



Cognitive benefits of exercise interventions: an fMRI activation likelihood estimation meta-analysis

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Abstract

Despite a growing number of functional MRI studies reporting exercise-induced changes during cognitive processing, a systematic determination of the underlying neurobiological pathways is currently lacking. To this end, our neuroimaging meta-analysis included 20 studies and investigated the influence of physical exercise on cognition-related functional brain activation. The overall meta-analysis encompassing all experiments revealed physical exercise-induced changes in the left parietal lobe during cognitive processing. Subgroup analysis further revealed that in the younger-age group (< 35 years old) physical exercise induced more widespread changes in the right hemisphere, whereas in the older-age group (≥ 35 years old) exercise-induced changes were restricted to the left parietal lobe. Subgroup analysis for intervention duration showed that shorter exercise interventions induced changes in regions connected with frontoparietal and default mode networks, whereas regions exhibiting effects of longer interventions connected with frontoparietal and dorsal attention networks. Our findings suggest that physical exercise interventions lead to changes in functional activation patterns primarily located in precuneus and associated with frontoparietal, dorsal attention and default mode networks.

Keywords Exercise · Cognition · Brain health · Network

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Introduction

Cognition covers a wide range of mental abilities that allow us to perceive, process and store information (Bostrom 2009; Herold et al. 2018) although no uniform definition has yet been reached (Allen 2017; Bayne et al. 2019). Consequently, cognitive performance is crucial to successfully interact with our environment and thus determine several aspects of successful everyday life functioning (MacNeill and Lichtenberg 1997; Salthouse 2011). Notably, there is increasing evidence in the literature showing that specific cognitive abilities such as processing speed and episodic memory gradually decline with aging (Harada et al. 2013; Hedden and Gabrieli 2004; Park et al. 2002; Salthouse 2011), and poor cognitive performance leads to impairments in several aspects of everyday life, including walking, financial management, and driving (Cohen et al. 2016; Montero-Odasso et al. 2012; Morris et al. 2016). Moreover, poorer cognitive performance has been shown to predict neurological diseases such as dementia (Boraxbekk et al. 2015) and has been associated with a higher mortality risk (Shipley et al. 2006). It is thus of general importance to preserve cognitive functions across the lifespan and particularly in aging populations.

Current approaches to preserve cognitive functions in the aging population focus on lifestyle changes and emphasize the role of regular physical activity and exercise (Bherer et al. 2013; Hillman et al. 2008; Kivipelto et al. 2018; Liu-Ambrose et al. 2018). An increase in physical activity level is usually achieved through regular physical exercise (also referred to as physical training). Indeed, accumulating evidence indicates that both acute bout of physical exercise (Chang et al. 2012; Ludyga et al. 2016; Pontifex et al. 2019) and chronic physical exercise intervention (hereafter referred to as physical training) (Falck et al. 2019; Ludyga et al. 2016) can influence cognitive performance positively. However, the underlying neurobiological processes which lead to an increase in cognitive performance after physical interventions are not fully understood.

According to Stillman (Stillman et al. 2016), physical exercise interventions induce changes on different levels of analysis which, in turn, promote the improvement of cognitive performance. In particular, physical exercise and physical training lead to changes on (i) molecular and cellular levels (e.g., brain-derived neurotrophic factors); (ii) structural and/or functional levels (e.g., hippocampus volume and hippocampal activity), and (iii) socioemotional level (e.g., sleep quality, well-being, self-efficacy) (Stillman et al. 2016). Currently, there are systematic reviews and meta-analysis available which summarize the beneficial effects of acute physical exercise (a single bout of

exercise) on changes: (i) molecular and cellular level (i.e., brain-derived neurotrophic factor) (Coelho et al. 2013; de Assis and de Almondes 2017; Dinoff et al. 2017; Knaepen et al. 2010); (ii) structural level (Chen et al. 2020; Zheng et al. 2019) and (iii) socioemotional level (i.e., sleep) (Kelley and Kelley 2017; Kovacevic et al. 2018). In contrast, the effects of physical exercise on functional brain changes that accompany acute physical exercise and physical training are currently less well understood. In this context, previous qualitative reviews have summarized the effects of acute physical exercises (Herold et al. 2020) and physical training (Herold et al. 2019; Voelcker-Rehage 2013) on functional brain changes, but did not perform a systematic quantitative meta-analyses. Moreover, given that some theories of cognitive aging emphasize the importance of compensatory brain activation patterns in distinct functional neural networks (Park and Reuter-Lorenz 2009; Reuter-Lorenz and Park 2014), a deeper understanding of physical exercise-induced functional brain activation changes can help us to better tailor physical exercise interventions to individuals. Hence, this meta-analysis addresses this gap in the literature and investigates the influence of physical exercise interventions on cognition-related changes of functional brain activation.

Methods

This study followed the recommendations outlined in the Preferred Reporting Items for Systematic Review and Meta-Analysis (PRISMA) guidelines and has been registered on OSF Registries (Registration <https://doi.org/10.17605/OSF.IO/674HF>).

Data sources

The literature search was conducted on April 19th, 2020, through four electronic databases (Pubmed, Web of Science, PsycINFO, and Embase). Search terms related to the object of this meta-analysis (for details see the supplementary material 1) were used to identify eligible studies based on a search in title and abstract. Furthermore, reference lists of included articles were manually searched for relevant articles that were captured through the database searches.

Inclusion criteria and study selection

The screening for relevant studies was conducted adhering to the PICOS-principles which stands for participants (P), intervention (I), comparisons (C), outcomes (O), and study design (S) (Harris et al. 2014; Moher et al. 2009). We included peer-review journal article published in English when they met the following inclusion criteria: (P) no

restrictions were applied and we included all age groups regardless of pathologies; (I) only studies performing physical exercise/physical training were considered as eligible; (C) only studies with a pre/post-intervention experimental designs and at least one group assigned to physical exercise/physical training intervention were included; (O) the relevant studies needed to assess cognition-related brain activation patterns via fMRI, PET or SPECT (task-based imaging studies) and needed to report retrievable data in standard Talairach or Montreal Neurologic Institute (MNI) coordinates; (S) we included interventional studies, but excluded cross-sectional studies. Based on the above-mentioned inclusion criteria, two independent researchers (QY and LZ) first screened article titles and abstracts to identify eligible articles. Afterwards and as recommended, a more detailed screening using the full-text of the article was conducted to ensure that all inclusion criteria were met.

Data extraction

The extraction of the relevant data was performed by two independent reviewers (QY and LZ) and the following information were extracted: (i) name of the lead author; (ii) imaging modality; (iii) population characteristics (e.g., health status and age); (iv) intervention characteristics (e.g., the number of participants, type of physical exercise, exercise duration, exercise intensity, training frequency, training duration, and control condition); (v) cognitive task paradigms employed to assess the effects of the intervention, and (vi) functional brain activation results (e.g., the number of foci). Risk of bias was independently assessed (by QY and LY) using the PEDro scale including 11 items (de Morton 2009).

Activation likelihood estimation (ALE)

ALE is a coordinate-based meta-analytical method that has been used to determine the consistent locations of brain activation in included studies that use similar experimental conditions. In ALE, activation foci are regarded as probability distributions centered on the reported coordinates. ALE maps are constructed after calculating each space voxel's likelihood to be activated. The null-distributions acquired from independent studies' ALE values is usually used to test the reliability of ALE map and this process have similarities with permutation test across experiments. The contribution of each study is weighted by its sample size, and each study is considered as a random effect. The observed values in the ALE distribution are compared to the null distribution in order to assign probability estimates to the observed experimental data. For ALE meta-analysis, Eickhoff et al. (2016) recommended that 17 or more experiments are needed to control excessive

contribution of a single experiment for the cluster-level thresholding; and 14 experiments are necessary for voxel-level thresholding (Eickhoff et al. 2016). Usually, each study will include one or more experiments; each experiment only includes the foci in the same condition (i.e., activation status, intervention type).

In this study, we performed ALE analyses with GingerALE v3.0.2 (<http://www.brainmap.org/ale/>) (Eickhoff et al. 2009) in MNI space, and cluster-based family-wise error (FWE) was used with a threshold of $p < 0.05$ (permuted 1000 times) (Eickhoff et al. 2009). The p -value accounts for the proportion of the random spatial relation between the various experiments under the null distribution. Coordinates reported in Talairach space in the original studies were initially transformed to the MNI space using the Lancaster transform `icbm2tal` software procedure as implemented in the `Convert Foci` tool of GingerALE (Laird et al. 2010). The ALE maps were imported into Mango Version 4.1 (<http://ric.uthscsa.edu/mango/mango.html>) software and overlaid on an anatomical template in MNI space for visualization and comparison.

Anatomical connectivity, functional connectivity and functional characterization of the identified brain regions

In this study, if a meta-analytic identified cluster included more than one peak, we considered the area where the peak was centered at as sub-region (radius = 3 mm). Anatomical and functional connection patterns of each sub-region were further determined by the Brainnetome Atlas and visualized by the Brainnetome Atlas Viewer (V1.0) (Fan et al. 2016; Liu et al. 2013). The functional characterizations of each sub-region are illustrated through probabilistic maps reflecting the behavioral domain and paradigm class according to meta-data labels of the BrainMap database (<http://www.brainmap.org/taxonomy>). Overlapping behavioral domain(s) or paradigm(s) across sub-regions were subsequently selected (Fan et al. 2016; Liu et al. 2013). Brainnetome Atlas offers a fine-grained and cross-validated atlas providing structural information of more than 200 sub-regions. Brainnetome Atlas also maps the brain structure and function to mental processes by reference to the BrainMap database. Thus, Brainnetome Atlas provides an effective way for researchers to explore the complex relationship between anatomy, connectivity and function.

We also used the DPABI, a surface-based fMRI data analysis toolbox, to explore associations between regions/sub-regions identified in both overall and subgroup analyses and the 7 networks proposed by Yeo et al. (Yan et al. 2016; Yeo et al. 2011). DPABI yoked between images were obtained from ALE analysis and DPABI template.

Subgroup analysis

Given that physical exercise-induced changes on the brain functional and behavioral level might be influenced by both individual characteristics (e.g., health status, age and regular level of physical activity) and intervention characteristics (e.g., training duration), the included studies were categorized into the following subgroups (as recommended by previous reviews Piercy et al. 2018; Stillman et al. 2020): (i) health status (healthy vs. patients)—the patient groups included studies conducted in individuals with social anxiety disorder, major depressive disorder, cognitive impairment, fibromyalgia, Parkinson’s disease, and bipolar disorder; (ii) age (“younger-age group: < 35 years old” vs “older-age group: \geq 35 years old”). The cut-off of 35 years was employed to achieve a relatively balanced number of studies for each age-related subgroup analysis and was additionally based on evidence that cognitive function typically peaks around 35 years (Hartshorne and Germine 2015); (iii) training duration (“shorter-term: < 12 weeks” vs “longer-term: \geq 12 weeks”). Based on previous meta-analysis, we used a 12-week training duration as the cut-off point for subgroup analysis (Piercy et al. 2018); (iv) although physical activity is characterized by three important components (intensity, frequency, and duration), it is reported that total amount of physical activity (i.e., minutes of exercise per week) is the most important factor for achieving health benefits (Piercy et al. 2018). Hence, weekly total minutes of exercise were used to categorize studies into two groups, including physically inactive group (PIG) and physically active group (PAG). Specifically, we followed the criteria from the Physical Activity Guidelines for Americans (2nd Edition) to determine the group of each experiment-based study with the age-specific cut-off values: (a) 180 min per week in children and adolescents aged 6 to 17 years (PIG: < 180 min per week; PAG: \geq 180 min per week); (b) 150 min (moderate-intensity aerobic exercise) per week in adults (PIG: < 150 min per week; PAG: \geq 150 min per week). Notably, individuals aged over 65 years who kept on exercising were considered as physically active, as suggested by the aforementioned guideline (Piercy et al. 2018).

Data and code availability statement

This ALE analysis was conducted with GingerALE v3.0.2 (<http://www.brainmap.org/ale/>) and Mango Version 4.1 (<http://ric.uthscsa.edu/mango/mango.html>) software were used for presentation. Anatomical and functional connection patterns of each sub-region were further determined by the Brainnetome Atlas (<http://atlas.brainnetome.org/>) and visualized by the Brainnetome Atlas Viewer (<http://atlas.brainnetome.org/download.html>). DPABI software (<http://rfmri.org/dpabi>) was used to explore associations among

identified regions/sub-regions. Study data used for the meta-analysis are provided in Table 1.

Results

Study selection

The systematic literature search returned a total of 42,302 records and 7870 duplicates were removed (see Fig. 1). The remaining 34,432 records were initially screened by examining the article titles and abstracts, with 34,195 records being excluded due to their failure to meet the pre-determined inclusion criteria (e.g., no original research or case reports, no cognition-related outcomes). Full-text assessment of 237 was further conducted by two independent authors (QY and LZ), which resulted in 20 eligible studies (26 experiments); as shown in Fig. 1, 1217 records were excluded according to our selection criteria (review and conference abstract = 24, no pre-to-post imaging assessment = 10, no cognition-related outcomes = 168, no coordinates of whole-brain analysis = 3, non-cognitive task = 2; non-exercise intervention = 10).

Characteristics of included studies

There are 260 foci of activation within 20 studies including a total 745 participants with mean age of 47.41 (SD = 22.04): 283 patients (mean age = 44.01, SD = 20.28) and 463 healthy people (mean age = 49.49, SD = 22.83). Study characteristics are detailed in Table 1.

Overall analysis: exercise-induced brain activation associated with cognition

Twenty studies that investigated the effects of physical exercise on cognition-associated functional brain activation were included in this meta-analysis (Baeck et al. 2012; Boa Sorte Silva et al. 2020; Chen et al. 2016; Duchesne et al. 2016; Goldin et al. 2012; Gourgouvelis et al. 2017; Hsu et al. 2018; Krafft et al. 2014; Li et al. 2019; Liu-Ambrose et al. 2012; Martinsen et al. 2018; Metcalfe et al. 2016; Nishiguchi et al. 2015; Pensel et al. 2018; Schmitt et al. 2019; Smith et al. 2013; Wagner et al. 2017; Wriessnegger et al. 2014; Wu et al. 2018). Overall, 260 foci from all 26 experiments converged onto a 2200 mm³ cluster centered at (−26.2, −57.9, 45.3) with 4 peaks (Fig. 2a). All 4 peaks were located at the parietal lobe of left cerebrum. More specifically, the cluster encompassed regions in the precuneus (69.7%), inferior parietal lobule (27.3%), and superior parietal lobule (3%) and covered regions the following Brodmann areas 7, 19, 39, and 40.

Examining the direction of the effects in terms of increased activation, 184 foci from 17 experiments were

Table 1 Information of included studies

No.	Study	Mode	Population	Age	Intervention of experimental group	Intensity	Duration	Task/stimuli	Control	Total foci number and experiments (Exp.)
1	Goldin et al. (2012a)	fMRI	Patients with social anxiety disorder	32.88 ± 7.97	Cardiovascular exercise (<i>n</i> = 18)	n.a	2.5 h/week * 8 weeks (weekly at least two individual AE sessions and one group AE session, nearly 2.5 h)	Self-referential encoding task (self-referent)	Mindfulness-based stress reduction (<i>n</i> = 24)	15 Exp.1: 1 + Exp.2: 14-
2	Goldin et al. (2012b)	fMRI	Patients with social anxiety disorder	32.88 ± 7.97	Cardiovascular exercise (<i>n</i> = 19)	n.a	2.5 h/week * 8 weeks (weekly at least two individual AE sessions and one group AE session, nearly 2.5 h)	Cognitive reappraisal task (negative self-belief)	Mindfulness-based stress reduction (<i>n</i> = 23)	6 Exp.3: 6 +
3	Gourgouvelis et al. (2017)	fMRI	Patients with major depressive disorder	37.25 ± 8.00	Cardiovascular exercise and resistance exercise for patients (<i>n</i> = 8)	Cardiovascular exercise: 60–80% of their age-predicted maximum heart rate Resistance exercise: 95% of the 10-repetition maximum	60 min/session * 1 session/week * 8 weeks	Associative memory task (memory)	Cardiovascular exercise and resistance exercise for healthy people (<i>n</i> = 8)	10 Exp.4: 10-
4	Schmitt et al. (2019)	fMRI	Athletes	27.20 ± 4.20	Low- vs high-intensity cardiovascular exercise (<i>n</i> = 21)	Low-intensity: 35% under lactate threshold High-intensity: 20% above lactate threshold	40 min for low-intensity exercise 40 min for high-intensity exercise	fMRI-based face-matching paradigm (fear/happiness)	–	4 Exp.5: 4-
5	Boa Sorte Silva et al. (2020)	fMRI	Older adults	67.50 ± 7.30	Multiple-modality exercise + mind-motor training (<i>n</i> = 63)	65–85% of their predicted maximum heart rate (in multiple modality exercise)	60 min/day * 3 days/week * 24 weeks	Computer-based memory tasks (working & episodic memory)	Multiple-modality exercise + others (<i>n</i> = 64)	38 Exp.6: 38 +
6	Hsu et al. (2018)	fMRI	Older adults with vascular cognitive impairment	72.60 ± 8.90	Aerobic cardiovascular (<i>n</i> = 10)	60% to 70% of heart rate reserve	60 min/session * 3 sessions/week * 24 weeks	Eriksen flanker task (executive function)	Usual care (<i>n</i> = 11)	16 Exp.7: 16 +

Table 1 (continued)

No.	Study	Mode	Population	Age	Intervention of experimental group	Intensity	Duration	Task/stimuli	Control	Total foci number and experiments (Exp.)
7	Li et al. (2019)	fMRI	Female college students	25.50 ± 0.67	Acute cardiovascular exercise (n = 12)	Target heart rate was calculated using Karvonen formula with an intensity of 60–69%	30 min	N-back task (executive function)	Acute aerobic exercise for low-fitness students (n = 12)	5 Exp. 8: 5 +
8	Martinsen et al. (2018)	fMRI	Patients with fibromyalgia	25–64y	Resistance exercise for patients (n = 19)	Progressively increased intensity (~60–80% 1-repetition maximum)	60 min/session * 2 sessions/week * 15 weeks	Stroop colour word test (executive function)	Resistance exercise for healthy people (n = 20)	3 Exp. 9: 3 +
9	Nishiguchi et al. (2015)	fMRI	Older adults	73.00 ± 4.80	Dual-task group exercise (n = 24)	n.a	90 min/session * 1 session/week * 12 weeks	N-back task (working memory)	Usual care (n = 24)	7 Exp. 10: 7-
10	Smith et al. (2013)	fMRI	Patients with mild cognitive impairment	78.70 ± 7.50	Cardiovascular exercise for mild cognitive impairment patients (n = 17)	50–60% of heart rate reserve	30 min/session * 44 sessions, for 12 weeks	Famous name discrimination task (semantic memory)	Cardiovascular exercise for healthy people (n = 18)	26 Exp. 11: 26-
11	Krafft et al. (2014)	fMRI	Overweight children	9.70 ± 0.80	Cardiovascular exercise (n = 24)	n.a	40 min/session * 138 sessions, for 8 months	Antisaccade task Flanker task (cognitive control)	Attention control (n = 19)	10 Exp. 12: 4 + Exp. 13: 6-
12	Duchesne et al. (2016)	fMRI	Patients with Parkinson's disease	59.00 ± 7.11	Cardiovascular exercise for Parkinson's disease patients (n = 19)	Increased from 60 to 80% of maximal aerobic power output	20 min/session (+5 min/week up to 40 min) * 3 sessions/week * 12 weeks	Serial reaction time task (procedural memory)	Aerobic exercise for healthy people (n = 19)	8 Exp. 14: 4 + Exp. 15: 4+
13	Chen et al. (2016)	fMRI	Preadolescent children	10y	Acute cardiovascular exercise (n = 9)	60–69% of the predicted maximal heart	35 min	N-back task (working memory)	–	5 Exp. 16: 5 +
14	Pensel et al. (2018)	fMRI	Middle-aged sedentary males	49.00 ± 5.32	Cardiovascular exercise (n = 23)	n.a	90 min/session * 3 sessions/week * 24 weeks	Flanker paradigm (Executive control processing)	Sedentary lifestyle (n = 14)	32 Exp. 17: 32 +
15	Liu-Ambrose et al. (2012)	fMRI	Community-dwelling senior women	69.3 ± 3.2y	Once-weekly resistance training (n1 = 20) Twice-weekly resistance training (n2 = 15)	2 sets of 6 to 8 repetitions of 7 repetition maximum	1 month (once-weekly/twice weekly)	Modified Eriksen flanker task (selective attention and conflict resolution)	Twice-weekly balance and tone training (n = 17)	12 Exp. 18: 12 +

Table 1 (continued)

No.	Study	Mode	Population	Age	Intervention of experimental group	Intensity	Duration	Task/stimuli	Control	Total foci number and experiments (Exp.)
16	Wagner et al. (2017)	fMRI	Young healthy male	25.0 ± 3.3y	Cardiovascular exercise (n = 17)	~77 ± 9% of VO ₂ max at beginning then individual progressed	60 min/session * 3 sessions/week * 6 weeks	Paired-associated learning task (Memory)	No additional specific training (n = 17)	36 Exp.19: 13 + Exp.20: 23 +
17	Wu et al. (2018)	fMRI	Healthy people	64.9 ± 2.8y	Yang-style Tai Chi Chuan (n = 16)	~65.4 ± 1.1% of individual participant's age-predicted maximal heart rate	60 min/week * 12 weeks	Numerical Stroop paradigm (Executive function: task-switching performance)	Telephone consultation (n = 10)	5 Exp.21: 5 +
18	Wriessneger et al. (2014)	fMRI	Healthy people	28.4 ± 4.3y	Playing soccer and tennis via "Kinect" (n = 23)	n.a	20 min	Imagery task (Motor imagery)	–	8 Exp.22: 8 +
19	Metcalfe et al. (2016)	fMRI	Adolescents with bipolar disorder	16.8 ± 1.4y	Acute cardiovascular exercise (n = 30)	60–80% of the predicted maximal heart rate	27 min	Sustained attention to response task (Attention)	Acute aerobic exercise for healthy people (n = 20)	Exp.23: 3 + Exp.24: 4- Exp.25: 1-
20	Baeck et al. (2012)	fMRI	Healthy people	27.2 ± 7.9	Shooting training (n = 18)	n.a	90 min/session * 3 sessions/week * 20 weeks	Motor imagery paradigm (Motor imagery)	–	6 Exp.26: 6 +

¹In the column of Intensity, *n.a.* means "not applicable"

²In the column of foci, "+ " means the number of positively activated foci, "–" means the number of negatively activated foci. If more than one "+ " or "–" appears in a study, it means that the activated/deactivated foci were divided into different experiments due to the intervention and participant type)

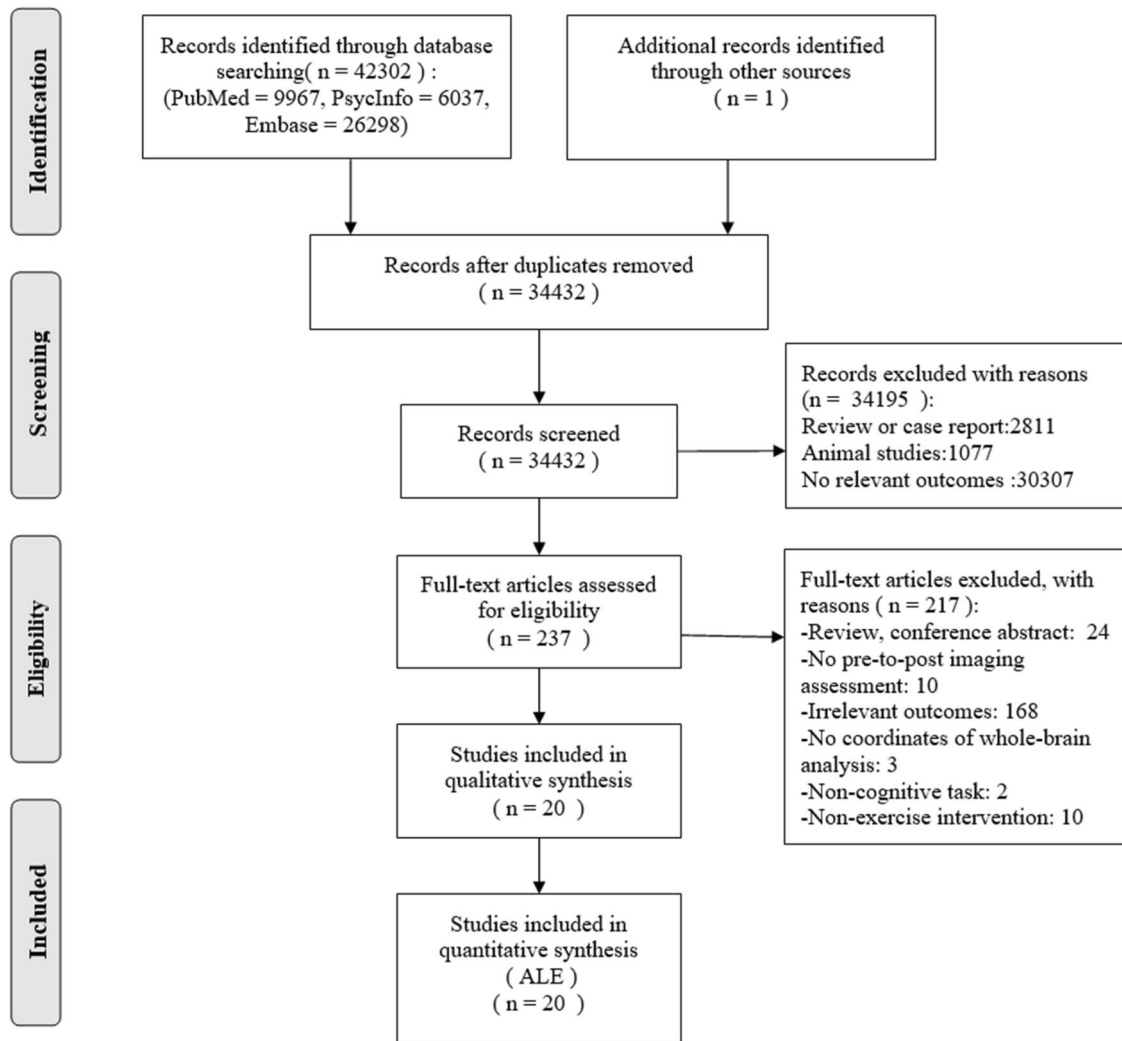


Fig. 1 Flowchart of literature searching, screening and assessment

included and converged on a 1616 mm³ cluster centered at (38.1, -21.9, 4.9) with 3 peaks (Fig. 2b) covering sub-lobar and temporal lobe regions in the right brain hemisphere. The cluster was primarily located in the lentiform nucleus (71.2%) and spread in the adjacent claustrum (17.8%), superior temporal gyrus (6.8%), as well as the insula (4.1%). Accordingly, the activated cluster comprises the Brodmann areas 13, 22 and 41 as well as the putamen. With respect to decreased activation reported in 9 experiments, no region of convergently decreased activation was identified.

Subgroup analysis for health status

For the healthy population (Baeck et al. 2012; Boa Sorte Silva et al. 2020; Chen et al. 2016; Krafft et al. 2014; Li et al. 2019; Liu-Ambrose et al. 2012; Nishiguchi et al. 2015; Pensel et al. 2018; Schmitt et al. 2019; Wagner et al. 2017; Wriessnegger et al. 2014; Wu et al. 2018), 168 foci from 14

experiments were included for the subgroup analysis; and for the patient population (Duchesne et al. 2016; Goldin et al. 2012a,b; Gourgouvelis et al. 2017; Hsu et al. 2018; Martinsen et al. 2018; Metcalfe et al. 2016; Smith et al. 2013), 92 foci from 12 experiments were selected. There was no activated cluster for either analysis.

Subgroup analysis for age

In the younger-age group (Baeck et al. 2012; Chen et al. 2016; Goldin et al. 2012a, b; Krafft et al. 2014; Li et al. 2019; Metcalfe et al. 2016; Schmitt et al. 2019; Wagner et al. 2017; Wriessnegger et al. 2014), 103 activation foci from 15 experiments were included and converged on 2112 mm³ cluster centered at (14.1, -62.2, 30.8) with 3 peaks (Fig. 2c) located at the occipital, parietal and limbic lobe. The identified cluster primarily encompassed the precuneus (78.5%), spreading into the adjacent cuneus (10.3%), posterior

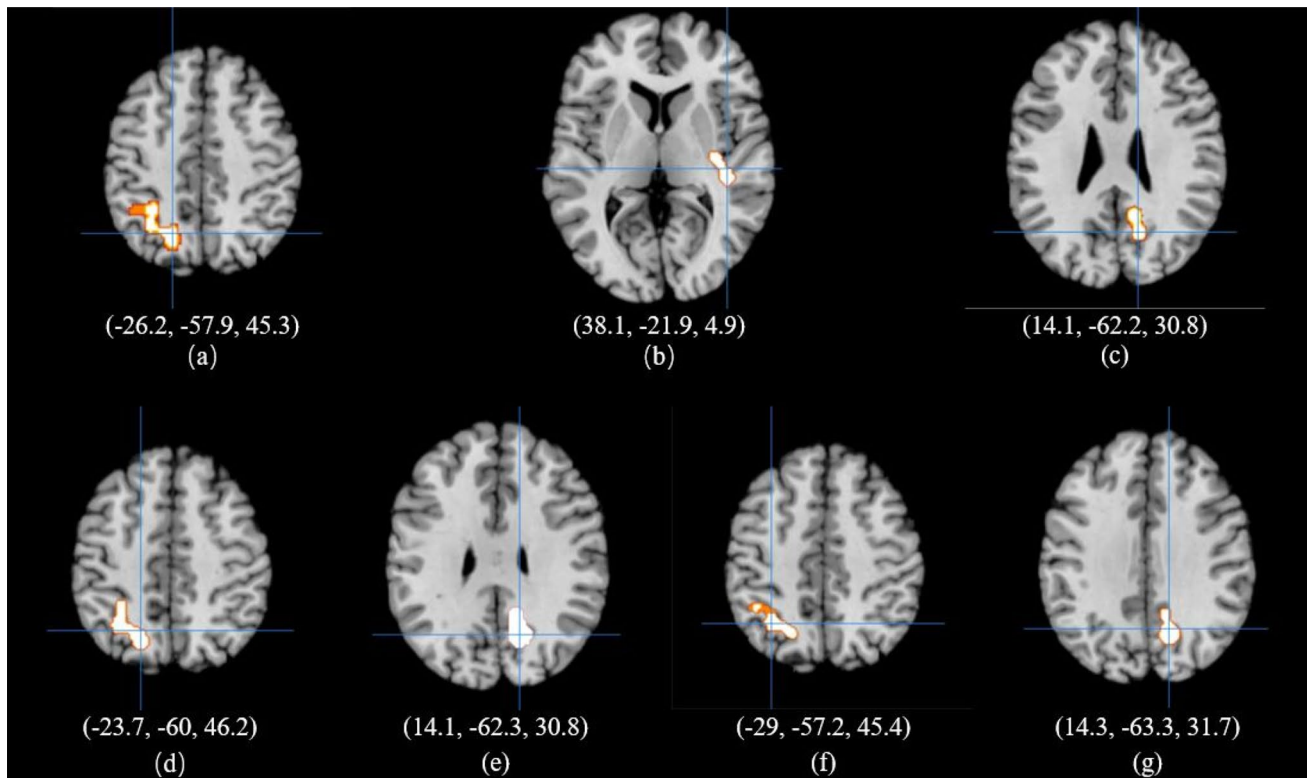


Fig. 2 Activated clusters in overall analysis and subgroup analyses. Notes: **a** activation for overall analysis; **b** increased activation; **c** activation for <35-year-old subgroup; **d** activation for \geq 35-year-old sub-

group; **e** activation for <12-week subgroup; **f** activation for \geq 12-week subgroup; **g** activation for physically inactive subgroup

cingulate (6.5%), and cingulate gyrus (4.7%). The cluster encompassed regions located in Brodmann areas 7 and 31.

In the older-age group (Boa Sorte Silva et al. 2020; Duchesne et al. 2016; Gourgouvelis et al. 2017; Hsu et al. 2018; Liu-Ambrose et al. 2012; Martinsen et al. 2018; Nishiguchi et al. 2015; Pensel et al. 2018; Smith et al. 2013; Wu et al. 2018), 157 foci from 11 experiments were included and converged on a 2360-mm³ cluster spanning from (−36, −72, 40) to (−10, −44, 56), with the center at (−23.7, −60, 46.2) (Fig. 2d). The 4 peaks of this cluster were located in the parietal lobe of the left hemisphere. The identified cluster was mainly located in the precuneus (82.2%), and additionally encompassed the inferior parietal lobule (6.7%), superior parietal lobule (6.7%), and angular gyrus (4.4%). In addition, the cluster included regions located in Brodmann areas 7, 19 and 39.

Subgroup analysis for intervention duration

For the shorter-term duration interventions (Chen et al. 2016; Goldin et al. 2012a,b; Gourgouvelis et al. 2017; Li et al. 2019; Liu-Ambrose et al. 2012; Metcalfe et al. 2016; Schmitt et al. 2019; Wagner et al. 2017; Wriessnegger et al. 2014), 109 foci from 14 experiments converged on a 2008-mm³ cluster

centered at (14.1, −62.3, 30.8) with 3 peaks in right hemisphere (Fig. 2e). Among these peaks, 1 peak with the maximum value was located in the occipital lobe and 2 peaks were situated in the limbic lobe. The cluster was primarily located in the precuneus (80.2%), spreading into the cuneus (10.9%), posterior cingulate (5%), and cingulate gyrus (4%). At the brain region level, Brodmann areas 31 and 7 contributed to 73.3% and 25.7% to this cluster, respectively. For the longer-term duration interventions (Baeck et al. 2012; Boa Sorte Silva et al. 2020; Duchesne et al. 2016; Hsu et al. 2018; Krafft et al. 2014; Martinsen et al. 2018; Nishiguchi et al. 2015; Pensel et al. 2018; Smith et al. 2013; Wu et al. 2018), 151 foci from 12 experiments converged on a 1968-mm³ cluster centered at (−29, −57.2, 45.4) with 4 peaks in the parietal lobe of left cerebrum (Fig. 2f). The cluster encompassed the precuneus (45.5%), as well as the inferior parietal lobule (45.5%), angular gyrus (6.1%), and supramarginal gyrus (3%) corresponding to Brodmann areas 7, 19, 39 and 40.

Subgroup analysis for total amount of physical activity

In the PIG (Chen et al. 2016; Duchesne et al. 2016; Gourgouvelis et al. 2017; Hsu et al. 2018; Krafft et al.

2014; Liu-Ambrose et al. 2012; Martinsen et al. 2018; Metcalfe et al. 2016; Nishiguchi et al. 2015; Schmitt et al. 2019; Wriessnegger et al. 2014; Wu et al. 2018), 85 foci from 16 experiments converged on a 1792-mm³ cluster centered at (14.3, -63.3, 31.7) with 2 peaks (Fig. 2g). Peaks 1 and 2 were located in the right occipital lobe and in the right limbic lobe, respectively. The cluster covered the occipital lobe, parietal lobe and limbic lobes. The cluster primarily included the precuneus (83.8%), and cuneus (33.3%) with additional engagement of the cingulate gyrus (5.1%), which correspond to Brodmann area 31 (72.7%) and Brodmann area 7 (27.3%). In the PAG (Baeck et al. 2012; Boa Sorte Silva et al. 2020; Goldin et al. 2012a, b; Hsu et al. 2018; Pensel et al. 2018; Smith et al. 2013; Wagner et al. 2017), 175 foci from 10 experiments did not converge on a robust cluster.

Anatomical connectivity, functional connectivity and functional characterizations of activated brain regions

In this study, each activated cluster includes more than one peak with each one that can generate a sub-region (radius = 3 mm). Anatomical and functional connectivity of activated sub-regions (4 peaks) in the overall analysis and subgroup analyses are shown in Fig. 3. Functional characterizations of activated sub-regions in both overall analysis and subgroup analyses are summarized in Tables 2, 3, and revealed that the identified sub-regions in the overall analysis exhibited a strong positive coupling with the entire frontoparietal control network and were functionally characterized by engagement in core cognitive domains, including attention, executive functions and working memory. For the functional characterizations, sub-regions were connected with spatial cognition in the overall analysis. In the subgroup analyses, activated sub-regions were associated with explicit memory in the younger-age group, shorter-term group, and PIG, whereas activated sub-regions were associated with working memory in the older-age group and longer-term intervention group. In addition, in both overall analysis and subgroup analysis (older-age group), activated sub-regions were associated with paradigms measuring mental rotation.

In the overall analysis, the frontoparietal and dorsal attention networks were involved in exercise-induced changes in functional activation patterns. For subgroup analyses, associated networks varied across subgroups: (1) younger-age group (frontoparietal network) and older-age group (dorsal attention network); (2) shorter-term group (frontoparietal and default networks) and longer-term group (frontoparietal and dorsal attention networks); (3) PIG (frontoparietal and default networks).

Risk of bias assessment

Total score across included studies ranged from 3 to 7 ($M = 4.80$ and $SD = 1.40$) that correspond to poor to good quality. Notably, only two studies scored 7. Points in the majority of included studies were mainly deducted due to their study design such as lack of random allocation ($n = 9$), concealed allocation ($n = 14$), assessor blinding ($n = 18$), intention-to-treat analysis ($n = 18$). Detailed information is displayed in Supplementary data 2.

Discussion

Exercise-induced brain activation associated with cognition

The overall meta-analysis encompassing data from all original studies demonstrated robust exercise-induced changes in cognition-related functional activation of the left parietal lobe, primarily covering the precuneus (69.7%) and spreading into inferior (27.3%) and superior (3%) parietal lobe. The precuneus plays a key role in a range of highly integrated mental processes, ranging from basic cognitive processes to regulatory control over performance under stress (Cavanna 2006; Zhao et al. 2020). The integrative function of the precuneus is further reflected in the functional connectivity profiles of these regions, which include separable interactions with networks engaged in sensorimotor and cognitive processes (e.g., executive function, episodic memory, visuospatial processing) (LJ 2006). Sporns and Bullmore proposed that the precuneus plays a key role in the frontoparietal network by interconnecting parietal and prefrontal regions (“small-world network” hub), which provides an explanation for the aforementioned activation during cognitive tasks (Bullmore 2009). Moreover, the precuneus is a brain region which is commonly affected in individuals with mild cognitive impairment and early stages of Alzheimer’s disease (Jacobs et al. 2012). In light of this observation, our findings suggest that regular physical exercise can be a valuable approach to prevent cognitive decline (Alty et al. 2020; Bherer et al. 2013; Kivipelto et al. 2018) by enhancing the functional integration of the frontoparietal control network via effects on the precuneus.

The activated sub-regions were connected with dorsal attention network and frontoparietal network, and the functional characterizations revealed a strong engagement in core cognitive domains, including attention as well as executive functions. The frontoparietal network is a functional hub sharing connectivity with diverse brain networks and plays an essential role in modulating cognitive control (Power et al. 2013). Moreover, the degree of frontoparietal network’s coupling with other brain networks (especially

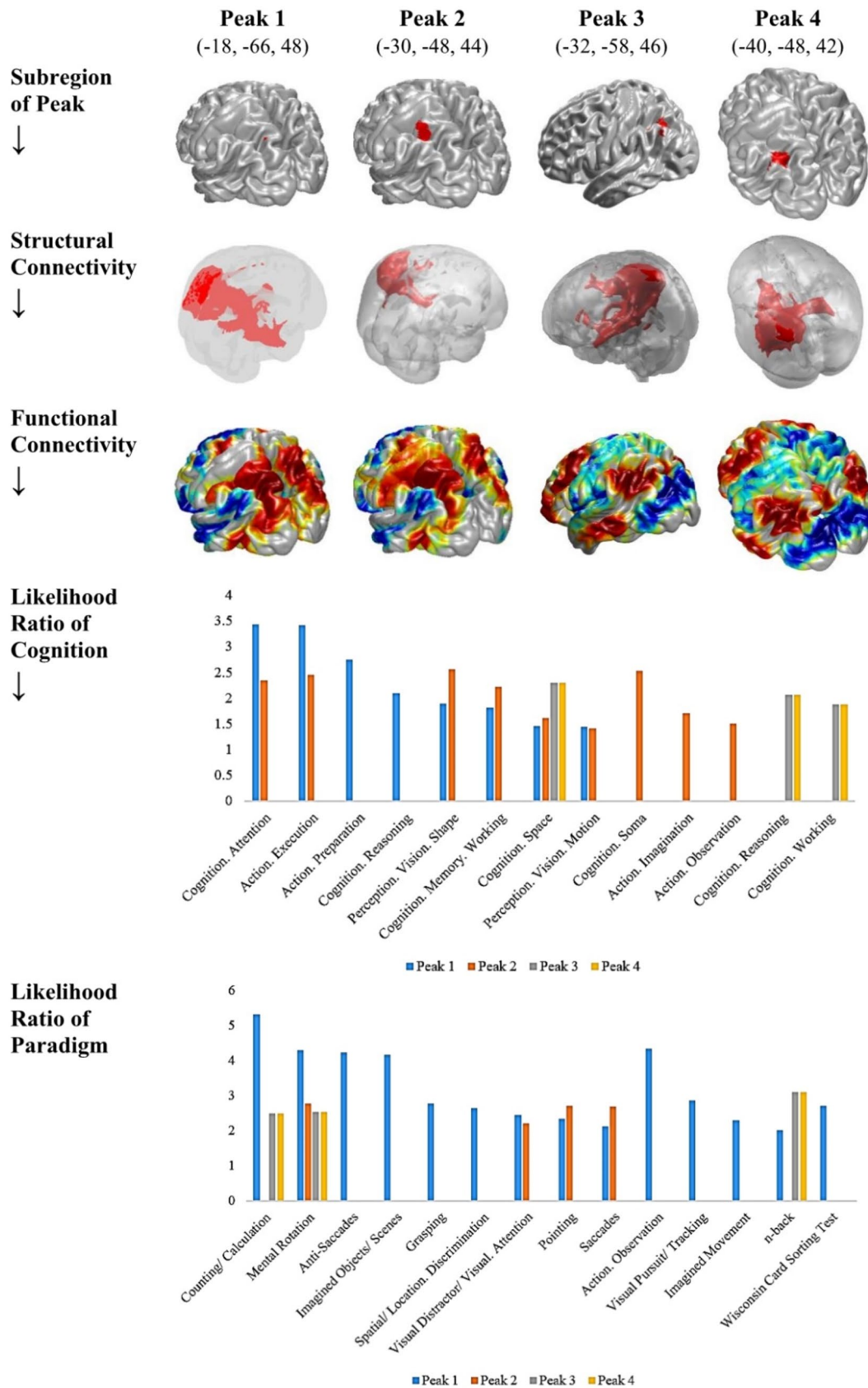


Fig. 3 Anatomical connectivity, functional connectivity and functional characterizations of activated brain sub-regions in the overall analysis

Table 2 Characterizations of activated brain regions in overall analysis

	Coordinate			Location	Likelihood ratio of behavioral domain	Likelihood ratio of paradigm		
	x	y	z					
Peak 1	-18	-66	48	Left Cerebrum. Parietal Lobe. Precuneus. Gray Matter. Brodmann area 7	Cognition. Attention	3.44	Counting/calculation	5.31
					Action. Execution	3.41	Mental Rotation©	4.28
					Action. Preparation	2.75	Anti-Saccades	4.23
					Cognition. Reasoning	2.10	Imagined Objects/scenes	4.15
					Perception. Vision. Shape	1.90	Grasping	2.77
					Cognition. Memory. Working	1.82	Spatial/Location. Discrimination	2.65
					Cognition. Space ®	1.46	Visual Distractor/Visual. Attention	2.45
					Perception. Vision. Motion	1.45	Pointing	2.33
Peak 2	-30	-48	44	Left Cerebrum. Parietal Lobe. Superior Parietal Lobule. Gray Matter Brodmann area 7	Perception. Vision. Shape	2.56	Action. Observation	4.34
					Cognition. Soma	2.53	Visual Pursuit/Tracking	2.85
					Action. Execution	2.45	Mental Rotation©	2.77
					Cognition. Attention	2.35	Pointing	2.70
					Cognition. Memory. Working	2.22	Saccades	2.69
					Action. Imagination	1.71	Imagined Movement	2.29
					Cognition. Space ®	1.61	Visual Distractor/Visual. Attention	2.20
					Action. Observation	1.51	n-back	2.01
Peak 3	-32	-58	46	Left Cerebrum. Parietal Lobe. Inferior Parietal Lobule. Gray Matter Brodmann area 39	Cognition. Space ®	2.29	n-back	3.1
					Cognition. Reasoning	2.06	Wisconsin Card Sorting Test	2.71
					Cognition. Working	1.88	Mental Rotation©	2.53
							Counting/Calculation	2.49
Peak 4	-40	-48	42	Left Cerebrum. Parietal Lobe. Inferior Parietal Lobule. Gray Matter Brodmann area 40	Cognition. Space ®	2.29	n-back	3.1
					Cognition. Reasoning	2.06	Wisconsin Card Sorting Test	2.71
					Cognition. Working	1.88	Mental Rotation©	2.53
							Counting/Calculation	2.49

® and © indicate overlapping behavioral domain(s) or paradigm(s) across sub-regions, respectively

default mode network) is positively correlated with fluid intelligence (e.g., problem solving/executive function and visuospatial ability) and overall cognitive ability (Cole et al. 2015; Sheffield et al. 2015), which is consistent with our results in overall analysis for global cognition (all cognitive components were pooled together for the overall analysis). The dorsal attention network, centered around the intraparietal sulcus and frontal eye fields, participates in top-down control of attention as well as sensory–motor information integration, which support several higher level cognitive domains (Fox et al. 2006).

Moderators have impact on exercise-induced cognitive improvement

In this review, age and training duration considerably moderated exercise effects on brain activation that is associated with anatomical and functional networks. For instance, in the younger-age group, the identified sub-regions are associated with explicit memory. This result is strongly supported by

a recent meta-analysis (including 25 experimental studies) demonstrating robust effects of exercise on episodic memory function (Loprinzi et al. 2019). Furthermore, the recruited participants in 23 studies were reported with age range of 18 to 30 years old, which coincides with the younger-age group (less than 35 years old) in the current review. In the older-age group, identified sub-regions are related to executive functions, including working memory, spatial cognition and mental rotation. Such improvements induced by exercise have been well-documented in previous meta-analytical studies at the behavioral level (Chen et al. 2020b; Falck et al. 2019; Northey et al. 2018).

With regard to the moderating effect of training duration on exercise-induced activation, shorter-term exercise interventions induced primarily changes in occipital, parietal and limbic regions and the peaks/sub-regions are generally linked to explicit memory, whereas the longer-term training protocols induced changes in activation in the parietal lobe. Of note, the shorter-term group included 14 experiments with 7 using acute physical exercise, which is partially

Table 3 Characterizations of activated brain regions in subgroup analyses

	Coordinate			Location	Likelihood ratio of behavioral domain	Likelihood ratio of paradigm	
	x	y	z				
<35 Years Old Subgroup							
Peak 1	14	-64	32	Right Cerebrum. Occipital Lobe. Precuneus. Gray Matter. Brodmann area 31	Perception. Vision. Motion Cognition. Memory. Explicit®	2.06 1.94	Saccades 2.51
Peak 2	12	-52	32	Right Cerebrum. Limbic Lobe. Cingulate Gyrus. Gray Matter. Brodmann area 31	Cognition. Social Cognition Cognition. Memory. Explicit®	3.11 2.45	Cued Explicit. Recognition Episodic Recall Theory of Mind Task 5.88 3.48 2.46
Peak 3	12	-54	22	Right Cerebrum. Limbic Lobe. Posterior Cingulate. Gray Matter. Brodmann area 23	Perception. Vision. Motion Cognition. Memory. Explicit®	2.06 1.94	Saccades 2.51
≥35 Years Old Subgroup							
Peak 1	-18	-68	48	Left Cerebrum. Parietal Lobe. Precuneus. Gray Matter. Brodmann area 7	Cognition. Attention Action. Execution Action. Preparation Cognition. Reasoning Perception. Vision. Shape Cognition. Memory. Working® Cognition. Space® Perception. Vision. Motion	3.44 3.41 2.75 2.10 1.90 1.82 1.46 1.45	Counting/Calculation Mental Rotation Anti-Saccades Imagined Objects/Scenes Grasping Spatial/Location. Discrimination Visual Distractor/Visual. Attention Pointing Saccades 4.34 2.11
Peak 2	-30	-48	44	Left Cerebrum. Parietal Lobe. Superior Parietal Lobule. Gray Matter. Brodmann area 7	Perception. Vision. Shape Cognition. Soma Action. Execution Cognition. Attention Cognition. Memory. Working® Action. Imagination Cognition. Space® Action. Observation Perception. Vision. Motion	2.56 2.53 2.45 2.35 2.22 1.71 1.61 1.51 1.41	Action. Observation Visual Pursuit/Tracking Mental Rotation Pointing Saccades Imagined Movement Visual Distractor/Visual. Attention n-back 2.01
Peak 3	-32	-58	46	Left Cerebrum. Parietal Lobe. Inferior Parietal Lobule. Gray Matter. Brodmann area 39	Cognition. Space® Cognition. Reasoning Cognition. Memory. Working®	2.29 2.06 1.88	n-back Wisconsin Card Sorting Test Mental Rotation Counting/Calculation 2.49
Peak 4	-12	-64	54	Left Cerebrum. Parietal Lobe. Precuneus. Gray Matter. Brodmann area 7	–	–	–
<12 weeks subgroup							
Peak 1	14	-64	32	Right Cerebrum. Occipital Lobe. Precuneus. Gray Matter. Brodmann area 31	Perception. Vision. Motion Cognition. Memory. Explicit®	2.06 1.94	Saccades 2.51
Peak 2	12	-52	32	Right Cerebrum. Limbic Lobe. Cingulate Gyrus. Gray Matter. Brodmann area 31	Cognition. Social Cognition Cognition. Memory. Explicit®	3.11 2.45	Cued Explicit. Recognition Episodic Recall Theory of Mind Task 5.88 3.48 2.46
Peak 3	12	-54	22	Right Cerebrum. Limbic Lobe. Posterior Cingulate. Gray Matter. Brodmann area 23	Perception. Vision. Motion Cognition. Memory. Explicit®	2.06 1.94	Saccades 2.51

supported by results from a previous meta-analysis which investigated the beneficial effects of acute exercise on episodic memory as a type of explicit memory (Tulving 1972).

Meanwhile, researchers emphasized that the timing of acute exercise plays an important role in the interaction of exercise and memory; such that cognitive improvement was observed

Table 3 (continued)

	Coordinate			Location	Likelihood ratio of behavioral domain	Likelihood ratio of paradigm		
	x	y	z					
≥ 12 weeks subgroup								
Peak 1	-32	-58	46	Left Cerebrum. Parietal Lobe. Inferior Parietal Lobule. Gray Matter. Brodmann area 39	Cognition. Space®	2.29	n-back	3.10
					Cognition. Reasoning®	2.06	Wisconsin Card Sorting Test	2.71
					Cognition. Memory. Working®	1.88	Mental Rotation	2.53
							Counting/Calculation	2.49
Peak 2	-40	-48	42	Left Cerebrum. Parietal Lobe. Inferior Parietal Lobule. Gray Matter. Brodmann area 40	Cognition. Space®	2.29	n-back	3.10
					Cognition. Reasoning®	2.06	Wisconsin Card Sorting Test	2.71
					Cognition. Memory. Working®	1.88	Mental Rotation	2.53
							Counting/calculation	2.49
Peak 3	-20	-64	46	Left Cerebrum. Parietal Lobe. Precuneus. Gray Matter. Brodmann area 7	Cognition. Reasoning®	3.25	Saccades	3.54
					Cognition. Memory. Working®	2.92	Delayed Match To Sample	3.37
					Cognition. Attention	2.81	Stroop Task	2.73
					Perception. Vision. Motion	2.37	Mental Rotation	2.68
					Cognition. Space®	1.72	Wisconsin Card Sorting Test	2.45
					Perception. Vision. Shape	1.68	Task Switching	2.44
Peak 4	-12	-64	54	Left Cerebrum. Parietal Lobe. Precuneus. Gray Matter. Brodmann area 7	–	–	–	–
Physically inactive subgroup								
Peak 1	14	-64	32	Right Cerebrum. Occipital Lobe. Precuneus. Gray Matter. Brodmann area 31	Perception. Vision. Motion	2.06	Saccades	2.51
					Cognition. Memory. Explicit®	1.94		
Peak 2	12	-52	32	Right Cerebrum. Limbic Lobe. Cingulate Gyrus. Gray Matter. Brodmann area 31	Cognition. Social Cognition	3.11	Cue Explicit. Recognition	5.88
					Cognition. Memory. Explicit®	2.45	Episodic Recall	3.48
							Theory of Mind Task	2.46

® and © indicate overlapping behavioral domain(s) or paradigm(s) across sub-regions, respectively

with acute exercise occurring before memory encoding, during early memory consolidation and during late memory consolidation. Positive results on explicit memory function may be attributed to improved pattern separation (Suwabe et al. 2018; Voss et al. 2019) and attenuated memory interference (Crawford et al. 2020) following a relatively short intervention duration. For the longer-term group, the activated cluster was located in the parietal lobe and peak areas were associated with spatial processing, reasoning and working memory. For longer-term exercise, various positive effects on cognition have been reported at the molecular and cellular levels (e.g., brain-derived neurotrophic factor) (Hotting et al. 2016), structural level (e.g., increased gray matter volume in frontal and hippocampal regions) (Chaddock-Heyman et al. 2014), and behavioral level (e.g., improvement in executive function) (Fernandes et al. 2017; Kramer et al. 2005).

Of note, the analyses separately examining healthy subjects and patient groups did not converge on a specific brain region that exhibited changes. However, when lowering the statistical threshold (uncorrected) convergent activation in the parietal lobe emerged for both, healthy subjects

and patients (detailed information can be found in supplementary data 3), indicating that the lack of convergent activation in the subgroups might be due to the reduced number of included experiments in each subgroup analysis. Besides, previous reviews have shown that both healthy people (Kramer and Colcombe 2018) and individuals with neurological, non-neurological, and psychiatric illnesses (Eggermont et al. 2006; Heyn et al. 2004) can benefit from physical exercise on the behavioral level. For instance, a meta-analysis by Dauwan et al., focusing on individuals with chronic disorders (Alzheimer's disease, Huntington's disease, multiple sclerosis, Parkinson's disease, schizophrenia, and unipolar depression), suggested that exercise interventions can improve several cognitive domains (attention, working memory and executive function) with small but significant effect sizes (Dauwan et al. 2016). In addition, an activated cluster was only observed in PIG, but not in PAG. Such results might be explained by physiological and psychological (cognitive) adaptation, which refers to that cellular stress and the resultant metabolic signals have reached relatively stable status (MacInnis and Gibala 2017). It is well known that fMRI measures brain activation by detecting

blood-oxygen-level dependent response to cognitive tasks. Regarding the total amount of exercise, if cognitive benefits reach a peak at 150-min exercise or below, activation may no longer last as the amount of exercise increases due to blood-oxygen metabolism and neural adaptation (Logothetis et al. 2001).

In addition, currently there is no sufficient number of studies available, which would allow to investigate how other important moderators, namely type of exercise and exercise intensity, can influence functional brain activation patterns. In this context, behavioral data suggest that coordinative exercise leads to greater benefits than purely aerobic exercises (Chen et al. 2020a; Ludyga et al. 2020). Hence, it seems a promising area for further research do investigate the underlying neural processes (e.g., functional brain activation patterns) driving this difference in behavioral findings. With regard to exercise intensity, the evidence from behavioral data is not univocal. In physical training studies no clear effect of exercise intensity could be observed (Ludyga et al. 2020), whereas in acute exercise studies the moderating effect of exercise intensity was clearly demonstrated as an exercising at higher intensity improves cognitive performance to a greater extent (Chang et al. 2012; Oberste et al. 2019). Hence, further studies are necessary to investigate whether the difference in behavioral findings, caused by the moderators type of exercise and exercise intensity, are mirrored in altered cognition-related functional brain activation patterns.

Implications

This is the first study to systematically examine the relationship between functional brain activation and cognition as a function of exercise practice. Therefore, the present results can be used to guide future research on exercise effects on brain health and cognition. Firstly, in the subgroup analyses of age and training duration, activation was observed in different brain regions. Follow-up studies should explore the associations between hemispheric lateralization and moderators (age and duration). Secondly, only the improvement of spatial cognition was observed in the overall analysis without consideration of moderators, while the benefits on various cognitive components like explicit memory, working memory and reasoning were found in the subgroup analyses. It can be inferred that the cognitive benefits from participating in physical exercise interventions are specifically influenced by age and training duration, which is consistent with previous behavioral meta-analytical reviews (Verburgh et al. 2014; Xue et al. 2019). Thirdly, because of the limited number of studies available for analysis, we cannot determine the influences of sex, exercise type, exercise intensity, and disease type on cognitive improvement. Given the high heterogeneity in the approaches to describe and report exercise

intensity in the reviewed studies, a more standardized prescription of exercise intensity would be highly desirable to facilitate a meta-analytical data analysis and to simplify the reproducibility of the studies (Gronwald et al. 2019; Herold et al. 2020), although the optimal approach to describe exercise intensity is still debated (Herold et al. 2019, 2020; Herold 2020; Jamnick et al. 2020). However, these moderators should be examined in future well-designed exercise–cognition studies using neuroimaging techniques. Fourthly, the small number of included experiments has limited researchers to focus on two age groups only for subgroup analysis. Furthermore, original imaging studies on this topic that focused on children and adolescent, middle-aged people, young-old (55–65 years old) group, oldest-old population are still in its infancy, which requires further investigation. Fifth, results in this current meta-analysis are generated from pre-to-post experiment, instead of between-group contrast. As a result, observed positive changes could be attributed to a variety of other factors that interact with the physical exercise interventions. For example, when people start an physical exercise program, they may change their diet and/or have more social interaction that are highly associated with improved cognitive function (Bailey et al. 2019; Hardman et al. 2020; Yu et al. 2020). Thus, future imaging studies should include active and/or passive control group (s) in order to draw a firm conclusion about the cognitive benefits of physical exercise intervention (Herold et al. 2020). Lastly, when using the well-recognized PEDro scale for risk of bias assessment, the majority of included studies scored 7 below (blinding of participants and instructor are relatively difficult in physical exercise intervention, leading to 9 points in total). Moreover, the most of the reviewed has a relatively small sample size which can result in a low power influencing effect size estimation and reproducibility negatively (Button et al. 2013). Thus, more well-designed randomized controlled trials with an adequate sample size should be conducted on this topic to enhance the robustness of the findings.

Conclusions

The evidence for exercise effects on cognition is extensive but still growing. Combined with structural brain effects and behavioral data from previous studies, this article demonstrates that exercise-induced changes in functional brain activation in parietal regions (precuneus, superior and inferior parietal lobule, cingulate gyrus and posterior cingulate) and associated networks (frontoparietal network, dorsal attention network and default mode network) may neutrally mediate exercise-induced cognition enhancement. Furthermore, the present findings emphasize that the brain functional effects of exercise vary as a function of age and duration.

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Compliance with ethical standards

Declaration of interest The authors declare that they have no conflict of interest.

Research involving human participants and/or animals and informed consent This manuscript does not report on any findings from studies that enrolled human participants or animals. Rather, it synthesizes information gathered from existing literature and the informed consent is not applicable.

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