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41	Abstract

Accumulating research suggests that, as a result of reduced neural activity in the prefrontal 42 cortex (PFC), higher-order cognitive function may be compromised while engaging in high-43 intensity acute exercise, with this phenomenon referred to as the transient hypofrontality effect. 44 However, findings in this field remain unclear and lack a thorough synthesis of the evidence. 45 Therefore, the purpose of this meta-analysis was to evaluate the effects of in-task acute exercise 46 47 on cognitive function, and further, to examine whether this effect is moderated by the specific type of cognition (i.e., PFC-dependent vs. non-PFC-dependent). Studies were identified by 48 49 electronic databases in accordance with the PRISMA guidelines. In total, twenty-two studies met 50 our inclusion criteria and intercept only meta-regression models with robust variance estimation were used to calculate the weighted average effect sizes across studies. Acute exercise at all 51 intensities did not influence cognitive function ( $\beta = -0.16, 95\%$  CI = [-0.58, 0.27], p = .45) when 52 exercise occurred during the cognitive task, and no significant moderation effects emerged. 53 However, there was evidence that cognitive task type (PFC-dependent vs. non-PFC-dependent) 54 55 moderated the effect of high-intensity acute exercise on a concomitant cognitive performance ( $\beta$ = -0.81, 95% CI = [-1.60, -0.02], p = .04). Specifically, our findings suggest that PFC-dependent 56 cognition is impaired while engaging in an acute bout of high-intensity exercise, providing 57 58 support for the transient hypofrontality theory. We discuss these findings in the context of a cognitive-energetic perspective. 59

60

Keywords: Cognitive decline, intense physical activity, mental resources, prefrontal activation,
transient hypofrontality theory.

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- 64

#### Introduction

Cognitive functions in the brain are mental processes enabling individuals to receive, 65 select, store, transform, develop, and remember information that originated from external stimuli 66 (Zhang, 2019). There are various types of cognitions, including attention, memory, and executive 67 function, which play a significant role in optimal daily functioning across the lifespan, for 68 instance, in making decisions and completing tasks demanding complex reasoning and 69 70 information processing (Aretouli & Brandt, 2010; Warren et al., 1989). Neuroimaging studies on healthy participants have shown that a critical brain region involved in such higher-order 71 72 cognitive functions is the prefrontal cortex (PFC), and these results provided evidence of 73 increased activation in a wide range of prefrontal regions during task performance (Blumenfeld & Ranganath, 2006; D'Esposito et al., 1999; Petrides, 2000). The role of PFC is to provide the 74 infrastructure to compute executive processing, which is strongly associated with the ability to 75 engage in goal-consistent behaviors and inhibit goal-inconsistent behaviors (Funahashi, 2017; 76 77 Goethals et al., 2004). Taken together, several cognitive tasks, such as the Stroop task or the 78 Digit Symbol Substitution test, which depend largely on frontal-prefrontal network and require substantial prefrontal engagement, are generally regarded as PFC-dependent tasks. In contrast to 79 80 the PFC-dependent tasks, non-PFC-dependent cognition encompasses relatively automatized, 81 simple decisional tasks (e.g., Choice Reaction Time [RT] and Visual Recognition task) that rely less on cognitive effort (Dienes & Perner, 1999). Using brain imaging techniques, it is suggested 82 83 that such tasks are primarily dependent on early sensory and late motor processes rather than 84 prefrontal cognitive processes (Grèzes et al., 2003). 85 Over the past few decades, exercise psychologists have been interested in the immediate

effect of acute exercise on multiple subdomains of cognition and, in particular, attempted to evaluate whether the type of cognitive task (e.g., PFC-dependent vs. non-PFC-dependent)

influences the acute exercise-cognition relationship in protocols in which the cognitive task is 88 carried out during exercise (i.e., concomitance protocols). With this protocol, some studies have 89 indicated that acute exercise has the selective potential to enhance cognitive function (Audiffren 90 et al., 2008; Davranche et al., 2005; Lambourne et al., 2010), whereas others have suggested a 91 particular pattern of cognitive impairment occurred during a single bout of exercise (Del Giorno 92 93 et al., 2010; Dietrich & Sparling, 2004; Komiyama et al., 2020; Loprinzi et al., 2019). This facilitation or impairment effect is likely to be influenced based on the exercise intensity and 94 95 type of cognitive task. Various theoretical accounts have been developed to explain these 96 contradicting effects, such as arousal, attention, and cognitive-energetic models (see Tomporowski & Qazi, 2020). Central to the present review is the potential impairment effects of 97 cognitive function during an acute bout of exercise. We evaluate this phenomenon within a 98 cognitive-energetic model, namely the transient hypofrontality theory (Dietrich, 2003, 2006). 99 100 The transient hypofrontality theory posits that in dual-task conditions where acute 101 exercise at higher intensity and cognitive tasks are performed simultaneously, the neural activation in non-motor areas of the PFC may be reduced as more metabolic and cognitive 102 resources may be allocated toward sensory and motor cortices to maintain physical movement. 103 104 Given that the brain operates on a limited amount of such resources (Miller & Cohen, 2001), the widespread activation of motor and sensory cortices while exercising at higher intensity may 105 106 come at the expense of activity in other neural structures that are not essential for controlling 107 motor movement. As such, this may result in a temporary deactivation of prefrontal structures 108 involved in higher-order cognitive processing, and ultimately, compromise task performance 109 (Audiffren, 2016; Dietrich, 2003). This basic assumption has been supported in experimental 110 studies in animals showing that exercise at a high intensity ( $\geq 85\%$  VO<sub>2max</sub>) increases neural

activity in areas (e.g., motor cortex) of the frontal lobe involved in motor control (39%  $\Delta$ 111 (increase) from baseline, p = .001), with no such changes in the PFC (11%  $\Delta$  from baseline, p 112 113 > .05) or frontal cortex (6%  $\Delta$  from baseline, p > .05) (Vissing et al., 1996). Moreover, in a human sample, exercise above 80% of VO<sub>2max</sub> significantly decreased cerebral oxygenation in 114 the right frontal cortex (Ando et al., 2011), which plays an important role in planning-based 115 116 cognition (Henson et al., 1999). Although speculative, the reduction in cerebral oxygenation of the PFC induced by this exercise protocol may impair PFC-dependent cognition. For example, 117 118 not only does high-intensity acute exercise decrease PFC oxygenation, but research also shows 119 that PFC-dependent cognition is compromised while engaging in acute high-intensity exercise (Mekari et al., 2015). Further, prior work shows that premotor time of the Eriksen Flanker task 120 did not improve while engaging in exercise above 80% VO<sub>2max</sub>, but improved at 60% VO<sub>2max</sub> of 121 exercise, compared to a resting condition (Ando et al., 2011). Thus, high-intensity exercise may 122 123 decrease regional levels of cerebral oxygenation (e.g., frontal cortex), which may fail to meet 124 cerebral metabolic demands and cause a concomitant transient inhibition of the PFC function, leading to PFC-dependent cognitive decline (Dietrich & Audiffren, 2011; Subudhi et al., 2007). 125 Collectively, the results from these animal and human studies suggest that high-intensity acute 126 127 exercise increases neural activity in the motor cortex, but either plateaus or downregulates neural activity and oxygenation in other areas of the PFC not specifically involved in motor control. 128 129 As stated previously, per the tenets of the transient hypofrontality theory, such 130 detrimental effects may be moderated by cognitive task types according to their dependence on 131 prefrontal functioning (i.e., PFC-dependent task vs. non-PFC-dependent task). For example, Bue-Estes et al. (2008) showed that working memory (PFC-dependent task) was impaired during 132 133 short-term maximal incremental treadmill exercise. Schmit et al. (2015) reported that the error

rate of the modified version of the Eriksen Flanker task (PFC-dependent task) was higher during 134 cycling at 85% of maximal aerobic power compared to a non-exercise condition. On the other 135 136 hand, an experimental study by Rattray and Smee (2016) found that response time in the Speed Match task (non-PFC-dependent task) was faster during exercise at 90% VO<sub>2peak</sub> than at rest. 137 Summarily, during high-intensity acute exercise, performance on tasks demanding PFC-138 139 dependent cognition may be potentially compromised, whereas cognitive tasks requiring less prefrontal activity may be enhanced or unaffected. A few near-infrared spectroscopy studies, 140 however, showed deleterious effects of acute low- and moderate-intensity exercise on a 141 142 concomitant PFC-dependent cognitive performance measured by the Eriksen Flanker task and the Simon task (Davranche & McMorris, 2009; Pontifex & Hillman, 2007). In addition to these 143 findings, other studies showing positive impacts of acute in-task exercise on PFC-dependent 144 cognition provided suggestive evidence that other potential moderators (e.g., too short duration 145 146 of exercise, too light-intensity exercise, and too high level of participants' physical fitness) 147 would influence the relationship between acute exercise and PFC-dependent cognitive task performance under dual-task conditions (Lucas et al., 2012; Martins et al., 2013; Pesce & 148 Audiffren, 2011). Accordingly, additional research is needed to fully understand the transient 149 150 hypofrontality effect and whether this is moderated by multiple characteristics, in particular, by different types of cognition. 151

Although recent experiments (Chang et al., 2017; Loprinzi et al., 2019; Siddiqui & Loprinzi, 2018), narrative reviews (Loprinzi et al., 2017; Netz, 2019), and meta-analytic reviews (Chang et al., 2012; Jung et al., 2020; Lambourne & Tomporowski, 2010; Loprinzi et al., 2019; Roig et al., 2013) have demonstrated that acute exercise can influence cognitive function, we have less knowledge as to whether cognitive function is impaired during an acute bout of

exercise and whether the specific type of cognition (i.e., PFC-dependent vs. non-PFC-dependent) 157 moderates this effect. In fact, while a number of past meta-analytic publications have confirmed 158 the moderating effects of cognitive task type on the acute exercise-cognition association by 159 grouping this into six categories (i.e., information processing, reaction time, attention, crystalized 160 intelligence, executive function, and memory) using a general approach (Chang et al., 2012, Jung 161 162 et al., 2020), we divided the task type into PFC-dependent and non-PFC-dependent tasks, which is a novel approach in this field. Therefore, the purpose of this meta-analytic review was to 163 investigate the potential differential effects of acute exercise on PFC-dependent and non-PFC-164 165 dependent cognition. In this meta-analysis, we specifically focused on articles that directly compared a non-exercise control group to acute exercise that occurred during the cognitive task 166 and evaluated studies separately based upon exercise intensity (i.e., all studies from comparisons 167 168 with low-, moderate-, and high-intensity acute exercise [Aim 1] and only studies from 169 comparisons with high-intensity acute exercise [Aim 2]). Parallel to the predictions of the 170 transient hypofrontality effect, we hypothesized that only PFC-dependent cognition will be impaired while engaging in high-intensity acute exercise. 171 **Methods** 172

173 Data Sources and Search Strategy

Studies were identified by electronic databases in accordance with the Preferred
Reporting Items for Systematic Reviews and Meta-Analyses (PRISMA) guidelines: PubMed
(1951–present), Scopus (2004–present), PsycINFO (1981–present), Google Scholar (2004–
present), and SPORTDiscus (1985–present). All documents were retrieved from inception to
March 14<sup>th</sup>, 2020. The search terms, including their combinations, were: acute exercise, physical
activity, cognitive function, prefrontal cortex, and transient hypofrontality. To minimize the

possibility of errors during searching articles, we used database-appropriate syntax for each
database, in combination with the selected terms, based on a recently developed systematic
search strategy (Bramer, et al., 2018). Table 1 represents the database-appropriate syntax for
each database.

184

#### [Insert Table 1 Here]

#### 185 Study Selection

Two separate authors (Jung and Ryu) independently employed the computerized searches 186 to determine the number of eligible studies. Each of the searches in each respective database was 187 imported into EndNote, and then, duplicate references were removed in EndNote from March 188 16<sup>th</sup> to 18<sup>th</sup>, 2020. Agreement on eligible studies was reached from these two independent 189 reviews. In addition to the aforementioned databases, to identify additional relevant articles for 190 191 inclusion, two authors reviewed key review papers on acute exercise and cognition (Chang et al., 2012; Loprinzi et al., 2019; Roig et al., 2013; Tomporowski, 2003) as well as transient 192 hypofrontality papers (Dietrich, 2003, 2006; Dietrich & Audiffren, 2011; Dietrich & Sparling, 193 2004). Notably, the Scopus database was used to identify all citations of these four transient 194 hypofrontality papers. Any studies that cited any of these four transient hypofrontality papers 195 196 were reviewed for possible inclusion in our meta-analysis. All studies appearing to meet the inclusion criteria were reviewed and cross-checked at the full text level from March 20<sup>th</sup> to 22<sup>nd</sup>, 197 198 2020. If any disagreement occurred, a third review author was invited to reach consensus through 199 discussion.

200 Inclusion Criteria

Studies were included if they: (1) employed an experimental design with a comparison to
a control group/visit (i.e., no exercise), (2) included human participants, (3) assessed exercise as

an independent variable, (4) performed acute exercise (defined as a single bout of exercise), (5)
evaluated cognitive function as a primary outcome, (6) conducted cognitive tasks during an acute
bout of exercise, (7) provided sufficient data (e.g., sample size, mean, and standard deviation
[SD]) for computing an effect size (ES) estimate, and (8) were published in English.

207 Methodological Quality of Evaluated Studies

208 Two authors independently reviewed the included studies for methodological quality using the modified Downs and Black checklist (Downs & Black, 1998). This checklist was 209 210 developed for the assessment of the methodological quality of randomized and nonrandomized 211 studies and was based on 27 criteria across 4 domains (e.g., reporting, external validity, internal validity, and power), providing a total maximum score of 28 (1 point per question except 212 question five [2 points]). All disagreements in quality ratings between reviewers were solved by 213 consensus. To avoid confusion of power calculation for users, the last question was revised from 214 a 5-point to a 1-point rating, where 1 was scored if a power or sample size computation was 215 216 reported, and 0 was scored when there was no power computation or indication of whether the number of subjects was appropriate for the study design. Of the 27 items, 3 items, "Have the 217 characteristics of patients lost to follow-up been described?", "In trials and cohort studies, do the 218 219 analyses for different lengths of follow-up of patients, or in case-control studies, is the time period between the intervention and outcome the same for cases and controls?" and "Were losses 220 of patients to follow-up taken into account?" were removed from the checklist as they did not fit 221 222 the inclusionary criteria of our meta-analysis. Thus, we utilized the modified version of the Downs and Black checklist including 24 items; scoring ranging from 0 to 25, with a higher score 223 224 indicating a higher level of methodological quality.

225 Data Extraction of Included Studies

Detailed data from each of the included studies were extracted and coded, including the following information: (1) author, date, and country of study, (2) sample size and characteristics, (3) study design, (4) exercise protocol (e.g., exercise type, exercise intensity, and exercise duration), (5) type of cognitive task (e.g., PFC-dependent task and non-PFC-dependent task), and (6) mean and SD of cognitive function between performing tasks under control (no exercise) and exercise conditions.

#### 232 Categorization of Moderators

233 The evaluated moderators included age, sex, exercise protocol, and specific cognitive 234 task type. These evaluated moderators were selected since they have been shown to potentially affect cognitive performance or influence the effects of acute exercise on cognitive function 235 (Chang et al., 2012). Sex was categorized as males, mixed samples, and predominantly female. 236 Predominantly female was defined as a study including > 71% females (Barha et al., 2017). Age 237 was categorized as young adult (18–30 years), middle-aged adult (31–60 years), and older adult 238 239 (> 60 years) (Chang et al., 2012). Exercise protocol included exercise intensity, exercise duration, and exercise modality. Exercise intensity was based on thresholds suggested by the 240 American College of Sports Medicine (Garber et al., 2011). For instance, according to maximum 241 242 heart rate estimates, low-, moderate-, and high-intensity exercise were defined as < 64%, 64%-76%, and > 76%, respectively. Exercise duration was defined as short (< 20 min), medium (20– 243 244 40 min), and long duration (> 40 min) (Roig et al., 2013). Exercise modality was defined as 245 treadmill-based walking/running and cycling. Lastly, in alignment with other reviews (Chang et 246 al., 2012), cognitive task type was categorized as PFC-dependent task and non-PFC-dependent 247 task. For example, the Wisconsin Card Sorting Task is commonly associated with increased 248 neural activity in the PFC, whereas Basic Choice RT and/or Visual Recognition Task are

considered tasks that require little prefrontal activation (see Supplementary Tables 1 and 2,

250 respectively, for PFC-dependent and non-PFC-dependent cognitive tasks).

#### 251 Data Analyses

The ESs were calculated as Hedges' *g* indices and expressed as a standardized mean difference (*g*) between the exercise and control groups. Hedges' *g* was used as the ES for analysis, given that it is a relatively unbiased estimate of the population standardized mean difference ES, while Cohen's *d* is a biased estimate. In the below formula,  $\overline{X_1}$  and  $\overline{X_2}$  are the means for the exercise group (EG) and the control group (CG), respectively.  $SD *_{pooled}$  is the weighted and pooled standard deviation of EG and CG.  $SD_1$  and  $SD_2$  are the standard deviations for EG and CG, respectively, and  $n_1$  and  $n_2$  are the sample sizes for EG and CG, respectively.

Hedges'g = 
$$\frac{\overline{X_1} - \overline{X_2}}{SD *_{pooled}}$$

260 
$$SD *_{pooled} = \sqrt{\frac{(n_1 - 1)SD_1^2 + (n_2 - 1)SD_2^2}{(n_1 + n_2 - 2)}}$$

To calculate ES measures, mean differences in cognitive function between performing 261 tasks during acute exercise  $(\overline{X_1})$  and non-exercise  $(\overline{X_2})$  were divided by the weighted and pooled 262 SD (SD \*pooled). Standardized mean differences adjusted for sampling error were subsequently 263 computed as a measure of individual ES by assigning more weight to studies with larger sample 264 265 sizes. The ESs were estimated so that a negative effect indicated the existence of a deleterious acute exercise effect on cognitive performance. When we calculated the ESs for RT,  $\overline{X_1} - \overline{X_2}$ 266 was replaced with  $\overline{X_2} - \overline{X_1}$ , so that a negative effect also indicated cognitive impairment. Effect 267 sizes values were classified as small effect (0.2), moderate effect (0.5), and large effect (0.8)268 (Hedges & Olkin, 2014). 269

Inverse variance weighted random-effects models were used to meta-analyze the ESs 270 since they make inferences about the effects of acute exercise on cognitive function across 271 various procedures and settings. Most research addresses multiple outcomes due to the 272 multifaceted nature of cognitive function. Multiple outcomes pose a problem for traditional 273 274 meta-analytic approaches, as averaging ESs through studies without considering their 275 correlations can lead to unreliable ES estimates. We meta-analyzed all qualifying ESs in each 276 sample, enabling studies to contribute several ESs to preserve as much data as possible (i.e., multiple ESs based on the same sample in a study). Thus, we used the robust variance estimation 277 278 in the meta-analytic technique, a random-effects meta-regression that can account for the dependence between ES estimates (Hedges et al., 2010). By correcting the within-study standard 279 errors for correlations between ESs, this estimation approach enables clustered data (i.e., ESs 280 281 nested within samples) to be meta-analyzed. This approach includes estimating the mean correlation between all pairs of within-study ESs ( $\rho$ ), which is then used to modify the within-282 study sampling variance  $(\tau^2)$  to account for these statistical dependencies. We conducted a 283 284 sensitivity analysis by setting  $\rho = 0.8$  based on Tanner-Smith and Tipton (2014), and the findings were consistent across various rational values of  $\rho$  (ranging from 0 to 1). We also reported  $I^2$ , 285 which quantifies the proportion of ES variance due to between-sample heterogeneity. Notably,  $I^2$ 286 values of 25%, 50%, and 75% indicate low, moderate, and high levels of heterogeneity, 287 respectively (Higgins et al., 2003). 288 The 'robumeta' package in R was used to conduct inverse variance weighted random-289

effects meta-analyses with Hedges et al. (2010) robust variance estimation (Fisher & Tipton,
2015). We ran the test twice, once with all of the studies included and then with only the studies
from the high-intensity exercise comparisons. An intercept-only meta-regression model was

fitted to assess the overall effect of acute exercise on a concomitant cognitive performance. The 293 constant coefficient in this model has the interpretation of the weighted average effect of acute 294 295 in-task exercise on cognitive function. Exercise-induced cognitive enhancement is represented by a positive constant coefficient, while the exercise-induced cognitive impairment is 296 represented by a negative constant coefficient. Next, to test for the possibility that sex, exercise 297 298 intensity, exercise duration, exercise modality, and cognitive task type explain between-study differences in the weighted average effect of acute in-task exercise on cognitive function (Aim 299 300 1), we added several covariates to our intercept-only meta-regression model. Sex was a 301 categorical variable with three levels (males, mixed samples, and predominantly female) and thus, we included two dummy covariates for sex. The first, males, reflected the males versus 302 others contrast (coded males = 1, predominantly female and mixed samples = 0) while the 303 second, predominantly female, reflected the predominantly female versus others contrast (coded 304 predominantly females = 1, males and mixed samples = 0). Mixed samples was the reference 305 306 group when these dummy variables were inserted into the meta-regression model. We added two dummy covariates for exercise intensity as it was a categorical variable with three levels (low-, 307 moderate-, and high-intensity). The first, low-intensity, reflected the low-intensity versus others 308 309 contrast (coded low = 1, moderate- and high-intensity = 0) while the second, moderate-intensity, reflected the moderate-intensity versus others contrast (coded moderate = 1, low- and high-310 311 intensity = 0). The reference group was high-intensity when these dummy variables were entered 312 to the meta-regression model. We included two dummy covariates for exercise duration because 313 it was a categorical variable with three levels (short duration, medium duration, and long 314 duration). The first, medium duration, reflected the medium duration versus others contrast 315 (coded medium = 1, short duration and long duration = 0) while the second, long duration,

reflected the long duration versus others contrast (coded long = 1, short duration and medium
duration = 0). The short duration was the reference group when these dummy variables were
entered into the meta-regression model. Exercise modality (0 = treadmill-based walking/running,
1 = cycling) and cognitive task type (0 = non-PFC-dependent task, 1 = PFC-dependent task)
were categorical variables with two levels. We had also intended to use age as a moderator for
Aim 1, but we were unable to include this moderator in the analysis due to a lack of studies
(described below).

The 'robumeta' function utilizes the method of moments estimator to estimate  $\tau^2$ 323 (Thompson & Sharp, 1999). This estimator and its degrees of freedom were modified for small 324 sample sizes, as suggested by Tipton (2015). Nevertheless, robust variance estimation with small 325 326 sample adjustment remains biased (i.e., increased type I error rate) when the adjusted degrees of 327 freedom are less than 4 (Tanner-Smith & Tipton, 2014). Consequently, despite our plans to look 328 at sex, age, exercise duration, and exercise modality as moderators of the ESs for Aim 2, we 329 were unable to perform moderator analyses due to a lack of studies in each category (i.e., fewer than five studies). Thus, in Aim 2, only the cognitive task type was included in the moderator 330 331 analyses. Finally, using the 'metafor' package in R (Viechtbauer, 2010), Egger's regression test 332 for funnel plot asymmetry (Egger et al., 1997) and Duval and Tweedie's trim and fill method 333 (Duval & Tweedie, 2000) were conducted to evaluate potential risk of publication bias across 334 studies. First, Egger's regression test was performed to examine the relationship between the observed ESs and their corresponding standard errors. If a significant result for the Egger's 335 336 regression test indicates funnel plot asymmetry, publication bias may be present. Next, in the presence of publication bias, Duval and Tweedie's trim and fill analysis is generally carried out 337 to obtain the estimated number of missing studies from a meta-analysis in the funnel plot. This 338

339	method augments the observed data, which in turn makes the funnel plot more symmetric and
340	adjusts the observed average ES (Rodgers & Pustejovsky, 2020). A two-sided $p < .05$ was
341	considered statistically significant.
342	Results
343	Retrieved Articles
344	Figure 1 displays the flow chart of the literature search process. The computerized
345	searches yielded 585 articles. Furthermore, 13 additional articles were identified in reference lists
346	of key review articles on acute exercise and cognition (Chang et al., 2012; Loprinzi et al., 2019;
347	Roig et al., 2013; Tomporowski, 2003) as well as transient hypofrontality papers (Dietrich, 2003,
348	2006; Dietrich & Audiffren, 2011; Dietrich & Sparling, 2004). Among the 598 articles, 176 were
349	eliminated due to duplication and 422 articles were screened. After initial screening of 422 titles
350	and abstracts, 69 full text articles were reviewed. Among these 69 articles, 47 were ineligible as
351	they did not meet the inclusion criteria (e.g., acute exercise not occurring during the cognitive
352	task, not directly comparing a non-exercise control group to an exercise group, and/or not
353	providing sufficient information for an ES calculation). As a result, 22 studies met our eligibility
354	for the systematic review via the computerized searches, and thus, a total of 22 studies were
355	chosen for the meta-analysis.
356	[Insert Figure 1 Here]
357	Article Synthesis
358	Detailed information on the study characteristics is displayed in Table 2. Sample sizes
359	ranged from 8 to 79 participants, with the majority of studies (95%) testing young adults (18-30
360	years). Among the 22 studies, 18 (82%) employed a within-subject design.
361	[Insert Table 2 Here]

#### 362 Study Quality

Based on the modified Downs and Black checklist, the methodological quality of the included studies was fair to good ( $18.74 \pm 1.82$ , mean  $\pm$  SD), ranging from 16 to 23 (Hooper et al., 2008). All studies were scored within an acceptable range of methodological quality and thus, were included in the quantitative meta-analysis.

#### 367 **Quantitative Analysis**

Figure 2 displays the weighted average ESs with robust variance estimation for the 368 studies comparing cognitive task performance during acute exercise of all intensities with 369 370 performance during a control condition. Among these 22 studies, 75 ESs were calculated. As illustrated by Figure 2 and Table 3, there was no statistically significant effect of acute in-task 371 exercise of all intensities on cognitive function ( $\beta = -0.16, 95\%$  CI: -0.58, 0.27, p = .45). 372 Between-study heterogeneity was 0.66 ( $\tau^2$ ) with approximately 87% ( $I^2 = 87.18$ ) of variance 373 374 attributable to systematic error. Table 3 shows the results of the moderation analyses for the 375 studies comparing acute exercise of all intensities during a cognitive task vs. control group. No significant moderation effects were observed, indicating that sex, exercise intensity, exercise 376 377 duration, exercise modality, and cognitive task type did not play a moderating role in the acute 378 exercise-cognition associations under the dual-task protocols. The Egger's regression test for 379 funnel plot asymmetry was not statistically significant (z = 0.45, p = .65), indicating that there 380 was no evidence of publication bias across studies.

381

# [Insert Figure 2 Here]

382

#### [Insert Table 3 Here]

Figure 3 displays the weighted average ESs with robust variance estimation for the studies comparing cognitive task performance during high-intensity acute exercise with

385	performance during a control condition. Among these 12 studies, 27 ESs were calculated. As
386	illustrated by Figure 3 and Table 4, there was no statistically significant effect of high-intensity
387	acute in-task exercise on cognitive function ( $\beta = 0.03$ , 95% CI: -0.41, 0.46, $p = .89$ ). Between-
388	study heterogeneity was 0.46 ( $\tau^2$ ) with approximately 80% ( $I^2 = 79.82$ ) of variance attributable
389	to systematic error. Table 4 shows the results of the moderation analyses for the studies
390	comparing high-intensity acute exercise during a cognitive task vs. control group. Significant
391	moderation effect was observed for cognitive task type ( $\beta = -0.81, 95\%$ CI = -1.60, -0.02, p
392	= .04), indicating that PFC-dependent cognition was impaired during high-intensity acute
393	exercise compared to non-PFC-dependent cognition. The Egger's regression test for funnel plot
394	asymmetry was statistically significant (z = 3.51, $p < .001$ ), suggesting that there was evidence of
395	publication bias across studies. After applying the Duval and Tweedie's trim and fill methods
396	(see Figure 4), all four estimated missing studies were located to the left of the average ES and
397	the observed outcomes were distributed symmetrically in the funnel plot ( $z = 0.95$ , $p = .34$ ).
398	Further, a negative adjustment to the ES was shown and decreased to $-0.2$ (SE = 0.13).
399	[Insert Figure 3 Here]
400	[Insert Table 4 Here]
401	[Insert Figure 4 Here]
402	Discussion
403	The present meta-analysis evaluates the transient hypofrontality theory by investigating
404	the impact of acute exercise on PFC-dependent cognition. In alignment with our hypothesis as
405	well as the transient hypofrontality theory, our meta-analysis demonstrates high-intensity acute
406	exercise had a selective effect on cognition, in that PFC-dependent cognition was compromised
407	while exercising at a high intensity compared to non-PFC-dependent cognition.

Several theories have been proposed to test these effects, namely arousal, attention, and 408 cognitive-energetic theories (Tomporowski & Qazi, 2020). Although there is some theoretical 409 410 overlap among these theories, each theory highlights a specific concept. First, arousal theory hypothesizes that in a dose-dependent inverted U-shaped fashion (Yerkes & Dodson, 1908), 411 exercise-induced arousal influences cognition and cognitive task performance is improved with 412 413 moderate, but not high, levels of arousal (McMorris & Hale, 2015). On the basis of hypotheses drawn from the arousal theory, a meta-analysis by Lambourne and Tomporowski (2010) reported 414 415 that the negative ESs on information-processing tasks were observed while engaging in exercise 416 protocols designed to assess the impacts of the inverted-U hypothesis. Second, attention theory dictates that attention is a focusing process that plays an important role in cognitive function 417 (Jonides et al., 2008). The basic assumption of attention theory is that attention acts as a gate, 418 which determines what information enters into consciousness (Tomporowski & Qazi, 2020). In 419 420 the dual-task paradigm, motor-related cognitive interference (i.e., while motor performance 421 remains stable, cognitive performance deteriorates; Plummer et al., 2013) may occur due to greater attentional allocation toward physical movement. Next, cognitive-energetic theory 422 suggests that when multicomponent tasks compete for available energetical resources, more 423 424 metabolic resources may be allocated to one task that optimizes behavioral actions (e.g., exercise) and the other(s) is likely to be impaired or unaffected (e.g., cognitive performance) 425 426 depending on the dual-task workloads.

Potential mechanisms of the exercise-induced cognitive impairment effect have been
discussed elsewhere (Audiffren, 2016; Dietrich & Audiffren, 2011), which align with the
cognitive-energetic theory discussed above. Briefly, acute changes in cerebral blood flow (CBF)
induced by exercise may be closely associated with cognitive fluctuations as a result of

alterations in regional neuronal activation and metabolism in the brain. For example, during 431 acute moderate-intensity exercise, CBF increases in response to neuronal activity and 432 433 metabolism, whereas during acute high-intensity exercise, regional levels of CBF (e.g., frontal cortex) progressively decrease despite global levels of blood flow, metabolism, and oxygen 434 uptake to the brain remaining stable (Ide & Secher, 2000; Ogoh & Ainslie, 2009). This high-435 436 intensity exercise-induced decrease in CBF lowers partial pressure of arterial carbon dioxide and the total cardiac output rate to the brain, and thus, metabolic demands of the brain may not be 437 438 fulfilled (Ogoh & Ainslie, 2009; Smith & Ainslie, 2017). Moreover, based on the neurovascular 439 coupling, local neural activation of brain structures involved in physical exercise facilitates an elevation of CBF in the motor cortex, whereas neural deactivation in other local brain areas, not 440 involved in exercise, leads to a reduction of CBF in the PFC. As such, reduced CBF may 441 attenuate cerebral metabolism in the prefrontal areas while exercising at a high intensity and 442 443 ultimately may compromise PFC-dependent cognition (Mekari et al., 2015). This, however, is in 444 contrast to a recent publication (Komiyama et al., 2020), which demonstrated that a decline in CBF is not a major factor in impairment of executive function during acute vigorous-intensity 445 exercise. Specifically, CBF restoration through  $CO_2$  inhalation did not prevent executive 446 447 degradation due to acute intense exercise. Although it was different from their expected results, we cannot rule out the plausibility that CBF is not related to exercise-induced cognitive 448 449 impairment given the methodological limitations of this study (e.g., they did not directly measure 450 changes in CBF in the prefrontal regions); thus, this is an area where future research is needed. 451 Alternatively, from a neurochemical perspective, it is possible that the mechanisms of 452 such a debilitating effect may be due to modulation of select neurotransmitters, such as 453 dopamine (DA) and noradrenaline (NA). Plasma concentrations of catecholamine

neurotransmitters in the brain released under high arousal conditions play a significant role in 454 brain networks that are particularly involved in cognitive function (McMorris et al., 2016). 455 456 Studies in human (Dalsgaard et al., 2004) and animals (Hattori et al., 1994; Kitaoka et al., 2010) provide suggestive evidence that arousal induced by moderate-intensity exercise induces an 457 acute increase in firing of the high affinity  $\alpha_{2A}$ -adrenoreceptors by NA, which helps to strengthen 458 459 neuronal signals in the target stimuli by inhibiting cyclic adenosine monophosphate (cAMP) activity (Deutch & Roth, 1990; Roth et al., 1988). Likewise, the high affinity  $D_1$ -receptors by 460 461 DA reduces neuronal noise by inhibiting the firing of non-target stimuli (Finlay et al., 1995). 462 This improvement of signal-to-noise ratio may help individuals to facilitate effective encoding of the stimuli, discrimination, and decision-making processes (Berridge & Waterhouse, 2003). In 463 consequence, modest elevations in DA and NA via moderate-intensity exercise may activate 464 prefrontal neuronal networks and further enhance PFC-dependent cognition (McMorris, 2016). 465 On the other hand, excessively elevated levels of DA and NA may deteriorate cognitive 466 467 ability during a single bout of heavy exercise. Extreme stimulation of  $\alpha_1$ -adrenoreceptors may lead to reduced neuronal firing in the PFC and too high levels of  $D_1$ -receptors and  $\beta$ -468 adrenoreceptors may responsible for greater cAMP activation (Arnsten, 2011). These may cause 469 470 hyperpolarization, which inhibits action potentials by closing voltage-gated Na<sup>+</sup> channels and opening of nearby  $K^+$  channels (Arnsten, et al., 2012). While hyperpolarized, the neuron is in a 471 472 period of physiological refractory, which may prevent the neuron from generating subsequent 473 action potentials (Becker et al., 2009; Pack, 2011). Hence, these effects may lead to synaptic inhibition that lessens the likelihood that a postsynaptic neuron will fire and may contribute to a 474 475 temporal reduction in neural activity in the prefrontal areas, and thereby PFC-dependent 476 cognition may be impaired (Arnsten, 2009, 2011; Cooper, 1973; McMorris et al., 2016). Another

notable plausibility linking altered catecholamines and cognitive impairment during exposure to 477 acute stress (e.g., a single bout of strenuous exercise) can be explained from Arnsten's Dynamic 478 479 Network Connectivity (DNC) mechanisms (Arnsten et al., 2010). Arnsten and her colleagues posited that DNC is a form of neuroplasticity that can rapidly vary the strength of PFC network 480 connections depending on momentary alterations in arousal state. Under maximal arousal 481 482 conditions, excessive release of DA and NA through  $D_1$ -receptors and  $\beta$ -adrenoreceptors, respectively, may induce activation of cAMP signaling (Ramos et al., 2005; Vijayraghavan et al., 483 2007). These actions may increase opening HCN channels<sup>1</sup>, thereby weakening PFC network 484 connectivity and PFC-dependent cognition (Wang et al., 2007). In a similar manner, high levels 485 of NA release by  $\alpha_1$ -adrenoreceptors may activate Ca<sup>2+</sup>/PKC<sup>2</sup> signaling, which could facilitate 486 the loss of dendritic spines in the PFC, and thus, suppress PFC neural activity (Birnbaum et al., 487 2004). As a result, precise control of DNC in prefrontal regions is likely to play a key part in 488 prevention of cognitive deficits. As most of these findings came from animal studies, future 489 490 research in human subjects calls on us to identify novel neurophysiological mechanism(s) to explain as to why PFC-dependent cognitive dysfunction is usually observed during higher 491 intensity acute exercise. 492 493 Furthermore, these impairment effects may be explained from neurophysiological-based arousal and cognitive-energetic theories as discussed above, which also may impact 494 495 psychological attention-related mechanisms. Exercise-induced arousal has long been regarded as 496 a means of improving attention. The amount of available attentional resources depends on the

497 arousal level, which in turn, is determined by the intensity of exercise and task demand

<sup>&</sup>lt;sup>1</sup> Hyperpolarization-activated cyclic nucleotide-gated channels are integral membrane proteins, which act as nonselective voltage-gated cation channels in the plasma membranes of brain cells.

<sup>&</sup>lt;sup>2</sup> Protein kinase C is a family of enzymes that their activity is controlled by  $Ca^{2+}$  or diacylglycerol. It works for controlling signal transduction pathway of other proteins, playing a critical part in intracellular signaling.

(Kahneman, 1973). In the inverted U-shape fashion, if arousal reaches its peak through high-498 intensity acute exercise, this may elicit peripheral and central mental fatigue (McMorris & Hale, 499 2015). Further, this may result in a narrowing of attention, and thus, even task-related cues may 500 be missed, ultimately impairing cognitive performance (Easterbrook, 1959). Further, in spite of 501 PFC-dependent tasks requiring higher levels of attention capability, additional resources (i.e., 502 503 mental effort) may not be allocated toward top-down processing in the PFC, but toward the supplementary motor cortex for maintaining dynamic physical movement. Thus, and although 504 505 speculative, reduced attention may mediate impaired PFC-dependent cognition during high-506 intensity acute exercise. In contrast to the PFC-dependent cognition, maximal arousal level induced by high-intensity acute exercise is unlikely to impair non-PFC-dependent cognition 507 because stimulus-driven and automatized tasks may be less sensitive to attentional resources 508 (i.e., low attentional cost and low mental workload) and less dependent on the prefrontal 509 510 functioning (Audiffren et al., 2008; Dietrich & Audiffren, 2011).

511 The current study has several strengths. This is the first meta-analytic review that evaluates the cognitive impairment effect observed during acute exercise based on the transient 512 hypofrontality theory. This meta-analysis is significant in that it will help future work to 513 514 reconcile existing inconsistencies as to how and why acute exercise impacts cognitive decline in dual-task protocols. Second, we also tested multiple moderating variables to identify the cause of 515 516 heterogeneity regarding the detrimental effects of acute in-task exercise on cognition. Third, our 517 meta-analytic approach addressed the issue of non-independent ESs that can be found in 518 outdated meta-analytic techniques by employing robust variance estimation in meta-regression 519 model. Multiple ESs that emanate from the same study can be problematic because the correct 520 sampling variance for that aggregate is unknown for each study, and thus, it makes the standard

error calculations underestimated and loses statistical power particularly for moderators (Hedges
et al., 2010). Although there are appropriate techniques for handling this problem, several
previous meta-analysis publications that investigated the relationship between acute exercise and
cognitive function have relied on traditional meta-analytic methods without consideration of
such measurement errors (Chen et al., 2020; Jung et al., 2020). Given the characteristics of
cognitive function research (i.e., multiple ESs are generated from a single study), future metaanalysis on this topic should handle this issue carefully.

Despite these strengths, there are some limitations worth considering. First, several 528 moderation effects should be carefully interpreted due to the limited number of studies within 529 each category. In future research, the identification of additional moderators (e.g., mood, 530 exercise preferences, hypoxia, hypoxia with exercise, and contextual factors related to cognition 531 encoding) will provide insight as to under what circumstances such detrimental effects of in-task 532 exercise on cognitive function occurs. Relatedly, it would be also noteworthy for subsequent 533 534 studies to consider the degree of PFC activation in each PFC-dependent cognitive task as one of the moderators. Some tasks classified as "prefrontal-dependent" require higher demands on 535 prefrontal activity while others do only mildly so. In spite of being aware of this, it was difficult 536 537 to evaluate how much PFC activation occurs across the different PFC-dependent tasks due to limited research directly comparing the degree of PFC activation across different tasks within the 538 539 same individuals. As such, more work is needed in this area. Second, considering that the 540 threshold for high-intensity exercise varies substantially across research protocols (McMorris, 541 2015), further work should assess participant's maximum heart rate, oxygen consumption or power output using metabolic measurements to better individualize exercise intensity. Third, we 542 543 found evidence of publication bias for high-intensity acute exercise studies; however, we

adjusted for this using the trim and fill method. It typically occurs when the publication of 544 research results depend not only on the hypothesis tested but also the significance and direction 545 546 of effects detected (Dickersin, 1990). Accordingly, findings showing evidence of publication bias in meta-analyses should be cautiously interpreted and require special attention in future 547 work. Fourth, other cognitive performance indexes not used in this study may play an important 548 549 role in validating the transient hypofrontality theory. As an example, the interference cost 550 (incongruent RT – congruent RT) in select PFC-dependent tasks (e.g., Flanker task, Simon task, 551 or Stroop task) may be more sensitive to inhibitory control than raw RT (reaction time) data. 552 Given that employing raw RT contains a larger variance, which is more related to speed of information processing rather than executive functioning, additional studies on this topic should 553 be careful in choosing the appropriate performance indexes. Another limitation of this field (not 554 555 our meta-analysis) is a lack of neural mechanistic measures for the observed PFC-dependent cognitive impairment effects during acute high-intensity exercise. Future research should 556 557 evaluate which prefrontal areas are activated when high-intensity acute exercise occurs during PFC-dependent and non-PFC-dependent cognitive tasks. Functional near-infrared spectroscopy 558 or electroencephalogram (EEG) can measure the temporal changes in neuronal activation within 559 560 broad regions of the PFC. For instance, with EEG, P3 and/or N2 amplitude can be altered across frontal and lateral electrode sites during stimulus engagement (e.g., stationary cycling) (Grego et 561 562 al., 2004). If these event related potentials observed through EEG data can provide useful 563 information on how exercise-induced prefrontal activation/deactivation influences the cognitive 564 task type (PFC-dependent vs. non-PFC-dependent), this may shed additional insight into the 565 neurophysiological mechanisms of the exercise-related cognitive impairment effect.

566	In conclusion, our meta-analysis demonstrates that PFC-dependent cognition is likely to
567	be impaired when high-intensity acute exercise occurs during a cognitive task, which provides
568	support for the transient hypofrontality theory. Further work is needed to determine which
569	mechanism(s) underlies PFC-dependent cognitive impairment observed during high-intensity
570	acute exercise (e.g., Dietrich's prefrontal deactivation, Arnsten's DNC, or a combination of both
571	mechanisms).
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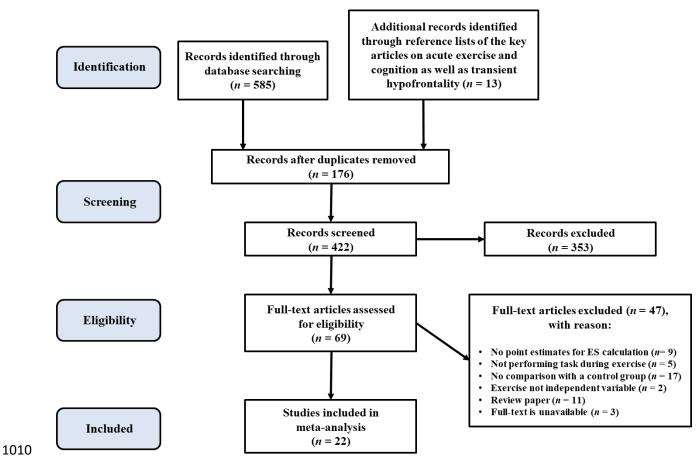


Figure 1. PRISMA flowchart of the evaluated studies and final number of included studies in a 

meta-analysis. 

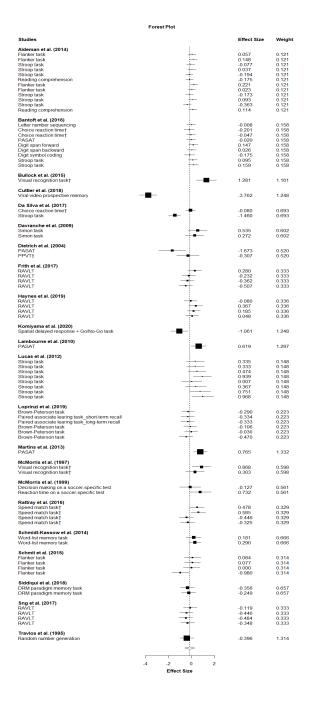


Figure 2. Forest plot depicting the effect size estimates for acute exercise studies including all
intensities. The black boxes symbolize the point estimates from each study. The white diamond
symbolize s the pooled estimates result. The horizontal lines symbolize the length of the 95%
confidence intervals of the study result. † Non-PFC-dependent cognitive task.

## Forest Plot

Studies		Effect Size	Weight
Bullock et al. (2015) Visual recognition task†		1.281	1.550
Da Silva et al. (2017)			
Choice reaction time†	<b></b>	-0.080	0.971
Stroop task	_ <b>_</b>	-1.460	0.971
Frith et al. (2017)			
RAVLT		0.280	0.459
RAVLT		-0.232	0.459
RAVLT		-0.362	0.459
RAVLT		-0.507	0.459
Komiyama et al. (2020) Spatial delayed response + Go/No-Go task	— <b>—</b>	-1.061	1.712
	—		
<b>Lambourne et al. (2010)</b> PASAT	<b>↓■</b>	0.619	1.782
Lucas et al. (2012)			
Stroop task		0.333	0.389
Stroop task	<b>-</b>	0.939	0.389
Stroop task		0.367	0.389
Stroop task		0.968	0.389
Loprinzi et al. (2019)			
Brown-Peterson task		-0.290	0.465
Paired associate learning task_short-term recall		-0.334	0.465
Paired associate learning task_long-term recall Brown-Peterson task		-0.333 -0.470	0.465 0.465
DIOWIFFEIEISON lask	-	-0.470	0.405
McMorris et al. (1997)	-	0.969	1 507
Visual recognition task†		0.868	1.597
McMorris et al. (1999)	_		
Decision making on a soccer-specific test		-0.127 0.732	0.753 0.753
Reaction time on a soccer-specific test	-	0.732	0.755
Rattray et al. (2016)	_		
Speed match task†		0.478	0.902
Speed match task†		-0.446	0.902
Schmit et al. (2015)			
Flanker task	<b>}_</b>	0.084	0.427
Flanker task Flanker task		0.077 0.000	0.427 0.427
Flanker task	<b>-</b> ]	-0.980	0.427
Travilag at al. (4005)			
Travlos et al. (1995) Random number generation		-0.396	1.808
		-0.550	1.000
	-2 -1 0 1 2 3		
	Effect Size		

- **Figure 3.** Forest plot depicting the effect size estimates for high-intensity acute exercise studies.
- 1022 The black boxes symbolize the point estimates from each study. The white diamond symbolize s
- the pooled estimates result. The horizontal lines symbolize the length of the 95% confidence
- 1024 intervals of the study result. † Non-PFC-dependent cognitive task.

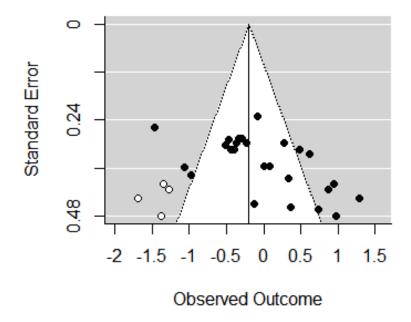


Figure 4. Funnel plot obtained with the Duval and Tweedie's trim and fill methods of the effect
size estimates for high-intensity acute exercise studies. Closed circles represent the observed
studies included in the meta-analysis. Open circles represent the estimated, imputed studies by
suggested by trim and fill analysis. The reference line indicated the adjusted mean effect size.

Database	Search Query				
PubMed	(("exercise"[mh] OR (exercise*[tiab])) AND (("Cognitive function"[mh] OR				
Fubilited	cognitive*[tiab])) AND (("hypofrontality"[mh] OR (hypofrontality*[tiab]))				
Saapua	TITLE-ABS-KEY(((Exercise OR exercise*))) AND (((Cognitive function OR				
Scopus	cognitive*))) AND ((((Hypofrontality OR hypofrontality*)))				
PsycINFO	(Exercise / OR exercise*).ab,kf,ti.) AND (Cognitive function / OR cognitive*))).ab,kf,				
FSycinfO	ti.) AND (hypofrontaility / OR hypofrontality*))).ab,kf,ti.)				
Google	Exercise   Acute exercise   Cognitive function   Hypofrontality				
Scholar	Exercise   Acute exercise   Cognitive function   Hypotrontanty				
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SPORTDiscus	function+ OR TI (cognitive*)) OR AB (cognitive*))) AND (MH Hypofrontality OR				
	TI (hypofrontality*)) OR AB (hypofrontality*)))				

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Study	Subject Characteristics	Study Design	Exercise Protocol	Cognitive Function Assessment	Mean (SD)	Results
Travlos & Marisi (1995)	<i>n</i> = 20, Age = 23.0 (3.0)	Within-subject	Cycle ergometer exercise for 50-min at high-intensity of 80% of heart rate reserve (HRR)	Random number generation	<sup>ns</sup> EX: 68.77 (5.81) Ctrl: 70.86 (4.46)	There were no differences in cognitive performance across the control and exercise conditions.
McMorris & Graydon (1997)	<i>n</i> = 12, Age = 20.8 (1.34)	Within-subject	Cycle ergometer exercise with progressive load (0%, 70%, 100% MPO)	Visual recognition task† (1. 100% MPO_RT) (2. 70% MPO_RT)	1.* EX: 776 (167) Ctrl: 923 (160) 2.* EX: 872 (165) Ctrl: 923 (160)	Performance during maximal exercise was significantly better than in the other two conditions.
McMorris et al. (1999)	n = 9, Age = 22.0 (2.4)	Within-subject	Cycle ergometer exercise with progressive load (0%, anaerobic threshold, 100% MPO)	Decision making on a soccer- specific test (1. ACC) Reaction time on a soccer- specific test <sup>†</sup> (2. RT)	1. <sup>ns</sup> EX: 8.22 (1.48) Ctrl: 8.44 (1.81) 2.* EX: 1.32 (0.20) Ctrl: 1.47 (0.19)	There was no significant effect of exercise on accuracy, but speed of decision to be significantly affected by exercise.
Dietrich & Sparling (2004)	n = 8, Age = 25.1 (6.3)	Within-subject	Running for 65-min at 70- 80% of maximum heart rate	PASAT (1. ER) PPVT <sup>†</sup> (2. ER)	1.* EX: 52 Ctrl: 32 2. <sup>ns</sup> EX: 13 Ctrl: 12	For the PASAT, exercise condition committed more errors than control group. For the PPVT, exercise did not affect cognitive performance.
Davranche & McMorris (2009)	<i>n</i> = 12, Age = 32 (9)	Within-subject	Cycle ergometer exercise for 21 minutes at moderate- intensity	Simon task (1. RT congruent) (2. RT incongruent)	1.* EX: 305 (33) Ctrl: 323 (32) 2.* EX: 344 (42) Ctrl: 355 (36)	Response time was better when the cognitive task is performed simultaneously with exercise.
Lambourne et al. (2010)	<i>n</i> = 19, Age = 21.1 (1.7)	Within-subject	Cycle ergometer exercise for 40-min at high-intensity of 90% VO <sub>2peak</sub>	PASAT	<sup>ns</sup> EX: 0.94 (0.02) Ctrl: 0.93 (0.01)	PASAT scores did not change when comparing performance during exercise and non-exercise conditions.
Lucas et al. (2012)	n = 22 (13 young adults) (9 older adults), Age = 24 (5), Age = 62 (3)	Within-subject	Two 8-min bouts of cycle ergometer exercise at 30% followed by 70% HRR	Stroop task (1. Younger: Simple task RT at 30% HRR) (2. Younger: Simple task RT at 70% HRR) (3. Younger: Difficult task RT at 30% HRR) (4. Younger: Difficult task RT at 70% HRR) (5. Older: Simple task RT at 30% HRR) (6. Older: Simple task RT at 70% HRR) (7. Older: Difficult task RT at 30% HRR) (8. Older: Difficult task RT at 70% HRR)	1. <sup>ns</sup> EX: 633 (119) Ctrl: 677 (135) 2. <sup>ns</sup> EX: 635 (108) Ctrl: 677 (135) 3.* EX: 1060 (215) Ctrl: 1157 (180) 4.* EX: 978 (189) Ctrl: 1157 (180) 5. <sup>ns</sup> EX: 830 (172) Ctrl: 829 (84) 6. <sup>ns</sup> EX: 787 (129) Ctrl: 829 (84) 7.* EX: 1434 (176) Ctrl: 1603 (247) 8.* EX: 1371 (208) Ctrl: 1603 (247)	Difficult-task response times on the Stroop task improved during exercise, with the enhancement greater at high intensity than at rest and low intensity.
Martins et al. (2013)	<i>n</i> = 24, Age = 20.5 (0.89)	Between-subject	Cycle ergometer exercise at moderate-intensity	PASAT	* EX: 89 (9) Ctrl: 82 (9)	For the PASAT, the exercise group outperformed the control group.

## **Table 2.** Extraction table of evaluated studies

Alderman et al. (2014)	<i>n</i> = 66, Age = 21.06 (1.6)	Within-subject	Self-paced treadmill walking for 15 minutes.	Flanker task (1. ACC congruent) (2. ACC incongruent) (3. RT congruent) (4. RT incongruent) Stroop task (5. ACC neutral) (6. RT neutral) (7. ACC interference word) (8. RT interference word) (9. ACC interference color) (10. RT interference color) (10. RT interference color) Reading comprehension (11. ACC) (12. RT)	$\begin{array}{c} 1.* \ {\rm EX:} \ 98.73 \ (2.82) \\ {\rm Ctrl:} \ 97.58 \ (6.76) \\ 2.* \ {\rm EX:} \ 93.29 \ (5.63) \\ {\rm Ctrl:} \ 93.13 \ (8.14) \\ 3.* \ {\rm EX:} \ 318.39 \ (59.31) \\ {\rm Ctrl:} \ 322.10 \ (69.31) \\ 4.* \ {\rm EX:} \ 380.40 \ (60.16) \\ {\rm Ctrl:} \ 390.63 \ (76.13) \\ 5.* \ {\rm EX:} \ 95.14 \ (5.43) \\ {\rm Ctrl:} \ 95.97 \ (4.03) \\ 6.* \ {\rm EX:} \ 672.19 \ (115.84) \\ {\rm Ctrl:} \ 663.34 \ (111.39) \\ 7.* \ {\rm EX:} \ 94.08 \ (10.84) \\ {\rm Ctrl:} \ 92.91 \ (13.91) \\ 8.* \ {\rm EX:} \ 808.26 \ (195.11) \\ {\rm Ctrl:} \ 815.88 \ (213.17) \\ 9.* \ {\rm EX:} \ 83.60 \ (23.57) \\ {\rm Ctrl:} \ 90.96 \ (16) \\ 10.* {\rm EX:} \ 932.56 \ (215.98) \\ {\rm Ctrl:} \ 892.84 \ (189.98) \\ 11.* {\rm EX:} \ 76.72 \ (14.18) \\ {\rm Ctrl:} \ 75.22 \ (11.92) \\ 12.* {\rm EX:} \ 3310.85 \ (757.89) \\ {\rm Ctrl:} \ 3452.12 \ (849.13) \\ \end{array}$	There were no significant differences in response speed or accuracy between walking and sitting conditions for any of the cognitive tests.
Schmidt- Kassow et al. (2014)	<i>n</i> = 49, Age = 21.7 (2.7)	Within-subject	Self-selected walking pace for 30-min	Recall memory task using 40- item (unfamiliar) word list (1. Experiment 1) (2. Experiment 2)	1.* EX: 5.5 (3.3) Ctrl: 4.8 (4.2) 2.* EX: 5.3 (4.6) Ctrl: 4.1 (3.5)	Experiment 1 & 2: Words recalled during walking was higher than non-exercise.
Bullock et al. (2015)	<i>n</i> = 12, Age = 20 (1.08)	Within-subject	Cycling for 50-min at low and high intensities	Visual recognition task <sup>†</sup> (RT)	* EX: 505 Ctrl: 525	Behavior target detection was faster during high-intensity exercise compared to rest.
Schmit et al. (2015)	<i>n</i> = 15, Age = 22.1 (0.6)	Within-subject	Cycling for 20-min at high- intensity of 85% MAP	Flanker task (1. RT congruent) (2. ER congruent) (3. RT incongruent) (4. ER incongruent)	1. <sup>ns</sup> EX: 410 (49) Ctrl: 414 (43) 2. <sup>ns</sup> EX: 0.7 (2.1) Ctrl: 0.7 (1.7) 3.* EX: 456 (76) Ctrl: 461 (46) 4.* EX: 20.6 (14.2) Ctrl: 9.9 (4.9)	The frequency of errors was lower during rest than when exercising for incongruent trials, but not for congruent trials.
Bantoft et al. (2016)	<i>n</i> = 45, Age = 22.6 (6.2)	Within-subject	Low intensity treadmill walking	Letter number sequencing (1. ACC) Choice reaction time† (2. ACC) (3. RT) PASAT (4. ACC) Digit span forward (5. ACC) Digit span backward	1. <sup>ns</sup> EX: 11.47 (2.55) Ctrl: 11.49 (2.35) 2. <sup>ns</sup> EX: 91.56 (4.75) Ctrl: 91.79 (4.87) 3. <sup>ns</sup> EX: 466.16 (60.44) Ctrl: 455.13 (47.36) 4. <sup>ns</sup> EX: 37.45 (14.83) Ctrl: 37.76 (15.23) 5. <sup>ns</sup> EX: 11.73 (2.20) Ctrl: 11.40 (2.24)	There were no significant differences in cognitive tasks across three conditions (sitting vs. standing vs. walking).

				<ul> <li>(6. ACC)</li> <li>Digit symbol coding</li> <li>(7. ACC)</li> <li>Stroop task</li> <li>(8. ER)</li> <li>(9. RT)</li> </ul>	6. <sup>ns</sup> EX: 7.28 (2.56) Ctrl: 7.22 (1.98) 7. <sup>ns</sup> EX: 88.91 (15.46) Ctrl: 91.60 (14.99) 8. <sup>ns</sup> EX: 0.38 (0.68) Ctrl: 0.49 (0.69) 9. <sup>ns</sup> EX: 17.39 (5.94) Ctrl: 17.92 (5.09)	
Rattray & Smee (2016)	<i>n</i> = 20, Age = 26.6 (5.2)	Within-subject	Cycle ergometer exercise for 50-min at low- and high-intensity of 50% VO <sub>2max</sub> and 90% VO <sub>2max</sub> .	Speed match task <sup>†</sup> (1. Low-intensity EX ACC) (2. Low-intensity EX RT) (3. High-intensity EX ACC) (4. High-intensity EX RT)	1.* EX: 93.6 (5.9) Ctrl: 95.3 (4.2) 2.* EX: 643 (52) Ctrl: 672 (45) 3.* EX: 93.1 (5.4) Ctrl: 95.3 (4.2) 4.* EX: 648 (53) Ctrl: 672 (45)	The exercise conditions facilitated response time, but reduced accuracy scores compared to control condition.
Da Silva et al. (2017)	n = 37 Age = 25.1 (4.6)	Within-subject	Cycle ergometer exercise for 20 minutes at high- intensity	Choice reaction time† (1. RT) Stroop task (2. ER)	1.* EX: 322.2 (261.7) Ctrl: 305.2 (138.2) 2.* EX: 1.7 (1.2) Ctrl: 0.3 (0.6)	Exercise at higher intensities was negatively associated with reaction time and inhibitory control.
Frith et al. (2017)	<i>n</i> = 88, Age = 21.9 (2.4)	Between-subject	15 minutes treadmill walking/running of progressive high-intensity aerobic exercise	RAVLT (1. Trial 1 recall) (2. 20 min delayed recall) (3. 24h recognition recall) (4. 24h attribution recall)	1.* EX: 6.55 (1.14) Ctrl: 6.18 (1.44) 2.* EX: 8.86 (2.51) Ctrl: 9.41 (2.13) 3.* EX: 35.23 (4.0) Ctrl: 36.59 (3.36) 4.* EX: 31.68 (5.09) Ctrl: 33.95 (3.57)	High-intensity exercise before memory encoding was effective in improving long-term memory, 20- min delayed recall, and 24-h delayed recall.
Sng et al. (2017)	<i>n</i> = 88, Age = 23.3 (3.7)	Between-subject	15 minutes self-paced treadmill walking at moderate-intensity	RAVLT (1. Trial 1_recall) (2. 20 min delay recall) (3. 24h recognition recall) (4. 24h attribution recall)	1.*         EX: 6.45 (1.63)           Ctrl: 6.64 (1.50)           2. <sup>ns</sup> EX: 10.23 (2.20)           Ctrl: 11.23 (2.20)           3.*         EX: 34.09 (3.44)           Ctrl: 35.95 (4.08)           4.*         EX: 31.73 (4.36)           Ctrl: 33.36 (4.84)	Exercising prior to memory encoding was superior for improving learning, 24-h memory recognition, and 24-h memory attribution.
Cuttler et al. (2018)	<i>n</i> = 120, Age = 19.77 (0.15)	Between-subject	30-min self-paced treadmill walking/running at moderate-intensity	Viral video prospective memory	* EX: 14 (0.94) Ctrl: 17.59 (0.95)	The control group had better prospective memory than the exercise group.
Siddiqui & Loprinzi (2018)	<i>n</i> = 20, Age = 21.1 (1.0)	Within-subject	20-min treadmill bout of self-paced brisk walking (moderate-intensity exercise)	Deese-Roediger-McDermott (DRM) paradigm (1. Immediate memory recall) (2. Delayed memory recall)	1.* EX: 8.20 (1.6) Ctrl: 8.80 (1.7) 2.* EX: 5.90 (1.7) Ctrl: 6.40 (2.2)	For both short-term and long-term memory, the exercise group during memory encoding resulted in the worst memory performance.
Haynes et al. (2019)	n = 24, Age = 20.9 (1.9)	Within-subject	Self-selected brisk walking pace for 15 minutes at moderate-intensity	RAVLT (1. Trial 1 recall) (2. 20 min delayed recall) (3. 24h recognition recall) (4. 24h attribution recall)	1.* EX: 6.0 (1.7) Ctrl: 6.13 (1.5) 2. <sup>ns</sup> EX: 11.2 (2.9) Ctrl: 10.04 (3.3) 3. <sup>ns</sup> EX: 22.48 (4.2) Ctrl: 22.26 (4.8)	Short-term memory was superior in the visit that involved exercise before the memory task. Similar outcomes occurred for long-term memory, but there were no

					4. <sup>ns</sup> EX: 19.08 (5.0) Ctrl: 17.96 (6.8)	exercise effects on prospective memory.
Loprinzi et al. (2019)	<i>n</i> = 48, Age = 21.9 (1.9)	Within-subject	Treadmill exercise at low (30% HRR), moderate (50% HRR), and high- intensity (80% HRR)	Experiment 1 Brown-Peterson task (1. Memory recall) Paired associate learning task (2. Short-term recall) (3. Long-term recall) Experiment 2 Brown-Peterson task (4. Memory recall at low- intensity EX) (5. Memory recall at moderate-intensity EX) (6. Memory recall at high- intensity EX)	1.* EX: 10.75 (3.41) Ctrl: 11.67 (2.79) 2.* EX: 3.67 (2.88) Ctrl: 4.71 (3.23) 3.* EX: 3.50 (2.84) Ctrl: 4.50 (3.06) 4. <sup>ns</sup> EX: 9.41 (3.9) Ctrl: 9.83 (3.9) 5. <sup>ns</sup> EX: 9.70 (4.5) Ctrl: 9.83 (3.9) 6.* EX: 7.66 (5.1) Ctrl: 9.83 (3.9)	Experiment 1: Both working memory and episodic memory were compromised during high- intensity exercise when compared to rest. Experiment 2: Similar results occurred for working memory, but memory function was not impaired during low- and moderate-intensity exercise.
Komiyama et al. (2020)	<i>n</i> = 17, Age = 22.1 (1.7)	Within-subject	Cycle ergometer exercise for 15 minutes at high- intensity of 80% VO <sub>2peak</sub>	Spatial delayed response + Go/No-Go task	* EX: 84.1 (13.3) Ctrl: 95.1 (5.3)	Accuracy of the cognitive tasks was impaired during high-intensity exercise in the exercise group relative to the control group.

*Note*. ACC = accuracy, ER = error rate, EX = exercise condition, Ctrl = control, HRR = heart rate reserve, MAP = maximal aerobic power, MPO = maximum power output, ms = millisecond, PASAT = paced auditory serial addition task, PPVT = peabody picture vocabulary test, RAVLT = rey auditory verbal learning task, RT = reaction time, SD = standard deviation,  $VO_{2max}$  = maximum oxygen uptake, † non-PFC-dependent cognitive task. All reaction time units are milliseconds. \* significantly different, <sup>ns</sup> not significantly different

Variable	k	0	b	SE	95% CI		р	<i>t</i> ( <i>df</i> )	Heterogeneity	
					LL	UL			$ au^2$	$I^2$
Cognitive function	22	75								
Intercept only									0.66	87.18
Constant			-0.16	0.20	-0.58	0.27	.45	-0.77 (20.9)		
Moderators									0.77	87.46
Constant			0.21	0.61	-1.17	1.59	.74	0.34 (9.17)		
Males			-0.40	0.69	-2.05	1.25	.58	-0.57 (6.70)		
Predominantly female			-0.38	0.52	-1.89	1.14	.51	-0.72 (4.63)		
Low-intensity exercise			0.58	0.51	-0.60	1.75	.29	1.14 (7.58)		
Moderate-intensity exercise			-0.01	0.51	-1.16	1.14	.98	-0.02 (8.71)		
Medium duration exercise		-0.87	0.88	-3.00	1.26	.36	-0.99 (6.27)			
Long duration exercise		-0.44	0.72	-2.19	1.31	.56	-0.61 (6.37)			
Cycling		0.95	0.61	-0.44	2.34	.16	1.56 (8.41)			
PFC-dependent cognitive task			-0.66	0.33	-1.48	0.16	.10	-1.99 (5.77)		

**Table 3**. Weighted average effects with robust variance estimation and moderation analyses for acute exercise at all intensities during cognitive task vs. control

*Note*. k = number of studies, o = total number of comparisons, b = coefficient in the meta-regression model, SE = standard error of the coefficient, 95% CI = 95% confidence interval for the coefficient, LL = lower limit of the 95% confidence interval for the coefficient, UL = upper limit of the 95% confidence interval for the coefficient, t = t-statistic calculated based on the predicted mean, df = small sample corrected degrees of freedom of the distribution of the *t*-statistic,  $\tau^2 =$  between-study sampling variance,  $I^2 =$  proportion of effect size variance due to between-sample heterogeneity, PFC = prefrontal cortex.

Variable	k	0	b	SE	95% CI		р	<i>t</i> ( <i>df</i> )	Heterogeneity	
					LL	UL			$ au^2$	$I^2$
Cognitive function	12	27								
Intercept only									0.46	79.82
Constant			0.03	0.20	-0.41	0.46	.89	0.14 (10.9)		
Moderators									0.37	76.01
Constant			0.57	0.29	-0.27	1.41	.13	1.97 (3.60)		
PFC-dependent cognitive task			-0.81*	0.33	-1.60	-0.02	.04	-2.46 (6.71)		

**Table 4**. Weighted average effects with robust variance estimation and moderation analyses for high-intensity acute exercise during cognitive task vs. control

*Note.* k = number of studies, o = total number of comparisons, b = coefficient in the meta-regression model, SE = standard error of the coefficient, 95% CI = 95% confidence interval for the coefficient, LL = lower limit of the 95% confidence interval for the coefficient, UL = upper limit of the 95% confidence interval for the coefficient, t = t-statistic calculated based on the predicted mean, df = small sample corrected degrees of freedom of the distribution of the *t*-statistic,  $\tau^2 =$  between-study sampling variance,  $I^2 =$  proportion of effect size variance due to between-sample heterogeneity, PFC = prefrontal cortex. \*p < .05.