# Somatotopy of cervical dystonia in motor-cerebellar networks: Evidence from resting state fMRI

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# Somatotopy of Cervical Dystonia in motor-cerebellar networks: evidence from resting state functional connectivity

## Running title: Resting state fMRI in Cervical Dystonia

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## Abstract

Background

Cervical dystonia is the most frequent form of isolated focal dystonia. It is often associated with a dysfunction in brain networks, mostly affecting the basal ganglia, the cerebellum, and the somatosensory cortex. However, it is unclear if such a dysfunction is somato-specific to the brain areas containing the representation of the affected body part, and may thereby account for the focal expression of cervical dystonia.

In this study, we investigated resting state functional connectivity in the areas within the motor cortex and the cerebellum containing affected and non-affected body representation in cervical dystonia patients.

Methods

Eighteen patients affected by cervical dystonia and 21 healthy controls had resting state fMRI. The functional connectivity between the motor cortex and the cerebellum, as well as their corresponding measures of gray matter volume and cortical thickness, were compared between groups. We performed seed-based analyses, selecting the different body representation areas in the precentral gyrus as seed regions, and all cerebellar areas as target regions.

Results

Compared to controls, patients exhibited increased functional connectivity between the bilateral trunk representation area of the motor cortex and the cerebellar vermis 6 and 7b, respectively. These functional abnormalities did not correlate with structural changes or symptom severity.

Conclusions

Our findings indicate that the abnormal function of the motor network is somato-specific to the areas encompassing the neck representation. Functional abnormalities in discrete relevant areas of the motor network could thus contribute to the focal expression of CD.

## Introduction

Cervical dystonia (CD) is the most common form of isolated focal dystonia, characterized by involuntary muscle contractions in the neck, which results in abnormal head posture and movements.[1] It has been associated with various brain dysfunctions, such as maladaptive neuroplasticity, abnormal sensorimotor processing and integration,[2] and its pathophysiological mechanisms are still unclear.[3]

CD is considered a network disorder arising from abnormal communication among different brain areas.[2] Neuroimaging studies have evidenced functional and structural abnormalities in the basal ganglia,[2, 4] the sensorimotor and frontoparietal regions, the insula, the cerebellum, and the brainstem.[5-7] Animal and human research indicate that both the cerebello-thalamo-cortical network and basal ganglia–thalamo-cortical network project into the motor cortex, where the motor output is generated,[8] and may contribute to the abnormal movements.[9] Other studies focusing on the cerebellum have reported loss of Purkinje cells, areas of focal gliosis,[10] as well as increased gray matter (GM) volume of cerebellar flocculus, in CD patients compared to healthy controls (HC).[11] Altogether, these findings point to the motor cortex and the cerebellum as critical structures for the pathogenesis of CD.

It is mostly unknown whether these network abnormalities represent a general marker of the pathogenesis of dystonia, irrespectively of the affected body part, or if the pathogenesis of different types of dystonia affects relevant discrete areas within the motor areas and cerebellum. Distinct patterns of altered microstructures within regions of basal ganglia and cerebellar circuits have been associated with different phenotypes of focal dystonia,[12] and regional patterns of functional connectivity within the striatum and a sensorimotor-parietal network, as opposed to global network dysfunction, may contribute to focal dystonia.[13, 14] So far, there is no detailed study of the cerebellar somatotopy in CD, since a high-definition functional cerebellar atlas has been developed only recently.[15] Such an investigation is not trivial, as it has been shown that the representations of multiple body parts are organized in an orderly manner in the cerebellar lobes, mirroring the functional specialization of the motor cortex.[15, 16]

In this study, we investigated the specificity of motor-cerebellar networks in CD using resting state functional connectivity (RS-FC). RS-FC is a widely used non-invasive technique, based on functional magnetic resonance imaging (fMRI), where the time courses of predefined regions of interest (ROIs) are extracted from the brain at rest, and correlated with each other, under the assumption that functionally connected areas show high correlation.[17] We compared patterns of RS-FC in the different body representation areas of the motor cortex and the cerebellum between patients affected by CD and

HC, and performed a morphometric analysis on the same areas to study differences in GM volumes and cortical thickness (CT) between groups. We tested the hypothesis that network dysfunctions in CD are somato-specific, i.e., CD patients exhibit abnormal RS-FC and structural differences only in the head and neck representation areas, in both the motor cortex and the cerebellum.

## Materials and Methods

### ***Subjects and general procedure***

We recruited 18 CD patients (7 male, mean age=46.9±8.7 years) and 21 sex- and age-matched HC (9 male, mean age=45.3±10.6 years). Patients were recruited at the Pitié-Salpêtrière Hospital, Paris, FR. Inclusion criteria for patients were: a diagnosis of CD, no botulin toxic injection within 3 months prior to the examination, and stable pharmacological treatment in the month preceding inclusion. Exclusion criteria common to both HC and CD patients were: i) any other neurological sign, and ii) incompatibility with MR acquisition. Severity of CD was assessed at the time of inclusion with the Toronto Western Spasmodic Torticollis Rating Scale (TWSTRS), subscale for severity.[18] Between-group differences in age were assessed with independent-sample t-tests, whereas differences in the ratio between male and female participants were assessed with $\chi^2$ tests.

The study was carried out in accordance with the latest version of the Declaration of Helsinki and approved by the local Ethics Committee (approval number: C17-04 - AU 1360, ClinicalTrial.gov ID: NCT03351218). All participants gave written informed consent prior to the study.

### ***Neuroimaging acquisition parameters and pre-processing***

During the MR session, participants lied in the scanner while fixating a cross displayed on a screen. Their gaze was monitored with eye-tracking. Neuroimaging data were acquired using a 3T Magnetom Prisma (Siemens, DE) with a 64-channel head coil. Resting state fMRI and structural images were acquired in one session. Structural images were acquired with a T1-weighted MP2RAGE sequence with Repetition Time (TR)=5 s, Inversion Time (TI)=700/2500 ms, field of view (FOV)=232×256 in plane ×176 slices, 1 mm isotropic, Ipat acceleration factor=3. FMRI data were acquired with an echo-planar imaging (EPI) sequence performed with a multi-slice, multi-echo acquisition, TR=1.9 s, Echo Time (TE)=17.2/36.62/56.04 ms, Ipat acceleration factor=2, Multi-band=2, isotropic voxel size=3 mm,

dimensions=66×66 in plane ×46 slices, 350 volumes, duration=11 min. The first 10 time points were not recorded to ensure magnetization equilibrium.

Image preprocessing was done as follows: structural images were background denoised[19, 20] in order to improve the quality of the subsequent steps, segmented and normalized to the Montreal Neurological Institute (MNI) space using the Computational Anatomy Toolbox (CAT12)[21] extension for SPM12.[22] Functional images were pre-processed according to standard pipelines (despiking, slice timing correction and realignment to the volume with the minimum outlier fraction driven by the first echo) using AFNI.[23] A brain mask was computed on the realigned shortest echo temporal mean using FSL BET[24] in order to increase the robustness against signal bias intensity. Afterwards, the TEDANA toolbox[25] version 0.0.7 was used to optimally combine the realigned echoes, to apply principal component analysis and reduce the dimensionality of the data, and to perform an independent component analysis (ICA) decomposition to separate BOLD and non-BOLD components.[26] This step ensured robust artefact removal of non-BOLD signals, such as movement, respiration or heartbeat, and has already shown to be superior over standard denoising techniques in regressing out motion.[25] Framewise displacement (FD) was further computed according to standard methods,[27] and compared between CD patients and HC. The quality of the signal was verified as head movement amplitude was minimal, and the FD did not statistically differ between CD patients (0.018±0.016 mm) and HC (0.016±0.008 mm) [$t(37)=0.49$, $p=0.629$]. Finally, using SPM12, functional images were co-registered to the T1-weighted image, normalized to MNI space, and smoothed with a Gaussian kernel with full width at half maximum of 4x4x4 mm, as previously suggested.[28]

After pre-processing, the CONN toolbox[29] implemented in Matlab r2018a (The MathWorks Inc. USA) was used to parcellate the brain images into 274 functional regions, based on the Brainnetome Atlas,[28] and to extract the region-averaged time series. Motion parameters obtained during the realignment, as well as the average signal of white matter and cerebrospinal fluid obtained during the segmentation, were regressed out with aCompCor.[30] This step reduced spatial correlations resulting from physiological noise. Time series were finally band-pass filtered at $0.01<f<0.1$ Hz, according to previous research.[28]

***Analysis of resting state functional connectivity***

We entered all the time series extracted from the 274 functional regions into a first level general linear model (GLM), where we performed a ROI-based analysis, for each participant, to determine significant

resting state connections at individual level: In particular, we used bivariate correlation coefficients between all pairs of ROIs as indicators of their functional connectivity. Next, we converted the correlation coefficients to z-scores using Fisher-Z transformation, in order to normalize them to a Gaussian distribution. We then implemented a second level GLM testing seed-based ROI-to-ROI differences between CD and HC. For the latter, as we were only interested in the connectivity between the motor cortex and the cerebellum, we isolated the regions of the Brainnetome Atlas that were associated with the body representation areas within the precentral gyrus (PrG) and paracentral lobule (PCL), as well as all cerebellar regions without a priori, i.e. irrespective of body representation (Table 1). We then selected the bilateral motor areas as seed ROIs and the cerebellar ROIs as target regions. False positive control for multiple comparisons was implemented using false discovery rate (FDR)-corrected p-values with a threshold of $p_{FDR}<0.050$.

A correlation analysis was performed by computing Pearson correlation's coefficients between the severity of dystonia (TWSTRS scores) and the connectivity values between the motor cortex and the cerebellum. A FDR-corrected threshold of $p_{FDR}<0.050$ was applied.

Table 1. List of seed and target ROIs used in the analysis of functional connectivity. The Brainnetome Atlas was used to isolate all regions associated with the body representation areas within the motor cortex, as well as all cerebellar regions. R = right, L = left, PrG = precentral gyrus. PCL = paracentral lobule.

| **Seed regions (Motor cortex)** | **MNI coordinates (x,y,z)** | **Target regions (Cerebellum)** | **MNI coordinates (x,y,z)** |
|---|---|---|---|
| R / L PrG, area 4 (head and face region) | R: 55, -2, 33<br>L: -49, -8, 39 | R / L Lobules 1, 2, 3, 4 | R: 10 -43 -18<br>L: -7 -44 -17 |
| R / L PrG, caudal dorsolateral area 6 | R: 33, -7, 57<br>L: -32, -9, 58 | R / L Lobule 5 | R: 14 -51 -19<br>L: -13 -50 -19 |
| R / L PrG, area 4 (upper limb region) | R: 34, -19, 59<br>L: -26, -25, 63 | R / L Lobule 6 | R: 24 -58 -25<br>L: -23 -59 -25 |
| R / L PrG, area 4 (trunk region, including neck) | R: 15, -22, 71<br>L : 15, -22, 71 | R / L Lobule 7b | R: 28 -66 -51<br>L: -26 -66 -51 |
| R / L PrG, area 4 (tongue and larynx region) | R : 54, 4, 9<br>L: -52, 0, 8 | R / L Lobule 8a | R: 26 -58 -53<br>L: -24 -57 -53 |

| | | | |
|---|---|---|---|
| R / L PrG, caudal ventrolateral area 6 | R: 51, 7, 30<br>L: -49, 5, 30 | R / L Lobule 8b | R: 18 -51 -55<br>L: -17 -50 -55 |
| R / L PCL, areas 1, 2, 3 (lower limb region) | R: 10, -34, 54<br>L: -8, -38, 58 | R / L Lobule 9 | R: 7 -53 -49<br>L: -7 -53 -48 |
| R / L PCL, area 4 (lower limb region) | R: 5, -21, 61<br>L: -4, -23, 61 | R / L Lobule 10 | R: 22 -37 -46<br>L: -21 -37 -45 |
| | | R / L Crus 1 | R: 38 -68 -32<br>L: -36 -68 -32 |
| | | R / L Crus 2 | R: 26 -76 -41<br>L: -26 -75 -42 |
| | | Vermis 6 | 1 -70 -21 |
| | | Vermis / Crus 1 | -4 -78 -27 |
| | | Vermis / Crus 2 | 1 -75 -31 |
| | | Vermis 7b | 0 -68 -31 |
| | | Vermis 8a | 0 -67 -38 |
| | | Vermis 8b | 0 -63 -42 |
| | | Vermis 9 | 0 -56 -37 |
| | | Vermis 10 | 1 -48 -35 |

***Analysis of structural data***

We performed a morphometric analysis to study differences in GM volume between CD patients and HC. After preprocessing, the normalized GM volumes obtained during the segmentation were smoothed using a 4 mm full breadth at half maximum kernel, in line with our functional analysis. Volumes from the ROIs in the motor cortex were extracted with CAT12[21] using the "ROI tool" option, whereas volumes in the cerebellar ROIs were extracted with the Spatially Unbiased Infratentorial toolbox (SUIT).[31] For the latter, we first isolated the infratentorial structures using *suit_isolate_seg*, we then performed an affine alignment to the specific SUIT template and applied a normalisation using *suit_normalize_dartel*, and finally we extracted the GM volumes using *suit_reslice_dartel*.

We also studied differences in CT between patients and controls, by extracting CT in the investigated ROIs with DL+DiReCT.[32] This method already showed high accuracy compared to standard instruments.[32] Due to the lack of available tools to reliably measure CT in the cerebellum, this analysis was performed only for the somatotopic regions of the motor cortex.

The extracted GM volumes and CTs were compared between groups with independent-sample t-tests implemented in SPSS 25 (IBM Inc., USA). A threshold of $p<0.050$ was selected.

## Results

No significant differences were found in age and sex ratio between male and female participants (Table 2). CD patients showed symptom severity of 18.3 ± 4.4, as assessed with the TWSTRS.

Table 2. Clinical and demographic information. Comparison of clinical and demographic scores between groups. TWSTRS = Toronto Western Spasmodic Torticollis Rating Scale, HC = healthy controls, CD = cervical dystonia,

| | **HC (N = 21)** | **CD (N = 18)** | **Statistics** |
|---|---|---|---|
| **Sex [male / female]** | 9 / 12 | 7 / 11 | $\chi(1) = 0.06$, $p = 0.802$ |
| **Age [years, mean ± SD]** | 45.3 ± 10.6 | 46.9 ± 8.7 | $t(37) = 0.53$, $p = 0.600$ |
| **TWSTRS** | - | 18.3 ± 4.4 | - |
| **Overall medication [N (% of CD)]** | - | 18 (100%) | |
| - Botox [N (% of TD)]* | - | 17 (94.4%) | |
| - Others (Tramadol, Levothyroxine, Diazepam, Escitalopram – [N (% of CD)] ) | - | 3 (16.7%) | |

* last Botox injection administered at least 3 months prior to the experiment.

***Resting state functional connectivity and structural analysis***

The analysis of RS-FC revealed increased functional connectivity, in CD patients compared to HC, of the bilateral trunk representation area 4 (PrG) with the cerebellar Vermis 6 [$F(2,36)=10.78$, $p_{FDR}=0.039$]

and Vermis 7b [$F(2,36)=10.33$, $p_{FDR}=0.039$], respectively (Figure 1). Detailed values of RS-FC in the two groups are displayed in Figure 2. No significant differences in the connectivity between the other seed ROIs and the cerebellum (Table 1) were found across groups ($p_{FDR}>0.050$ – Supplementary Figure 1).

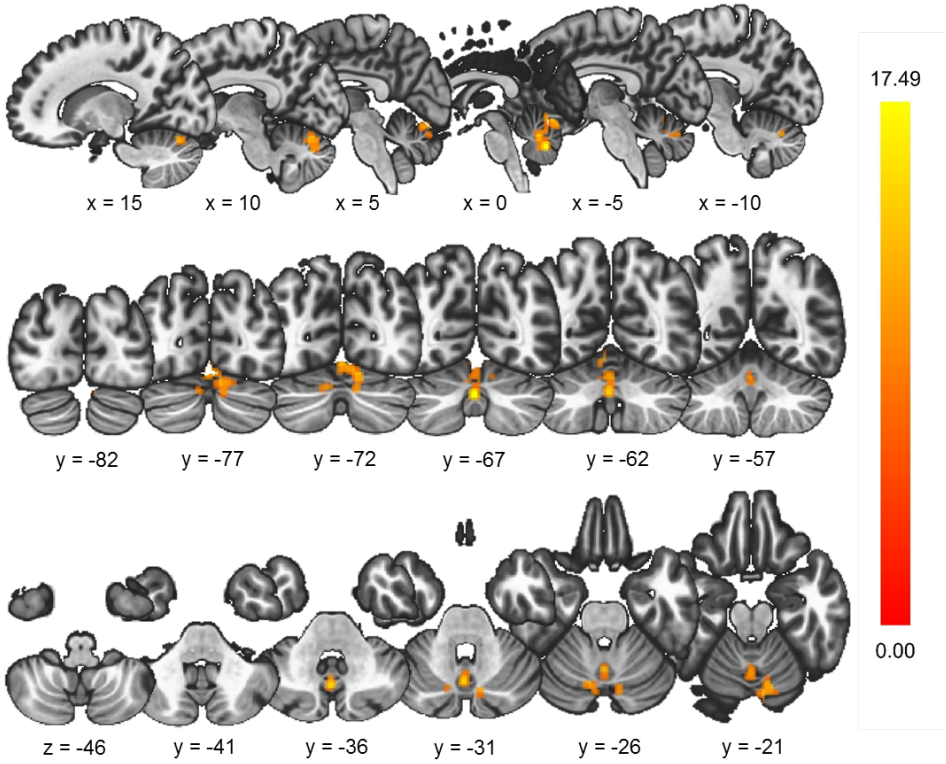


Figure 1. Between-group differences in the connectivity of the bilateral trunk representation area 4 of the PrG (seed) with the cerebellar Vermis 6 and 7b (target). Color bar represents the F-values assessed with the general linear model. For visualization purposes, the significant connectivity values are displayed irrespectively of the cerebellar ROIs.

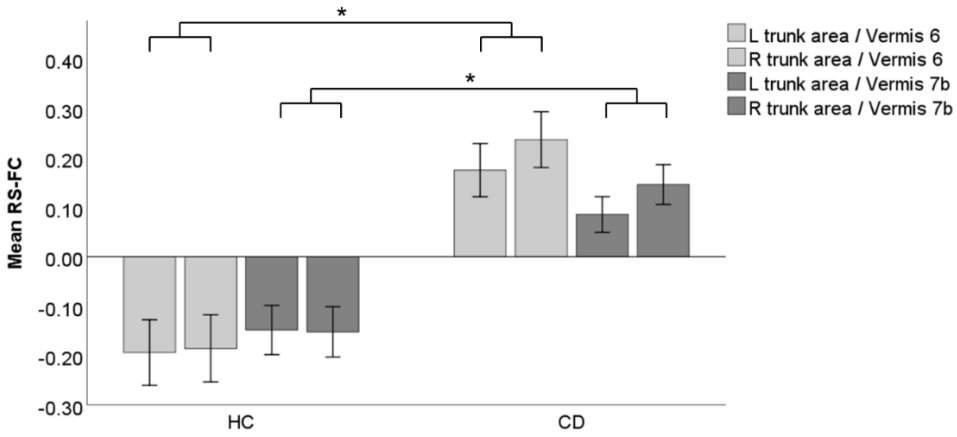


Figure 2. Mean values of resting state functional connectivity between ROIs. Error bars represent the standard error of the mean. * depicts significant differences ($p_{FDR}$<0.050) between CD patients and HC.

No correlation was found between any of the investigated connections and symptoms' severity in CD ($p_{FDR}$>0.050). Neither differences in GM volume, nor in CT, were found between the studied groups, in any of the investigated ROIs (p>0.050).

## Discussion

Compared to HC, patients with CD showed increased RS-FC between the neck and trunk representation in area 4 of the precentral gyrus (PrG) and the cerebellar Vermis 6 and 7b. This pattern was neither associated with differences in GM volumes or CT between groups, nor with symptoms' severity in CD. Our findings indicate that the abnormal function of the motor network is somato-specific to the areas encompassing the neck and trunk representation. Functional abnormalities in discrete relevant areas of the motor network could thus contribute to the focal expression of CD.

The use of resting state fMRI was a strength of this study, as it allowed us to investigate task-free patterns of functional connectivity, hence, to avoid a potential bias due to adaptive/compensatory processes associated with task execution. This is relevant, as dystonic symptoms tend to worsen during voluntary motor activity, and compensatory mechanisms are more likely to occur during task execution than rest.[33] However, neuroimaging analyses can hardly distinguish causes from effects, therefore the causal mechanisms here discussed are speculative.

The cerebellum is of particular importance for the pathophysiology of CD, as the topographical expression of dystonia depends on the extent of cerebellar dysfunction in a dedicated mouse model.[34]

In this model, a dysfunction of the entire cerebellum induces a phenotype similar to generalized dystonia, whereas dysfunctions in limited cerebellar regions restrict abnormal movements to isolated body parts, like in focal dystonia. Hence, the extent, but not the location of the cerebellar dysfunction, seems to be linked to the severity of dystonia. Altered connectivity between the cerebellum and the sensorimotor areas has also been associated with the pathogenesis of CD.[3, 35, 36] However, reported changes were not limited to the somatotopic representation of the affected body parts,[3, 37, 38] and the potential link between critical areas within these regions and the location of dystonic manifestations is still poorly understood. Our results challenge this view, and provide evidence of well-defined regions in both the precentral gyrus (PrG) and the cerebellum specifically relevant to the clinical expression of CD, namely the trunk representation area 4 of the PrG and the cerebellar Vermis 6 and 7b. No other areas within the PrG and the cerebellum showed any group differences in RS-FC between patients and HC, thereby reinforcing the hypothesis of a selective area, with a critical role in motor control of head and neck, responsible for the pathogenesis of CD. More specifically, the trunk representation area 4 of the PrG contains neuronal populations responsible for the sternocleidomastoid muscle,[39, 40] typically affected in CD. Likewise, the cerebellar Vermis 6 has been related to saccadic eye movements in healthy subjects,[41, 42] and clinical studies have confirmed a link between abnormal cerebellar output and an impairment in saccadic adaptation,[43] vestibule-ocular reflex,[44] and eye-hand coordination[45] in CD. This may reflect cerebellar-related maladaptive plasticity associated with an attempt to compensate the abnormal head posture with a modulation of eye movements.[46, 47]

Abnormal connectivity of brain networks linking the cerebellum and the motor cortex has been largely investigated for its role in the pathogenesis of dystonia.[4] In non-human primates[48] and mice,[34] dystonic movements can be provoked by manipulations of the cerebellum, abnormal firing of the Purkinje cells has been found in DYT1 torsin1-knock-in mice,[49] and acute knock-down of Sgce in the cerebellum produces motor symptoms close to myoclonus dystonia in mice.[50] In humans, abnormal anatomical cerebello-thalamo-cortical connectivity can play a role in the clinical expression of dystonia,[51] as well as in the loss of cerebellar control over sensorimotor plasticity.[52] CD patients with a sensory trick show a differential ability to modulate the connectivity of the sensorimotor network, likely through a cerebellar mediation,[36] and modulation of the cerebello-cortical connectivity has been associated with the clinical improvement following botulinum toxin injections in these patients.[53]

We hypothesize that the abnormal connectivity between the motor cortex and the cerebellum found in our study reflects a cerebellar dysfunction mediating sensorimotor integration and maladaptive plasticity. This interpretation is consistent with the head neural integrator model,[54, 55] by which changes in any of the inputs of the integrator affect the communication between the cerebellum and cortical areas, and potentially lead to the manifestation of CD.[7, 56] In healthy subjects, proprioceptive input from the neck can substantially change the way cerebellar output influences plasticity at the level of the motor cortex,[57] and in CD, an abnormal integration of the neck proprioceptive information could drive an atypical functioning of the integrator,[57] which may generate head twists. The specific involvement of Vermis 6 and 7b found in our study provides further insights into the role of cerebellar areas associated with saccadic eye movements in the integrator dysfunction of CD.[11, 46]

The investigated differences in functional connectivity were not associated with structural changes in CD, as evidenced by our analysis of GM volume and CT, and this supports the vision that CD mainly affects brain functions. Research on this topic has shown controversial results, and while the classic assumption is that CD is not associated with structural changes in the brain,[58] some studies have reported altered GM concentration in CD patients in the cerebellar flocculus, as well as in the basal ganglia, the thalamus and the motor cortex.[11, 59] Such differences from our findings may be explained by different methodological approaches: While the previously used whole-brain approaches have identified differences in various structures, regardless of their functional roles, we focused on areas associated with specific functions relevant for CD, such as neck movements, and therefore opted for a ROI approach, which grouped together voxels belonging to the same functional areas. Even though our choice may have been more conservative than other approaches, it suggests that CD-related brain dysfunctions are not linked to structural changes. Future research should further investigate this topic, for instance by applying multimodal imaging techniques or ultra-high field MR, which could reveal subtle structural changes undetectable with conventional MRI.

In conclusion, our results point to an impairment in the communication of somato-specific cerebello-cortical networks related to head position and saccadic eye movements.[4] This impairment might be the consequence of abnormal processing of proprioceptive input from the neck, which affects the functioning of the head neural integrator, and in turn generates abnormal head posture, as well as related compensatory eye movements.

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## Authors' roles

Research project. Conception and organization: ER, YW. Execution: CT, AE, CH, SS, CB, CD, AM, BD, FB, MAB, PJ.

Statistical analysis. Design and execution: GAZ, CG, BB, MR, YW. Review and critique: GAZ, CG, BB, MR, ER, YW.

Manuscript. Writing of the first draft: GAZ, YW. Review and critique: GAZ, CG, BB, MR, CT, AE, CH, SS, AMB, CD, AM, BD, FB, MAB, PJ, ER, YW.

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## Figures' captions

Figure 1. Between-group differences in the connectivity of the bilateral trunk representation area 4 of the PrG (seed) with the cerebellar Vermis 6 and 7b (target). Color bar represents the F-values assessed with the general linear model. For visualization purposes, the significant connectivity values are displayed irrespectively of the cerebellar ROIs.

Figure 2. Mean values of resting state functional connectivity between ROIs. Error bars represent the standard error of the mean. * depicts significant differences ($p_{FDR}<0.050$) between CD patients and HC.