Gray Matter Volume of the Lingual Gyrus Mediates the Relationship between Inhibition Function and Divergent Thinking

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Although previous research provides converging evidence for the role of posterior regions of the brain (including temporal, occipital, and parietal regions) involved in inhibition on creative thinking, it remains unclear as to how these regions influence individual differences in creative thinking. Thus, we explored the relationship between posterior regions (i.e., hippocampal, parahippocampal, lingual gyrus, precuneus, and cuneus), inhibition function, and divergent thinking (DT) in 128 healthy college students. The results revealed that lower inhibition was associated with larger gray matter volume (GMV) in the lingual gyrus, which in turn was associated with higher DT. In addition, GMV in the lingual gyrus mediated the association between inhibition and DT. These results provide new evidence for the role of inhibition in creative thinking. Inhibition may affect the amount of information stored in long-term memory, which, in turn influences DT.

Keywords: creativity, divergent thinking, fluency, gray matter volume, inhibition, lingual gyrus, long-term memory, originality

INTRODUCTION

Creativity can be defined as the production of a novel, useful, and surprising solution to a complex problem (Sternberg and Lubart, 1996) and is crucial to several aspects of human life (Simonton, 2000). DT, the essence of creative thinking (Guildford, 1987; Zhu et al., 2013), is a complex process that is regulated by multiple cognitive and neural factors (Ward et al., 1999; Benedek et al., 2014; Boccia et al., 2015b). Alpha synchronization in creative ideation was linked to prefrontal and right parietal and occipital sites. Increases in prefrontal alpha activity are usually observed in cognitive tasks involving high top-down control (e.g., inhibition function). In contrast, alpha synchronization at posterior parietal and occipital sites appears to be more specific to cognitive processes involved in creativity (Fink and Benedek, 2014). Further, Dietrich (2004) described that task-relevant memory stored in the TOP areas is pulled by the prefrontal cortex which acts as a search engine and then temporarily represent it in working memory. The prefrontal cortex’s cognitive flexibility allows for retrieved information to be combined. The functions of posterior regions, including TOP areas, differ from those of the prefrontal cortex (Dietrich, 2004). The TOP

Abbreviations: DT, divergent thinking; GMV, gray matter volume; ROIs, regions of interest; TOP, temporal, occipital, and parietal; VBM, voxel based morphometry; MRI, magnetic resonance imaging.
nerve cells are applied mainly to sensory feelings and long-term memory. However, the mechanism of creativity-related cognition remains unclear. Therefore, we adopted an inhibition-TOP-DT approach to investigate the links between inhibition, posterior brain regions, and DT.

Recent investigations have consistently linked inhibition function to DT. For instance, creative people with fluency of ideas and associations (Mednick et al., 1964; Benedek et al., 2012b), are characterized by a lack of inhibition function to filter task-irrelevant information (Eysenck, 1995; Martindale, 1999), and appear to show over-inclusive thinking (Carson et al., 2003). Moreover, the frequency and originality of ideas is facilitated when resources for inhibition are lacking (Radel et al., 2015). Studies in the field of psychology and psychiatry show that several mental states associated with impaired executive functioning can lead to creative performance (Dietrich, 2004; Healey and Rucklidge, 2006; White and Shah, 2006; Keri, 2009; Morgan et al., 2010; Wieth and Zacks, 2011; Chrysikou et al., 2013; Radel et al., 2015). However, contrary positions on the relationship between the creativity and inhibition exist (Golden, 1975; Groborz and Necka, 2003; Benedek et al., 2014; Edl et al., 2014). For instance, a positive correlation between inhibition – measured by random motor generation tasks – and ideational fluency was observed (Benedek et al., 2012a; Zabelina et al., 2012). Obviously, the contradiction exists across abovementioned studies. Therefore, some researchers in the creative field speculated that creative cognition may involve a two-stage process including idea generation and idea evaluation (Finke et al., 1992). These researchers commonly think idea generation as a bottom-up process which is associated with diffuse attention. By contrast, idea evaluation is regarded as a top-down process with focused attention and cognitive control (Jung et al., 2013). Furthermore, it has been suggested by a model that idea generation and idea evaluation are separately depend on the default and the control networks (Jung et al., 2013). A series of studies suggest that creative thought involves cooperation between idea generation and idea evaluation or cooperation between the default and the control networks (Dietrich and Kanso, 2010; Ellamil et al., 2012; Benedek et al., 2013; Beaty et al., 2014b, 2016). For example, it is indicated that both effects of semantic distance (related with ideas generation) and executive function (related with ideas evaluation) variables predict the DT in the same structural equation model (Beaty et al., 2014b).

On the one hand, functional imaging studies have repeatedly demonstrated inhibition control cortex plays a crucial role in the creative thinking (Fink et al., 2009; Abraham et al., 2012a,b; Kleibuker et al., 2013a,b; Beaty et al., 2014a; Pinho et al., 2014; Zhang et al., 2014; Fink et al., 2015; Mayeless et al., 2015). Importantly, inferior frontal gyrus (IFG) is classically associated with inhibitory control (Aron et al., 2004, 2014; Houdé et al., 2010). On the other hand, structural and neuroscience evidence indicated the role of TOP in promoting DT. TOP regions are the primary sensory cortices of all sense modalities. Sensory information initially decoded in the primary cortex is further assembled and assimilated the TOP regions responsible for association. It is widely accepted that TOP regions are the sites of long-term memory storage (Gilbert, 2001; Dietrich, 2004). In addition, hippocampal and parahippocampal regions are involved in verbal DT (Fink et al., 2009). Furthermore, the middle occipital gyrus and parietal lobes which are drawn into mental imagery, such as (Sack et al., 2005; Belardinelli et al., 2009; Boccia et al., 2015a), may be involved during creative thinking. Jauk et al. (2015) discovered that temporo-lateral gyrus and cuneus was negatively correlated with DT. Zhang et al. (2014) also discovered that the left lingual gyrus (BA 18) underlying relevant visual imagery is involved in the process of semantic relatedness. They inferred that the lingual gyrus may take part in visual imagery in this process. According to Fink et al. (2014), enhanced gray matter density in the right precuneus and the right cuneus may be indicative of vivid imaginative abilities in more creative individuals. The volumetric approach is preferable in studying inhibition function, as structural changes may reveal more stable individual differences relative to task-related activations identified in functional neuroimaging research (DeYoung et al., 2010).

Though little empirical evidence suggests a direct link between inhibition function and TOP regions, it is noteworthy that the inhibition function – a component of executive function – can filter task-irrelevant information that can therefore disrupt goal completion (Dempster, 1992; Miyake et al., 2000; Radel et al., 2015). In other words, a lack of inhibition function result in a broader range of information penetrating working memory space (Radel et al., 2015). A broader range of information in working memory space contributes to more information stored in long-term memory (Norman, 2013). More importantly, Dietrich (2004) proposed that the frontal lobe, which is responsible for executive functions, allows cognitive freedom and flexibility, thereby freeing us from the confines of direct environmental information or memory stored in TOP regions; this is at the core of creativity theories such as Guilford’s (1967) concept of DT.

Collectively, the literature indicates a three-way relationship between inhibition function, TOP, and DT, although the direction of influences is unclear. Therefore, the mediation hypothesis in the present study states that individuals with poorer inhibition function would perform better at DT tasks. Poorer inhibition function may increase the amount information stored in brain regions associated with long-term memory, and the demand for large information storage contributes to increased volumes in these brain regions. In addition, evidence suggests that brain structures represent perceptual- and cognitive-level variables to some degree (Kana and Rees, 2011; Montag et al., 2013; Gilai-Dotan et al., 2014). More information helps form new combinations. Based on previous evidence linking inhibition function and the three posterior cortices (TOP) to DT (Boccia et al., 2015b; Liu et al., 2015; Lustenberger et al., 2015) and inhibition function to DT (Radel et al., 2015), the present study investigated the associations among GMV of classical regions located in TOP, inhibition function, and DT. In addition, we examined the potential mediating role of GMV in these ROIs in the relationship between inhibition function and DT. Furthermore, IFG-a classical brain region associated with inhibition function is related to originality and appropriateness.
of the ideas proposed by the participants (Aron et al., 2004, 2014; Houdé et al., 2010). We expected that (a) the volume of ROIs located in TOP would be positively associated with DT and negatively associated with inhibition function. Furthermore, inhibition function would be negatively associated with DT; (b) the volume of ROIs located in TOP would mediate the relationship between the inhibition and DT; (c) the volume of IFG is related with inhibition function and DT. The multidimensional approach employed in the present study has the potential to advance our understanding of the cognitive and neural factors influencing DT.

MATERIALS AND METHODS

Subjects

Participants comprised 128 right-handed, healthy subjects (49 males; age: Mean = 19.69 years; range = 17–22) who took part in our project to investigate the relationships among creativity, mental health, and brain imaging (Li et al., 2014; Wei et al., 2014). All of them were undergraduates from the Southwest University, China. Based on a self-report questionnaire administered before the scan, we found that none experienced neurological or psychiatric illness, and substance abuse. All participants signed the written informed consent. The study was granted by the Institutional Review Board of Southwest University Imaging Center for Brain Research.

Assessing Inhibition

The Stroop color–word-interference task was used to assess the inhibition of prepotent responses (Stroop, 1935). It is generally thought to be a classical inhibition task (Miyake et al., 2000). The task involves presenting four words (“red”, “green”, “blue,” or “yellow”) which denote any a color name in the middle of a black computer screen. All words are segmented into congruent trials or in congruent trials. Congruent trials signify that the color of each word was consistent with its meaning. On the contrary, incongruent trials imply that the color of each word was inconsistent with its meaning. In this study, the task included one practice block, and four task blocks, each comprising 24 trials: 12 incongruent, and 12 congruent trials. Incongruent and congruent trials are presented in a random order within each block. Participants were requested to identify the color of word as quickly as possible within 4 s by pressing one of four appointed keys (D, F, J, K). After an incorrect response feedback will be presented to participants to ensure high accuracy. In the presence of incongruent trials participants require suppressing the dominant process of word naming. Mean response time difference between incongruent trials and congruent trials in each block is taken as the interference score. However, the interference score in the Stroop task is reverse to inhibitory control. Thus, the negative interference score was used to the next step analysis.

Assessing Creativity

The Torrance (1974) Tests of Creative Thinking – based on DT – is the most commonly used test (Cramond et al., 2005; Krumm et al., 2014a,b). We used three modified product-improvement tasks, which are a kind of verbal DT tasks from Torrance’s tests. This task requires participants to improve a product, particularly a toy, such that they would enjoy the activity. It differs considerably from extremely popular, Alternate uses test (Christensen et al., 1960). Differing from traditional verbal evaluations of the Torrance Tests of Creative Thinking, we only focus on fluency and originality of DT: they reflect quantity of ideas in a limited time and quality, respectively (Jauk et al., 2014).

Participants were given 3 min per task to write all the novel and appropriate responses that they could think of on a personal computer. A consensual assessment technique was used to assess the ideas (Amabile et al., 1990). All responses were assessed for originality and fluency by four experienced professionals using a similar 10-point scale. Originality means the degree of originality of the responses (i.e., the quality of ideas) and fluency reflects the number of ideas produced by a participant (i.e., quantity of ideas). For each task, originality and fluency scores reflected the average originality and fluency rating of the three ideas assessed by the raters. In the next step analysis, we analysis the two scores as two dependent variables.

Assessing Intelligence

To control the impact of general intelligence on brain structure (Colom et al., 2006; Jung and Haier, 2007; Takeuchi et al., 2011a), we measured participants’ general intelligence with the Combined Raven’s Test-Rural in China (revised by the Psychology Department of East China Normal University in 1994; Takeuchi et al., 2010, 2011b). This scale is adopted to test intelligence in a wide range for good reliability and validity (Wang, 2007). The Revised Combined Raven’s Test-Rural in China contains the Raven’s colored progressive matrix (A, B, and AB sets) and Raven’s standard progressive matrix (C, D, and E sets). This scale includes 72 non-verbal items. In every item, participants are asked to select the best answer from six or eight alternatives to fill a matrix with a missing piece.

Image Data Acquisition and Preprocessing

Magnetic resonance imaging (MRI) data were acquired with a 3T scanner (Siemens Magnetom Trio TIM, Erlangen, Germany). A magnetization-prepared rapid gradient echo (MPRAGE) sequence was used to collect the high-resolution T1-weighted anatomical images (TR = 1900 ms; TE = 2.52 ms; TI = 900 ms; flip angle = 9; 176 slices; slice thickness = 1.0 mm; FOV = 250 mm × 250 mm; matrix size = 256 × 256).

Structural magnetic resonance images were processed with SPM8 software (Welcome Department of Cognitive Neurology, London, UK). Specifically, artifacts or anatomical abnormalities were first screened in SPM8. Subsequently, images were manually set along the anterior-posterior commissure (AC-PC) line to get better registration. Next, the new segmentation tool in SPM8 was employed to segment images into gray
matter, white matter, and cerebrospinal fluid. The segmented images were submitted for registration, normalization, and modulation using SPM8 plus DARTEL (Ashburner, 2007). After that, Jacobian determinants were used to modulate images in order to conserve the absolute amount of gray matter (Good et al., 2002). Finally, the modulated gray matter images were smoothed with a 10 mm full-width at half-maximum (FWHM) Gaussian kernel to improve the signal-to-noise ratio.

Region of Interest Extraction
We extracted six cortical ROIs from the Automated Anatomical Labeling atlas using the WFU PickAtlas Toolbox (Tzourio-Mazoyer et al., 2002). The ROIs (hippocampal, parahippocampal, lingual gyrus, precuneus, cuneus, IFG) were identical to those used in previous studies of structural correlation networks (Fink et al., 2009; Jung et al., 2010; Fink et al., 2014; Zhang et al., 2014; Boccia et al., 2015b; Jauk et al., 2015). These ROIs were resampled to the dimension of the voxel-based morphometry images. Subsequently, the ROI images were used as masks to extract the GMV within each ROI from the modulated, normalized gray-matter images using the REX toolbox (Massachusetts Institute of Technology, Cambridge, MA, USA). In order to eliminate the influence of non-interest covariates, subjects’ age, gender, and total brain volume (including gray matter, white matter, and cerebrospinal fluid) were submitted to a linear regression analysis for each ROI. The residuals of each regression were used for subsequent analysis to replace the raw ROI volume values (He et al., 2007; Bernhardt et al., 2011; Hosseini et al., 2012).

Mediation Analysis
Before performing a mediation analysis, all data were normalized. The non-interest effect of age, gender, and general intelligence was removed from behavioral data (i.e., DT and Stroop-effect scores). We used Stroop-effect scores as the independent variable, DT as the dependent variable, and GMV of ROIs as mediators. To further investigate the effects of inhibition and GMV on facets of DT, we set up several models for originality and fluency, wherein originality and fluency of DT tasks were used as dependent variables.

A simple mediation analysis was employed using SPSS macro with a 0.95 confidence level and 5000-bootstrap sample (Preacher and Hayes, 2004, 2008). In typical mediation analyses, several paths between the variables are estimated. The chief estimated path is the total effect of an independent variable X on a dependent variable Y (path c). This path comprises the direct effect of X on Y after controlling for mediator M (path c’) and the indirect effect of X on Y through M (i.e., path ab: product of path a → M and M → Y: Figure 1). After accounting for M, the mediation analysis was determined by assessing whether there is a significant distinction between the total (path c) and direct effects (path c’). When the zero is not encompassed within the confidence interval (CI), a significant difference between the indirect effect (path a b) and zero exists, which indicates that regional structural volume accounts for a statistically significant portion of the relationship between inhibition functions and DT.

RESULTS
Increased ROIs Volume Linked to Decreased Inhibition Function and Increased DT (Originality and Fluency) Performance
All behavioral data were analyzed using SPSS 18.0 (SPSS Inc., Chicago, IL, USA). Partly confirming our first prediction, only lingual gyrus volume was positively associated with originality (r = 0.276, p < 0.01, corrected for multiple comparisons, shown in Table 1) and fluency (r = 0.269, p < 0.01, corrected for multiple comparisons, shown in Table 1), and negatively associated with inhibition function (r = −0.247, p < 0.05, corrected for multiple comparisons, shown in Table 1).

Descriptive statistics are presented in Table 2 and correlations between the behavioral variable and ROIs volume are presented in Table 1. Since only the relationship between behavioral data and lingual gyrus was significant (Figure 1), only lingual gyrus data were considered for further analysis (Figure 2).

Mediation Results
Inhibition was selected as the independent variable after controlling for intelligence, gender, and age. The right and left lingual gyrus were used separately as mediators to investigate a concrete neural basis for originality and fluency of DT. It was found that (1) the GMV of the right lingual gyrus accounts for a significant portion of the relationship between inhibition functions and originality (path a = −0.231, p = 0.005; path b = 0.3, p = 0.002; path a × b = −0.069, bootstrapped 95% CI = −0.154, −0.02; Figure 2); (2) GMV of the right lingual gyrus accounts for a significant portion of the relationship between inhibition functions and fluency (path a = −0.231, p = 0.005; b = 0.289, p = 0.003; a × b = −0.067, bootstrapped 95% CI = −0.159, −0.02; Figure 3); (3) GMV of the left lingual gyrus accounts for a significant portion of the relationship
### TABLE 1 | Correlations between behavioral and volumetric measures.

| Region of interest | Inhibition | Parahippocampal | Cuneus | Precuneus | Hippocampal | Lingual | Inferior frontal gyrus (IFG) |
|--------------------|------------|-----------------|--------|-----------|-------------|---------|-----------------------------|
| Inhibition          | 0.009      | −0.182          | −0.042 | 0.072     | −0.247∗     | −0.031  |                              |
| Originality         | −0.022     | 0.101           | 0.023  | 0.050     | 0.276**     | 0.001   |                              |
| Fluency             | −0.028     | 0.072           | −0.016 | −0.014    | 0.269**     | −0.006  |                              |

∗p < 0.05 (two-tailed), **p < 0.001 (two-tailed), corrected for multiple comparisons. DT, divergent thinking. The values in the blackest represent the P value.

### TABLE 2 | Descriptive statistics of subject demographics and psychological measures (N = 128; males = 49, females = 79).

| Measure      | M    | SD   | Range |
|--------------|------|------|-------|
| Age          | 19.7 | 0.97 | 17–22 |
| CRT          | 65.63| 3.76 | 49–72 |
| DT           | 77.06| 28.76| 24–176|
| Stroop       | 0.13 | 0.06 | 0.02–0.32|

CRT, Combined Raven’s Test; DT, divergent thinking.

**FIGURE 2 | Increased lingual gyrus volume is associated with better DT performance and poorer inhibition.** Scatterplots show a significant positive correlation between the GMV of the lingual gyrus and DT (A); and a significant negative correlation between the GMV of the lingual gyrus and inhibition (B).

**FIGURE 3 | Mediation analysis for right lingual gyrus volume and fluency.** Path c’ is the direct effect of DT on inhibition functions after controlling for the GMV of lingual gyrus; the product of the paths a and b (ab) is the indirect effect of inhibition functions through the GMV of right lingual gyrus on fluency. The GMV of lingual gyrus mediated the association between inhibition functions and fluency (bootstrap estimate 95% confidence interval [−0.159, −0.02]). ∗∗p < 0.001.

**FIGURE 4**

**DISCUSSION**

The present study used mediation analysis to classify the associations between GMV of classical regions located in TOP areas, inhibition function, and DT. The results showed two main findings: poorer inhibition function was associated with increased lingual gyrus volume, which in turn was associated with better DT performance; and the lingual gyrus GMV mediated the relationship between inhibition function and DT.

The association between lingual gyrus GMV and DT advances previous structural and functional neuroimaging evidence that suggests that greater lingual gyrus GMV volume is associated with better ideational fluency (Jauk et al., 2015) and activation of the left lingual gyrus is associated with better performance in processing of semantic relatedness (Zhang et al., 2014). Specifically, research consistently shows that the lingual gyrus is associated with visual memory (Bogousslavsky et al., 1987), vivid visual imagery (Belardinelli et al., 2009), and motion imagery (Malouin et al., 2003). The creative process necessitates the generation of new visual imagery (Anderson and Helstrup, 1993; LeBoutillier and Marks, 2003). Moreover, visual imagery is an important strategy for DT (Gilhooly et al., 2007; Fink et al., 2014). As suggested by Guilford (1967), the prefrontal cortex draws task-relevant information from long-term memory stored in the TOP areas and temporarily represents it in the working memory space. This allows the prefrontal cortex cognitive flexibility when combining the retrieved information to form new combinations. Lingual gyrus GMV may be one important neural marker of long-term memory (specifically visual memory) storage.

The current findings demonstrate a negative association between lingual gyrus volume and inhibition function. Although there is little experimental evidence for the direct relationship...
between inhibition function and lingual gyrus, Dietrich (2004) proposed that inhibition function would modulate the associative activity among knowledge nodes in TOP regions. Furthermore, some researchers (Radel et al., 2015) suggest that the inhibition function may automatically modulate the spreading of concept activation. Impaired inhibition function would activate a greater number of concepts in working memory, with several weak links to the initial concept. Further, the working memory buffer – with its ability to sustain online processing in real time – may be a prerequisite access to long-term memory (Dietrich, 2004). Importantly, it is widely accepted that TOP regions is the site of long-term memory storage (Gilbert, 2001; Dietrich, 2004). Therefore, the negative association between inhibition function and lingual gyrus volume found in our study may indicate that poorer inhibition function is associated with better visual memory storage, and thus, greater GMV in TOP regions. However, according to Mechelli et al. (2000), they used positron emission tomography to investigate the effect word length and visual contrast in the visual cortex during reading. Results indicate that the effect of lingual gyrus enhanced by low-contrast words reflects global shape processing when reading. In other words, the GMV of lingual gyrus may be related to the global shape processing ability in incompatible trials in Stroop task instead of the amount of memory storage. But task-related brain activity is different from VBM. Task-related brain activity is the correlation between task phase and brain activity, which might be a kind of momentary effect, while VBM provide structural information, which may be a kind of accumulative effect. So it is not obvious that the GMV of lingual gyrus is associated with inhibition interference in the Stroop task.

Inconsistent with previous studies showing that inhibition function is positively or negatively associated with creative cognition, we find that inhibition function is not directly associated with DT. Creative thought has been variably associated with focused attention and effective inhibition control (Beaty et al., 2014b, 2016), disinhibition and defocused attention (Radel et al., 2015), or a flexible adaption of inhibition control (Cassotti et al., 2016a). While some researchers suggested that inhibitory control hindered the creative potential (Radel et al., 2015), however, other studies reported that generating creative solutions to a problem require the inhibition of past inappropriate ideas that induce fixation phenomena (Cassotti et al., 2016a,b). In addition, many famous scientists indicated when original thoughts enter into their brain they are not deliberately aimed at solving the problem they were trying to solve. The possible interpretation is that creative cognition involves a flexible adaptation of inhibition function, as a line of studies that creative cognition may involve a two-stage process that include idea generation and idea evaluation (Finke et al., 1992). Researchers often think idea generation as a bottom-up process which is associated with diffuse attention. By contrast, idea evaluation is regarded as a top-down process with focused attention and cognitive control (Jung et al., 2013). In the idea generation stage, diffuse attention and disinhibition may contribute to ideas generation, as concepts used for generating uncommon ideas are dispersedly stored in semantic memory (Tyler and Moss, 2001; Thompson-Schill et al., 2006). Therefore, in this stage, it may show a negative correlation pattern between inhibition and creativity. While in the idea evaluation stage, inhibition control plays a key role to screen creative ideas/associations by suppressing prepotent and common ideas. Thus, a positive correlation pattern between inhibition and creativity would occur. In the present study, the results indicated that inhibition is negatively associated with DT in an indirect way. Inhibition function is negatively associated with the GMV of lingual gyrus—a brain region supporting for visual memory (Bogousslavsky et al., 1987), vivid visual imagery (Belardinelli et al., 2009), and motion imagery (Malouin et al., 2003). And then, the GMV of lingual gyrus is positively associated with DT. This may conform to the suggestion that diffuse attention and disinhibition contribute to ideas generation by using concepts dispersedly stored in semantic memory (Tyler and Moss, 2001; Dietrich, 2004).
Improvement, Unusual Uses of Cardboard Boxes, and Just
verbal activity on The Torrance Tests of Creative Thinking
possible relationships between the performance in each
improvement performance. Future research should examine
the relationship between inhibition function and product-
this process. Therefore, lingual gyrus volume may mediate
which a product could be changed to be more interesting
in the present study required participants to list ways in
et al., 2009). Further, the product improvement tasks used
be because the lingual gyrus is associated with visual memory
parahippocampal, precuneus, and cuneus regions. This may
function and DT, compared to the volume of hippocampal,
stimulated by this DT task. These results are consistent
in prefrontal regions (Martindale and Hasenfus, 1978; Mölle
impaired executive functioning, including ADHD (Healey and
White and Shah, 2006), fronto-executive disfunctioning/schizophrenia (Keri, 2009), lateral frontal lesions
(Chrysikou et al., 2013), and cannabis intoxication (Morgan
et al., 2010) can lead to creative performance. Obviously, the
contradiction exists across abovementioned studies. Therefore,
some researchers in the creative field speculated that creative
cognition may involve a two-stage process including idea
generation and idea evaluation (Finke et al., 1992). These
researchers commonly think idea generation as a bottom-up
process which is associated with diffuse attention. By contrast,
idea evaluation is regarded as an top-down process with focused
attention and cognitive control (Jung et al., 2013). Furthermore,
it has been suggested by a model that idea generation and idea
evaluation are separately depend on the default and the control
networks (Jung et al., 2013). In the present study, we find that
the GMV of IFG is not significantly related with DT, but the
GMV of lingual gyrus is related with DT. IFG is classically
associated with inhibitory control (Aron et al., 2004, 2014;
Houdé et al., 2010). Lingual gyrus is a brain region supporting
visual memory (Bogousslavsky et al., 1987) and vivid visual
imagery (Belardinelli et al., 2009), which is a region important
for idea generation. In other words, idea generation and idea
evaluation (also the default and the control networks) may show
little cooperation in this DT task. These results are consistent
with the EEG studies indicating that successful performance
at ideational fluency tasks do not depend on cortical arousal
in prefrontal regions (Martindale and Hasenfus, 1978; Mölle
et al., 1999). Moreover, individuals with PFC damage solve
insight problems better than the healthy counterparts (Reverberi
et al., 2005). Importantly, a fMRI study found that Uncommon-
Use-task reliably activated regions of occipito-temporal cortex
without regions of lateral prefrontal cortex compared with the
Common-Use-task (Chrysikou and Thompson-Schill, 2011). In
addition, previous studies usually found IFG is activated when
individuals are performing a creativity task. Task-related brain
activity is the correlation between task phase and brain activity,
which might be a kind of momentary effect, while VBM provide
structural information, which may be a kind of accumulative
effect. Although previous studies indicate that the prefrontal
cortex is associated with inhibition function, at the same time,
prefrontal cortex is activated during idea generation process, it is
not obvious that the GMV of IFG is associated with inhibition
function and DT.

However, there are some limitations in the present study.
First, it is not sufficient to attribute the link between inhibition
and DT to memory processes due to a lack of working memory
and long-term memory assessment. In future research, memory
measurements should be involved to provide a substantial
evidence to examine the mechanism underlying the association between inhibition and lingual gyrus. Second, the Stroop task used in this study may not be an appropriate measurement to test inhibition function, as this version of Stroop task lacks a neural condition. Because of this, we are not sure the difference is due to inhibition function or attention control. Modified task is required to test the relationship between inhibition function and DT in future research.

In summary, the present findings – lingual gyrus volume mediated the relationship between inhibition function and DT – are important because they potentially explain the influence of inhibition on DT through the posterior cortex. Limited research has simultaneously examined inhibition, brain structure, and DT behavior to identify their interrelationships and the mechanisms through which inhibition influences DT. The results provide novel evidence for the role of inhibition function in creative thinking by its influence on the amount of information stored in long-term memory. However, longitudinal studies are needed to establish the long-term impact of lingual gyrus volume and the direction of influence in our model.

REFERENCES

Abraham, A., Beudt, S., Ott, D. V., Yves, and von Cramon, D. (2012a). Creative cognition and the brain: dissociations between frontal, parietal-temporal and basal ganglia groups. Brain Res. 1482, 55–70. doi: 10.1016/j.brainres.2012.09.007

Abraham, A., Pieritz, K., Thybusch, K., Rutter, B., Kroger, S., Schweckendiek, J., et al. (2012b). Creativity and the brain: uncovering the neural signature of creative expansion. Neuropsychologia 50, 1906–1917. doi: 10.1016/j.neuropsychologia.2012.04.015

Amabile, T. M., Goldfarb, P., and Brickfield, S. C. (1990). Social influences on creativity: evaluation, coaction, and surveillance. Creat. Res. J. 3, 6–21. doi: 10.1080/10400419009534330

Anderson, R. E., and Helstrup, T. (1993). Visual discovery in mind and on paper. Mem. Cognit. 21, 283–293. doi: 10.3758/BF03208261

Aron, A. R. (2007). The neural basis of inhibition in cognitive control. Neuroscientist 13, 214–228. doi: 10.1177/1073858407299288

Aron, A. R., Robbins, T. W., and Poldrack, R. A. (2004). Inhibition and the right inferior frontal cortex. Trends Cogn. Sci. 8, 170–177. doi: 10.1016/j.tics.2004.02.010

Aron, A. R., Robbins, T. W., and Poldrack, R. A. (2014). Inhibition and the right inferior frontal cortex: one decade on. Trends Cogn. Sci. 18, 177–185. doi: 10.1016/j.tics.2013.12.003

Ashburner, J. (2007). A fast diffeomorphic image registration algorithm. Neuroimage 38, 95–113. doi: 10.1016/j.neuroimage.2007.07.007

Beatty, R. E., Benedek, M., Silvia, P. J., and Schacter, D. L. (2016). Creative cognition and brain network dynamics. Trends Cogn. Sci. 20, 87–95. doi: 10.1016/j.tics.2015.10.004

Beatty, R. E., Benedek, M., Wilkins, R. W., Jauk, E., Fink, A., Silvia, P. J., et al. (2014a). Creativity and the default network: a functional connectivity analysis of the creative brain at rest. Neuropsychologia 64, 92–98. doi: 10.1016/j.neuropsychologia.2014.09.019

Beatty, R. E., Silvia, P. J., Nusbaum, E. C., Jauk, E., and Benedek, M. (2014b). The roles of associative and executive processes in creative cognition. Mem. Cognit. 42, 1186–1197. doi: 10.3758/s13421-014-0428-8

Belardinelli, M. O., Palmiero, M., Sestieri, C., Nardo, D., Di Matteo, R., Londei, A., et al. (2009). An fMRI investigation on image generation in different sensory modalities: the influence of vividness. Acta Psychol. 132, 190–200. doi: 10.1016/j.actpsy.2009.06.009

Benedek, M., Franz, F., Heene, M., and Neubauer, A. C. (2012a). Differential effects of cognitive inhibition and intelligence on creativity. Pers. Individ. Dif. 53, 334–480. doi: 10.1016/j.paid.2012.04.014

AUTHOR CONTRIBUTIONS

LZ, LQ, QC, JQ, and DY programmed the experiments and analyzed the data; LZ and WY drafted the manuscript and LZ, LQ, QC, WY, MX, and XY provided critical revisions; LZ prepared the figures.

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Benedek, M., Jauk, E., Fink, A., Koschutnig, K., Reishofer, G., Ebner, F., et al. (2013). To create or to recall? Neural mechanisms underlying the generation of creative new ideas. Neuroimage 88, 125–133. doi: 10.1016/j.neuroimage.2013.11.021

Benedek, M., Jauk, E., Sommer, M., Arendasy, M., and Neubauer, A. C. (2014). Intelligence, creativity, and cognitive control: the common and differential involvement of executive functions in intelligence and creativity. Intelligence 46, 73–83. doi: 10.1016/j.intell.2014.05.007

Benedek, M., Kön, T., and Neubauer, A. C. (2012b). Associative abilities underlying creativity. Psychol. Aesthet. Creat. Arts 6, 273–281. doi: 10.1037/00207059

Benedek, M., Chen, Z. J., He, Y., Evans, A. C., and Bernasconi, N. (2011). Graph-theoretical analysis reveals disrupted small-world organization of cortical thickness correlation networks in temporal lobe epilepsy. Cereb. Cortex 21, 2147–2157. doi: 10.1093/cercor/bhq291

Boccia, M., Piccardi, L., Palermo, L., Nemmi, F., Sulpizio, V., Galati, G., et al. (2015a). A penny for your thoughts? Patterns of fMRI activity reveal the content and the spatial topography of visual mental images. Hum. Brain Mapp. 36, 945–958. doi: 10.1002/hbm.22678

Boccia, M., Piccardi, L., Palermo, L., Nori, R., and Palmiero, M. (2015b). Where do bright ideas occur in our brain? Meta-analytic evidence from neuroimaging studies of domain-specific creativity. Front. Psychol. 6:1195. doi: 10.3389/fpsyg.2015.01195

Bogousslavsky, J., Miklossy, J., Deruaz, J. P., Assal, G., and Regli, F. (1987). Lingual and fusiform gyri in visual processing: a clinico-pathologic study of superior altitudinal hemianopia. J. Neurol. Neurosurg. Psychiatry 50, 607–614. doi: 10.1136/jnnp.50.5.607

Carson, S. H., Peterson, J. B., and Higgins, D. M. (2003). Decreased latent inhibition is associated with increased creative achievement in high-functioning individuals. J. Pers. Soc. Psychol. 85, 499–506. doi: 10.1037/0022-3514.85.3.499

Cassotti, M., Agogué, M., Camarda, A., Houdé, O., and Borst, G. (2016a). Inhibitory control as a core process of creative problem solving and idea generation from childhood to adulthood. New Dir. Child Adolesc. Dev. 2016, 61–72. doi: 10.1002/cad.20153

Cassotti, M., Camarda, A., Poirel, N., Houdé, O., and Agogué, M. (2016b). Flexibility in creative ideas generation: Opposite impacts of example in children and adults. Think. Skills Creat. 19, 146–152. doi: 10.1016/j.tsc.2015.10.008

Christensen, P. R., Guilford, J. P., Merrifield, P. R., and Wilson, R. C. (1960). Alternate Uses. Beverly Hills, CA: Sheridan Psychological Services.

Chrysikou, E. G., Hamilton, R. H., Coslett, H. B., Datta, A., Bikson, M., and Thompson-Schill, S. L. (2013). Noninvasive transcranial direct current stimulation over the left prefrontal cortex facilitates cognitive flexibility in tool use. Cogn. Neurosci. 4, 81–89. doi: 10.1016/j.paid.2012.04.014

Christensen, P. R., Guilford, J. P., Merrifield, P. R., and Wilson, R. C. (1960). Alternate Uses. Beverly Hills, CA: Sheridan Psychological Services.
Martinodale, C. (1999). "Biological bases of creativity," in *Handbook of Creativity*, ed. R. J. Sternberg (Cambridge: Cambridge University Press), 137–152.

Martinodale, C., and Hasenfus, N. (1978). EEG differences as a function of creativity, stage of the creative process, and effort to be original. *Biol. Psychol.* 6, 157–167. doi: 10.1016/0301-0511(78)90118-2

Mayseless, N., Erao, A., and Shmay-Tsoory, S. G. (2015). Generating original ideas: the neural underpinning of originality. *Neuroimage* 116, 232–239. doi: 10.1016/j.neuroimage.2015.05.030

Mechelli, A., Humphreys, G. W., Mayall, K., Olson, A., and Price, C. J. (2000). Differential effects of word length and visual contrast in the fusiform and lingual gyri during. *Proc. R. Soc. Lond. B Biol. Sci.* 267, 1909–1913. doi: 10.1098/rspb.2000.1229

Mednick, M. T., Mednick, S. A., and Jung, C. C. (1964). Continual association as a function of level of creativity and type of verbal stimulus. *J. Abnorm. Soc. Psychol.* 69, 511–515. doi: 10.1037/h0041086

Miyake, A., Friedman, N. P., Emerson, M. J., Witzki, A. H., Howarter, A., and Wagner, T. D. (2000). The unity and diversity of executive functions and their contributions to complex "frontal lobe" tasks: a latent variable analysis. *Cognit. Psychol.* 41, 49–100. doi: 10.1016/cogpsych.1999.0734

Mölle, M., Marshall, L., Wolf, R., Fehm, H. L., and Born, J. (1999). EEG complexity and performance measures of creative thinking. *Psychophysiology* 36, 95–104. doi: 10.1016/0048-5772(99)99616-9

Montag, C., Reuter, M., Jurkiewicz, M., Markett, S., and Pankepp, J. (2013). Imaging the structure of the human anxious brain: a review of findings from neuroscientific personality psychology. *Rev. Neurosci.* 24, 167–190. doi: 10.1515/revre-2012-0085

Morgan, C. J., Rothwell, E., Atkinson, H., Mason, O., and Curran, H. V. (2013). Preacher, K. J., and Hayes, A. F. (2004). SPSS and SAS procedures for estimating indirect effects in simple mediation models. *Behav. Res. Methods Instrum. Comput.* 36, 717–731. doi: 10.3758/BF03206553

Preacher, K. J., and Hayes, A. F. (2008). SPSS and SAS procedures for estimating indirect effects in multiple mediation models. *Behav. Res. Methods* 40, 879–891. doi: 10.3758/BRM.40.3.879

Radel, R., Davranche, K., Fournier, M., and Dietrich, A. (2015). The role of (dis)inhibition in creativity: decreased inhibition improves idea generation. *Cognition* 134, 110–120. doi: 10.1016/j.cognition.2014.09.001

Reverberi, C., Toraldo, A., D’Agostini, S., and Skrap, M. (2005). Better without inhibition in verbal creative thinking to cerebral structures: an optimal voxel-based morphometry study. *Neuroimage* 26, 213–218. doi: 10.1016/j.neuroimage.2008.09.002

Norman, D. A. (2013). *Models of Human Memory*. Amsterdam: Elsevier.

Simonton, D. K. (2000). Creativity: cognitive, personal, developmental, and social aspects. *Am. Psychol.* 55, 151–158. doi: 10.1037/0003-066X.55.1.151

Sternberg, R. J., and Lubart, T. I. (1996). Investing in creativity. *Am. Psychol.* 51, 677–688. doi: 10.1037/0003-066X.51.7.677

Stroop, J. R. (1935). Studies of interference in serial verbal reactions. *J. Exp. Psychol.* 18, 643–648. doi: 10.1002/160054651

Takeuchi, H., Taki, Y., Hashizume, H., Sassa, Y., Nagase, T., Nouchi, R., et al. (2011a). Failing to deactivate: the association between brain activity during a working memory task and creativity. *Neuroimage* 55, 681–687. doi: 10.1016/j.neuroimage.2010.11.052

Takeuchi, H., Taki, Y., Sassa, Y., Hashizume, H., Sekiguchi, A., Fukushima, A., et al. (2011b). Working memory training using mental calculation impacts regional gray matter of the frontal and parietal regions. *PLoS ONE* 6:e23175. doi: 10.1371/journal.pone.0023175

Takeuchi, H., Taki, Y., Sassa, Y., Hashizume, H., Sekiguchi, A., Fukushima, A., et al. (2010). Regional gray matter volume of dopaminergic system associate with creativity: evidence from voxel-based morphometry. *Neuroimage* 51, 578–585. doi: 10.1016/j.neuroimage.2010.02.078

Thompson-Schill, S. L., Kan, I. P., and Oliver, R. T. (2006). "Functional neuroimaging of semantic memory," in *Handbook of Functional Neuroimaging of Cognition*, Vol. 2, eds R. Cabeza and A. Kingston (Cambridge, MA: MIT Press), 149–190.

Tyler, L. K., and Moss, H. E. (2001). Towards a distributed account of conceptual knowledge. *Trends Cogn. Sci.* 5, 244–252. doi: 10.1016/S1364-6613(00)01651-X

Zhang, H., Liu, J., and Zhang, Q. (2014). Neural representations for the generation of inventive conceptions inspired by adaptive feature optimization of biological species. *Cortex* 51, 1909–1913. doi: 10.1016/j.cortex.2013.09.004

White, H. A., and Shah, P. (2006). Uninhibited imaginations: creativity in adults with attention-deficit/hyperactivity disorder. *Pers. Individ. Differ.* 40, 1121–1131. doi: 10.1016/j.paid.2005.11.007

Werbel, M. A., and Thackre, R. (2014). Time of day effects on problem solving: When the non-optimal is optimal. *Think. Reason.* 17, 387–401. doi: 10.1080/13546783.2011.625663

Zabelina, D. L., Robinson, M. D., Council, J. R., and Breslin, K. (2012). Patterning and nonpatterning in creative cognition: insights from performance in a random number generation task. *Psychol. Aesthet. Creat. Arts* 6, 137–145. doi: 10.1037/a0025452

Zhang, H., Liu, J., and Zhang, Q. (2014). Neural representations for the generation of inventive conceptions inspired by adaptive feature optimization of biological species. *Cortex* 50, 162–173. doi: 10.1016/j.cortex.2013.01.015

Zhu, F., Zhang, Q., and Qiu, J. (2013). Relating inter-individual differences in verbal creative thinking to cerebral structures: an optimal voxel-based morphometry study. *PLoS ONE* 8:e79272. doi: 10.1371/journal.pone.0079272

Conflict of Interest Statement: The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

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