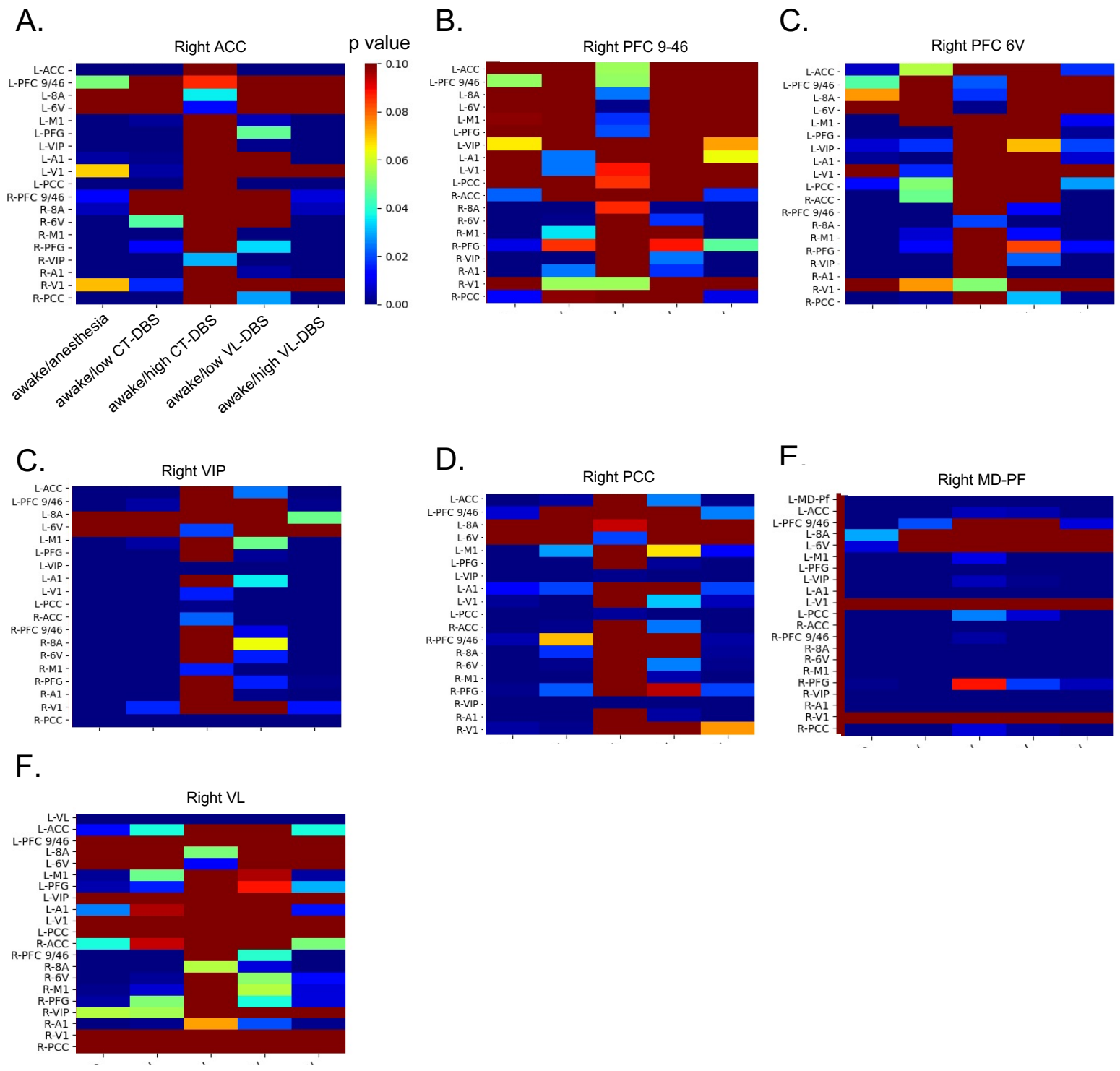


Figure S6: p values matrices for the ANOVA comparison of the awake state versus different experimental conditions of the static functional correlations within the macaque Global Neuronal Workspace (GNW) nodes and sensori-motor areas.

Statistical p values obtained by ANOVA to compare static functional correlations within the macaque GNW nodes and sensori-motor areas in the different experimental conditions. X-axis displays the experimental conditions (awake, anesthesia, low central thalamic (CT) DBS, high CT-DBS, low ventral-lateral thalamic (VL) DBS and high VL-DBS states), y-axis represents the macaque GNW areas correlated to the seed ($p < 0.001$, FDR corrected). For each region, the matrix stands for the p value for the comparison of the awake state versus anesthesia, low CT-DBS, high CT-DBS, low VL-DBS and high VL-DBS between the seed and the rest of the macaque GNW nodes and sensori-motor areas.

Anterior cingulate cortex (ACC) ; Prefrontal cortex (area 9/46, 8A, 6 Ventral) ; Primary motor cortex (M1) ; Parietal cortex (area PFG) ; Ventral Intraparietal sulcus (VIP) ; Primary auditory cortex (A1) ; Primary visual area (V1) ; Posterior cingulate cortex (PCC).

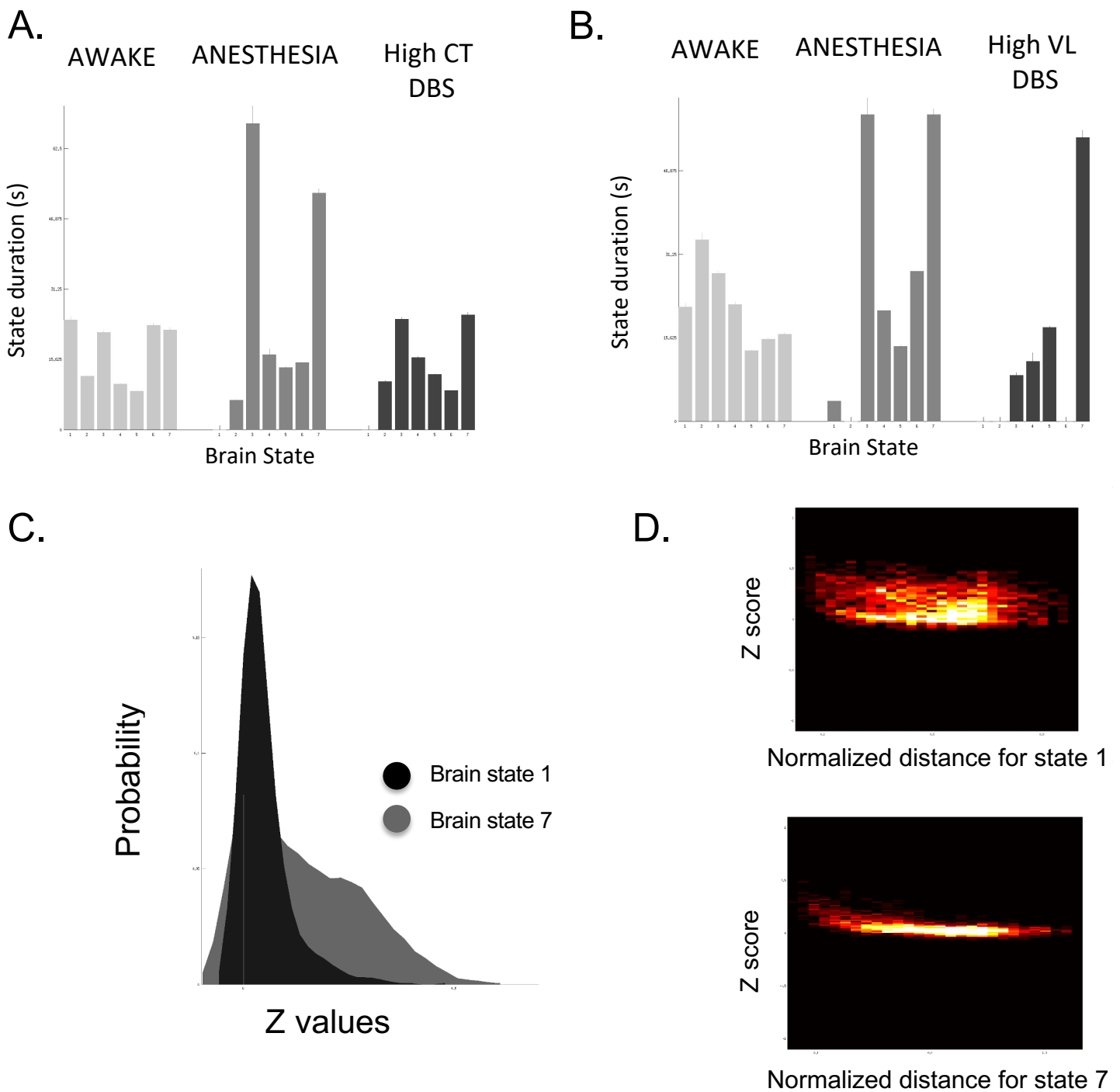


Figure S8: Average life time of brain states in the awake state, under anesthesia and during high central thalamic (CT) DBS (left) or high ventral-lateral thalamic (VL) DBS (right); normalized probability distribution and two-dimensional normalized histograms

Average life time of brain states for the awake, anesthesia and high central thalamic (CT) DBS condition (A) or high ventral-lateral thalamic (VL) DBS (B) for 7 brain states obtained by k-means clustering. Error bars stand for 1 SEM. Normalized probability distribution of all Z values for the functional brain state 1 (the least similar to the structural brain connectivity) and functional brain state 7 (the most similar to the structural brain connectivity) for awake, anesthesia and high CT-DBS resting state pooled together. Similar results were obtained regardless the inputs conditions for the clustering (C). Two-dimensional normalized histograms for functional brain state 1 and functional brain state 7 for the clustering of awake, anesthesia and high CT-DBS condition. (D) Z values as a function of distance between pairs of regions of interest for brain state 1 (upper right) and brain state 7 (lower right) for the clustering of awake, anesthesia and high CT-DBS condition.

fMRI activations during the auditory “Local-global” experiment

Local effect: High CT- DBS > Anesthesia

Individual results: Monkey N

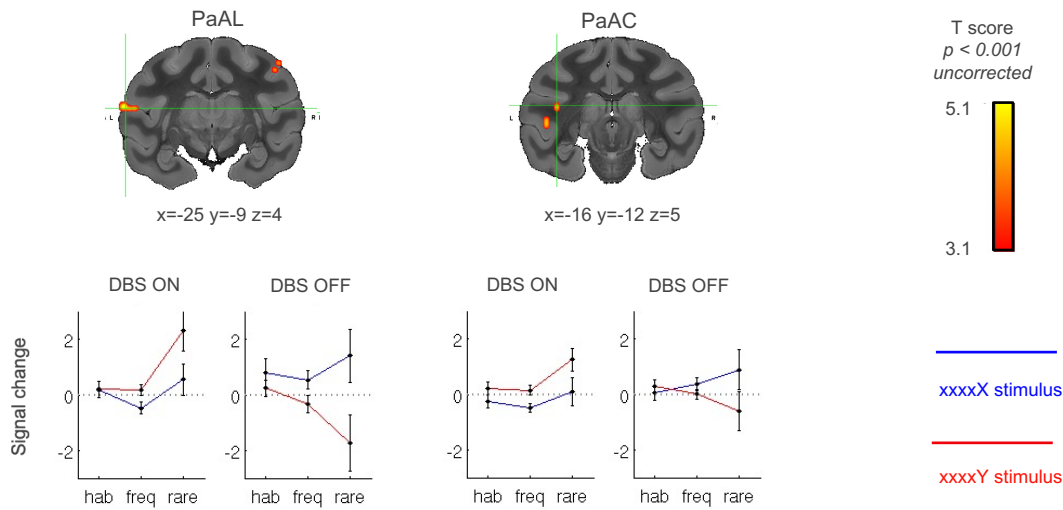
A.



Left Hemisphere

Right Hemisphere

B.



C.

Individual results: Monkey T



Left Hemisphere

Right Hemisphere

D.

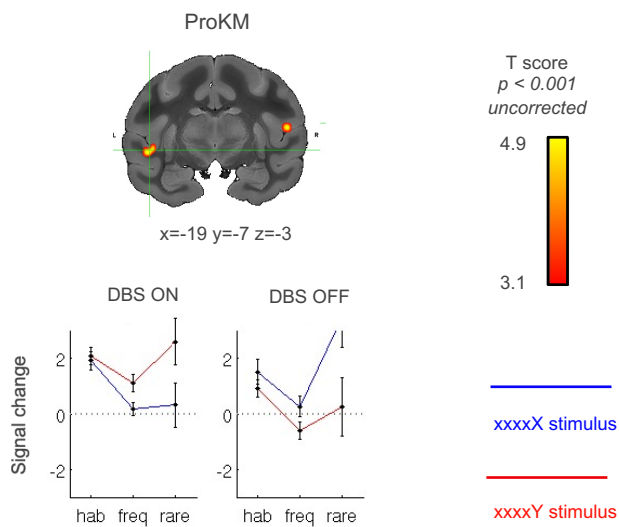


Figure S10: Activations maps for the local effect showing stronger activations under high central thalamic (CT) DBS compared to anesthesia state

Activation maps for the local effect showing stronger activations under high CT-DBS compared to anesthesia state (A, C). fMRI signal changes in areas responsive to the local effect (green cursor) (B,D). Individual results for monkey N (top panel - A,B) and monkey T (lower panel - C,D), $p < 0.001$, uncorrected.

PaAL, parauditory cortex, lateral part; *PaAc*, parauditory cortex, caudal part; *ProKM*, auditory cortex, prokonio cortex, medial part.

fMRI activations during the auditory “Local-global” experiment

Global effect: High CT-DBS > Anesthesia

Individual results: Monkey N

A.



Left Hemisphere

Right Hemisphere

B.

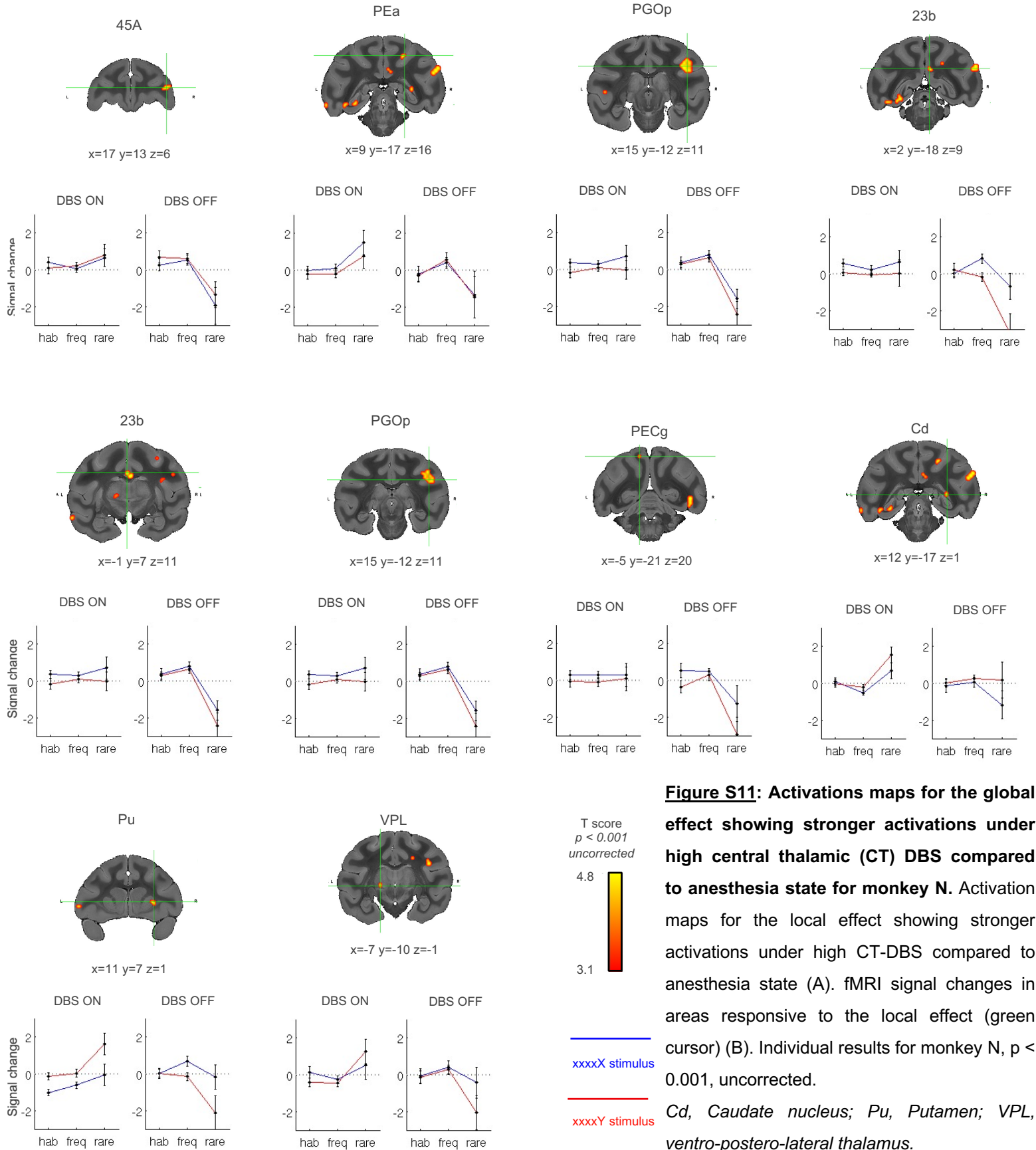


Figure S11: Activations maps for the global effect showing stronger activations under high central thalamic (CT) DBS compared to anesthesia state for monkey N. Activation maps for the local effect showing stronger activations under high CT-DBS compared to anesthesia state (A). fMRI signal changes in areas responsive to the local effect (green cursor) (B). Individual results for monkey N, $p < 0.001$, uncorrected.

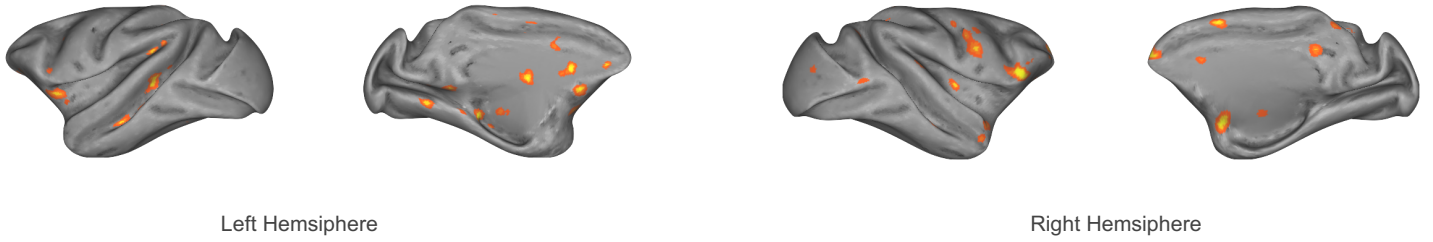
Cd, Caudate nucleus; *Pu*, Putamen; *VPL*, ventro-postero-lateral thalamus.

fMRI activations during the auditory "Local-global" experiment

Global effect: High CT-DBS > Anesthesia

Individual results: Monkey T

A.



B.

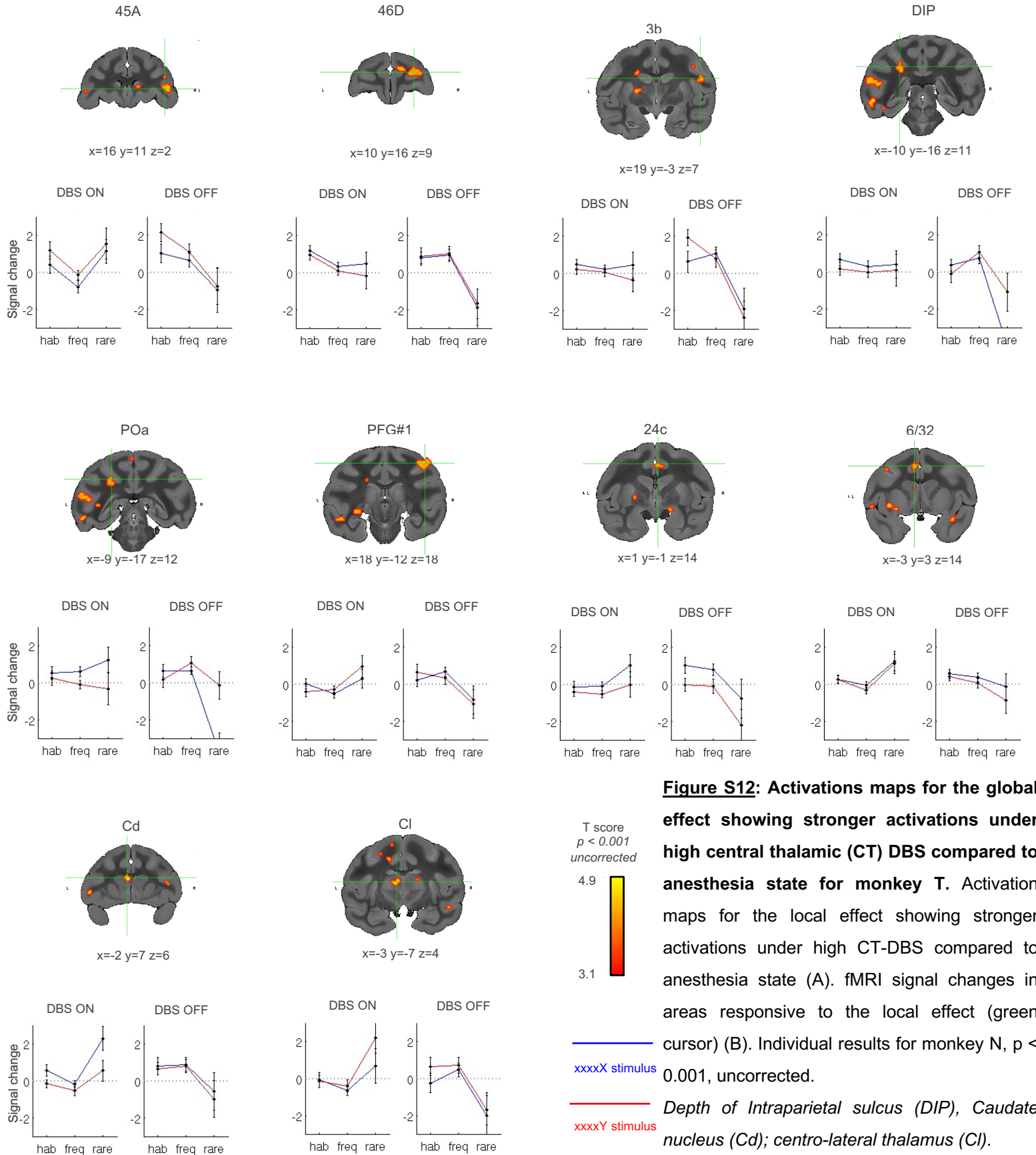


Figure S12: Activations maps for the global effect showing stronger activations under high central thalamic (CT) DBS compared to anesthesia state for monkey T. Activation maps for the local effect showing stronger activations under high CT-DBS compared to anesthesia state (A). fMRI signal changes in areas responsive to the local effect (green cursor) (B). Individual results for monkey N, $p < 0.001$, uncorrected. Depth of Intraparietal sulcus (DIP), Caudate nucleus (Cd); centro-lateral thalamus (CI).

Thalamic nuclei	Abbreviation	CT-DBS				VL-DBS			
		low		high		low		high	
		monkey N	monkey T	monkey N	monkey T	monkey N	monkey T	monkey N	monkey T
Ventral-Postero-Medial	VPM	X	X	X	X		X	X	X
Centro-Median, Medial part	CMM	X	X	X	X				X
Centro-Median, Lateral part	CML	X	X	X	X				X
Medio-Dorsal, Central part	MDC	X	X	X	X		X	X	X
Ventral-Postero-Lateral	VPL			X	X	X	X	X	X
Medio-Dorsal, Lateral part	MDL			X	X	X	X	X	X
Ventral-Lateral, Medial part	VLM			X	X	X	X	X	X
Medio-Dorsal-Medial part	MDM			X	X				X
Inter-Medio-Dorsal	IMD			X	X				
Centro-Lateral	CL					X	X	X	X
Medio-Dorsal, Dorsal part	MDD					X		X	
Lateral-Dorsal, Superficial part	LDSF							X	
Ventral-Anterior, Lateral part	VAL							X	X
Ventral-Lateral, Lateral part	VLL							X	

Table S1: Simulation of the stimulated thalamic nuclei around the DBS lead using the LEAD DBS macaque toolbox

Estimation of the thalamic nuclei that were included in the volume of activated tissue around the DBS lead active contact(71) for monkey N and monkey T across the four experimental conditions (low central thalamic (CT) DBS, high CT-DBS, low ventral thalamic (VL) DBS and high VL-DBS).

Physiological data during the resting state fMRI experiments

Condition	Animal	Heart Rate (HR in bpm)	Oxygen Saturation (SpO ₂ in %)	Blood pressure			Respiration Rate (RR in breath/min)	End-tidal CO ₂ (EtCO ₂ in mmHg)	Temperature (T in °C)
				Systolic (SBP in mmHg)	Diastolic (DBP in mmHg)	Mean (MBP in mmHg)			
Anesthesia	monkey N	111 ± 9	98 ± 2	116 ± 10	59 ± 7	87 ± 9	20 ± 1	41 ± 2	37.0 ± 0.7
	monkey T	116 ± 17	97 ± 3	95 ± 8	46 ± 7	68 ± 9	18 ± 2	37 ± 2	38.3 ± 0.7
Low CT-DBS	monkey N	134 ± 13	99 ± 1	128 ± 13	70 ± 12	101 ± 13	21 ± 2	43 ± 2	37.1 ± 0.3
	monkey T	123 ± 18	99 ± 1	98 ± 12	47 ± 6	71 ± 8	18 ± 2	39 ± 2	38.6 ± 0.5
High CT-DBS	monkey N	164 ± 16	98 ± 2	129 ± 15	71 ± 15	99 ± 14	21 ± 1	45 ± 3	37.3 ± 0.4
	monkey T	181 ± 19	98 ± 2	133 ± 12	74 ± 12	104 ± 13	19 ± 2	43 ± 2	39.6 ± 0.4
Low VL-DBS	monkey T	145 ± 3	98 ± 2	113 ± 14	54 ± 12	81 ± 12	19 ± 2	38 ± 1	39.0 ± 0.5
High VL-DBS	monkey T	163 ± 14	98 ± 1	127 ± 15	71 ± 19	99 ± 19	18 ± 2	42 ± 2	39.1 ± 0.6

Table S2: Physiological data during the fMRI resting-state experiments.

Oxygen saturation (SpO₂); systolic, diastolic and mean blood pressure (respectively SBP, DBP, MBP); respiration rate (RR); end-tidal CO₂ (EtCO₂) and temperature (T) for each animal (monkey N and T) under general anesthesia, general anesthesia plus low central thalamic (CT) DBS, general anesthesia plus high CT-DBS, general anesthesia plus low ventral-lateral thalamic (VL) DBS and general anesthesia plus high VL-DBS.

fMRI activations during the DBS block-design experiment

CT-DBS

VL-DBS

low

high

low

high

Area	Abbreviation	Hemisphere	T score	p value	T score	p value	T score	p value	T score	p value
<i>Orbitofrontal cortex</i>										
Area 13 of cortex	13	Right		<i>n.s.</i>	19.15	$p_{FWE} < 1 \times 10^{-12}$		<i>n.s.</i>		<i>n.s.</i>
Area 13a of cortex	13a	Left	5.11	$p_{FW} = 7.4 \times 10^{-3}$	5.11	$p_{FWE} = 7.4 \times 10^{-3}$		<i>n.s.</i>		<i>n.s.</i>
Area 14o	14o	Right		<i>n.s.</i>	5.91	$p_{FWE} = 1.21 \times 10^{-4}$		<i>n.s.</i>		<i>n.s.</i>
Area 25 of cortex	25	Left	5.48	$p_{FWE} = 9.66 \times 10^{-4}$	5.48	$p_{FWE} = 9.66 \times 10^{-4}$		<i>n.s.</i>		<i>n.s.</i>
Orbitofrontal cortex	OPro	Right		<i>n.s.</i>	19.15	$p_{FWE} < 1 \times 10^{-12}$		<i>n.s.</i>		<i>n.s.</i>
<i>Frontal cortex</i>										
Area 4 of cortex (primary motor)	4#1	Left		<i>n.s.</i>	18.47	$p_{FWE} < 1 \times 10^{-12}$		<i>n.s.</i>		<i>n.s.</i>
		Right		<i>n.s.</i>	18.47	$p_{FWE} < 1 \times 10^{-12}$		<i>n.s.</i>	9.94	$p_{FWE} < 1 \times 10^{-12}$
Area 6 of cortex, Medial part (supplementary motor)	6M	Left		<i>n.s.</i>	19.15	$p_{FWE} < 1 \times 10^{-12}$		<i>n.s.</i>		<i>n.s.</i>
		Right		<i>n.s.</i>	19.15	$p_{FWE} < 1 \times 10^{-12}$		<i>n.s.</i>	5.77	$p_{FWE} = 2.67 \times 10^{-4}$
Area 6 of cortex, DorsoCaudal part (Matellis F2)	6DC	Right		<i>n.s.</i>	18.47	$p_{FWE} < 1 \times 10^{-12}$		<i>n.s.</i>	12.81	$p_{FWE} < 1.0 \times 10^{-12}$
Area 6 of cortex, DorsoRostral part (Matellis F7)	6DR	Right		<i>n.s.</i>	19.15	$p_{FWE} < 1 \times 10^{-12}$		<i>n.s.</i>	12.81	$p_{FWE} < 1.0 \times 10^{-12}$
Area 6 of cortex, Ventral part, Caudal subdivision (Matellis F4)	6VC	Left		<i>n.s.</i>	6.54	$p_{FWE} = 2.37 \times 10^{-6}$		<i>n.s.</i>		<i>n.s.</i>
		Right		<i>n.s.</i>	18.47	$p_{FWE} < 1 \times 10^{-12}$		<i>n.s.</i>		<i>n.s.</i>
Area 6 of cortex, Ventral part, Rostral subdivision (Matellis F5)	6VR	Left		<i>n.s.</i>	6.16	$p_{FWE} = 2.54 \times 10^{-5}$		<i>n.s.</i>		<i>n.s.</i>
		Right		<i>n.s.</i>	19.15	$p_{FWE} < 1 \times 10^{-12}$		<i>n.s.</i>	5.61	$p_{FWE} = 6.87 \times 10^{-4}$
Area 6/32 of cortex	6 32	Left		<i>n.s.</i>	19.15	$p_{FWE} < 1 \times 10^{-12}$		<i>n.s.</i>		<i>n.s.</i>
		Right	4.96	$p_{FWE} = 1.56 \times 10^{-2}$	19.15	$p_{FWE} < 1 \times 10^{-12}$		<i>n.s.</i>	5.77	$p_{FWE} = 2.67 \times 10^{-4}$
Area 8A of cortex	8A	Left		<i>n.s.</i>	5.79	$p_{FWE} = 2.41 \times 10^{-4}$		<i>n.s.</i>		<i>n.s.</i>
		Right		<i>n.s.</i>	18.47	$p_{FWE} < 1 \times 10^{-12}$		<i>n.s.</i>	12.81	$p_{FWE} < 1.0 \times 10^{-12}$
Area 8of cortex, AnteroDorsal part	8AD	Left		<i>n.s.</i>	7.10	$p_{FWE} = 5.04 \times 10^{-8}$		<i>n.s.</i>		<i>n.s.</i>
		Right	5.92	$p_{FWE} = 7.47 \times 10^{-5}$	18.47	$p_{FWE} < 1 \times 10^{-12}$		<i>n.s.</i>	12.81	$p_{FWE} < 1.0 \times 10^{-12}$
Area 8 of cortex, AnteroVentral part	8AV	Right		<i>n.s.</i>	19.15	$p_{FWE} < 1 \times 10^{-12}$		<i>n.s.</i>	12.81	$p_{FWE} < 1.0 \times 10^{-12}$
Area 8B of cortex	8B	Left		<i>n.s.</i>	7.10	$p_{FWE} = 5.04 \times 10^{-8}$		<i>n.s.</i>		<i>n.s.</i>
		Right		<i>n.s.</i>	18.47	$p_{FWE} < 1 \times 10^{-12}$		<i>n.s.</i>		<i>n.s.</i>
Area 8/32 of cortex	8 32	Right		<i>n.s.</i>	19.15	$p_{FWE} < 1 \times 10^{-12}$		<i>n.s.</i>		<i>n.s.</i>
Area 9 of cortex, Medial part	9M	Left		<i>n.s.</i>	18.47	$p_{FWE} < 1 \times 10^{-12}$		<i>n.s.</i>		<i>n.s.</i>
		Right		<i>n.s.</i>	14.54	$p_{FWE} < 1 \times 10^{-12}$		<i>n.s.</i>		<i>n.s.</i>
Area 9/32 of cortex	9 32	Right	7.72	$p_{FDR} = 2.84 \times 10^{-10}$		$p_{FDR} = 2.84 \times 10^{-10}$		<i>n.s.</i>		<i>n.s.</i>
Area 9/46 of cortex	9 46	Right	6.98	$p_{FWE} = 7.08 \times 10^{-8}$	19.15	$p_{FWE} < 1 \times 10^{-12}$		<i>n.s.</i>	9.94	$p_{FWE} < 1 \times 10^{-12}$
Area 44 of cortex	44	Left		<i>n.s.</i>	6.66	$p_{FWE} = 1.03 \times 10^{-6}$		<i>n.s.</i>		<i>n.s.</i>
		Right		<i>n.s.</i>	19.15	$p_{FWE} < 1 \times 10^{-12}$		<i>n.s.</i>	9.94	$p_{FWE} < 1 \times 10^{-12}$
Area 45A of cortex	45A	Right	7.30	$p_{FWE} = 7.23 \times 10^{-9}$	19.15	$p_{FWE} < 1 \times 10^{-12}$		<i>n.s.</i>	9.94	$p_{FWE} < 1.0 \times 10^{-12}$
Area 45B of cortex	45B	Left		<i>n.s.</i>	6.66	$p_{FWE} = 1.03 \times 10^{-6}$		<i>n.s.</i>		<i>n.s.</i>
		Right	7.30	$p_{FWE} = 7.23 \times 10^{-9}$	19.15	$p_{FWE} < 1 \times 10^{-12}$		<i>n.s.</i>	9.94	$p_{FWE} < 1.0 \times 10^{-12}$
Area 46D of cortex	46D	Right	6.98	$p_{FWE} = 7.08 \times 10^{-8}$		$p_{FWE} = 7.08 \times 10^{-8}$		<i>n.s.</i>		<i>n.s.</i>
Area 46V of cortex	46V	Left		<i>n.s.</i>	5.27	$p_{FWE} = 4.68 \times 10^{-3}$		<i>n.s.</i>		<i>n.s.</i>
		Right	6.98	$p_{FWE} = 7.08 \times 10^{-8}$		$p_{FWE} = 7.08 \times 10^{-8}$		<i>n.s.</i>		<i>n.s.</i>
Area 47 (old 12) of cortex, Lateral part	47L	Left	6.27	$p_{FWE} = 8.28 \times 10^{-6}$		$p_{FWE} = 8.28 \times 10^{-6}$		<i>n.s.</i>		<i>n.s.</i>
Area 47 (old 12) of cortex, Orbital part	47O	Left		<i>n.s.</i>	6.66	$p_{FWE} = 1.03 \times 10^{-6}$		<i>n.s.</i>		<i>n.s.</i>
Area ProM (promotor)	ProM#1	Left		<i>n.s.</i>	11.36	$p_{FWE} < 1 \times 10^{-12}$		<i>n.s.</i>		<i>n.s.</i>

fMRI activations during the DBS block-design experiment

low CT-DBS high

low VL-DBS high

Area	Abbreviation	Hemisphere	low CT-DBS		high CT-DBS		low VL-DBS		high VL-DBS	
			T score	p value	T score	p value	T score	p value	T score	p value
<i>Parietal cortex</i>										
Area 1 of cortex (somatosensory)	1	Left		n.s	18.47	$p_{FWE} < 1x10^{-12}$		n.s		n.s
		Right		n.s	18.47	$p_{FWE} < 1x10^{-12}$		n.s	9.94	$p_{FWE} < 1x10^{-12}$
Area 2 of cortex (somatosensory)	2#1	Left		n.s	18.47	$p_{FWE} < 1x10^{-12}$		n.s		n.s
		Right		n.s	18.47	$p_{FWE} < 1x10^{-12}$		n.s	12.81	$p_{FWE} < 1x10^{-12}$
Area 2 of cortex, Vesticular part	2Ve			n.s	18.47	$p_{FWE} < 1x10^{-12}$		n.s		n.s
Area 3a of cortex (somatosensory)	3a	Left		n.s	18.47	$p_{FWE} < 1x10^{-12}$		n.s		n.s
		Right		n.s	18.47	$p_{FWE} < 1x10^{-12}$		n.s	9.94	$p_{FWE} < 1x10^{-12}$
Area 3b of cortex (somatosensory)	3b	Right		n.s	18.47	$p_{FWE} < 1x10^{-12}$		n.s	16.05	$p_{FWE} < 1.0x10^{-12}$
Depth IntraParietal area	DIP	Right		n.s	18.47	$p_{FWE} < 1x10^{-12}$		n.s		n.s
Dorsal parietal area	Dpt	Left		n.s	7.27	$p_{FWE} = 1.53x10^{-8}$		n.s	7.23	$p_{FWE} = 2.09x10^{-8}$
		Right	5.73	$p_{FWE} = 2.32x10^{-4}$	18.47	$p_{FWE} < 1x10^{-12}$		n.s		n.s
OccipitoParietal area	OPt	Left		n.s	12.59	$p_{FWE} < 1x10^{-12}$		n.s		n.s
		Right		n.s	18.47	$p_{FWE} < 1x10^{-12}$		n.s		n.s
Parietal area PE	PE	Right	6.51	$p_{FWE} = 1.72x10^{-6}$	18.47	$p_{FWE} < 1x10^{-12}$		n.s	7.26	$p_{FWE} = 1.72x10^{-8}$
Parietal area PE, Caudal part	PEC	Right		n.s	5.39	$p_{FWE} = 2.3x10^{-3}$		n.s	7.85	$p_{FWE} = 2.08x10^{-10}$
Parietal area PEa	PEa	Right	6.30	$p_{FWE} = 7.07x10^{-6}$	18.47	$p_{FWE} < 1x10^{-12}$		n.s	7.85	$p_{FWE} = 2.08x10^{-10}$
Parietal area PF (cortex)	PFCx	Right		n.s		n.s		n.s	9.94	$p_{FWE} < 1x10^{-12}$
Parietal area PF, Opercular part	PFOp	Left		n.s	7.27	$p_{FWE} = 1.53x10^{-8}$		n.s		n.s
		Right		n.s	18.47	$p_{FWE} < 1x10^{-12}$		n.s		n.s
Parietal area PG	PG	Right		n.s	12.62	$p_{FWE} < 1x10^{-12}$		n.s		n.s
Parietal area PG, Opercular part	PGOp	Left		n.s	7.27	$p_{FWE} = 1.53x10^{-8}$		n.s		n.s
		Right	7.38	$p_{FWE} = 3.90x10^{-9}$	18.47	$p_{FWE} < 1x10^{-12}$		n.s	5.95	$p_{FWE} = 9.51x10^{-5}$
Parietal area POa, External part	POaE	Right		n.s	18.47	$p_{FWE} < 1x10^{-12}$		n.s		n.s
Parietal area POa, Internal part	POaI	Right		n.s	18.47	$p_{FWE} < 1x10^{-12}$		n.s		n.s
ParietoOccipital associated area in the intraparietal sulcus	POa	Left		n.s	7.27	$p_{FWE} = 1.53x10^{-8}$		n.s		n.s
		Right		n.s	18.47	$p_{FWE} < 1x10^{-12}$		n.s	9.94	$p_{FWE} < 1x10^{-12}$
Posterior Parietal area	PPT	Right		n.s	18.47	$p_{FWE} < 1x10^{-12}$		n.s		n.s
Retrolsular area, Parietal part	RelP	Left		n.s	7.27	$p_{FWE} = 1.53x10^{-8}$		n.s		n.s
		Right		n.s	12.62	$p_{FWE} < 1x10^{-12}$		n.s		n.s
Secondary somatosensory cortex	S2	Left		n.s	7.27	$p_{FWE} = 1.53x10^{-8}$		n.s		n.s
		Right		n.s	18.47	$p_{FWE} < 1x10^{-12}$		n.s	5.66	$p_{FWE} = 5.12x10^{-4}$
Secondary somatosensory cortex, External part	S2E	Right		n.s		n.s		n.s	12.81	$p_{FWE} < 1x10^{-12}$
Secondary somatosensory cortex, Internal part	S2I	Right		n.s	18.47	$p_{FWE} < 1x10^{-12}$		n.s		n.s
Visual area 4, Dorsal part	V4D	Right		n.s	18.47	$p_{FWE} < 1x10^{-12}$		n.s		n.s
Visual area 4, Transitional part	V4T	Right		n.s	18.47	$p_{FWE} < 1x10^{-12}$		n.s		n.s
Visual area 4A	V4A	Left		n.s	7.27	$p_{FWE} = 1.53x10^{-8}$		n.s	7.23	$p_{FWE} = 2.09x10^{-8}$
		Right		n.s	18.47	$p_{FWE} < 1x10^{-12}$		n.s		n.s
<i>Cingulate cortex</i>										
Area 23 of cortex	23	midline	6.84	$p_{FWE} = 1.82x10^{-7}$		n.s		n.s		n.s
Area 23a of cortex	23a	midline		n.s	5.60	$p_{FWE} = 7.36x-4$		n.s		n.s
Area 23b of cortex	23b			n.s	18.47	$p_{FWE} < 1x10^{-12}$		n.s		n.s
		Left		n.s	18.47	$p_{FWE} < 1x10^{-12}$		n.s		n.s
Area 23c of cortex	23c	Right		n.s	18.47	$p_{FWE} < 1x10^{-12}$		n.s		n.s
		Right		n.s	18.47	$p_{FWE} < 1x10^{-12}$		n.s		n.s
Area 24/23b of cortex	24 23b	Right		n.s	18.47	$p_{FWE} < 1x10^{-12}$		n.s		n.s
Area 24/23c of cortex	24 23c	Left		n.s	18.47	$p_{FWE} < 1x10^{-12}$		n.s		n.s
		Right		n.s		n.s		n.s	5.77	$p_{FWE} = 2.67x10^{-4}$
Area 24b of cortex	24b	Left		n.s	18.47	$p_{FWE} < 1x10^{-12}$		n.s		n.s
		Right		n.s	18.47	$p_{FWE} < 1x10^{-12}$		n.s		n.s
Area 24c of cortex	24c	Left		n.s	18.47	$p_{FWE} < 1x10^{-12}$		n.s		n.s
		Right	6.98	$p_{FWE} = 7.08x10^{-8}$	18.47	$p_{FWE} < 1x10^{-12}$		n.s		n.s
Area 24d of cortex	24d	Left		n.s	18.47	$p_{FWE} < 1x10^{-12}$		n.s		n.s
Area 31 of cortex	31	Left		n.s	18.47	$p_{FWE} < 1x10^{-12}$		n.s		n.s
		Right		n.s	18.47	$p_{FWE} < 1x10^{-12}$		n.s		n.s
Area PGM/31 of cortex	PGM 31	midline	5.41	$p_{FWE} = 1.4x10^{-3}$		n.s		n.s		n.s
Parietal area PE, Cingulate part	PECg	Right		n.s	18.47	$p_{FWE} < 1x10^{-12}$		n.s		n.s

fMRI activations during the DBS block-design experiment

CT-DBS

VL-DBS

low

high

low

high

Area	Abbreviation	Hemisphere	CT-DBS low		CT-DBS high		VL-DBS low		VL-DBS high	
			T score	p value	T score	p value	T score	p value	T score	p value
<i>Temporal cortex</i>										
Area PG associated, region of superior temporal sulcus	PGa	Right		<i>n.s.</i>	18.47	$p_{FWE} < 1 \times 10^{-12}$		<i>n.s.</i>		<i>n.s.</i>
Auditory Koniocortex, Lateral part	AKL	Left		<i>n.s.</i>	7.27	$p_{FWE} = 1.53 \times 10^{-8}$		<i>n.s.</i>		<i>n.s.</i>
		Right		<i>n.s.</i>	18.47	$p_{FWE} < 1 \times 10^{-12}$		<i>n.s.</i>		<i>n.s.</i>
Auditory Koniocortex, Medial part	AKM	Left		<i>n.s.</i>	7.27	$p_{FWE} = 1.53 \times 10^{-8}$		<i>n.s.</i>		<i>n.s.</i>
		Right		<i>n.s.</i>	18.47	$p_{FWE} < 1 \times 10^{-12}$		<i>n.s.</i>		<i>n.s.</i>
Fundus of Superior Temporal sulcus	FST	Left		<i>n.s.</i>	6.48	$p_{FWE} = 3.37 \times 10^{-6}$		<i>n.s.</i>		<i>n.s.</i>
		Right		<i>n.s.</i>	18.47	$p_{FWE} < 1 \times 10^{-12}$		<i>n.s.</i>		<i>n.s.</i>
Medial Superior Temporal area	MST	Left		<i>n.s.</i>	7.27	$p_{FWE} = 1.53 \times 10^{-8}$		<i>n.s.</i>		<i>n.s.</i>
		Right		<i>n.s.</i>	18.47	$p_{FWE} < 1 \times 10^{-12}$		<i>n.s.</i>		<i>n.s.</i>
Middle Temporal area (visual area 5)	MT	Left		<i>n.s.</i>	7.27	$p_{FWE} = 1.53 \times 10^{-8}$		<i>n.s.</i>		<i>n.s.</i>
		Right		<i>n.s.</i>	18.47	$p_{FWE} < 1 \times 10^{-12}$		<i>n.s.</i>	6.08	$p_{FWE} = 4.26 \times 10^{-5}$
ParaAuditory area, Caudal part	PaAC	Left		<i>n.s.</i>	7.27	$p_{FWE} = 1.53 \times 10^{-8}$		<i>n.s.</i>		<i>n.s.</i>
		Right		<i>n.s.</i>	20.85	$p_{FWE} < 1 \times 10^{-12}$		<i>n.s.</i>		<i>n.s.</i>
ParaAuditory area, lateral part	PaAL	Right		<i>n.s.</i>	18.47	$p_{FWE} < 1 \times 10^{-12}$		<i>n.s.</i>		<i>n.s.</i>
ParaAuditory area, Rostral part	PaAR	Right		<i>n.s.</i>	12.62	$p_{FWE} < 1 \times 10^{-12}$		<i>n.s.</i>		<i>n.s.</i>
ProKoniocortex	ProK	Right		<i>n.s.</i>	18.47	$p_{FWE} < 1 \times 10^{-12}$		<i>n.s.</i>		<i>n.s.</i>
Retroinsular area	Rel	Right		<i>n.s.</i>	18.47	$p_{FWE} < 1 \times 10^{-12}$		<i>n.s.</i>		<i>n.s.</i>
Retroinsular area, Temporal part	RelT	Left		<i>n.s.</i>	7.27	$p_{FWE} = 1.53 \times 10^{-8}$		<i>n.s.</i>		<i>n.s.</i>
		Right		<i>n.s.</i>	18.47	$p_{FWE} < 1 \times 10^{-12}$		<i>n.s.</i>		<i>n.s.</i>
Superior Temporal sulcus area 1	ST1	Right		<i>n.s.</i>	8.83	$p_{FWE} < 1 \times 10^{-12}$		<i>n.s.</i>		<i>n.s.</i>
Superior Temporal area, gyral part	ST2g	Left		<i>n.s.</i>	11.36	$p_{FWE} < 1 \times 10^{-12}$		<i>n.s.</i>		<i>n.s.</i>
Superior Temporal area, sulcal part	ST2s	Left		<i>n.s.</i>	11.36	$p_{FWE} < 1 \times 10^{-12}$		<i>n.s.</i>		<i>n.s.</i>
Temporal area TAa	TAa	Right		<i>n.s.</i>	18.47	$p_{FWE} < 1 \times 10^{-12}$		<i>n.s.</i>		<i>n.s.</i>
Temporal area TE, Medial part	TEM	Right		<i>n.s.</i>	18.47	$p_{FWE} < 1 \times 10^{-12}$		<i>n.s.</i>		<i>n.s.</i>
Temporal area TE, OccipitoMedial part	TEOM	Right		<i>n.s.</i>	18.47	$p_{FWE} < 1 \times 10^{-12}$		<i>n.s.</i>		<i>n.s.</i>
Temporal area TEa	TEa	Right		<i>n.s.</i>	18.47	$p_{FWE} < 1 \times 10^{-12}$		<i>n.s.</i>		<i>n.s.</i>
Temporal ParietoOccipital associated area in sts	TPO	Right	7.38	$p_{FWE} = 3.90 \times 10^{-9}$	18.47	$p_{FWE} < 1 \times 10^{-12}$		<i>n.s.</i>		<i>n.s.</i>
Temporal ParietoOccipital associated area in sts, Caudal part	TPOC	Left		<i>n.s.</i>	7.27	$p_{FWE} = 1.53 \times 10^{-8}$		<i>n.s.</i>		<i>n.s.</i>
		Right		<i>n.s.</i>	18.47	$p_{FWE} < 1 \times 10^{-12}$		<i>n.s.</i>		<i>n.s.</i>
Temporoparietal cortex	Tpt	Left	5.02	$p_{FWE} = 1.15 \times 10^{-2}$		<i>n.s.</i>		<i>n.s.</i>		<i>n.s.</i>
		Right	7.38	$p_{FWE} = 3.90 \times 10^{-9}$	18.47	$p_{FWE} < 1 \times 10^{-12}$		<i>n.s.</i>		<i>n.s.</i>
<i>Occipital cortex</i>										
Visual area 1 (primary visual cortex)	V1	Left		<i>n.s.</i>		<i>n.s.</i>		<i>n.s.</i>	9.22	$p_{FWE} < 1 \times 10^{-12}$
		Right	7.22	$p_{FWE} = 1.26 \times 10^{-8}$	18.47	$p_{FWE} < 1 \times 10^{-12}$		<i>n.s.</i>	7.44	$p_{FWE} = 4.52 \times 10^{-9}$
Visual area 2	V2	Left		<i>n.s.</i>	5.70	$p_{FWE} = 4.08 \times 10^{-4}$		<i>n.s.</i>		<i>n.s.</i>
		Right	8.65	$p_{FWE} < 1 \times 10^{-12}$	18.47	$p_{FWE} < 1 \times 10^{-12}$	7.97	$p_{FWE} = 3.94 \times 10^{-11}$	9.27	$p_{FWE} < 1 \times 10^{-12}$
Visual area 3, Dorsal part	V3D	Right		<i>n.s.</i>	18.47	$p_{FWE} < 1 \times 10^{-12}$		<i>n.s.</i>		<i>n.s.</i>
Visual area 3A	V3A	Left		<i>n.s.</i>	7.27	$p_{FWE} = 1.53 \times 10^{-8}$		<i>n.s.</i>		<i>n.s.</i>
		Right		<i>n.s.</i>		<i>n.s.</i>		<i>n.s.</i>	5.63	$p_{FWE} = 6.10 \times 10^{-4}$
<i>Insular cortex</i>										
Dysgranular Insular cortex	DI	Left		<i>n.s.</i>	6.18	$p_{FWE} = 2.29 \times 10^{-5}$		<i>n.s.</i>		<i>n.s.</i>
		Right		<i>n.s.</i>	12.62	$p_{FWE} < 1 \times 10^{-12}$		<i>n.s.</i>		<i>n.s.</i>
Granular Insular cortex	GI	Left		<i>n.s.</i>	7.52	$p_{FWE} = 2.54 \times 10^{-9}$		<i>n.s.</i>		<i>n.s.</i>
		Right		<i>n.s.</i>	18.47	$p_{FWE} < 1 \times 10^{-12}$		<i>n.s.</i>	5.32	$p_{FWE} = 1.54 \times 10^{-2}$
Insular Proisocortex	IPro	Right		<i>n.s.</i>	12.32	$p_{FWE} < 1 \times 10^{-12}$		<i>n.s.</i>		<i>n.s.</i>
<i>Striatum</i>										
Caudate nucleus	Cd	Left		<i>n.s.</i>	18.47	$p_{FWE} < 1 \times 10^{-12}$		<i>n.s.</i>		<i>n.s.</i>
		Right	6.34	$p_{FWE} = 5.50 \times 10^{-6}$	18.47	$p_{FWE} < 1 \times 10^{-12}$		<i>n.s.</i>		<i>n.s.</i>
Putamen	Pu	Left	5.30	$p_{FWE} = 2.6 \times 10^{-3}$	18.47	$p_{FWE} < 1 \times 10^{-12}$		<i>n.s.</i>		<i>n.s.</i>
		Right	13.68	$p_{FWE} < 1 \times 10^{-12}$	18.47	$p_{FWE} < 1 \times 10^{-12}$		<i>n.s.</i>	5.30	$p_{FWE} = 3.7 \times 10^{-3}$

fMRI activations during the DBS block-design experiment

Area	Abbreviation	Hemisphere	CT-DBS				VL-DBS			
			low		high		low		high	
			T score	p value	T score	p value	T score	p value	T score	p value
<i>Thalamus</i>										
Lateral Geniculate Nucleus	LGN	Right		<i>n.s.</i>	18.47	$p_{FWE} < 1 \times 10^{-12}$		<i>n.s.</i>		<i>n.s.</i>
Lateral pulvinar	Lpul	Left		<i>n.s.</i>	5.20	$p_{FWE} = 1.44 \times 10^{-2}$		<i>n.s.</i>		<i>n.s.</i>
Medial Geniculate nucleus, Ventral part	MGV	Right		<i>n.s.</i>	5.87	$p_{FWE} = 1.55 \times 10^{-4}$		<i>n.s.</i>		<i>n.s.</i>
Medial pulvinar	Mpul	Right	5.59	$p_{FWE} = 5.17 \times 10^{-4}$	11.07	$p_{FWE} < 1 \times 10^{-12}$		<i>n.s.</i>		<i>n.s.</i>
MedioDorsal thalamic nucleus, Central part	MDC	Left	5.65	$p_{FWE} = 3.56 \times 10^{-4}$	11.07	$p_{FWE} < 1 \times 10^{-12}$		<i>n.s.</i>		<i>n.s.</i>
MedioDorsal thalamic nucleus, Dorsal part	MDD	Left		<i>n.s.</i>	11.07	$p_{FWE} < 1 \times 10^{-12}$		<i>n.s.</i>		<i>n.s.</i>
MedioDorsal thalamic nucleus, Medial part	MDM	Left		<i>n.s.</i>	11.07	$p_{FWE} < 1 \times 10^{-12}$		<i>n.s.</i>		<i>n.s.</i>
Paraventricular thalamic nucleus	PV	Left		<i>n.s.</i>	11.07	$p_{FWE} < 1 \times 10^{-12}$		<i>n.s.</i>		<i>n.s.</i>
		Right		<i>n.s.</i>	11.07	$p_{FWE} < 1 \times 10^{-12}$		<i>n.s.</i>		<i>n.s.</i>
Paraventricular Thalamus	PVT	Left		<i>n.s.</i>	11.07	$p_{FWE} < 1 \times 10^{-12}$		<i>n.s.</i>		<i>n.s.</i>
		Right		<i>n.s.</i>	11.07	$p_{FWE} < 1 \times 10^{-12}$		<i>n.s.</i>		<i>n.s.</i>
Reticular thalamic nucleus	R#4	Right		<i>n.s.</i>	5.91	$p_{FWE} = 1.22 \times 10^{-4}$		<i>n.s.</i>		<i>n.s.</i>
<i>Hypothalamus</i>										
Hypothalamus	Hy	Left		<i>n.s.</i>	14.54	$p_{FWE} < 1 \times 10^{-12}$		<i>n.s.</i>		<i>n.s.</i>
<i>Pallidum</i>										
External Globus Pallidus	EGP	Left		<i>n.s.</i>	5.25	$p_{FWE} = 4.9 \times 10^{-3}$		<i>n.s.</i>		<i>n.s.</i>
<i>Paraseptal subpallium</i>										
Accumbens nucleus, Core	AcbC	Left		<i>n.s.</i>	18.47	$p_{FWE} < 1 \times 10^{-12}$		<i>n.s.</i>		<i>n.s.</i>
		Right		<i>n.s.</i>	18.47	$p_{FWE} < 1 \times 10^{-12}$		<i>n.s.</i>		<i>n.s.</i>
Accumbens nucleus, Shell	AcbSh	Left		<i>n.s.</i>	18.47	$p_{FWE} < 1 \times 10^{-12}$		<i>n.s.</i>		<i>n.s.</i>
		Right		<i>n.s.</i>	18.47	$p_{FWE} < 1 \times 10^{-12}$		<i>n.s.</i>		<i>n.s.</i>
Basal nucleus, Meynert	BM	Left		<i>n.s.</i>	14.54	$p_{FWE} < 1 \times 10^{-12}$		<i>n.s.</i>		<i>n.s.</i>
Substantia Innominata	SI	Left		<i>n.s.</i>	15.09	$p_{FWE} < 1 \times 10^{-12}$		<i>n.s.</i>		<i>n.s.</i>
<i>Subpallial amygdala</i>										
Anterior Amygdaloid area	AA	Left		<i>n.s.</i>	14.54	$p_{FWE} < 1 \times 10^{-12}$		<i>n.s.</i>		<i>n.s.</i>
Bed nucleus of the Stria Terminalis IntraAmygdaloid	BSTIA	Left		<i>n.s.</i>	15.09	$p_{FWE} < 1 \times 10^{-12}$		<i>n.s.</i>		<i>n.s.</i>
Central amygdaloid nucleus, Lateral division	CeL	Left		<i>n.s.</i>	14.54	$p_{FWE} < 1 \times 10^{-12}$		<i>n.s.</i>		<i>n.s.</i>
Central amygdaloid nucleus, Medial division	CeM	Left		<i>n.s.</i>	14.54	$p_{FWE} < 1 \times 10^{-12}$		<i>n.s.</i>		<i>n.s.</i>
<i>Ventral pallium</i>										
BasoMedial amygdaloid nucleus	BM	Left		<i>n.s.</i>	14.54	$p_{FWE} < 1 \times 10^{-12}$		<i>n.s.</i>		<i>n.s.</i>
Medial amygdaloid nucleus	Me	Left		<i>n.s.</i>	14.54	$p_{FWE} < 1 \times 10^{-12}$		<i>n.s.</i>		<i>n.s.</i>
<i>Midbrain</i>										
Midbrain	MD	Left	5.83	$p_{FWE} = 1.25 \times 10^{-4}$	8.32	$p_{FWE} < 1 \times 10^{-12}$		<i>n.s.</i>		<i>n.s.</i>
		Right		<i>n.s.</i>	11.07	$p_{FWE} < 1 \times 10^{-12}$		<i>n.s.</i>		<i>n.s.</i>
<i>Cerebellum</i>										
Cerebellum	Cb	Left	8.38	$p_{FWE} < 1 \times 10^{-12}$	13.74	$p_{FWE} < 1 \times 10^{-12}$		<i>n.s.</i>	7.53	$p_{FWE} = 2.29 \times 10^{-9}$
		Right	8.41	$p_{FWE} < 1 \times 10^{-12}$	14.03	$p_{FWE} < 1 \times 10^{-12}$		<i>n.s.</i>	5.31	$p_{FWE} = 1.8 \times 10^{-3}$

Table S3: fMRI activations during low central thalamic (CT), high CT-DBS, low ventral-lateral thalamic (VL) and high VL-DBS

Thalamic DBS-induced fMRI activity during the electrical stimulation block-design experiment, $p < 0.05$, FWE corrected, ns: non significant.

Interpretation of the Bayes Factors

Value	BF10	BF01
<i>>100</i>	Obvious evidence for H1	Obvious evidence for H0
<i>30 to 100</i>	Very strong evidence for H1	Very strong evidence for H0
<i>10 to 30</i>	Strong evidence for H1	Strong evidence for H0
<i>3 to 10</i>	Substantial evidence for H1	Substantial evidence for H0
<i>1 to 3</i>	Anecdotal evidence for H1	Anecdotal evidence for H0
<i>1</i>	No evidence for H1 or H0	
<i>1 to 0.33</i>	Anecdotal evidence for H0	Anecdotal evidence for H1
<i>0.33 to 0.10</i>	Substantial evidence for H0	Substantial evidence for H1
<i>0.10 to 0.03</i>	Strong evidence for H0	Strong evidence for H1
<i>0.03 to 0.01</i>	Very strong evidence for H0	Very strong evidence for H1
<i><0.01</i>	Obvious evidence for H0	Obvious evidence for H1

Table S4: Interpretation of the Bayes Factors.

Value of the Bayes Factor BF10 and BF01 to interpret statistical evidence in favor of the H1 or H0 hypothesis. A BF greater than 3 significantly support the evidence of the tested hypothesis.

fMRI activations during the auditory “Local-Global” experiment

Local effect

Group

Area	Abbreviation	Hemisphere	Anesthesia		High CT-DBS		High CT-BS > Anesthesia	
			T score	p value	T score	p value	T score	p value
<i>Orbitofrontal cortex</i>								
Orbital Proisocortex	OPro	Left	4.44	$p_{FDR} = 0.047$		<i>n.s.</i>		<i>n.s.</i>
<i>Parietal cortex</i>								
Area 3b of cortex (somatosensory)	3b	Left	4.59	$p_{FDR} = 0.039$		<i>n.s.</i>		<i>n.s.</i>
Parietal area PG#1	PG#1	Right		<i>n.s.</i>		<i>n.s.</i>	4.24	$p_{FDR} = 0.044$
Visual area 4, Ventral part	V4V	Left		<i>n.s.</i>	4.80	$p_{FDR} = 0.049$		<i>n.s.</i>
<i>Cingulate cortex</i>								
Parietal area PE, Cingulate part	PECg	Left	4.80	$p_{FDR} = 0.037$		<i>n.s.</i>		<i>n.s.</i>
<i>Temporal cortex</i>								
Fundus of Superior Temporal sulcus	FST	Left	4.27	$p_{FDR} = 0.048$		<i>n.s.</i>		<i>n.s.</i>
ProKoniocortex, Medial part	ProKM	Left		<i>n.s.</i>		<i>n.s.</i>	4.59	$p_{FDR} = 0.040$
Temporal ParietoOccipital associated area in STS	TPO	Right		<i>n.s.</i>		<i>n.s.</i>	4.42	$p_{FDR} = 0.040$
Temporoparietal cortex	Tpt	Right		<i>n.s.</i>		<i>n.s.</i>	4.38	$p_{FDR} = 0.040$
<i>Occipital cortex</i>								
Visual area 1 (primary visual cortex)	V1	Left	4.41	$p_{FDR} = 0.047$		<i>n.s.</i>		<i>n.s.</i>
<i>Striatum</i>								
Caudate nucleus	Cd	Right	4.28	$p_{FDR} = 0.048$		<i>n.s.</i>		<i>n.s.</i>
<i>Midbrain</i>								
Midbrain	MB	Right		<i>n.s.</i>		<i>n.s.</i>	4.13	$p_{FDR} = 0.044$

Table S5: Cerebral activations for the local effect

fMRI activations for the local effect under anesthesia, high central thalamic (CT) DBS and comparison between high CT-DBS > anesthesia. Group results, $p < 0.05$, FDR corrected, ns: non significant.

Global effect

Group

Awake

Anesthesia

High CT-DBS

High CT-DBS
> Anesthesia

Area	Abbreviation	Hemisphere	Tscore	p value	Tscore	p value	Tscore	p value	Tscore	p value
<i>Frontal cortex</i>										
Area 6 of the cortex, DorsoRostral part (Matellis F7)	6DR	Right		n.s		n.s	3.66	$p_{FDR} = 0.023$		n.s
Area 6 of the cortex, Ventral part, Rostral subdivision (Matellis FS)	6VR	Left	3.88	$p_{FDR} = 0.024$		n.s		n.s		n.s
Area 8B of cortex	8B	Right	4.61	$p_{FDR} = 0.014$		n.s		n.s		n.s
Area 8B of cortex	9/46V	Right	3.82	$p_{FDR} = 0.026$		n.s		n.s		n.s
Area 44 of cortex	44	Right	3.58	$p_{FDR} = 0.033$		n.s	4.75	$p_{FDR} = 0.002$	4.57	$p_{FDR} = 0.018$
				n.s		n.s	3.48	$p_{FDR} = 0.031$		n.s
Area 45B of cortex	45B	Left	3.92	$p_{FDR} = 0.024$		n.s	3.94	$p_{FDR} = 0.014$		n.s
		Right		n.s		n.s		n.s	4.57	$p_{FDR} = 0.018$
Area 47 (old 12) of cortex, Orbital part	47O	Left	3.91	$p_{FDR} = 0.024$		n.s	3.83	$p_{FDR} = 0.017$		n.s
		Right	4.09	$p_{FDR} = 0.018$		n.s		n.s		n.s
Area ProM (promotor)	ProM#1	Left		n.s		n.s	3.80	$p_{FDR} = 0.018$		n.s
<i>Parietal cortex</i>										
Area 3a of cortex (somatosensory)	3a	Right		n.s		n.s	5.06	$p_{FDR} = 0.002$		n.s
				n.s		n.s	3.50	$p_{FDR} = 0.030$		n.s
				n.s		n.s	3.47	$p_{FDR} = 0.031$		n.s
Area 3b of cortex (somatosensory)	3b	Left		n.s	4.73	$p_{FDR} = 0.042$		n.s		n.s
Parietal area PFG	PFG#1	Right	4.20	$p_{FDR} = 0.017$		n.s		n.s	4.29	$p_{FDR} = 0.018$
		Left		n.s		n.s		n.s	4.06	$p_{FDR} = 0.028$
Depth IntraParietal area	DIP	Right		n.s		n.s	4.11	$p_{FDR} = 0.010$		n.s
Parietal area PG	PG#1	Right	3.76	$p_{FDR} = 0.028$		n.s	3.98	$p_{FDR} = 0.013$		n.s
Dorsal parietal area	Dpt	Right		n.s		n.s	4.86	$p_{FDR} = 0.002$		n.s
Parietal area Poa, external part	POaE	Left	3.39	$p_{FDR} = 0.042$		n.s	4.51	$p_{FDR} = 0.005$		n.s
Parietal area Poa, internal part	POaI	Left	3.56	$p_{FDR} = 0.034$		n.s		n.s		n.s
Secondary somatosensory cortex	S2	Right		n.s		n.s	3.25	$p_{FDR} = 0.043$		n.s
Secondary soamatosensory cortex, External part	S2E	Right		n.s		n.s	3.70	$p_{FDR} = 0.021$		n.s
Visual area 4, Ventral part	V4V	Left		n.s		n.s	3.39	$p_{FDR} = 0.035$	4.32	$p_{FDR} = 0.018$
Visual area 4, Trnansitional part	V4T	left	4.37	$p_{FDR} = 0.017$		n.s		n.s		n.s
		Left		n.s		n.s	3.58	$p_{FDR} = 0.026$		n.s
		Right		n.s		n.s	3.76	$p_{FDR} = 0.019$		n.s
<i>Cingulate cortex</i>										
Area 23c of cortex	23c	Left		n.s		n.s	3.60	$p_{FDR} = 0.024$		n.s
Area 24b of cortex	23b	Left	3.45	$p_{FDR} = 0.039$		n.s		n.s		n.s
<i>Temporal cortex</i>										
Medial Superior Temporal area	MST	Left		n.s		n.s	3.24	$p_{FDR} = 0.044$		n.s
		Right	4.05	$p_{FDR} = 0.019$	4.46	$p_{FDR} = 0.042$		n.s		n.s
Fundus of Superior Temporal sulcus	FST	Left		n.s		n.s	3.33	$p_{FDR} = 0.039$		n.s
Superior Temporal sulcus area, gyral part	ST2g	Left		n.s		n.s	3.76	$p_{FDR} = 0.019$		n.s
retroinsular area, Temporal part	RelT	Right	3.76	$p_{FDR} = 0.028$		n.s		n.s		n.s
Temporal area TEa	TEa#1	Left	4.12	$p_{FDR} = 0.018$		n.s	4.87	$p_{FDR} = 0.002$	4.78	$p_{FDR} = 0.018$
Temporal area TE, Occipital part	TEO	Left		n.s		n.s	3.27	$p_{FDR} = 0.042$	4.32	$p_{FDR} = 0.018$
Temporal ParietoOccipital associated area in STS	TPO	Left		n.s		n.s	5.37	$p_{FDR} = 0.002$	4.62	$p_{FDR} = 0.018$
				n.s		n.s		n.s	4.27	$p_{FDR} = 0.018$
<i>Occipital cortex</i>										
Visual area 1 (primary visual cortex)	V1	Left		n.s	4.19	$p_{FDR} = 0.042$		n.s		n.s
		Right		n.s	4.33	$p_{FDR} = 0.042$	4.39	$p_{FDR} = 0.006$		n.s
				n.s	4.30	$p_{FDR} = 0.042$	3.90	$p_{FDR} = 0.015$		n.s
Visual area 2	V2	Left		n.s		n.s	3.57	$p_{FDR} = 0.026$		n.s
		Right	3.91	$p_{FDR} = 0.024$	4.53	$p_{FDR} = 0.042$	4.62	$p_{FDR} = 0.004$		n.s
			3.58	$p_{FDR} = 0.033$	4.46	$p_{FDR} = 0.042$		n.s		n.s
Visual area 3, Ventral part	V3V	Right		n.s		n.s	4.27	$p_{FDR} = 0.007$		n.s
Visual area 3, Dorsal part	V3D	Right		n.s		n.s	3.76	$p_{FDR} = 0.019$		n.s
Visual area 3A	V3A	Right		n.s		n.s	3.88	$p_{FDR} = 0.015$		n.s
				n.s		n.s	3.70	$p_{FDR} = 0.021$		n.s

Global effect

Group

Area	Abbreviation	Hemisphere	Awake		Anesthesia		High CT-DBS		High CT-DBS > Anesthesia	
			T score	p value	T score	p value	T score	p value	T score	p value
<i>Striatum</i>										
Caudate nucleus	Cd	Right	4.61	$p_{FDR} = 0.014$		<i>n.s.</i>	4.83	$p_{FDR} = 0.002$		<i>n.s.</i>
Putamen	Pu	Right	4.66	$p_{FDR} = 0.007$		<i>n.s.</i>	4.27	$p_{FDR} = 0.007$		<i>n.s.</i>
				<i>n.s.</i>		<i>n.s.</i>	3.77	$p_{FDR} = 0.019$		<i>n.s.</i>
<i>Thalamus</i>										
Ventral Posterolateral thalamic nucleus	VPL#1	Right			4.72	$p_{FDR} = 0.042$		<i>n.s.</i>		<i>n.s.</i>
Ventral Anterior thalamic nucleus, Medial part	VAM	Left		<i>n.s.</i>		<i>n.s.</i>	3.66	$p_{FDR} = 0.023$		<i>n.s.</i>
MedioDorsal thalamic nucleus, Lateral part	MDL	Left		<i>n.s.</i>		<i>n.s.</i>		<i>n.s.</i>	3.81	$p_{FDR} = 0.039$
Reticular thalamic nucleus	R#4	Left		<i>n.s.</i>		<i>n.s.</i>		<i>n.s.</i>	3.83	$p_{FDR} = 0.037$
Lateral Geniculate Nucleus	LGN	Left		<i>n.s.</i>		<i>n.s.</i>	4.08	$p_{FDR} = 0.011$		<i>n.s.</i>
		Right		<i>n.s.</i>		<i>n.s.</i>	3.49	$p_{FDR} = 0.030$		<i>n.s.</i>
				<i>n.s.</i>		<i>n.s.</i>	3.45	$p_{FDR} = 0.033$		<i>n.s.</i>
<i>Paraseptal subpallium</i>										
Accumbens nucleus	Acb	Left		<i>n.s.</i>		<i>n.s.</i>		<i>n.s.</i>	4.38	$p_{FDR} = 0.018$
<i>Midbrain</i>										
Midbrain	MB	Right		<i>n.s.</i>		<i>n.s.</i>	4.43	$p_{FDR} = 0.006$		<i>n.s.</i>
<i>Cerebellum</i>										
Cerebellum	Cb	Left	4.83	$p_{FDR} = 0.014$		<i>n.s.</i>	3.32	$p_{FDR} = 0.039$		<i>n.s.</i>
		midline		<i>n.s.</i>		<i>n.s.</i>		<i>n.s.</i>	4.21	$p_{FDR} = 0.020$
		Right	3.57	$p_{FDR} = 0.033$		<i>n.s.</i>	3.65	$p_{FDR} = 0.023$		<i>n.s.</i>
				<i>n.s.</i>		<i>n.s.</i>	3.64	$p_{FDR} = 0.023$		<i>n.s.</i>

Table S6: Cerebral activations for the global effect

fMRI activations for the global effect in the awake, anesthesia and high central thalamic (CT) DBS condition and comparison between high CT-DBS versus anesthesia. For the global effect, no regions are significantly different for the awake > high CT-DBS comparison. Group results, $p < 0.05$, FDR corrected, ns: non significant.

Cerebral areas labelling using the CIVM macaque brain atlas Revised version

Level	Labels CIVM	Abbreviations CIVM	Value CIVM		Labels CIVM_R	Abbreviations CIVM_R	Value CIVM_R left	Value CIVM_R
temporal cortex	area 35 of cortex	35	145		AREA 35 OF CORTEX	35	135	136
temporal cortex	area PG associated region of the superior temporal sulcus	PGa	146		REA PG, associated region of superior temporal sulcus	PGa	137	138
temporal cortex	area TF, medial part	TFM	147		AREA TF, medial part	TFM	139	140
temporal cortex	area TL, rostral part (area 36R)	TLR(R36)	148		AREA TL, rostral part (area 36R)	TLR(R36)	141	142
temporal cortex	auditory koniocortex, lateral part	AKL	149		AUDITORY KONIOCORTEX	AK	143	144
temporal cortex	auditory koniocortex, medial part	AKM	150					
temporal cortex	entorhinal cortex, caudal limited part	ECL	151					
temporal cortex	entorhinal cortex, caudal part	EC#2	152					
temporal cortex	entorhinal cortex, lateral part, caudal division	ELC	153					
temporal cortex	entorhinal cortex, lateral part, rostral division	ELR	154		ENTORHINAL CORTEX	ECL	145	146
temporal cortex	entorhinal cortex, rostral part	ER#1	155					
temporal cortex	entorhinal Cx, intermediate part	EI	156					
temporal cortex	entorhinal Cx, olfactory part	EOI	157					
temporal cortex	fundus of superior temporal sulcus	FST	158		FUNDUS OF SUPERIOR TEMPORAL SULCUS	FST	147	148
temporal cortex	intraparietal sulcus associated area in the superior temporal sulcus	IPa	159		INTRAPARIETAL SULCUS ASSOCIATED AREA	IPa	149	150
temporal cortex	medial superior temporal area	MST	160		MEDIAL SUPERIOR TEMPORAL AREA	MST	151	152
temporal cortex	middle temporal area (visual area 5)	MT(V5)	161		MIDDLE TEMPORAL AREA (visual area 5)	MT(V5)	153	154
temporal cortex	paraauditory area, caudal part	PaAC	162		PARAAUDITORY CORTEX	PaA	155	156
temporal cortex	paraauditory area, lateral part	PaAL	163					
temporal cortex	paraauditory cortex, rostral part	PaAR	164					
temporal cortex	parainsular cortex, lateral part	PaIL	165		PARAINSULAR CORTEX	PaI	157	158
temporal cortex	parainsular cortex, medial part	PaIM	166					
temporal cortex	prokoniocortex	ProK	167		PROKONIOCORTEX	ProK	159	160
temporal cortex	prokoniocortex, lateral part	ProKL	168					
temporal cortex	prokoniocortex, medial part	ProKM	169					
temporal cortex	retroinsular area	ReI	170		RETROINSULAR AREA	ReI	161	162
temporal cortex	retroinsular area, temporal part	ReIT	171					
temporal cortex	superior temporal sulcus area 1	ST1	172		SUPERIOR TEMPORAL SULCUS	ST1	163	164
temporal cortex	superior temporal sulcus area 3	ST3	173					
temporal cortex	superior temporal sulcus area, gyral part	ST2g	174					
temporal cortex	superior temporal sulcus area, sulcal part	ST2s	175					
temporal cortex	temporal area Taa	Taa	176		TEMPORAL AREA Taa	Taa	165	166
temporal cortex	temporal area TE, medial part	TEM	177					
temporal cortex	temporal area TE, occipital part	TEO	178					
temporal cortex	temporal area TE, occipitomedial part	TEOM	179					
temporal cortex	temporal area TE1	TE1#1	180		TEMPORAL AREA TE	TE	167	168
temporal cortex	temporal area TE2	TE2#1	181					
temporal cortex	temporal area TE3	TE3#1	182					
temporal cortex	temporal area TEa	TEa#1	183					
temporal cortex	temporal area TF	TF	184		TEMPORAL AREA TF	TF	169	170
temporal cortex	temporal area TF, lateral part	TFL	185					
temporal cortex	temporal area TF, occipital part	TFO	186					
temporal cortex	temporal area TH	TH	187		TEMPORAL AREA TH	TH	171	172
temporal cortex	temporal area TH, occipital part	Tho	188					
temporal cortex	temporal area TL	TL	189		TEMPORAL AREA TL	TL	173	174
temporal cortex	temporal area TL, occipital part (area 36O)	TLO(36O)	190					
temporal cortex	temporal parietooccipital associated area in sts	TPO	191		TEMPORAL PARIETOOCIPITAL ASSOCIATED AREA	TPO	175	176
temporal cortex	temporal parietooccipital associated area in sts, caudal part	TPOC	192					
temporal cortex	temporoparietal cortex	Tpt	193		TEMPOROPARIETAL CORTEX	Tpt	177	178
temporal cortex	temporopolar periallocortex	TPPAI	194		TEMPOROPOLAR PERIALLOCORTEX	TPPAI	179	180
temporal cortex	temporopolar proisocortex	TPPro	195		TEMPOROPOLAR PROISOCORTEX	TPPro	181	182
frontal cortex	area 4 of cortex (primary motor)	4#1	196		AREA 4 OF CORTEX (primary motor)	4#1	183	184
frontal cortex	area 6 of cortex, dorsocaudal part (Matellis F2)	6D(CF2)	197					
frontal cortex	area 6 of cortex, dorsostral part (Matellis F7)	6DR(F7)	198		AREA 6 OF CORTEX	6	185	186
frontal cortex	area 6 of cortex, medial (supplementary motor) part	6M	199					
frontal cortex	area 6 of cortex, ventral part, caudal subdivision (Matellis F4)	6VC(F4)	200					
frontal cortex	area 6 of cortex, ventral part, rostral subdivision (Matellis F5)	6VR(F5)	201					
frontal cortex	area 6/32 of cortex	10379	202		AREA 6/32 OF CORTEX	6/32	187	188
frontal cortex	area 8 of cortex, anterodorsal part	8AD	203					
frontal cortex	area 8 of cortex, anteroventral part	8AV	204					
frontal cortex	area 8/32 of cortex	10440	205		AREA 8 OF CORTEX	8	189	190
frontal cortex	area 8A of cortex	8A	206					
frontal cortex	area 8B of cortex	8B	207					
frontal cortex	area 9 of cortex, lateral part	9L	208		AREA 9 OF CORTEX	9	191	192
frontal cortex	area 9 of cortex, medial part	9M	209					
frontal cortex	area 9/32 of cortex	10471	210		AREA 9/32 OF CORTEX	9/32	193	194
frontal cortex	area 9/46 of cortex	15584	211					
frontal cortex	area 9/46 of cortex, dorsal part	9/46D	212		AREA 9/46 OF CORTEX	9/46	195	196
frontal cortex	area 9/46 of cortex, ventral part	9/46V	213					
frontal cortex	area 10 of cortex	10	214		AREA 10 OF CORTEX	10	197	198
frontal cortex	area 10 of cortex, dorsal part	10D	215					
frontal cortex	area 10 of cortex, medial part	10M	216					
frontal cortex	area 10 of cortex, ventral part	10V	217					
frontal cortex	area 44 of cortex	44	218		AREA 44 OF CORTEX	44	199	200
frontal cortex	area 45A of cortex	45A	219					
frontal cortex	area 45B of cortex	45B	220		AREA 45 OF CORTEX	45	201	202
frontal cortex	area 46D of cortex	46D	221					
frontal cortex	area 46V of cortex	46V	222		AREA 46 OF CORTEX	46	203	204
frontal cortex	area 47 (old 12) of cortex	47(12)	223					
frontal cortex	area 47 (old 12) of cortex, lateral part	47(12)L	224		AREA 47 OF CORTEX	47	205	206
frontal cortex	area 47 (old 12) of cortex, orbital part	47(12)O	225					
frontal cortex	area ProM (promotor)	proM#1	226		AREA PROM (promotor)	proM#1	207	208
frontal cortex	gustatory cortex	Gu	227		GUSTATORY CORTEX	Gu	209	210
orbitofrontal cortex	area 11 of cortex	11	228		AREA 11 OF CORTEX	11	211	212
orbitofrontal cortex	area 11 of cortex, lateral part	11L	229					
orbitofrontal cortex	area 11 of cortex, medial part	11m	230					
orbitofrontal cortex	area 13 of cortex	13	231					
orbitofrontal cortex	area 13 of cortex, lateral part	13L	232		AREA 13 OF CORTEX	13	213	214
orbitofrontal cortex	area 13 of cortex, medial part	13M	233					
orbitofrontal cortex	area 13a of cortex	13a	234					
orbitofrontal cortex	area 14 of cortex, medial part	14M	235					
orbitofrontal cortex	area 14o	14o	236		AREA 14 OF CORTEX	14	215	216
orbitofrontal cortex	area 25 of cortex	25	237					
orbitofrontal cortex	orbital periallocortex	OPAI	238		AREA 25 OF CORTEX	25	217	218
orbitofrontal cortex	orbital proisocortex	OPro	239		ORBITAL PERIALLOCORTEX	OPAI	219	220
white matter	cerebral white matter	cwm	240		ORBITAL PROISOCORTEX	OPro	221	222
white matter	external medullary lamina	eml	241				0	0
							0	0

Table S7: Whole brain areas labelling of the Center for In Vivo Microscopy atlas (CIVM) atlas Revised (CIVM_R) for functional correlations analysis.

Whole brain regions considered for the functional correlations analysis using the macaque CIVM atlas(76) that was revised into CIVM_R to match fMRI spatial resolution. Regions merged together (for instance dorsal, medial and ventral part of the medial geniculate nucleus in the original CIVM atlas into medial geniculate nucleus) share the same abbreviation and value in the CIVM_R. Brackets represent the biggest regions unified. Empty cells stands for deleted regions.