Imbalance between the caudate and putamen connectivity in obsessive–compulsive disorder

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\textbf{ABSTRACT}

Background: Compulsive behaviors in obsessive–compulsive disorder (OCD) have been suggested to result from an imbalance in cortico-striatal connectivity. However, the nature of this impairment, the relative involvement of different striatal areas, their imbalance in genetically related but unimpaired individuals, and their relationship with cognitive dysfunction in OCD patients, remain unknown.

Methods: In the current study, striatal (i.e., caudate and putamen) whole-brain connectivity was computed in a sample of OCD patients (OCD, n = 62), unaffected first-degree relatives (UFDR, n = 53) and healthy controls (HC, n = 73) by ROI-based resting-state functional magnetic resonance imaging (rs-fMRI). A behavioral task switch paradigm outside of the scanner was also performed to measure cognitive flexibility in OCD patients.

Results: There were significantly increased strengths (Z-transformed Pearson correlation coefficient) in caudate connectivity in OCD patients. A significant correlation between the two types of connectivity strengths in the relevant regions was observed only in the OCD patient group. Furthermore, the caudate connectivity of patients was negatively associated with their task-switch performance.

Conclusions: The imbalance between the caudate and putamen connectivity, arising from the abnormal increase of caudate activity, may serve as a clinical characteristic for obsessive–compulsive disorder.

1. Introduction

Obsessive-compulsive disorder (OCD) is surprisingly common in the general population (2.5%–3%) with a high genetic risk (Robbins et al., 2019). This disorder is characterized by persistent obsessions and compulsions. Specifically, compulsion (persistent behavior, despite negative consequences) is one of the core manifestations of the disorder, and may play an important role in the mechanism of OCD.

For understanding the pathomechanism of OCD behaviors, we need to know its cognitive and neural foundations (Figee et al., 2016; Gillan et al., 2016; Gillan and Robbins, 2014; Robbins et al., 2019; Simmler and Ozawa, 2019), in particular the cortico-striatal circuits (Dong et al., 2020; Heilbrunner et al., 2016; Morelli et al., 2011; Nagarajan et al., 2018; Pinhal et al., 2018). Previous studies proposed that the connectivity from cortex to dorsomedial (caudate) and dorsolateral striatum (putamen) constitute parallel loops, namely the associative and motor loops, respectively (Middleton, 2000; Milad and Rauch, 2012; O’Doherty et al., 2017; Seger, 2018; Yin and Knowlton, 2006). The associative loop connecting the caudate and prefrontal cortex is associated with goal-directed and flexible behaviors (Dong et al., 2020; Seger, 2018; Simmler and Ozawa, 2019). The motor loop connecting putamen with premotor and sensorimotor cortex is vital in habits (de Wit et al., 2012; Seger, 2018). Since compulsivity has been characterized as an imbalance between the brain’s goal-directed and

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habit-learning systems (Gillan et al., 2016), these loops are thought to mediate cognitive control of learning and behavior, and its automation into habits. It is understandable then that abnormal or excessive active avoidance has been perceived as a key impairment in OCD (Geramita et al., 2020).

Converging evidence indicates that compulsive behaviors may be associated with an imbalance between the associative and motor loops (Dolan and Dayan, 2013; Dong et al., 2020; Gillan et al., 2014b; Gillan and Robbins, 2014). When functioning properly, these two systems shift flexibly according to the external environment and feedback. Disruption of either of the two loops could result in an imbalance between them, and consequently the emergence of compulsive behavior (Banca et al., 2015; Dong et al., 2020; Gillan et al., 2014a, 2014b, 2011). However, it’s unclear which of the two loops (or both) drives the imbalance. Few studies have investigated the direct relationship between these two systems; however, such study could promote our understanding of their interaction in OCD (Geramita et al., 2020).

The risk of OCD is significantly higher for relatives of patients (Gottesman and Gould, 2003; Nestadt et al., 2010; Pauls, 2008; Pauls et al., 2014). There could thus be a genetic basis to OCD, although the specific gene and its relation with behavior remains elusive. The concept of endophenotype (Gottesman and Shields, 1973) was adopted to understand the gap between genetics and behavioral disease processes. Behaviorally, OCD patient probands and their unaffected first-degree relatives (UFDR) showed cognitive inflexibility (Chamberlain et al., 2007). Neurally, OCD patients and their UFDR exhibit an associated reduction or increase in brain functional and structural variation involved in the associative loop, compared to healthy controls (HC) (Menzies et al., 2007). Specifically, patients and their UFDR show leftward asymmetry of cortical thickness in the anterior cingulate cortex (ACC) (Peng et al., 2015). Another investigation identified reduced activation of orbitofrontal cortex, during reversal learning in patients and their UFDR (Chamberlain et al., 2008). A recent resting-state fMRI (rs-fMRI) study of our lab found that patients and UFDR showed greater effective connectivity between the left caudate and frontal cortex than HC (Dong et al., 2020). These results indicate that the abnormal activity or connectivity in the associative loop may be a hereditary risk factor for OCD. Considering that cognitive flexibility is an identified endophenotype of OCD (Robbins et al., 2019), it’s critical to clarify this loop as a whole and its relation with cognitive flexibility.

Currently, based on the previous study where we investigated the effective connectivity of specific areas only, we used rs-fMRI to measure whole-brain functional connectivity of caudate and putamen in OCD, UFDR and HC groups. First, we investigated the imbalance between caudate and putamen connectivity in OCD and measured whether it was also observed in UFDR of OCD patients, which could thus serve as a clinical characteristic for this disorder. Second, we explored whether the severity of obsessive–compulsive symptoms could influence the interaction between the two (caudate and putamen) systems. Finally, previous studies reported that goal-directed associative learning requires people to flexibly use feedback according to task demands. Cognitive inflexibility might prevent patients from shifting from one thought to another, thus rendering their behavior stimulus-driven and inflexible (Mayr and Keele, 2000; Monsell, 2003). Thus, we investigated whether cognitive inflexibility is related to these two loops.

2. Materials and Methods

2.1. Participants

Healthy controls (n = 73; males = 51), OCD patients (n = 62; males = 45) and their unaffected first-degree relatives (n = 53; males = 32) were recruited for the study from Guangzhou Psychiatric Hospital. All participants, aged 18 to 55, underwent the diagnosis performed by a clinical psychiatrist and an experienced psychologist. In accordance with the institutional research and ethics committee of Guangzhou Psychiatric Hospital, each subject gave written informed consent after understanding the complete study description. Both OCD patients and their UFDR were recruited in the hospital. All procedures contributing to this work comply with the ethical standards of the relevant national and institutional committees on human experimentation and with the Helsinki Declaration of 1975, as revised in 2008.

The patients with OCD received the Structured Clinical Interview (SCID) for DSM-IV-TR Axis I disorders (First et al., 2002), fulfilling Diagnostic and Statistical Manual of Mental Disorders, 4th edition (DSM-IV) criteria for OCD. Their unaffected first-degree relatives and healthy controls were assessed by the SCID for the DSM-IV-TR Axis I disorders, Research Version, Non-Patient edition (SCID-I/NP) (Spitzer et al., 2002). All participants had no history of traumatic brain injury or neurological disease and did not exhibit alcohol/substance abuse. The UFDR of patients and healthy controls were excluded if they reported any history of mental illness and/or treatment with any psychotropic medication. On this basis, healthy controls had an additional exclusion criterion of no family history of Axis I or Axis II psychiatric disorders.

Thirty-one OCD patients took normal psychotropic medications while scanning (see Supplementary Material, Table S1). Twenty patients had comorbid disorders such as anxiety or depression. It is worth noting that having only comorbid anxious and depressive symptoms were not considered as an exclusion criterion, provided that OCD was the primary clinical diagnosis.

2.2. Clinical assessments

The Yale-Brown Obsessive-Compulsive Scale (Y-BOCS) (Goodman et al., 1989) was administered to assess illness severity. The Obsessive-Compulsive Inventory-Revised (OCI-R) (Foa et al., 2002; Peng et al., 2011) was used for measuring the degree of distress or being disturbed with common OCD phenomena (Fernandez-Egea et al., 2018). The Beck Depression Inventory (BDI) (Beck and Steer, 1984) was used to estimate depressive symptoms, and the State-Trait Anxiety Inventory (STAI) (Spielberger, 1989) was used to measure anxiety symptoms. Individual clinical assessment measures were compared between or across groups using chi-square analyses and Analysis of Variance in SPSS version 20.0.

2.3. Imaging data acquisition

All MRI data were acquired on a 3.0-Tesla MR system (Philips Medical Systems Nederland B.V.) equipped with an eight-channel phased-array head coil. The resting-state functional MRI data were collected with the following parameters: gradient echo Echo-Planar Imaging (EPI) sequences; time repetition, TR = 2000 ms; echo time, TE = 30 ms; flip angle = 90°, 33 slices, field of view [FOV] = 220 mm × 220 mm, matrix = 64 × 64; slice thickness = 4.0 mm; voxel size = 3.4 × 3.4 × 4 mm³. For each participant, the fMRI scanning lasted for 480 s and generated 240 whole-brain volumes. During the scanning, participants were instructed to lie quietly with their eyes closed, and stay awake without moving. For spatial normalization and localization, the high-resolution T1-weighted anatomical images were obtained by using a magnetization prepared gradient echo sequence with the following parameters: TR = 8 ms, TE = 3.7 ms, flip angle = 7°, FOV = 240 mm × 240 mm, matrix = 256 × 256, slice thickness = 1.0 mm; voxel size = 1.0 × 1.0 × 1.0 mm³.

2.3.1. Functional imaging data preprocessing

The data was preprocessed using the Statistical Parametric Mapping toolbox (SPM12, https://www.fil.ion.ucl.ac.uk/spm), and Data Processing Assistant for Resting-State fMRI (DPARSFA version 4.4, http://rfMRI.org/dpabi). For image preprocessing, the first 10 time points were removed to denoise the signal. The remaining 230 volumes were corrected for slice timing and head motion, as all the subjects had no>1.5° of maximal rotation and 1.5 mm of maximal translation. After realignment with the corresponding T1-volume and visual inspection as
the image quality control method to exclude non-conforming images, the nuisance covariates (six head motion parameters, white matter signal and cerebrospinal fluid signal) were regressed in first-level analysis. Next, the functional data were normalized into the stereotactic space of the Montreal Neurological Institute and resampled at 3 × 3 × 3 mm³. The processed images were spatially smoothed with a 6-mm full-width half-maximum isotropic Gaussian kernel. Further preprocessing pipeline consisted of band-pass filtering (0.01–0.08 Hz) to reduce the effects of physiologic noise and micro-head-motion correction according to frame-wise displacement (FD) by replacing the rs-fMRI volume with FD > 0.5 mm (nearest neighbor interpolation).

2.3.2. Caudate connectivity and putamen connectivity construction

Two ROIs for functional connectivity were defined according to previous studies (Di Martino et al., 2008; Dong et al., 2020). The caudate ROI was constructed using the MNI-coordinates: X = ±13, Y = 15, Z = 9; the MNI-coordinates of the putamen ROI were: X = ±28, Y = 1, Z = 3. These ROIs were defined by spheres surrounding the central voxel with a radius of 3 mm. A previous study includes additional details about the anatomical delineation of these regions (Di Martino et al., 2008). We drew a plot to show the location of two spherical ROIs to demonstrate there is no overlap between them (see Supplementary Material, Fig. S3). All anatomical regions mentioned in the results of the study are identified by the automated anatomical labeling (AAL) atlas.

The averaged time course within each seed was extracted and correlated with all voxels in the entire brain to generate functional connectivity for the putamen and the caudate. Due to anatomical proximity, the caudate was regressed out as a covariate when analyzing the putamen, and vice versa. Then the Pearson correlation coefficients (r) between ROI activation and each voxel were subsequently Fisher-Z transformed: Z = 0.5 × ln((1 + r)/(1−r)). One-sample t tests in each group on functional connectivity maps for the caudate connectivity and putamen connectivity were conducted respectively, using the rest toolbox (https://restfmri.net/forum/REST_V1.8). Then, we obtained six connectivity maps as masks by False Discovery Rate used for cluster-level multiple comparisons correction (p < 0.001, FDR corrected). Afterwards, a union mask of the caudate connectivity including all significant connections in any of the three groups was produced, and another union mask of putamen connectivity was generated in the same way. Then each subject obtained the final connectivity maps of the two brain regions by multiplying the data of their two raw functional connectivity maps with the corresponding union mask respectively. Finally, the functional connectivity values (Z) were averaged for suprathreshold caudate and putamen connectivity separately. We referred to the mean Fisher-Z transformed value in caudate connectivity map as caudate connectivity strength; and the mean Fisher-Z transformed value in putamen connectivity map as putamen connectivity strength. Hence, each of the two ROIs (caudate, putamen) for each group, has a connectivity map (FDR corrected) and a mean functional connectivity strength value for further analysis that employed Analysis of Variance (ANOVA) and two-sample test at p < 0.001 corrected (cluster level p < 0.05, voxel level p < 0.001, corrected).

2.4. Task-switching paradigm

The task-switching paradigm was used to investigate cognitive flexibility (Gu et al., 2007). This paradigm was conducted outside the fMRI scanner (see Supplementary Material, Fig. S2), in the OCD patients. Participants were required to learn associations between stimuli and responses. A total of 264 trials were presented. Half of the trials were task-switching conditions, during which the subjects attended to a different dimension as on the previous trial (e.g., a square cue followed by a diamond cue); the other half of the trials were task-repeat conditions during which the dimensions were the same as on the previous trial. During the 10-minute training phase, participants were required to practice and learn the stimulus-action mapping. During the test phase, reaction time (RT) and accuracy were recorded. Switching cost (task-switch RT minus task-repeat RT) was used as a measure of cognitive flexibility, which meant the lower an individual’s switching cost, the better his cognitive flexibility. Only the correct responses are included in the following analyses. The experimental flow was the same as in a previous study which explored abnormal brain activity related to cognitive inflexibility in OCD (Gu et al., 2007).

2.5. Brain-behavior correlation analysis

Moderation analyses were performed between mean caudate connectivity, mean putamen connectivity, and Yale-Brown Obsessive-Compulsive Scale (YBOCS) scores to examine associations between functional connectivity and OCD symptom severity. All variables are zero-centered.

Correlation analyses were performed between mean strength of the ROI-based connectivity and task-switching performance in the OCD group, to examine associations between functional connectivity and cognitive flexibility (RT and accuracy data exceeding plus or minus three standard deviations are excluded). The moderation and correlation analyses were conducted using SPSS 20.0.

3. Results

3.1. Demographic and clinical analysis

The demographic and clinical variables of each participant group are presented in Table 1. There were no differences in age (F(2,185) = 2.4, p = 0.10), gender (χ² = 2.1, p = 0.34) or education level (F(2,185) = 0.1, p = 0.87) across the three groups. The head motion, the YBOCS, the OCI-R, the BDI, and the STAI of OCD patients were significantly higher compared to the HC group and the UFDR group. Since age, gender, education, head motion, BDI and STAI may play roles in differences

| Measures                      | OCD (n = 62) | UFDR (n = 53) | HC (n = 73) | F/χ² | p     |
|-------------------------------|--------------|---------------|-------------|------|-------|
| Demographic                  |              |               |             |      |       |
| Age (years) (SD)             | 26.8(8.3)    | 23.9(8.3)     | 27.2(9.4)   | 2.4  | 0.10  |
| Gender (Male: Female)        | 45.17        | 32.21         | 51.22       | 2.1  | 0.34  |
| Education (years) (SD)       | 12.77(3.5)   | 11.4(3.1)     | 11.6(3.3)   | 0.1  | 0.87  |
| Head motion (SD)             | 0.0673       | 0.0574        | 0.0517      | 4.0  | <0.05 |
|                          | (0.03954)    | (0.02799)     | (0.02756)   |      |       |
| Clinical                     |              |               |             |      |       |
| YBOCS total (SD)             | 29.0(4.9)    | 3.8(3.3)      | 0(0)        | 32.9 | <0.001|
| YBOCS obsessions (SD)        | 14.6(2.9)    | 1.8(2.0)      | 0(0)        | 27.8 | <0.001|
| YBOCS compulsions (SD)       | 14.4(3.1)    | 1.9(2.1)      | 0(0)        | 24.7 | <0.01 |
|                          | (0.03954)    | (0.02799)     | (0.02756)   |      |       |

OCD: Obsessive-compulsive disorder group; UFDR: unaffected first-degree relative group; HC: healthy control group. YBOCS: Yale-Brown Obsessive-Compulsive Scale; OCI-R: Obsessive Compulsive Inventory–Revised; BDI: Beck Depression Inventory; STAI: State-Trait Anxiety Inventory; SD: standard deviation.
between groups, they were treated as nuisance variables and controlled in all subsequent data analysis.

3.2. Caudate connectivity and putamen connectivity

We measured the caudate and putamen whole-brain connectivity patterns in all participants by one-sample t-tests, and found the caudate connectivity map for all three groups included the ventral medial prefrontal cortex (vmPFC), the orbitofrontal cortex (OFC), the ACC, the dorsolateral prefrontal cortex (DLPFC) and an area of the cerebellum (Fig. 1, upper row). The putamen connectivity map for all three groups included the supplementary motor area (SMA), the insular cortex, the primary motor cortex and an area of the cerebellum (Fig. 1, lower row). Based on visual inspection (i.e., magnitude and distribution of T-values), the pattern of UFDR group was more similar to that of HC group for both caudate and putamen connectivity.

Then, we conducted a mixed design two-factor repeated measures ANOVA analysis to investigate the differences in caudate and putamen connectivity strength among OCD, UFDR and HC groups. The two factors in this model are group (OCD, UFDR, and HC) and network (or map) type (caudate, putamen), and the dependent variable is the mean connectivity strength values in these two maps. We observed a significant group main effect, $F(2,104) = 22.357, p < 0.001, \eta^2 = 0.200$; and a network type main effect, $F(1,52) = 9.748, p < 0.001, \eta^2 = 0.181$. Specifically, for putamen connectivity strength, there was no difference between any of the groups ($F(2,104) = 1.573, p = 0.210, \eta^2 = 0.017$); but for caudate connectivity strength, there was a significant difference between groups ($F(2,104) = 41.808, p < 0.001, \eta^2 = 0.318$) (Fig. 2).

3.3. Moderation analysis

3.3.1. Group differences in correlation between caudate and putamen connectivity

We conducted ANOVA and subsequent t-tests based on the supra-threshold caudate and putamen functional connectivity maps between the three groups or between each pair of groups using the rest toolbox (Fig. 3). Combining the results of these two analyses (i.e., magnitude and distribution of F-values and T-values), the inter-group differences of the caudate connectivity maps in three groups were mainly contributed by the differences between the OCD group and the other two groups. The ANOVA results showed abnormal connectivities between caudate and several areas including the bilateral ACC, superior and inferior frontal gyri, inferior parietal gyri, inferior and middle temporal gyri, angular gyri and left inferior cerebellum (Fig. 3, upper row). In combination with the connectivity strength results mentioned above, the OCD group showed similar areas of abnormal activation compared to either the UFDR group or the HC group. The abnormal connectivities between the putamen and several areas were mainly concentrated in the bilateral insular cortex and hippocampus (Fig. 3, lower row).

In order to clarify whether the group modulated the correlation between caudate and putamen connectivity maps, a moderation analysis was performed to explore the influence of the group on the relationship between caudate and putamen connectivity strengths in the altered areas. The latter were defined as the regions obtained from the ANOVA between the three groups separately. The caudate connectivity strength significantly interacted with the group in predicting connectivity strength of the putamen in the OCD group (see Fig. 4, $R^2 = 0.075$, $F(1,179) = 24.1462, p < 0.001$). Since the moderator was a categorical variable, linear regressions of the independent variable (caudate strengths) on the dependent variable (putamen strengths) were further carried out separately for the three groups. It was found that the regression coefficients were significantly positive solely in the OCD group ($R^2 = 0.117, \beta = 0.342, p < 0.01$; UFDR: $R^2 = 0.060, \beta = 0.080, p > 0.05$; HC: $R^2 < 0.001, \beta = 0.012, p > 0.05$). In sum, the significant correlation between the two types of connectivity strengths in the altered regions was observed only in the patient group, which may reflect the imbalance between the two networks.

3.4. Task switching analysis

Here we acquired behavioral results of 62 OCD patients. The error rate and mean reaction time (RT) under task repeat condition and task switch condition appear in Supplementary Material (Table S2). Both the repeat error rate and the switch error rate were significantly correlated with the Y-BOCS scores ($r = 0.295, p = 0.030; r = 0.298, p = 0.029$). Switching cost, reflecting cognitive flexibility, was calculated as mentioned before (task-switch RT minus task-repeat RT).

3.4.1. Relationship between ROI-based connectivity and cognitive flexibility

We investigated the relationship between functional connectivity strength of whole brain and cognitive flexibility. Longer task-switch RT was significantly associated with increased caudate connectivity strength, while the correlation between switching cost and caudate connectivity strength was marginally significant (see Fig. 5a and b; $r = 0.308, p = 0.023; r = 0.267, p = 0.051$). There was no significant correlation between putamen connectivity strength and switching cost ($r = 0.062, p = 0.658$). Since only OCD patients participated in the behavioral experiment, we were not able to assess these relationships between connectivity strength and cognitive flexibility in the UFDR group and HC group.

4. Discussion

The present study investigated the balance between the caudate and
putamen connectivity in HC, OCD patients and their UFDR using rs-fMRI. Compared with HC and UFDR, OCD patients showed significantly increased caudate connectivity strength of the whole brain, but not putamen connectivity strength. These deficits represent trait rather than state impairments, that can exist in the absence of medication confounds and clinical phenotype (Chamberlain et al., 2007). The moderation analysis and linear regression for all three groups found a correlation between the two types of connectivity in OCD patients only, which may indicate that these two types of networks were imbalanced in this sense. In addition, there was a significant relationship between the caudate connectivity of patients and their RT in task-switch trials, suggesting that the abnormally activated caudate connectivity in OCD patients is associated with worse performance in the behavioral task.

Given the importance of caudate and putamen in goal-directed and habitual learning respectively (Balleine et al., 2007; Burton et al., 2015; Cabeza, 2002; Cox and Witten, 2019; Cushman and Morris, 2015; Di Martino et al., 2008; Dolan and Dayan, 2013; Dong et al., 2020), the abnormal correlations between two types of connections found in the OCD patients may support the hypothesis that compulsive symptoms are associated with an imbalance between the goal-directed and the habitual learning system. An increasing number of studies have found an imbalance between these two systems in OCD patients (Robbins et al., 2019; Simmler and Ozawa, 2019). Using a “slips-of-action” test, researchers found that the ability for understanding action outcome value changes was impaired in OCD patients compared to HC (Gillan et al., 2011). Furthermore, using a novel shock avoidance task designed to induce habits through goal-devaluation by overtraining, they observed that OCD patients have a tendency to develop excessive avoidance habits (Gillan et al., 2014b). Using fMRI, the authors further found that these increased avoidance habits in OCD patients were associated with
and found strong increases involving connectivity between the cerebellum and regions outside the CSTC (e.g., the frontal gyrus and temporal gyri) could be related to visuo-spatial and sensory-motor processing, which may indirectly impact cognitive flexibility in OCD (Wolff et al., 2017). Our findings of a positive correlation between Y-BOCS and error rates during both repeat and switch events may seem surprising, but were consistent with previous studies reporting that differences in adaptation during repeat trials also played an important role (Han et al., 2011; Page et al., 2009; Woolley et al., 2008). One possible explanation is that the severity of OCD may be beneficial to accuracy at the expense of prolonged responding during repetition (Remijnse et al., 2013).

There are several limitations to the current study. First, the demographics of participants were not completely matched among the three groups. The sample was young and predominantly male, indicating an imbalance in gender distribution. Intelligence level, drug use, disease duration, co-morbidity and distinct subtypes of OCD patients may serve as confounding factors. Future research should better control these factors and/or examine the influence of these factors on OCD. Second, because of the limited manpower, the switch task was not administered in healthy controls. Third, the length of resting-state connectivity data may be fairly limited in the range of 5–10 min, suggesting the length of the resting state scan should be extended appropriately to obtain stable results. Lastly, this is a cross-sectional study. A longitudinal study would be important to obtain data related to the trajectory of network changes in OCD patients.

5. Conclusions

We identified the abnormally increased caudate connectivity in OCD patients mainly resulted from the outward extension of CSTC regions, which may be a compensatory mechanism for impaired CSTC loop in the disorder. The correlation between caudate and putamen connectivity was observed significantly only in OCD patients. These findings suggest the imbalance between the goal-directed and habitual cortico-striatal connectivity may serve as a clinical characteristic for OCD.

CRediT authorship contribution statement

Ziwen Peng: Conceptualization, Methodology, Writing – original draft. Tingxin He: Data curation, Formal analysis, Writing – original draft. Ping Ren: Investigation, Writing – review & editing. Lili Jin: Data curation, Software, Writing – original draft. Qiong Yang: Project administration. Chuanyong Xu: Data curation, Visualization.
Rongzhen Wen: Investigation. Jierong Chen: Investigation. Zhen Wei: Validation, Resources, Investigation. Tom Vogruts: Writing – review & editing. Qi Chen: Supervision, Writing – review & editing.

Declaration of Competing Interest

The authors declare that they have no known competing financial interests or personal relationships that could have appeared to influence the work reported in this paper.

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