**Cortico-pallidal oscillatory connectivity in patients with dystonia**

Wolf-Julian Neumann,1,2 Ashwani Jha,3 Antje Bock,1 Julius Huebl,1 Andreas Horn,1 Gerd-Helge Schneider,4 Tillmann H. Sander,5 Vladimir Litvak2,* and Andrea A. Kühn1,6,7,*

*These authors contributed equally to this work.

Primary dystonia has been associated with an underlying dysfunction of a wide network of brain regions including the motor cortex, basal ganglia, cerebellum, brainstem and spinal cord. Dystonia can be effectively treated by pallidal deep brain stimulation although the mechanism of this effect is not well understood. Here, we sought to characterize cortico-basal ganglia functional connectivity using a frequency-specific measure of connectivity—coherence. We recorded direct local field potentials from the human pallidum simultaneously with whole head magnetoencephalography to characterize functional connectivity in the cortico-pallidal oscillatory network in nine patients with idiopathic dystonia. Three-dimensional cortico-pallidal coherence images were compared to surrogate images of phase shuffled data across patients to reveal clusters of significant coherence (family-wise error \( P < 0.01 \), voxel extent 1000). Three frequency-specific, spatially-distinct cortico-pallidal networks have been identified: a pallido-temporal source of theta band (4–8 Hz) coherence, a pallido-cerebellar source of alpha band (7–13 Hz) coherence and a cortico-pallidal source of beta band (13–30 Hz) coherence over sensorimotor areas. Granger-based directionality analysis revealed directional coupling with the pallidal local field potentials leading in the theta and alpha band and the magnetoencephalographic cortical source leading in the beta band. The degree of pallido-cerebellar coupling showed an inverse correlation with dystonic symptom severity. Our data extend previous findings in patients with Parkinson’s disease describing motor cortex-basal ganglia oscillatory connectivity in the beta band to patients with dystonia. Source coherence analysis revealed two additional frequency-specific networks involving the temporal cortex and the cerebellum. Pallido-cerebellar oscillatory connectivity and its association with dystonic symptoms provides further confirmation of cerebellar involvement in dystonia that has been recently reported using functional magnetic resonance imaging and fibre tracking.

1 Department of Neurology, Campus Virchow Klinikum, Charité–University Medicine Berlin, Augustenburger Platz 1, 13353 Berlin, Germany
2 The Wellcome Trust Centre for Neuroimaging, UCL Institute of Neurology, 12 Queen Square, London WC1N 3BG, UK
3 Sobell Department of Movement Disorders, UCL Institute of Neurology, Queen Square, London WC1N 3BG, UK
4 Department of Neurosurgery, Campus Virchow Klinikum, Charité–University Medicine Berlin, Augustenburger Platz 1, 13353 Berlin, Germany
5 Physikalisch-Technische Bundesanstalt, Institut Berlin, Abbestr. 2-12, 10587 Berlin, Germany
6 Berlin School of Mind and Brain, Charité - University Medicine Berlin, Unter den Linden 6, 10099 Berlin, Germany
7 NeuroCure, Charité - University Medicine Berlin, Charitéplatz 1, 10117 Berlin, Germany

Correspondence to: Prof. Dr. Andrea A. Kühn,
Department of Neurology, Charité,
University Medicine Berlin,
Campus Virchow Klinikum,
Augustenburger Platz 1,
13353 Berlin, Germany

© The Author (2015). Published by Oxford University Press on behalf of the Guarantors of Brain. All rights reserved.
For Permissions, please email: journals.permissions@oup.com
Cortico-pallidal connectivity in dystonia

Introduction

The dystonias are a heterogeneous group of syndromes that are characterized by involuntary twisting movements and abnormal posturing, sometimes with associated tremor (Fahn, 1988). Early neuropathological, CT and MRI studies investigating the pathophysiological mechanisms underlying this movement disorder have suggested that basal ganglia dysfunction plays a central role (Hallett, 2006). More recently, abnormal activity in a wider network of regions has been identified including the cortex, cerebellum, brainstem and spinal cord (Hendrix and Vitek, 2012). Both at rest and during voluntary movements, neuroimaging studies have demonstrated abnormal activation patterns in the putamen, pallidum, thalamus, cerebellum and motor and premotor cortices (Ceballos-Baumann et al., 1995; Eidelberg et al., 1998; Kumar et al., 1999; Detante et al., 2004; Lehericy et al., 2013). Electrophysiological evaluations of the sensorimotor and premotor cortices have suggested that there is a loss of intracortical inhibition, increased cortical plasticity and abnormal sensorimotor integration in patients with dystonia (Vidailhet et al., 2009; Quartarone and Hallett, 2013). Deep sites such as the basal ganglia have traditionally been difficult to electrophysiologically access in humans. Deep brain stimulation surgery for movement disorders now affords the opportunity to record from macroelectrodes in these areas via surgically placed deep brain stimulation (DBS) electrodes, before electrical stimulation (Brown and Williams, 2005). Such recordings from the target areas of DBS have revealed that subjects with Parkinson’s disease have a characteristic increase in local beta frequency activity whilst subjects with dystonia have increased lower frequency activity (Silberstein et al., 2003; Liu et al., 2008). Furthermore, pallidal low frequency oscillations temporally lead similar oscillations in dystonic muscle activity, suggesting that they may play a causal role (Sharott et al., 2008).

DBS of the globus pallidus internus (GPI) is a successful treatment for patients with generalized, segmental and cervical dystonia refractory to medical therapy (Vidailhet et al., 2005, 2007; Kupsch et al., 2006; Ostrem and Starr, 2008; Volkman et al., 2012, 2014). However, the therapeutic mechanism of DBS in dystonia is still not fully understood. One proposed hypothesis is that DBS suppresses or overrides the aforementioned pathological oscillatory network activity within cortico-basal ganglia circuits (Hammond et al., 2007; Brown and Eusebio, 2008; McIntyre and Hahn, 2010; Hendrix and Vitek, 2012). However, less is known about the wider network implications of such activity, in particular the physiological and pathological oscillatory activity ‘between’ distant brain regions, in patients with dystonia. A novel technique that records subcortical local field potentials (LFPs) from DBS electrodes and simultaneous cortical whole head magnetoencephalography (MEG) has made it possible to electrophysiologically characterize the cortico-basal ganglia network in subjects with Parkinson’s disease using coherence measures to characterize functional connectivity (Litvak et al., 2010, 2011a). Coherence describes the spectral distribution of oscillatory synchronization between concurrent signals and can be interpreted as interaction or communication between brain areas. Two spectrally and spatially distinct oscillatory networks were found in patients with Parkinson’s disease at rest involving the subthalamic nucleus and cortical sources: a frontal motor network for the beta band and a predominantly temporal network for the alpha band, which were modulated during movement and tremor episodes (Hirschmann et al., 2011, 2013; Litvak et al., 2011a, 2012; Oswal et al., 2013).

The aim of the present study was to use this novel technique of simultaneous MEG-LFP recording to investigate patterns of cortico-pallidal oscillatory connectivity in patients with dystonia. We hypothesized that such networks would again be spatially and spectrally segregated in dystonic patients but that key spatial and/or spectral features would be distinct from the abnormalities found in Parkinson’s disease and that such features would provide new insights into the brain networks involved into the pathophysiology of dystonia.

Materials and methods

Patients and surgery

Nine patients (six females; age 50.1 ± 3.9 years, range 24–68 years; disease duration 12.8 ± 2.5 years, range 3–23 years; mean ± standard error of the mean [SEM]) with either segmental dystonia (n = 2; Cases 2 and 5), cervical dystonia (n = 5; Cases 3, 4, 6, 7 and 8) or generalized dystonia (n = 2; Cases 1 and 9) who underwent bilateral implantation of DBS electrodes in the GPI were included in the study. All patients participated with informed consent and with the agreement of the ethical committee of the Charite–Universita¨tsmedizin, Berlin, Campus Virchow-Klinikum. The study was conducted according to the Declaration of Helsinki. Clinical details are summarized in Table 1. The DBS macroelectrode used was model 3389 (Medtronic) with four platinum-iridium...
cylindrical surfaces of diameter 1.27 mm, length 1.5 mm and centre-to-centre separations of 2 mm. Contacts 0 and 3 were the lowermost and uppermost contacts, respectively. Correct placement of the DBS electrodes was guided by intraoperative microelectrode recordings and confirmed by postoperative MRI. Postoperative 3D electrode localization was conducted using the Lead-DBS toolbox (Horn and Kühn, 2014; see Fig. 1 for an example). In brief, preoperative and postoperative magnetic resonance images were fused and normalized to Montreal Neurological Institute (MNI) space, before mapping the contacts to normalized subcortical atlases (BGHAT atlas, Prodoehl et al., 2008; Harvard-Oxford subcortical/cortical atlas, Frazier et al., 2005). This confirmed that 51/54 contact pairs had at least one contact in the internal pallidum, with the remaining three contact pairs from two patients (Case 2: R01, Case 7: R01/L01) lying below the edge of the target.

### Experiments and recordings

Simultaneous 125 channel MEG (YOKOGAWA ET 160) and pallidal LFP recordings were conducted 1 to 7 days after DBS electrode implantation (mean ± SEM = 3 ± 0.37 days), while the DBS leads were externalized. Rest recordings of 3–4 min were conducted, during which the patients were lying still with eyes open and instructed not to move. To avoid movement artefacts, patients with severe phasic head movements or head tremor were not included in the study. All data were recorded at a sampling rate of 2000Hz. Electro-oculographic, EMG and LFP recordings were recorded simultaneously through the integrated EEG system. Recordings from four DBS electrode contacts were referenced to contact 0 of the left electrode and later re-referenced offline (see below).

### Data analysis

Statistical parametric mapping (SPM; Litvak et al., 2011b) (http://www.fil.ion.ucl.ac.uk/spm/), Fieldtrip (Oostenveld et al., 2011) (http://www.ru.nl/neuroimaging/fieldtrip/) and Data Analysis in Source Space (DAiSS, https://code.google.com/p/spm-beamforming-toolbox/) toolboxes were used. To limit the effects of volume conduction to distant sources, LFP recordings were re-referenced to a bipolar montage between adjacent contact pairs (01, 12, 23). To avoid selection bias, all three contact pairs (01, 12, 23) from the DBS electrodes of all patients were included in the analysis. The data were then down-sampled to 300Hz. High-pass (>).1Hz) and notch filters (at 50, 100 and 150Hz) were applied to limit effects of line noise (fifth order zero-phase Butterworth filters). Continuous rest recordings were divided into arbitrary epochs of 3.41-s length (1024 samples). Epochs were rejected from further analysis if they contained artefacts in either MEG or LFP data. Artefacts in the LFP recordings were identified by thresholding the LFP amplitude at 30μV (Litvak et al., 2011a). This threshold was set by examining the typical values for clean data and artefactual deflections across epochs and subjects. The MEG data were contaminated by artefacts generated by ferromagnetic percutaneous connection wires. It was previously shown (Litvak et al., 2010) that such artefacts can be suppressed by beamforming. However, the Yokogawa system used in the present study has a lower dynamic range than the CTF-MEG system used previously, resulting in intermittent saturations in some of the channels close to the wires. These

### Table I  Clinical details of the patients

| Case | Age | Sex | Diagnosis       | TWSTRS | BFMDRS | Disease duration | Alpha peak location (mm x y z MNI) | Beta peak location (mm x y z MNI) | Theta peak location (mm x y z MNI) | Alpha peak location (mm x y z MNI) | Beta peak location (mm x y z MNI) | Theta peak location (mm x y z MNI) | Alpha peak location (mm x y z MNI) | Beta peak location (mm x y z MNI) |
|------|-----|-----|-----------------|--------|--------|------------------|----------------------------------|----------------------------------|---------------------------------|----------------------------------|---------------------------------|---------------------------------|---------------------------------|
| 1    | 48  | F   | Generalized dystonia | na     | 16     | 20               | 1/125                            | 54 – 24 6                        | -2 – 50 – 48                     | 40 – 2 24                       |                                 |                                 |                                 |
| 2    | 55  | M   | Segmental dystonia  | 20     | na     | 12               | 0/125                            | 32 – 28 – 28                     | 2 – 48 – 48                      | 4 – 46 64                      |                                 |                                 |                                 |
| 3    | 51  | F   | Cervical dystonia   | 23     | na     | 3                | 20/125                           | 56 – 44 – 16                     | 24 – 64 – 46                     | 14 – 8 70                      |                                 |                                 |                                 |
| 4    | 52  | F   | Cervical dystonia   | 22     | na     | 11               | 14/125                           | 46 – 14 – 22                     | 0 – 72 – 42                      | 48 – 4 28                      |                                 |                                 |                                 |
| 5    | 48  | F   | Segmental dystonia  | 25     | na     | 6                | 15/125                           | 54 – 32 – 18                     | 16 – 84 – 16                     | 46 – 30 52                     |                                 |                                 |                                 |
| 6    | 68  | M   | Cervical dystonia   | 16     | na     | 23               | 86/125                           | na                              | na                              | na                              |                                 |                                 |                                 |
| 7    | 47  | F   | Cervical dystonia   | 15     | na     | 4                | 18/125                           | 30 – 32 – 28                     | 0 – 52 – 40                      | 32 – 10 54                     |                                 |                                 |                                 |
| 8    | 58  | M   | Cervical dystonia   | 18     | na     | 20               | 26/125                           | 56 – 32 6                       | 8 – 62 – 40                      | 4 – 2 70                       |                                 |                                 |                                 |
| 9    | 24  | F   | Generalized dystonia| na     | 27     | 16               | 0/125                            | no peak                         | 38 – 78 – 34                     | 36 – 2 54                      |                                 |                                 |                                 |

Note that the TWSTRS was used to assess symptom severity in patients with clinical features involving primarily the neck, whereas the Burke Fahn Marsden Dystonia Rating Scale (BFMDRS) was used for the patients with generalized dystonia. aThis case was excluded from the analysis due to artefacts. bCase 2 had cervical dystonia with additional postural hand tremor. cCase 5 had cervical dystonia with mild contraction of the right abdominal wall.
saturated segments were defined as segments where the differences between adjacent samples stayed < 5 fT for 10 consecutive samples or more. Epochs with amplitude jumps of > 10 pT between consecutive samples were also excluded. If for a particular MEG channel > 20% of the epochs were contaminated by these artefacts, the channel was excluded from further analysis (see Table 1 for details). In one patient (Case 6), metal artefacts led to exclusion of 86/125 sensors; thus, data from this patient were omitted from further analysis. The mean number of excluded channels was 13 (range 0–26), concentrated in the area of the percutaneous connection wires on the left hemisphere (location corresponding to the area around electrode F3 in the 10–20 EEG system). After artefact rejection, a mean ± SEM of 216.9 ± 16.2 s (range 133–293 s) were analysed, resulting in an average of 62.9 ± 5.25 SEM trials (range 39–75 trials) in eight remaining patients. All data were visually inspected before and after automatic artefact rejection.

Characterization of the frequency distribution of significant LFP-MEG coherence

The principal correlate of functional connectivity investigated in this study was coherence, which provides a frequency domain measure of the linear phase and amplitude relationships between signals (Buzsaki and Draguhn, 2004; Litvak et al., 2010). Single-subject analysis at the sensor level was designed to investigate the frequency distribution of significant peaks of coherence before group analysis. We used the procedure originally suggested by Litvak et al. (2011a). MEG and bipolar GPi-LFP epoched data were converted to the frequency domain (range 5–45 Hz with a frequency resolution of 2.5 Hz, and 55–95 Hz with frequency resolution of 7.5 Hz) using the multi-taper method (Thomson, 1982). Coherence was then calculated between each GPi-LFP channel and each MEG channel. Scalp maps of coherence for each frequency bin were linearly interpolated to produce a 2D image (64 × 64 pixels). The resulting images were stacked to produce a 3D image with two spatial and one frequency dimension. Thus, 3D images for the low and high frequency ranges were created for each LFP channel in each patient. Significant regions of coherence within these images were determined by statistical comparison to 10 surrogate coherence images in which any coherence was destroyed. The surrogate coherence images were generated from the same MEG trials, but with the order of GPi-LFP trials shuffled. To ensure conformance to the assumptions of random field theory, all images were smoothed with a Gaussian kernel (10 mm × 10 mm × 2.5 Hz for low frequency, and 10 mm × 10 mm × 7.5 Hz for high frequency images). Standard SPM procedures were used for a two-sample t-test between the original image and the 10 surrogate images with a family-wise-error corrected threshold of P < 0.01 to identify significant regions in sensor space and frequency. For each original LFP-MEG coherence image in each patient, frequencies of the clusters of significant coherence were identified and taken into a histogram (Fig. 2B) to visualize the frequency distribution of significant sensor level coherence across patients.

Characterization of coherent source activity across patients

To identify the spatial topography and spectral properties of distinct frequency-specific cortico-pallidal coherence networks, we used MEG beamforming. For this method, a linear projection of sensor data is generated by using a spatial filter computed from the lead field of the source of interest and the data covariance in the time domain (Van Veen et al., 1997) or the cross-spectral density matrix in the frequency domain (Gross et al., 2001). A lead field computation, a perturbed sphere forward model (Nolte, 2003) was created based on an inner skull mesh obtained by inverse-normalizing a canonical mesh to the subject’s individual preoperative MRI image (Mattout et al., 2007). All MEG data sets were coregistered to the individual patient’s MRI by using MEG head position indicator coils attached to the nasion and left and right pre-auricular points. The DICS (dynamic imaging of coherent sources) beamforming method was used to locate cortical sources coherent to the GPi-LFP activity (Gross et al., 2001). The coherence values were computed on a 3D grid in Montreal Neurological Institute space, with spacing of 5 mm bounded by the inner skull surface. Values at the grid points were then linearly interpolated to produce volumetric images with 2 mm resolution. The images were stored in Neuroimaging Informatics Technology Initiative (NIfTI) format and smoothed with an 8 mm isotropic Gaussian kernel. The resulting 3D images of LFP-MEG coherence were compared to surrogate images from phase shifted data using SPM procedures (Friston et al., 1994; Litvak et al., 2011b). Original and surrogate coherence images were created for five frequency bands, theta (4–8 Hz), alpha (7–13 Hz), beta (13–30 Hz), low gamma (30–60 Hz) and high gamma (60–90 Hz) and taken into the analysis. Images derived from the left pallidal electrodes were reflected across the median sagittal plane, to allow comparison of sources ipsilateral and contralateral to the GPi regardless of the original hemisphere. All images were subjected to a fixed-effect ANOVA to test for a significant difference between real and surrogate coherence images, with subject and side as additional factors. A family-wise error (FWE) corrected threshold of P < 0.01 and a minimum voxel extent of 1000 voxels were chosen to identify significant regions of cortico-pallidal coherence. The MNI coordinates of the global maxima for each significant frequency band on the group level were further used for single patient analysis and source extraction. Single patient peak locations were analysed from averaged DICS images from all contact pairs by identifying local maxima in proximity to the group peak. Virtual electrode signals were extracted from group coherence peaks for each frequency band using a linearly constrained minimum variance beamformer (Van Veen et al., 1997) and regularization set at 0.01% of the mean of the diagonal of the channel covariance matrix (Litvak et al., 2010). Source and GPi-LFP power and coherence estimates were calculated from the resulting extracted source channels and are presented as the mean across all pallidal contact-pairs and patients. Individual peaks of GPi-LFP power spectra were investigated visually and reported. To investigate the effective directionality of functional coupling between the cortical sources and the pallidal LFP signal, we used a non-parametric variant of Granger causality (Brovelli et al., 2004) averaged for each frequency band. To identify
significant directional coupling, original data were compared
to time-reversed data (Haufe et al., 2013) across all patients
for each extracted source in the defined frequency range using
right-tailed paired t-tests. This procedure has proven to sup-
press the influence of weak data asymmetries not related to
time-lagged interactions in the data, while it is statistically
powerful in the detection of meaningful strong asymmetries
(Haufe et al., 2013). Hence, in the case of true directional
interaction a significantly higher Granger causality value is
expected for the original signals compared to the reversed sur-
rogate. Therefore, it was tested whether the results were
significantly greater across the group for the original data
when compared to the reversed surrogate data.

**Correlation with clinical symptoms**

To investigate a potential relation of LFP–source coherence
and dystonic symptom severity, the preoperative Torticollis
severity scale that is part of the Toronto Western Spasmodic
Torticollis Rating Scale (TWSTRS) was documented for patients
with cervical or segmental dystonia (all analysed patients,
except Cases 1 and 9). Note that patients with segmental
dystonia had mild additional symptoms that were not covered
by the TWSTRS (Case 2: postural hand tremor; Case 5: con-
traction of the right abdominal wall). Non-parametric
Spearman's correlations were used to investigate a potential
association of the TWSTRS severity score and the coherence
value in the contact pair, with the highest averaged frequency
band LFP-MEG coherence for the temporal theta (4–8 Hz),
cerebellar alpha (7–13 Hz) and motor-cortical beta band
(13–30 Hz) source.

**Results**

**Spectral distribution of cortico-pallidal coherence**

Significant peaks of cortico-pallidal coherence were present
in all patients at the sensor level. An example of sensor
level coherence with a significant beta peak and the result-
ing source coherence on an individual patient’s cortical
mesh is shown in Fig. 2. Frequencies of peaks were
Spatial distribution of frequency-specific cortico-pallidal coherence

Topographical mapping of LFP-MEG coherence revealed frequency-specific pallidal connectivity to sensorimotor cortical areas, temporal areas and the cerebellum. The coherence patterns were clearly lateralized to the ipsilateral hemisphere. For each frequency band, DICS images with their corresponding surrogate phase shifted DICS pairs (96 images from 48 contact pairs from 16 hemispheres from eight patients) were subjected to a fixed-effect ANOVA with subject and side as additional factors. Three significant clusters of cortico-pallidal coherence were revealed, in the theta (4–8 Hz), alpha (7–13 Hz) and beta (13–30 Hz) bands. The results are presented in Fig. 3.

In the theta band, a cluster extending from subcortical areas to the temporal cortex, with the coherence maximum in the inferior temporal gyrus, was identified (t = 8.87; P < 0.0001; cluster size: 4424; MNI coordinates x = 44, y = −26, z = −17). A cluster restricted to the central cerebellum was significant in the alpha band (t = 6.18; P < 0.0001; cluster size: 6688; MNI coordinates x = 2, y = −32, z = 13). This cluster overlapped peripherally with the brainstem. However, this overlap vanished when a more conservative FWE corrected P-value threshold (P < 0.00001) was chosen. In line with a role of beta oscillations in the sensorimotor network, cortico-pallidal coherence in the beta band (13–30 Hz) was significant in the sensorimotor areas (t = 8.09; P < 0.0001; cluster size: 24669; MNI coordinates x = 30, y = −14, z = 57).

For the low gamma band (30–60 Hz) a sensorimotor region and two subcortical clusters, and for the high gamma band (60–90 Hz) a small subcortical cluster reached significance (P < 0.05 FWE corrected), but did not survive our stringent thresholding criteria and were therefore not further analysed.

All other clusters remained significant, when the three contact pairs that were off target in the postoperative imaging were excluded. A direct comparison of hemispheres (left versus right) did not reveal significant differences in any of the frequency bands. Analysis of individual source locations confirmed the consistency of our findings in individuals within the patient group (Fig. 4). In the theta band, 7/8 patients had a local peak within an average distance of 19.07 ± 2.17 mm from the group mean in MNI space. Similar distances were found for the alpha (8/8 patients, mean distance ± SEM 21.38 ± 5.90 mm) and beta bands (8/8, 25.05 ± 4.16 mm).

Cortical and pallidal spectral power, coherence and directionality of information flow

In each subject, using a linearly constrained minimum variance beamformer, virtual channels of cortical activity were extracted from the locations of maximum coherence (group peak) in the significant clusters for each frequency band. Figure 5 shows the power spectra, LFP-source coherence and directionality measures for each of the three extracted sources. The cortical power spectra showed distinct low frequency peaks at ~10 Hz (including theta and alpha band) and additional increased power in the frequency range up to ~20 Hz for the motor cortical beta band source. For the pallidal LFP signals, power-spectra were averaged across all contact pairs and all patients for visualization. On an individual level, all patients had distinct low frequency peaks in 30/44 contact pairs overall (range 7–12 Hz; mean ± SEM 9 ± 0.5 Hz). A clear spectral
Figure 5 Source extraction and directionality analysis. Linearly constrained minimum variance beamformer (LCMV) source extraction allowed power, coherence and directionality analysis of the significant sources. The top first row shows the averaged power spectra and the second row the coherence spectra for the temporal theta (A), the cerebellar alpha (B) and the motor cortical beta (C) sources. The lowermost row shows the averaged Granger causality measure for the respective frequency bands [4–8 Hz (A), 7–13 Hz (B), 13–30 Hz (C)] of the three identified sources in comparison to reversed data. *p < 0.05 **p < 0.001.

Figure 4 Individual peak localization. Locations of individual local maxima closest to the respective group peak for the theta (yellow), alpha (blue) and beta band (red) were identified in the averaged source coherence images across all contact-pairs for each patient and projected to ‘glass brains’ (inner boundary of skull marked with grey mesh) viewed from the above (A), right (B) and front (C). All images from left hemispherical contact-pairs were mirrored along the sagittal plane prior to averaging.
distribution of LFP-MEG coherence was found for each cluster, with a focal theta peak (4–7 Hz) for the temporal cortex, a broader alpha band peak (6–13 Hz) for the cerebellar source, and two peaks for the motor cortical source (4–8 Hz + 10–30 Hz). Directionality analysis for the relevant frequency bands for each source revealed that the GPi drove the temporal cortex in the theta range (4–8 Hz; \( P < 0.01 \)) and the cerebellum in the alpha range (7–13 Hz; \( P = 0.01 \)). The motor cortical source drove the GPi in the beta range (\( P < 0.001 \)). Beta band activity in the pallidum has previously been shown to be driven by the motor cortex in LFP-EEG studies (Williams et al., 2002) and similar results were found for LFP-MEG and LFP-EEG coherence from the subthalamic nucleus in patients with Parkinson’s disease (Litvak et al., 2011a).

**Correlation of cortico-pallidal coherence and dystonic symptom severity**

A negative correlation was found between the degree of pallido-cerebellar connectivity and clinical symptom severity as measured by the TWSTRS. Averaged pallido-cerebellar alpha band coherence (7–13 Hz) negatively correlated with the preoperative TWSTRS score in patients with segmental or cervical dystonia (\( n = 6, \text{Rho} = -0.94, P = 0.017 \); Fig. 6), meaning that higher connectivity values were associated with less prominent dystonic motor symptoms in our patients. Detailed analysis revealed that higher scores due to additional symptoms in segmental dystonia patients (not covered in the TWSTRS) would improve (Case 2) or not change (Case 5) the rank based Spearman correlation. No significant correlation was found for the motor cortical beta and theta as well as the temporal theta source (Supplementary Fig. 1).

**Discussion**

We have demonstrated that the human internal pallidum is interconnected with the cerebral cortex in spatially and spectrally distinct functional oscillatory networks in patients with idiopathic dystonia. Three network connections with the pallidum were identified. Theta band coherence was identified with the temporal cortex, an alpha band coherence with a central cerebellar source, and the beta band coherence with motor and premotor cortical regions. Interestingly, the amount of coupling in the alpha band between the internal pallidum and the cerebellum was inversely correlated with the patients’ preoperative clinical state as measured by the TWSTRS. Although correlation does not prove causation, the demonstration of a significant correlation between cerebello-pallidal alpha band coherence and dystonic symptom severity suggests a pathophysiological role for alpha band oscillatory network activity in dystonia.

Before we proceed with a more detailed discussion of our findings, we should consider the potential limitations of our study. The methodological effort, combined with the sparseness of dystonic DBS patients suited for MEG recordings, led to a relatively small and heterogeneous cohort of patients; we tried to compensate for this by introducing conservative statistical thresholds and thorough validation of individual results, both in sensor and in source space. In this regard, it should be noted that the cerebellar alpha band group peak overlaps partially with the brainstem. This is of interest, because brainstem lesions can cause dystonic symptoms (Vidailhet et al., 1999) and altered structural basal ganglia–brainstem connectivity has been reported in dystonia (Blood et al., 2012). Thus, the observed alpha band coherence could be partially originated from the brainstem. However, the facts that all patients had the local maxima in the cerebellum and not in the brainstem, and that an even more conservative statistical threshold confined the source of alpha band coherence to the cerebellum, favours the cerebellum as the relevant source of coupling. Further, magnetic artefacts led to an exclusion of some of the channels near the DBS lead externalization in the left hemisphere, which may have compromised the reliability of coherence measures in this area. A potential influence of this channel exclusion on our results cannot be ruled out; however, no hemispherical differences in coherence patterns were revealed by the statistical analysis. Next, the correlation of alpha band coherence and clinical symptoms are obtained by investigating the severity of predominant cervical dystonic symptoms and we should be cautious not to generalize the findings to all forms of dystonia. Moreover, an inherent limitation of intracranial recordings in humans is the lack of healthy controls. Hence, we cannot predict the
pathophysiological influence of the disease on the reported networks, as our group consisted of patients with different types of dystonia with partially persistent motor symptoms at rest. With the intent to discuss the functional roles of the networks, we should note that the demonstrated results are derived without any functional modulation, and more likely reflect a resting or default mode network (Laufs et al., 2003). Finally, all intracranial recordings in human subjects lack histological confirmation of the origin of the recorded activity, thus the electrode placement remains presumptive. However, postoperative imaging and MRI-based electrode localization in all of our patients was consistent with at least two contacts of the DBS electrode placed in the Gpi.

The observed findings for the theta and the beta band are similar to previously described subthalamic LFP-MEG connectivity patterns from patients with Parkinson’s disease (Hirschmann et al., 2011; Litvak et al., 2011a; Neumann et al., 2014). Subcortico-cortical coherence with the temporal lobe has been described in these studies for the lower (yet alpha band) frequency range, whereas beta oscillations were primarily synchronized to sensorimotor regions. We describe a pallido-temporal network in the theta frequency range, which overlaps spatially with the hippocampus. The hippocampal theta rhythm is one of the most prominent oscillatory phenomena (Buzsaki and Draguhn, 2004) that has previously been described in MEG studies (Tesche, 1997; Tesche and Karhu, 2000a; Cornwell et al., 2012), and recently further supported by evidence from parallel hippocampal LFP-MEG recordings that has given first-line evidence for the feasibility of recording human hippocampal theta rhythms with MEG (Dalal et al., 2013). The hippocampus is anatomically interconnected with the pallidum via the nucleus accumbens and proposed to be simultaneously activated during learning and integration of limbic information on sensorimotor output (Thierry et al., 2000; Packard and Knowlton, 2002). Theta synchronization has been hypothesized to allow encoding of new information in hippocampo-cortical feedback loops (Klimesch, 1999), and theta band coherence between the hippocampus and the basal ganglia was previously described in rats, where the striatum has been reported to be driving information flow (DeCoteau et al., 2007). Thus, the hippocampus is a tempting candidate as a source for the described mid-temporal oscillatory theta network. However, given the low spatial resolution of the method and the relatively small size of the hippocampus and the larger extent of the described source, we propose a hippocampal network as one potential explanation of our findings, while keeping in mind that further studies would be needed to confirm this hypothesis through functional modulations of this oscillatory circuit to also elucidate its functional significance in dystonia.

The role of beta oscillatory activity in the cortico-basal ganglia circuit is comparatively much more studied, and beta network alterations have been reported in a variety of studies (Pfurtscheller and Aranibar, 1977; Pfurtscheller and Lopes da Silva, 1999; Kühn et al., 2004, 2008a, b; Lalo et al., 2007; Ray et al., 2008; Hirschmann et al., 2011; Litvak et al., 2011a; Brücke et al., 2012; Alegre et al., 2013). A suppression of beta band activity has frequently been reported during motor tasks in the motor cortex (Pfurtscheller and Aranibar, 1977; Pfurtscheller and Lopes da Silva, 1999; Lalo et al., 2007), Gpi (Brücke et al., 2008, 2012; Singh et al., 2011a, b; Herrojo Ruiz et al., 2014), the subthalamic nucleus (Kühn et al., 2004; Litvak et al., 2012; Alegre et al., 2013) and the motor thalamus (Paradiso et al., 2004; Kempf et al., 2009; Brücke et al., 2013). Thus, beta activity in the cortico-basal ganglia loops has been widely associated with movement initiation and termination in motor control (Jenkinson and Brown, 2011; Brittain and Brown, 2014). Stimulation of the motor cortex in the beta frequency range has been shown to slow movements, which is first-line evidence for a causal influence of this network activity on motor performance (Pogosyan et al., 2009; Joundi et al., 2012). More specifically, beta circuit synchronization may promote the status quo or generally tonic activity (Engel and Fries, 2010; Jenkinson and Brown, 2011), and premovement beta desynchronization could signal the likelihood that a new voluntary action will need to be performed (Jenkinson and Brown, 2011). Altered movement-related beta band desynchronization over motor cortical areas (Toro et al., 2000), and abnormal beta band functional connectivity across premotor–motor areas during movement (Jin et al., 2011) provides further evidence for motor cortical involvement in dystonia linked to alterations in beta band oscillatory activity. Our study demonstrates a cortico-pallidal beta network at rest that is spatially restricted to the sensorimotor area, and thus gives a framework for functional interactions of cortico-pallidal beta signalling in the cortex-basal ganglia feedback loops for motor control. Abnormal movement-related beta signalling may occur in the described motor network leading to impaired motor learning in dystonia. The oscillatory driving of information flow by the motor cortex could either be indirect via a corticostriatal connection or monosynaptic through the recently described super-direct pathway that links Area 4 and 6 with the internal pallidum (Milardi et al., 2014). It is interesting to note that both alpha and beta oscillatory activity is suppressed before and during movement, but demonstrate topographical and temporal differences that imply these rhythms as functionally distinct (Pfurtscheller and Lopes da Silva, 1999). PET studies have revealed remote effects of pallidal DBS on premotor and motor areas at rest (Detante et al., 2004) and during movement (Kumar et al., 1999). Moreover, neurophysiological parameters of motor cortex excitability, intracortical inhibition and induced plasticity tend to normalize in parallel with alleviation of dystonic symptoms after chronic DBS, as revealed with transcranial magnetic stimulation (Kühn et al., 2003; Tisch et al., 2006; Ruge et al., 2011a, b). Modulation of cortical activity patterns may be facilitated by a functional reorganization through
DBS-related modulation of oscillatory long range network synchronization.

Although there is a vast amount of literature that has established MEG recordings for the characterization of cortical surface activity, fewer studies have shown activity from deeper brain structures such as the cerebellum. Nevertheless, cerebellar signals picked up by MEG have been reported in association with somatosensory stimulation (Tesche and Karhu, 1997, 2000b; Stancak et al., 2011), motor tasks (Jousmaki et al., 1996; Gross et al., 2002; Pollok et al., 2005; Jerbi et al., 2007; Bourguignon et al., 2012), saccades (Ioannides et al., 2005) and pathological conditions, such as epileptic seizures (Mohamed et al., 2011), parkinsonian tremor (Timmermann et al., 2003; Pollok et al., 2009) and in patients with writer’s cramp during writing (Butz et al., 2006). Our results extend these findings, suggesting that the cerebellum is interconnected with the internal pallidum through 7–13 Hz alpha band oscillations. A cerebello-thalamo-cortical network at ~10 Hz has been proposed to reflect a fundamental network for the synchronization of spatially separated motor areas for the maintenance of intermittent motor control (Welsh et al., 1995; Gross et al., 2002). Alteration of functional connectivity in cerebellar pathways is consistent with recent findings of functional and structural abnormalities in patients with different types of dystonia; for example, a cerebellar hypermetabolism found during rest (Eidelberg tonia; for example, a cerebellar hypermetabolism found during rest (Eidelberg et al., 1998) and with pallidal DBS using PET (Detante et al., 2004), as well as a reduced cerebellar activation during motor execution (Carbon et al., 2008; Argyelan et al., 2009; Blood, 2013). While there is extensive evidence that the cerebellum and the basal ganglia are interconnected (Bostan et al., 2010, 2013; Bostan and Strick, 2010; Kipping et al., 2013), no direct anatomical connection between the internal pallidum and the cerebellum has been described, making a monosynaptic oscillatory connection that could explain our findings highly unlikely. However, a close connection between the GPi and cerebellum via the red nucleus (a relay nucleus for afferent connections to the cerebellum and in the cerebello-thalamic circuit) has previously been reported and implicated in dystonia (Blood, 2013). More specifically, alterations in structural white matter connectivity between the internal pallidum and the red nucleus have been reported in a diffusion tensor imaging study of patients with dystonia (Argyelan et al., 2009; Blood et al., 2012). A pathway via the red nucleus offers a feasible explanation for the pallido-cerebellar oscillatory connectivity presented in this study, which would be corroborated by the pallidal drive of information flow found in the directionality analysis. Alterations in structural connectivity are likely associated with alterations in functional connectivity, which could lead to a disease-specific signature of oscillatory pallido-cerebellar network activity. The inverse correlation between pallido-cerebellar coupling and dystonic symptom severity suggests that alpha band connectivity is necessary for physiological motor and posture maintenance, and is disturbed in dystonia. Similar to the reported low frequency peaks in our patients, previous studies have shown that dystonia patients show increased low frequency (4–12 Hz) synchrony in the pallidal LFP signature (Silberstein et al., 2003; Liu et al., 2008) that drives (Sharott et al., 2008) and correlates (Chen et al., 2006) with dystonic muscle activity and is suppressed by DBS (Barow et al., 2014). Whether increased pallidal low frequency synchronization is related to the loss of pallido-cerebellar coupling, e.g. as a compensatory mechanism, or may disrupt cortico-pallidal coupling itself will have to be investigated in future studies. Sensorimotor cortico-pallidal low frequency coherence was not correlated with dystonic symptoms at rest in our patients. Future studies should explore frequency-specific network changes during movement to further define its pathophysiological role for abnormal movement generation. In this context, it is of note that reduced cerebello-thalamic fibre tract integrity has been revealed in patients with dystonia (Argyelan et al., 2009; Carbon et al., 2011). Here, a larger amount of cerebellar outflow tract disruption was related not only to the occurrence of dystonic symptoms but associated with increased motor cortical activation at rest and during movement, suggesting interdependent network modulation (Carbon et al., 2011). Similarly, increased cerebellar-motor cortical functional connectivity has been described as a potential compensatory mechanism in patients with writer’s cramp using functional MRI that was progressively reduced with larger symptom severity, indicating symptom-related neuroplastic compensatory network changes in dystonia (Dresel et al., 2014). Moreover, recent findings suggest that modulation of cerebellar output by continuous theta burst transcranial magnetic stimulation modulates motor cortex activity via cerebello-pallido-thalamo-cortical pathways and thereby may improve dystonia or levodopa-induced dyskinesia in Parkinson’s disease (Koch et al., 2009). It could be speculated that theta burst stimulation may either enhance cerebello-pallidal low frequency connectivity or may transiently disrupt abnormal pallidal oscillations, thus leading to short-term improvement in dystonic symptoms. If so, cerebellar transcranial magnetic stimulation could be of potential interest to define optimal DBS candidates in the future.

In conclusion, we have demonstrated that cortico-pallidal circuits are interconnected via spatially and spectrally distinct oscillatory networks. Our findings suggest the existence of three segregated circuits that show a disease-related pattern. Deep brain stimulation may interfere with these networks, and could modulate abnormal motor circuit signalling in dystonia.

Acknowledgements

We would like to thank Lutz Trahms (Physikalisch Technische Bundesanstalt, Berlin, Germany), Vadim Nikulin (Neurophysics Group, Department of Neurology,
Charité - University Medicine Berlin, Germany) and Gabriel Curio (Neurophysics Group, Department of Neurology, Charité - University Medicine Berlin, Germany) for helpful discussions.

**Funding**

This work was supported by the German Federal Ministry of Education and Research (BMBF) project ‘dystrac’. W.J.N. was supported by a research grant by the German Society For Clinical Neurophysiology and Functional Imaging (DGKN). A.A.K. was supported by the German Research Foundation (DFG) grant KFO247.

**Supplementary material**

Supplementary material is available at Brain online.

**References**


Cortico-pallidal connectivity in dystonia

BRAIN 2015: 138: 1894–1906 | 1905


Mattout J, Henson RN, Friston KJ. Canonical source reconstruction for MEG. Comput Intell Neurosci 2007; 67613.


