

1 **Title:**

2 Preparation for mental effort recruits Dorsolateral Prefrontal Cortex: an fNIRS
3 investigation

4 **Abbreviated title:** DLPFC activity during preparation for mental effort

5

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17 arithmetic

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28 **Abstract**

29 Preparing for a mentally demanding task calls upon cognitive and motivational
30 resources. The underlying neural implementation of these mechanisms is receiving
31 growing attention, given the implications for professional, social, and medical
32 contexts. While several fMRI studies converge in assigning a crucial role to a cortico-
33 subcortical network including Anterior Cingulate Cortex (ACC) and striatum, the
34 involvement of Dorsolateral Prefrontal Cortex (DLPFC) during mental effort
35 anticipation has yet to be replicated. This study was designed to target DLPFC
36 contribution using functional Near Infrared Spectroscopy (fNIRS), as a more cost-
37 effective tool measuring cortical hemodynamics. We adapted a validated mental effort
38 task, where participants performed easy and difficult mental calculation, while
39 measuring DLPFC activity during the anticipation phase. As hypothesized, DLPFC
40 activity increased during preparation for a hard task as compared to an easy task.
41 Besides replicating a previous fMRI study, these results establish fNIRS as an
42 effective tool to investigate cortical contributions to preparation for effortful behavior.
43 This is especially useful if one requires testing large samples (e.g., to target individual
44 differences), populations with contraindication for functional MRI (e.g., infants or
45 patients with metal implants), or subjects in more naturalistic environments (e.g.,
46 work or sport).

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53 **Introduction**

54 Humans face cognitively challenging situations on a daily basis.

55 Accomplishing such tasks requires a great deal of cognitive resources, and typically

56 successful completion is facilitated by the possibility of gaining a reward. Preparing

57 for demanding tasks and anticipating possible rewards are core components of

58 motivation. Several studies investigating cost-benefit trade-offs in decision-making

59 showed that these motivational processes rely on a cortical-subcortical brain network,

60 involving the medial Prefrontal Cortex (MPFC, including dorsal Anterior Cingulate

61 Cortex, dACC) and striatum (Chong et al., 2017; Prévost, Pessiglione, Météreau,

62 Cléry-Melin, & Dreher, 2010; Westbrook & Braver, 2013, 2015). Interestingly, some

63 studies demonstrated that these regions are also implicated in preparing for effortful

64 performance (unconfounded by motor or decision-making factors), showing increased

65 neural activity when preparing for a harder task. For example, this is the case when

66 participants prepare for upcoming mentally demanding arithmetic problems (Vassena

67 et al., 2014) or perceptual discrimination (Krebs, Boehler, Roberts, Song, & Woldorff,

68 2012). This evidence is often interpreted as indexing proactive control, i.e. top-down

69 deployment of attentional control to ensure successful performance (Braver, 2012).

70 Recently, computational frameworks have been proposed where MPFC activity would

71 reflect the value of engaging in an effortful task to the extent that it can lead to a

72 reward (Holroyd & McClure, 2015; Holroyd & Yeung, 2012; Shenhav, Botvinick, &

73 Cohen, 2013; Verguts, Vassena, & Silvetti, 2015) or the monitoring processes

74 detecting the frequency of occurrence of motivationally relevant variables (Vassena,

75 Deraeve, & Alexander, 2017). Notwithstanding the different computational

76 implementation, all accounts agree in assigning to MPFC a crucial role in mechanisms

77 underlying effortful behavior.

78 The role of the Dorsolateral Prefrontal Cortex (DLPFC) in preparing for
79 cognitively demanding tasks is however less clear. DLPFC activity is generally
80 implicated in higher level cognitive processing (Miller & Cohen, 2001), such as
81 working memory updating, goal maintenance and task set representation. According
82 to recent theories, DLPFC indeed maintains abstract information about task-related
83 rules, instructions or context (Alexander & Brown, 2015; Badre, 2008; Koechlin &
84 Summerfield, 2007; Nee & Brown, 2012). One recent study showed that MPFC
85 coding of reward expectation seems to drive strategy selection in DLPFC, which in
86 turn regulates MPFC activity (Domenech, Redouté, Koechlin, & Dreher, 2017). This
87 dynamic provides theoretical support for DLPFC role in learning how to deploy
88 control when reward is available. Therefore, as DLPFC and MPFC interact in guiding
89 strategy-selection according to reward prospect, comparable dynamics may be
90 hypothesized in the context of preparing for a more effortful task.

91 In an earlier study using functional MRI (fMRI) we provided preliminary
92 evidence that preparing for an effortful task relies on a network also implicated in
93 reward expectation (Vassena et al., 2014). In this study we also showed that DLPFC
94 was more active when expecting to perform a mentally effortful task (as compared to
95 an easy task). The goal of the current study was to independently replicate
96 anticipation of effort in DLPFC with a novel and promising measurement technique.
97 The use of functional Near-Infrared spectroscopy (fNIRS) is rapidly growing in
98 cognitive and social neuroscience (Balconi & Vanutelli, 2017), as it allows measuring
99 cortical variations in regional blood oxygenation levels in a comparable way to fMRI,
100 but without a number of downsides that MRI has. In particular, fNIRS technology
101 does not involve a strong magnetic field nor gradients. As a consequence,
102 contraindications for participation due to the magnetic field do not apply.

103 Furthermore, the fNIRS machinery (and its use) has a much lower cost than an MRI
 104 machine. Because of these two reasons, one can test a larger sample of participants,
 105 with lower cost, including patients and other subjects with (non MR-compatible)
 106 metal implants, children, and babies (who normally do not undergo fMRI strictly for
 107 research purposes), and in more ecological context (as the equipment is portable,
 108 Ayaz et al., 2013; Balardin et al., 2017). Finally, motion artifacts are less problematic
 109 with fNIRS, which makes it an interesting tool to test hypotheses in domains where
 110 movement is required (Metzger et al., 2017; Pinti et al., 2015); a relevant example in
 111 the current context would be physical effort (where participants are normally required
 112 to move to exert force). One final noteworthy advantage is that subjects can be tested
 113 simultaneously and while interacting, making it an ideal tool for social neuroscience
 114 experiments (Balconi & Vanutelli, 2017).

115 Exploiting these advantages to investigate cortical contributions to preparation
 116 for mental effort requires establishing fNIRS as a reliable measurement method of
 117 cortical (prefrontal) activity, by replicating cortical hemodynamic effects observed
 118 with fMRI. We therefore adopted fNIRS to investigate the contribution of bilateral
 119 DLPFC in preparation for effortful tasks. We adapted a mental effort task from
 120 previous studies (Vassena et al., 2014; Vassena, Cobbaert, Andres, Fias, & Verguts,
 121 2015). Participants were presented with cues indicating if the upcoming task was
 122 going to be easy or hard. We measured oxygenated hemoglobin dynamics in 26
 123 measurement channels covering frontal cortex. Moreover, we tested whether DLPFC
 124 sensitivity to task demand during preparation was bilateral or lateralized to one
 125 hemisphere. For this purpose, we controlled participants' handedness, as previous
 126 studies show that left handers are more likely to present opposite or reduced

functional lateralization as compared to right-handers (e.g. in language, Mazoyer et al., 2014; and in face processing, Frässle, Krach, Paulus, & Jansen, 2016).

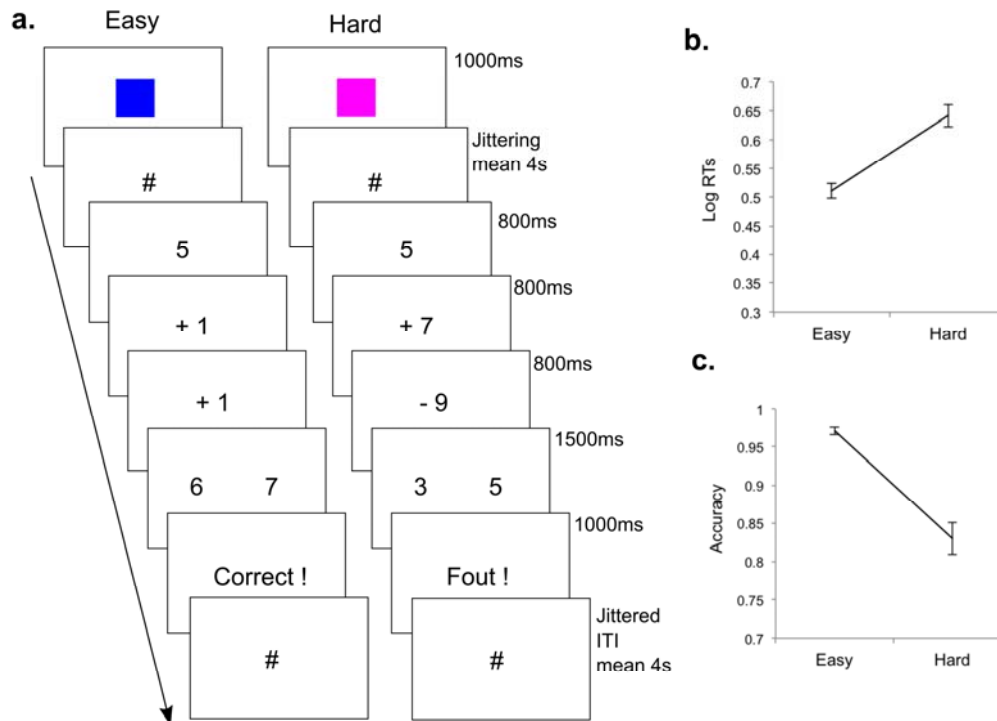
Materials and methods

Participants

Twenty undergraduate students from Ghent University participated in this study (mean age 20.1 ± 2.74 years, 13 females, 9 left handed), receiving one study credit as compensation to participate in the study. Written informed consent was obtained from all participants prior to participation. The study protocol was approved by the Local Ethics Committee of Ghent University. After data collection, one participant was excluded from further analysis due to technical failure. Sample size was determined based on previous studies using fNIRS to investigate cognitive function (Causse, Chua, Peysakhovich, Del Campo, & Matton, 2017; Ferreri et al., 2014; Nakahachi et al., 2008).

Experimental procedure

We examined difficulty-related hemodynamic cortical activation while participants performed a task consisting of easy and difficult arithmetic calculations (Figure 1).



146

147 **Figure 1.** Task and behavioral performance. a.Task structure. Each trial started with a cue
148 indicating if the upcoming task was going to be easy (blue square) or difficult (magenta
149 square). After a jittered interval, two subsequent operations (additions or subtractions) were
150 presented on the screen, followed by two possible results. Participants had to indicate the
151 response they thought to be correct by pressing right or left response button, and received
152 performance feedback. Subsequent inter-trial interval was also jittered. b. Log-transformed
153 reaction times for responses in easy as compared to hard trials. c. Accuracy of the responses
154 for easy as compared to hard trials. Error bars represent $\pm$ one standard error of the mean.
155 Responses to easy trials were significantly more accurate and faster as compared to hard trials.
156

157 The procedure consisted of one block with 130 trials, of which 65 were easy
158 trials and 65 were difficult trials. Easy and difficult trials were randomly intermixed.
159 At the beginning of every trial, a cue was presented for 1000 ms, indicating if the
160 upcoming trial was going to be easy (a blue square) or difficult (a magenta square),
161 followed by a screen showing the symbol # at fixation with a pseudo-exponentially
162 jittered duration (range 2.2 – 8 seconds, mean 4 seconds). Subsequently, the task was
163 presented. In an easy trial, the task consisted of a sequence of two arithmetic
164 operations, with three small numbers shown on subsequent screens (e.g., example 3 +
165 1 + 1). Each number remained on the screen for 800 ms, and first and second number

166 were followed by a blank screen (600 ms). In a difficult trial, the task consisted of a
167 sequence of more difficult arithmetic operations with three larger numbers shown on
168 subsequent screens (e.g. $8 + 15 - 6$, same timing as easy trials) and requiring carrying
169 and borrowing at each operation. We adapted this procedure from previous
170 experiments, as it elicits a reliable difficulty effect (Vassena et al., 2014, 2015). After
171 the arithmetic problem, two possible results were presented on the screen, and
172 participants had to select the result they thought to be correct, by pressing either a left
173 or a right button (F or J on the keyboard, response time limit 1500 ms). The response
174 was followed by a feedback screen, which could be correct (showing the Dutch word
175 “correct”), incorrect (showing the Dutch word “fout”) or too late (showing the Dutch
176 words “te laat”). The feedback was followed by a 500 ms blank screen, and a pseudo-
177 exponentially jittered inter-trial interval, with a screen showing the # symbol at
178 fixation (range 2.2 – 8 seconds, mean 4 seconds). Participants were instructed to be as
179 fast and accurate as possible. Before starting the experimental block, 8 training trials
180 were administered. During the training only, at the end of every trial participants were
181 asked to rate the trial on perceived difficulty and pleasantness. The questions were
182 presented on the screen one by one (randomized across participants) and participants
183 were asked to respond by pressing the number corresponding to their response on the
184 keyboard. The difficulty question asked how difficult that trial was for them (on a
185 visual 7-point scale, with 1 meaning very easy and 7 meaning very difficult). The
186 pleasantness question asked how much they liked to perform that trial (on a visual 7-
187 point scale, with 1 meaning not at all and 7 meaning very much). This procedure has
188 been used in previous studies to confirm subjective perception of difficult trials as
189 more difficult (Vassena et al., 2014, 2015).

190

191 *Questionnaires*

192 Participants filled in the Positive And Negative Affect Scales (PANAS,
193 (Watson, Clark, & Tellegen, 1988) twice, before and after the experimental session.
194 The goal of this procedure was to measure changes in affective state, and test whether
195 such changes may be related to task difficulty (by testing the correlation with
196 accuracy and reaction times at the task). At the end of the session, participants also
197 filled in the Need for Cognition scale (short version, (Cacioppo, Petty, & Kao, 1984),
198 assessing how much participants enjoy engaging in mentally demanding endeavors,
199 and the BisBas scale (Carver & White, 1994), assessing participants' behavioral
200 inhibition and activation tendencies.

201

202 *fNIRS methods*

203 We used the continuous-wave NIRS system (NIRScout; NIRx Medical
204 Technologies, Brooklyn, NY) utilizing two wavelengths of near-infrared light (760
205 and 850 nm). Data were acquired from the prefrontal cortex with 5 sources and 5
206 detectors per hemisphere, covering the lateral and medial PFC. The distance between
207 each source-detector pair was 3 cm, which provides an adequate compromise between
208 depth sensitivity and signal to noise ratio (G. E. Strangman, Li, & Zhang, 2013). An
209 a-priori DLPFC region-of-interest (ROI) was anatomically determined, by visual
210 inspection of optode locations projected on a 3D MNI atlas (Figure 2, Okamoto et al.,
211 2004), and by convergence of such locations with previously reported DLPFC activity
212 in a comparable task (Vassena et al., 2014). The DLPFC-ROI included the channels
213 F5-F3, F5-FC5, FC3-F3 and FC3-FC5 (for each hemisphere, note that in each pair the
214 first label is the sender, and the second label is the receiver). This approach had one
215 main limitation: our procedure did not include neuronavigation with the subject-

specific MRI scan, thus preventing from projecting specific MNI coordinates to the subject's adapted cortical coordinates. However, given the spatial resolution of the current fNIRS setup, and the large extent of DLPFC activity reported in above-mentioned studies, it seems plausible that a finer anatomical characterization would be difficult to achieve (and not necessary for the current purpose). Figure 2 shows the channel configuration.

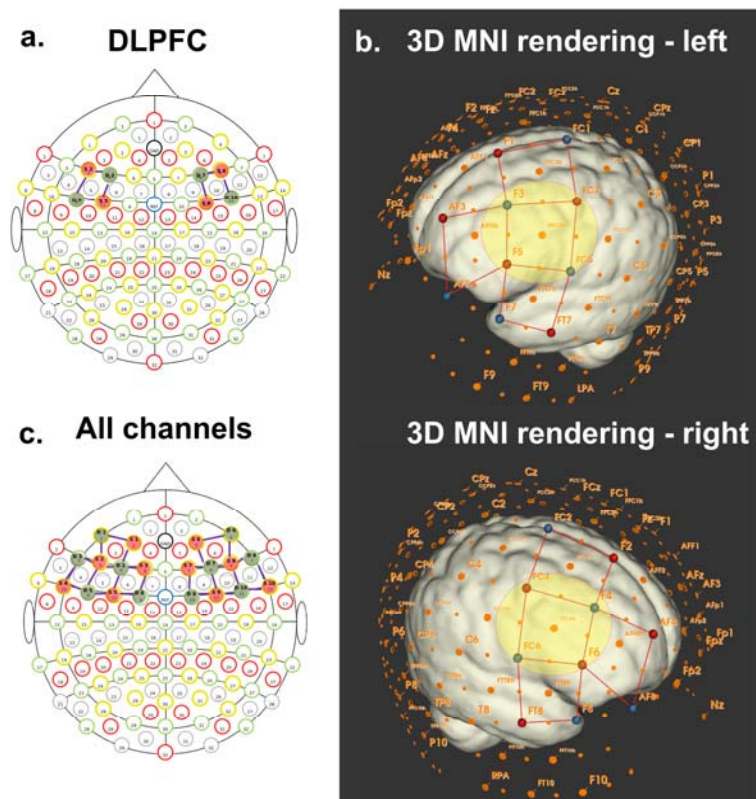


Figure 2. fNIRS setup a. fNIRS montage visualized on the 10-20 EEG template. Selected optodes and channels covering DLPFC. b. Whole montage visualized on a 3D rendering of MNI space, with optode coordinates projected on the cortex. The yellow circles highlight the channels included in the DLPFC-ROI. c. Whole montage visualized on the 10-20 EEG template.

233 **Data analysis**

234 *Behavioral data analysis*

235 A paired-sample t-test was performed on log-transformed reaction times
236 (RTs), comparing easy vs. difficult trials. A second paired-sample t-test was
237 performed, comparing accuracy in the easy vs. difficult trials. Reported significance
238 values are two-tailed.

239 We calculated the difference between positive and negative PANAS scores by
240 subtracting scores at the beginning of the session to the scores at the end of session.
241 Subsequently, these differences for positive and negative PANAS separately were
242 correlated with accuracy and RTs at the task, to test potential influences of difficulty
243 (as measured by indexes of task performance) on affective state. Performance was
244 also correlated with difficulty and liking ratings. Furthermore, we calculated Need for
245 Cognition and Bis and Bas scores, to test the relationship between task performance
246 and attitude towards mental effort and behavioral inhibition and activation. One
247 should note that these correlational analyses were exploratory in nature.

248

249 *fNIRS data preprocessing*

250 We analyzed the optical data using Homer2 NIRS processing package
251 functions (Huppert, Diamond, Franceschini, & Boas, 2009) based on MATLAB
252 (Mathworks, MA USA). For every participant, the raw optical intensity data series
253 were converted into changes in optical density (OD). Then PCA was performed,
254 which automatically adjusts the amount of variance to be removed from the data on a
255 subject-by-subject basis. A PCA parameter of 80% was chosen as it is more
256 conservative and removes only the variance supposed to account for the motion
257 artifacts (Brigadoi et al., 2014). Then a motion detection algorithm was applied to the

258 OD time series to identify residual motion artifacts (AMPthresh=0.5, SDThresh=50,
259 tMotion=0.5 s, tMask=1s). This means that if, over a temporal window of length 0.5 s,
260 the standard deviation increases by a factor exceeding 50, or the peak-to-peak
261 amplitude exceeds 0.5, then the segment of data of length 1 s starting at the beginning
262 of that window is defined as motion. Stimuli with artifacts from the HRF calculation
263 were excluded if any artifact appeared 5 seconds before the stimulus appearance, and
264 10 seconds after. Low-pass filtering with a cut-off frequency of 0.5 Hz was applied to
265 the data in order to remove variability due to the cardiac cycle. The OD data were
266 then converted into concentration changes using the modified Beer-Lambert law
267 (Cope & Delpy, 1988; Delpy et al., 1988) with the differential path length factors set
268 to 0.6. This method enabled us to calculate signals reflecting the oxygenated
269 hemoglobin (OxyHb), deoxygenated hemoglobin (DeoxyHb), and total hemoglobin
270 (Total Hb) signal changes. Afterwards, to recover the mean hemodynamic response
271 we solved the GLM based on ordinary least squares (Ye, Tak, Jang, Jung, & Jang,
272 2009), modeling the HRF with a modified gamma function convolved with its
273 derivative and 3rd order polynomial for drift correction. Statistics were done outside
274 Homer with in-house written scripts in Matlab. Note that we performed all further
275 analysis on the OxyHb signal. Most fNIRS studies focus on OxyHb, as previous
276 research showed that this signal correlates more robustly with the fMRI-BOLD signal
277 in several tasks, possibly due to a higher signal-to-noise ratio as compared to
278 DeoxyHb (Hoge et al., 2005; Huppert, Hoge, Diamond, Franceschini, & Boas, 2006;
279 Mehagnoul-Schipper et al., 2002; Okamoto, Dan, Shimizu, et al., 2004; G.
280 Strangman, Culver, Thompson, & Boas, 2002).

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fNIRS Statistical analysis

We used a mixed linear modeling (MLM) approach (Baayen, Davidson, & Bates, 2008, also) to analyze the relation between the peak OxyHb and task difficulty. All analyses were performed using the package lme4 (version 1.1-13) in R version 3.3.1.

Two sets of analyses were performed one on the data averaged within the DLPFC-ROI, and one on all channels. In all analyses, we followed a model building procedure. In a first step, we estimated a benchmark model (including only a random intercept for channels nested into participants, see Jasinska & Petitto, 2013 for a similar approach) to account for between-subjects variability in OxyHb concentration changes across channels. Next, three more complex models were created by expanding the baseline model with one of the fixed effects for Difficulty (Easy vs Difficult), Handedness (Left- vs Right-handed) or Hemisphere (Left vs Right). Each expanded model was compared to the benchmark model using a likelihood ratio test (significance level of 0.05). We introduced the factor Hemisphere to test whether preparation-related activity in PFC would be bilateral or lateralized; Handedness was introduced as a control factor.

In the second step, a new benchmark model was constructed by including the random effects of the original benchmark model, plus each statistically significant fixed effect from the first step. This benchmark model was then compared to the same model plus each two-way interaction effect.

In a third step, a third benchmark model was estimated, consisting of the previous benchmark model plus all significant two-way interaction effects from the second step. This benchmark model was then compared to the same model plus the three-way interaction effect.

311 As a control, all analyses were also conducted with the first benchmark model
312 including only a random intercept for participant (thus without nesting channels
313 within participants). This control analysis returned very similar results and therefore
314 will not be reported.

315 Finally, in order to explore brain-behavioral relationships, a difference score
316 was calculated for behavioral performance indexes (RTs and accuracy) and for
317 OxyHb peak within the DLPFC ROI (difficult – easy condition). Next we computed
318 Pearson’s correlations between these measures.

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321 **Results**

322 *Behavioural results*

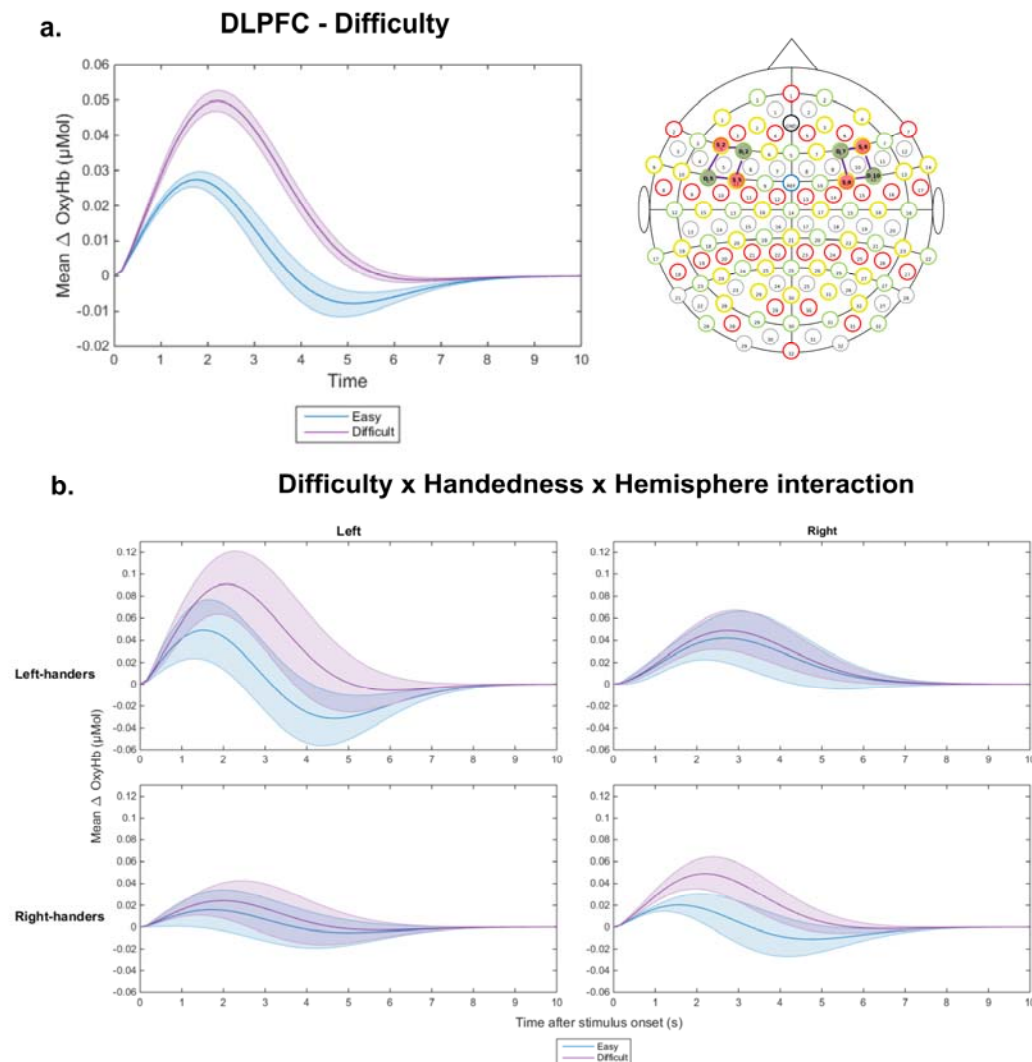
323 Prior to analysis, RTs were log-transformed. In line with previous reports,
324 participants responded faster to easy trials as compared to hard trials ($t_{(18)}=-6.56$,
325 $p<.0001$, mean difference $-.13$). Responses to easy trials were also more accurate as
326 compared to hard trials ($t_{(18)}=6.73$, $p<.0001$, mean difference $.14$), confirming
327 successful manipulation of task difficulty. Next we analyzed the ratings about task
328 difficulty and liking, given during the training.

329 Participants judged hard trials as more difficult ($t_{(18)}=2.31$, $p=.03$). Participants
330 who found hard trials more difficult also liked the easy trials more ($r=.51$, $p=.03$), and
331 showed larger RT differences between hard and easy trials performance ($r=.54$,
332 $p=.02$).

333 No significance difference was found in the liking ratings (how much
334 participants reported to like the easy as compared to the hard trials).

335 Next, we computed the results of the PANAS questionnaire, which was
 336 administered before and after the task to check participants' affective state. Both
 337 positive ($t_{(18)}=6.43$, $p<.001$) and negative affect scores ($t_{(18)}=3.29$, $p=.004$) were
 338 significantly lower after the task (possibly due to the long duration of the experiment,
 339 which lasted about a hour including set up time and task performance time).
 340 Subsequently, we performed an exploratory correlation analysis, correlating the
 341 difference between hard and easy condition in RTs and Accuracy with the difference
 342 in affective state pre- and post-task (both for negative and positive PANAS).
 343 However, no significant correlation surviving correction for multiple comparisons
 344 was observed. Performance also did not correlate with other questionnaires measures.

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 348 *fNIRS results*
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 351 Statistical significance of the results was assessed by likelihood ratio testing. χ^2 and p-
 352 values refer to comparisons between the benchmark model and the same model plus
 353 the fixed effect or interaction of interest. As the model residuals were right-skewed, a
 354 square root transformation was applied to the OxyHb.



355

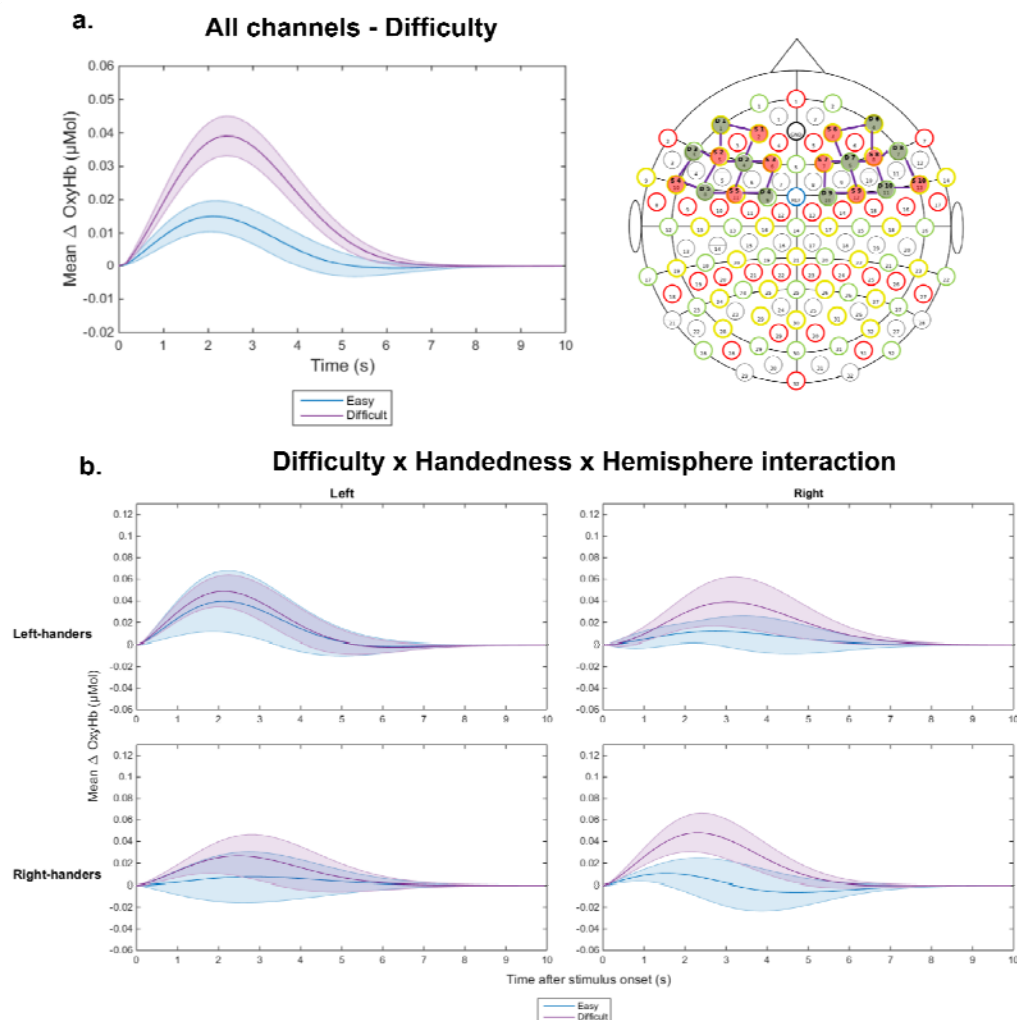
356 **Figure 3.** DLPFC fNIRS results. a. Cortical hemodynamic response of OxyHb within
 357 the DLPFC ROI time-locked with cue-onset (during task preparation) for easy (blue
 358 line) and difficult (pink line) trials. Shades around the lines represent $\pm$ standard error
 359 of the mean. b. Same as a, split by handedness (upper panels left-handers, lower
 360 panels right-handers) and hemisphere (left panels left hemisphere, right panels right
 361 hemisphere). In all plots shades around the lines represent $\pm$ one standard error of the
 362 mean.

363

364 The analysis on the DLPFC-ROI revealed a main effect of difficulty ($\chi^2(1)=6.63$,
 365 $p=.010$). Specifically, the average OxyHb peak was higher in the hard condition than
 366 in the easy condition (Figure 3). None of the other main effects demonstrated
 367 significance (Handedness: $\chi^2(1)=.07$, $p=.790$; Hemisphere: $\chi^2(1)=3.16$, $p=.075$). A

368 significant Handedness $\times$ Hemisphere interaction was also observed ($\chi^2(3)=8.85$,
369 $p=.031$), reflecting increased activity in the left DLPFC of left-handers. Again, no
370 other interaction effects were found to be significant (Difficulty $\times$ Handedness:
371 $\chi^2(2)=3.67$, $p=.159$; Difficulty $\times$ Hemisphere: $\chi^2(2)=.15$, $p=.923$). The three-way
372 interaction (Difficulty $\times$ Handedness $\times$ Hemisphere) showed a trend towards
373 significance ($\chi^2(3)=7.06$, $p=.070$). This reflects the fact that the difficulty effect was
374 more pronounced in right DLPFC for right-handers and in left DLPFC for left-
375 handers. This exploratory result is consistent with previous findings on lateralization
376 and handedness, demonstrating that left-handers tend to show effects in the opposite
377 hemisphere as opposed to right-handers. However, the sample size in each group was
378 small, so this finding should be further addressed and corroborated in future studies.
379 For the model including all the channels, a significant main effect for difficulty was
380 observed ($\chi^2(1)=20.59$, $p<.001$). The OxyHb activity was higher for difficult trials
381 compared to easy trials. (Figure 4). None of the other main effects reached
382 significance (Handedness: $\chi^2(1)=.88$, $p=.349$; Hemisphere: $\chi^2(1)=2.89$, $p=.089$. A
383 significant Handedness $\times$ Hemisphere interaction was also observed ($\chi^2(1)=14.90$,
384 $p=.002$), but no three-way interaction ($\chi^2(3)=2.41$, $p=.491$). No other two-way
385 interactions reached significance (Difficulty $\times$ Handedness: $\chi^2(2)=3.33$, $p=.189$;
386 Difficulty $\times$ Hemisphere: $\chi^2(2)=2.77$, $p=.251$).

387 Finally, for completeness in Figure 5 we plot cortical hemodynamic responses
388 for all channels across the whole montage, showing OxyHb, DeoxyHb and total Hb in
389 the hard (Figure 5a) and easy condition (Figure 5b).

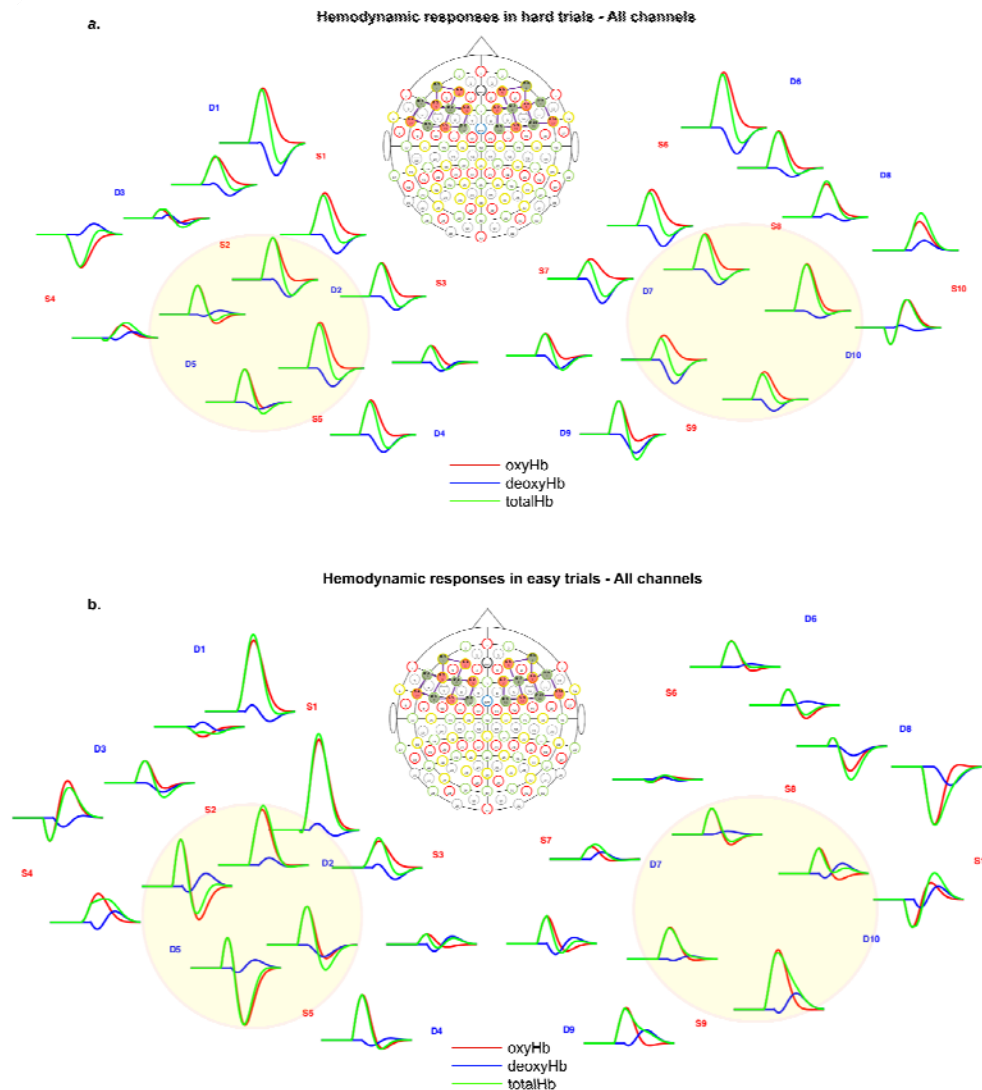


390

391 **Figure 3.** fNIRS results averaged over the whole montage (shown at top right). a.
 392 Cortical hemodynamic response of OxyHb averaged over all channels, time-locked
 393 with cue-onset (during task preparation) for easy (blue) and difficult (pink) trials. b.
 394 Same as a, split by handedness (upper panels left-handers, lower panels right-handers)
 395 and hemisphere (left panels left hemisphere, right panels right hemisphere). In all
 396 plots shades around the lines represent $\pm$ one standard error of the mean.
 397

398 Next, we correlated the difficulty effect on the OxyHb peak within the left and
 399 right DLPFC ROI (hard > easy) with difference scores of accuracy and RTs. Although
 400 no significant correlations were observed, there was a trend for a negative relationship
 401 between peak magnitude difference in Left DLPFC and RT difference ($r=-.400$,
 402 $p=.09$), and accuracy difference ($r=-.404$, $p=.086$). While this may indicate that

403 increased preparation-related activity in left DLPFC might reduce difficulty effects on
 404 performance, this relationship cannot be confirmed on the basis of this data and
 405 should be investigated in further research. No correlation was found for Right DLPFC
 406 ($r=.33$, $p=.16$).
 407



408
 409 **Figure 5.** Hemodynamic responses for all channels separately. a. OxyHb
 410 (red), DeoxyHb (blue) and total Hb (green) time-locked with cue onset in the hard
 411 trials. b. OxyHb (red), DeoxyHb (blue) and total Hb (green) time-locked with cue
 412 onset in the easy trials. In both panels the whole montage is shown. The yellow circles
 413 highlight the DLPFC-ROI channels.
 414

415 **Discussion**

416 This study investigated the role of DLPFC during preparation for a mentally
417 effortful task. Our results revealed an increase over DLPFC during expectation of a
418 mentally effortful task (difficult arithmetic problem). These results confirm a
419 contribution of DLPFC to anticipation of mental effort, as suggested by a previous
420 fMRI study (Vassena et al., 2014), and establishes optical imaging with fNIRS as a
421 non-invasive and cost-effective tool to investigate the role of DLPFC in motivation
422 for effortful behavior.

423 Growing neuroimaging evidence shows that preparing to exert mental effort is
424 associated with activity in medial PFC, especially dorsal ACC (e.g., Vassena et al.
425 2014, Parvizi, Rangarajan, Shirer, Desai, & Greicius, 2013). Few studies suggest a
426 crucial contribution of DLPFC to this process as well. For example, Vassena et al
427 (2014) and Sohn et al. (Sohn, Albert, Jung, Carter, & Anderson, 2007) found an effect
428 of task preparation prior to performing hard trials in DLPFC. McGuire & Botvinick
429 (2010) observed that MFC correlated with errors and RTs, but only DLPFC correlated
430 with avoidance ratings when performance (errors and RT) were factored out,
431 suggesting a more general role in strategic preparation.

432 Previous fNIRS studies have investigated DLPFC contribution to several
433 processes cognitive, such as word-encoding in a memory task (Ferreri et al., 2014),
434 inhibitory control in drug users (Roberts & Montgomery, 2015), dual motor and
435 cognitive task-performance (Mandrick et al., 2013). Relevant to the current results,
436 two studies targeted hemodynamic responses in DLPFC in effortful tasks, specifically
437 testing the effect of varying mental load in a working memory task (Molteni et al.,
438 2012), and comparing laboratory measures of load (executive function task) with real-
439 life effort (operating a flight-simulator Causse et al., 2017). Both studies confirm a

440 PFC contribution to the process. However, in both cases hemodynamic changes were
 441 measured *during* task performance, and not during task preparation as in our case.
 442 Interestingly, Causse and colleagues found no effects of performance on DLPFC
 443 activity, and conclude that this region may play a motivational role in sustaining effort
 444 exertion, rather than affecting task performance. Finally, a few studies also
 445 investigated the potential of fNIRS signal decoding in PFC as a Brain-Computer
 446 Interface, showing reliable decoding of brain activity during mental arithmetic as
 447 compared to rest (Bauernfeind, Steyrl, Brunner, & Muller-Putz, 2014; Bauernfeind et
 448 al., 2014; Herff et al., 2013).

449 The results of the current study thus relate to previous fNIRS evidence on
 450 DLPFC involvement during effortful tasks, showing for the first time with fNIRS a
 451 clear contribution of DLPFC to preparation for mental effort. It is however important
 452 to highlight a few limitations. First, the fNIRS montage used to measured cortical
 453 hemodynamics included only frontal channels. Although a similar approach has been
 454 adopted in several others studies (Bembich et al., 2014; Ernst et al., 2013; Laguë-
 455 Beauvais, Brunet, Gagnon, Lesage, & Bherer, 2013), other areas must be targeted in
 456 future work. In particular parietal cortex has often been found to be co-activated with
 457 DLPFC, including during preparation for mental effort (e.g., Boehler et al., 2011).
 458 Several studies suggest that DLPFC and parietal cortex form the fronto-parietal
 459 network (Dosenbach, Fair, Cohen, Schlaggar, & Petersen, 2008), implicated in
 460 attentional and top-down control, goal-directed behavior and translation of instruction
 461 to action (Buschman & Miller, 2007; Farooqui, Mitchell, Thompson, & Duncan,
 462 2012; Hartstra, Waszak, & Brass, 2012; Muhle-Karbe, Duncan, De Baene, Mitchell,
 463 & Brass, 2017). A second limitation is that the fNIRS technology only allows
 464 recording cortical activity; deeper regions such as dACC or subcortical structures

cannot be targeted. However, these regions have already been investigated in the context of preparation for mental effort (Kurniawan, Guitart-Masip, Dayan, & Dolan, 2013; Vassena et al., 2014), and therefore this caveat does not affect the relevance of the current findings. One last potential drawback concerns the interpretation of our results. We propose that DLPFC activity reflects the subject's preparation for the upcoming effortful task (i.e., as in Vassena et al., 2014). However, two alternative interpretations may be possible. This activity may reflect simply predicting the nature of the upcoming task (i.e., is it difficult or not, Vassena, Deraeve, et al., 2017), as opposed to playing a causal role in effort preparation (Brown & Alexander, 2017; Verguts et al., 2015). Alternatively, it may predict other variables related to task difficulty, such as increased error likelihood (Brown & Braver, 2005) or time-on-task (Grinband et al., 2011), although these models predict these effects in the dACC rather than DLPFC. Future designs should disentangle these possibilities, referring to available computational models of PFC function, which attempt to described the contribution of this region to task performance in a mechanistic fashion (Alexander, Vassena, Deraeve, & Langford, 2017; Koechlin, 2016). Future theorizing efforts should also consider the relative contribution of DLPFC, as opposed to dACC (Vassena, Holroyd, & Alexander, 2017) to develop novel frameworks able to describe the interaction between the two regions (Domenech et al., 2017; Kerns et al., 2004).

The conclusion that DLPFC activation is measurable via fNIRS opens up several avenues for research, given the portability and low price of fNIRS setups. In particular, it allows measurement while engaged in (or preparing for) active tasks, which is not possible with standard (e.g., EEG, fMRI) measurement protocols. This is especially of potential relevance in physical effort tasks, in which movement seems

489 almost by definition required, and for example in the emergent field of sport
490 psychology.

491 Another group of studies in which fMRI protocols are problematic, and hence
492 fNIRS is an interesting alternative, are those where measurement time is necessarily
493 long, either because a single session is long (e.g., studies on fatigue; Wang,
494 Trongnetrpunya, Samuel, Ding, & Kluger, 2016) or because several sessions must be
495 administered (e.g., studies on learning or memory). For example, concerning fatigue,
496 (Wang et al., 2016) conducted a Stroop task for 160 minutes using EEG. They
497 observed an anterior-frontal ERP that increased during the first 80 minutes of test-
498 taking; during that period, accuracy remained approximately constant. However, as
499 soon as the ERP dropped (after 80 minutes on the task), also accuracy dropped. They
500 thus interpreted the frontal ERP as a “compensation” signal, reflecting the investment
501 of cognitive effort to maintain task performance at an acceptable level (around 10%
502 errors). fNIRS could be fruitfully used to investigate the spatial localization of this
503 component more precisely. fNIRS also opens up interesting possibilities for studies
504 that require large groups. Examples include between-subject designs, studies on
505 individual differences, or studies where effect sizes are expected to be small. Due to
506 the cost of a single MRI scan, such large-group studies are typically not possible in
507 fMRI. Thus, fNIRS might provide an opportunity to better control Type-I and Type-II
508 errors in neuroimaging (Button et al., 2013). Finally, fNIRS also allows testing in
509 subjects with contra-indications for fMRI (e.g., pregnancy, non-removable ferro-
510 magnetic implants, or pacemakers, children). Based on our findings, one could
511 investigate the development of the difficulty effect in children across the life-span.
512 Finally, one could investigate preparation for effortful behavior in populations that
513 show clinically impaired motivation and effort exertion.

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798 **Conflict of interests statement:**

799 The authors have no competing interests to declare.

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