1 Overlapping attentional networks yield divergent behavioral predictions across tasks: 2 Neuromarkers for diffuse and focused attention? 3 4 Abbreviated title: Divergent predictions from attentional networks 5 Esther X.W. Wu^{a,b}, Gwenisha J. Liaw^b, Rui Zhe Goh^a, Tiffany T.Y. Chia^b, Alisia M.J. Chee^c, 6 Takashi Obana^{a,b,e}, Monica D. Rosenberg^f, B.T. Thomas Yeo^{b,c,d,g,h}, & Christopher L. 7 $Asplund^{a,b,d,e,g,h,*}$ 8 9 10 ^a Division of Social Sciences, Yale-NUS College, National University of Singapore, 16 College 11 Ave West, Singapore 138527 ^b N.1 Institute for Health, National University of Singapore, 28 Medical Drive, #05-COR, 12 13 Singapore 117456 ^c Department of Electrical and Computer Engineering, Faculty of Engineering, National 14 15 University of Singapore, Block E4, Level 5, Room 42, 4 Engineering Drive 3, Singapore 117583 ^d Clinical Imaging Research Centre, Yong Loo Lin School of Medicine, 14 Medical Drive, #B1-16 17 01, Singapore 117599 18 ^e Department of Psychology, Faculty of Arts and Social Sciences, National University of 19 Singapore, 5 Arts Link, Singapore, 117570 20 ^f Department of Psychology, University of Chicago, Chicago IL, USA 60637 21 ^g Centre for Cognitive Neuroscience, Duke-NUS Medical School, 8 College Road, Singapore 22 169857

^h Institute for Application of Learning Science and Educational Technology, National University 23 24 of Singapore, Singapore 119077 25 *To whom correspondence should be sent. Address: Yale-NUS College, 28 College Avenue 26 27 West #01-501, Singapore 138533. Tel: +65 6601 3327. Email: chris.asplund@yale-nus.edu.sg. 28 29 Acknowledgements: 30 The authors would like to thank Mike Esterman for the GradCPT experiment and analysis code. 31 This work was supported by grants from the Singapore Ministry of Defence, DSO National 32 Laboratories, and Singapore Ministry of Education and Yale-NUS College start-up, all to 33 Asplund. It was also supported by an NUS Cross-Faculty Research Grant to Yeo and Asplund. 34 The authors have no conflicts of interest, financial or otherwise, with respect to their authorship 35 or the publication of this article. 36 37 Number of pages for entire document (85), main text (52) 38 Number of figures (9), tables (3), multimedia (0) and 3D models (0) 39 Number of words for Abstract (250), Introduction (648), and Discussion (1497)

40 Abstract

41

42

43

44

45

46

47

48

49

50

51

52

53

54

55

56

57

58

59

60

61

62

Attention is a critical cognitive function, allowing humans to select, enhance, and sustain focus on information of behavioral relevance. Attention contains dissociable neural and psychological components. Nevertheless, some brain networks support multiple attentional functions. Connectome-based Predictive Models (CPM), which associate individual differences in task performance with functional connectivity patterns, provide a compelling example. A sustained attention network model (saCPM) successfully predicted performance for selective attention, inhibitory control, and reading recall tasks. Here we constructed a visual attentional blink (VAB) model (vabCPM), comparing its performance predictions and network edges associated with successful and unsuccessful behavior to the saCPM's. In the VAB, attention devoted to a target often causes a subsequent item to be missed. Although frequently attributed to attentional limitations, VAB deficits may attenuate when participants are distracted or deploy attention diffusely. Participants (n=73; 24 males) underwent fMRI while performing the VAB task and while resting. Outside the scanner, they completed other cognitive tasks over several days. A vabCPM constructed from these data successfully predicted VAB performance. Strikingly, the network edges that predicted better VAB performance (positive edges) predicted worse selective and sustained attention performance, and vice versa. Predictions from the saCPM mirrored these results, with the network's negative edges predicting better VAB performance. Furthermore, the vabCPM's positive edges significantly overlapped with the saCPM's negative edges, and vice versa. We conclude that these partially overlapping networks each have general attentional functions. They may indicate an individual's propensity to diffusely deploy attention, predicting better performance for some tasks and worse for others.

Significance statement

A longstanding question in psychology and neuroscience is whether we have general capacities or domain-specific ones. For such general capacities, what is the common function? Here we addressed these questions using the attentional blink (AB) task and neuroimaging. Individuals searched for two items in a stream of distracting items; the second item was often missed when it closely followed the first. How often the second item was missed varied across individuals, which was reflected in attention networks. Curiously, the networks' pattern of function that was good for the AB was bad for other tasks, and vice versa. We propose that these networks may represent not a general attentional ability, but rather the tendency to attend in a less focused manner.

73 Introduction

74

75

76

77

78

79

80

81

82

83

84

85

86

87

88

89

90

91

92

93

94

95

Attention is a critical cognitive function, allowing humans to select, enhance, and sustain focus on information of behavioral relevance. Visual attention plays numerous roles in different contexts, and it has been fractionated both behaviorally and neurally (Chun, Golomb, & Turk-Browne, 2011; Desimone & Duncan, 1995; Egeth & Yantis, 1997). In addition to such separable components, however, some brain networks support attentional processing across multiple domains (Asplund, Todd, Snyder, & Marois, 2010; Corbetta & Shulman, 2002; Duncan, 2010; Tamber-Rosenau, Dux, Tombu, Asplund, & Marois, 2013; Yeo, Krienen, et al., 2015). Recent studies using Connectome-based Predictive Models (CPM) support this idea. In a CPM approach, individual differences in behavioral performance are accounted for as a function of whole-brain functional connectivity patterns, after which performance for novel individuals can be predicted from fMRI data (Shen et al., 2017). Such predictions also apply across tasks. A sustained attention network model (saCPM) (Rosenberg, Finn, et al., 2016) could predict individual differences in performance for selective attention (Rosenberg, Hsu, Scheinost, Constable, & Chun, 2018), inhibitory control (Fountain-Zaragoza, Samimy, Rosenberg, & Prakash, 2019), and reading recall (Jangraw et al., 2018). Here we constructed a CPM for the visual attentional blink (VAB), aiming to test whether that model could predict performance on a variety of tasks and to compare its predictions and network features to the saCPM's. In a VAB paradigm, participants search for two items in a stream of distractors; they often fail to perceive the second item, but only when it closely follows the first (200-500 ms) (Raymond, Shapiro, & Arnell, 1992). The VAB is critically dependent on attention, as no deficit occurs when the first item is ignored. Individuals differ in their VAB severity (rate of second item detection failures), and these differences are

97

98

99

100

101

102

103

104

105

106

107

108

109

110

111

112

113

114

115

116

117

118

typically large and stable (Dale, Dux, & Arnell, 2013). It is unclear, however, which cognitive and neural factors underlie them. Numerous theoretical explanations have been proposed for the VAB, including a temporary loss of control (Di Lollo, Kawahara, Shahab Ghorashi, & Enns, 2005) or bottleneck-like processing limitations (Chun & Potter, 1995). VAB magnitude also correlates only weakly with most other attention tasks (Skogsberg et al., 2015). Intriguingly, VAB performance sometimes improves when attention to its primary detection task is reduced. Such reductions can be due to manipulation (Olivers & Nieuwenhuis, 2005, 2006) or dispositional factors (Dale & Arnell, 2010, 2015; Thomson, Ralph, Besner, & Smilek, 2015), and are thought to cause more diffuse attentional deployment. In particular, mindwandering is associated with better VAB performance, though it has the opposite association for many other attention tasks (Gonçalves et al., 2017; Hu, He, & Xu, 2012; Robertson, Manly, Andrade, Baddeley, & Yiend, 1997; Smilek, Carriere, & Cheyne, 2010). The VAB likely involves many factors (Dux & Marois, 2009), but which are reflected in individual differences of brain network function? To address this question using a CPM approach, we scanned 73 individuals while they performed the VAB task and while they rested. Resting state data allowed us to assess whether any predictive functional architecture persisted when participants were not engaged in attention-demanding tasks (Finn et al., 2015; Rosenberg, Finn, et al., 2016; Yoo et al., 2017). Outside the scanner, the same individuals completed cognitive tasks assessing sustained attention, selective attention, and fluid intelligence. We constructed a visual attentional blink CPM (vabCPM), from which we could make and assess predictions about the tasks. If attentional capacity predicts VAB performance, we would expect positive associations between vabCPM predictions for behavior and observed performance in other attention tasks. Conversely, if diffuse attentional tendencies predict VAB performance, we

might find significant *negative* associations between predicted and observed behavior for other attention tasks. For external validation, we made and assessed predictions about our tasks, including the VAB, using a sustained attention CPM (Rosenberg, Finn, et al., 2016). We then investigated and compared the networks associated with each model to better understand their relationship and potential psychological functions.

Materials and Methods

The present study included numerous tasks, with a primary focus on the visual attentional blink (VAB). Additional tasks provided critical context and points of comparison for understanding the individual differences in VAB performance and neural features. These additional tasks included those related to goal-directed attention and fluid intelligence. Many of these tasks are conceptually linked to the VAB (Dux & Marois, 2009), and many have also been studied themselves using Connectome-based Predictive Modeling (Finn et al., 2015; Rosenberg, Finn, et al., 2016; Rosenberg et al., 2018). To facilitate comparisons across these different tasks, we re-coded all behavioral performance measures such that positive numbers indicated better performance (e.g. higher accuracy or faster reaction times; see details below).

Experimental design

Participants. Eighty-two participants with self-reported normal or corrected-to-normal vision and normal hearing were recruited from the National University of Singapore (NUS) community. These individuals began a six-session study that included numerous behavioral and neuroimaging components, a subset of which are reported and analyzed here. Eight participants did not continue with the experiment after the first practice session (Session 0) and were excluded from the following analyses. One participant who did not achieve a target

143

144

145

146

147

148

149

150

151

152

153

154

155

156

157

158

159

160

161

162

163

164

discrimination score of 75% in the main VAB task (Session 1) was also excluded. Thus, unless otherwise stated, the following analyses included data from 73 participants (24 males) between the ages of 19-30 (M = 22.25, SD = 1.84). All participants provided written informed consent in accordance with a protocol approved by the NUS Institutional Review Board and received monetary compensation. Stimulus presentation. All sessions took place either inside the MR scanner or in the laboratory over a span of 3 weeks (see Table 1-1 of Extended Data for complete and detailed experimental protocol). Inside the scanner, stimuli were presented at a viewing distance of 91 cm on a 32-inch LCD monitor (NordicNeuroLab, Bergen, Norway) with a screen refresh rate of 60 Hz, connected to a MacBook Air (OS 10.12.1) running PyschoPy (Peirce, 2007). Participants made responses using an MR-compatible button box. In the laboratory, stimuli were presented at a distance of 57 cm on a 22-inch LCD monitor (Samsung SyncMaster 2233) with a screen refresh rate of 60 Hz using an NVIDIA Quadro FX 3450/4000 SD graphics card on Windows 7. Participants' responses were captured on a standard computer keyboard. Auditory stimuli were presented using PyschoPy (Peirce, 2007) binaurally through Creative headphones. Overview of task domains and specific tasks. The tasks in this study all investigated cognitive processing, primarily different forms of attention (Table 1). Each task is detailed below, organized by task domain. The task domains included the Attentional Blink, Sustained Attention, Selective Attention, and Fluid Intelligence. Table 1. Schedule of tasks and data used in analysis. Participants were encouraged to take breaks between the tasks to prevent fatigue. With the exception of the Raven's Progressive Matrices test, each task was performed twice on non-successive days. Task domains and tasks were as

follows. Attentional Blink: Visual Attentional Blink (VAB) and Auditory Attentional Blink (AAB). Sustained Attention: Visual Sustained Attention to Response Task (VSART), Auditory Sustained Attention to Response Task (ASART), and Gradual-onset Continuous Performance Task (GradCPT). Selective Attention: Attentional Network Task (ANT). Fluid Intelligence: Raven's Progressive Matrices test (Raven's).

Session	Days after previous session	Task	Data used in analysis Behavior	
0	0	VAB (practice)		
		Resting state	FMRI	
1	3 to 14	VAB (in scanner)	FMRI, Behavior	
		ANT, VSART	Behavior	
2	1	AAB, GradCPT, ASART	Behavior	
3	1	ANT, VSART, Raven's	Behavior	
4	1	AAB, GradCPT, ASART	Behavior	
5	1 to 10	VAB (in scanner)	Behavior	

Attentional Blink. Participants completed a visual attentional blink (VAB) and an auditory attentional blink (AAB) task. They were designed to be generally similar (Figure 1). Both versions were built around target discrimination and a probe detection, with each item embedded within a stream of distractors. Our main task of interest is the VAB, for which neuroimaging data was collected concurrently. For the AAB and all other tasks, only behavioral data were collected.

Visual Attentional Blink task

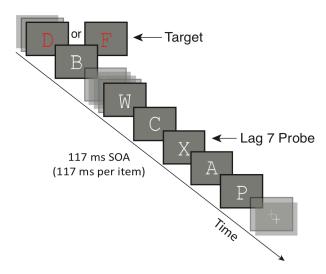


Figure 1. Experimental paradigm for the visual attentional blink (VAB) task. Participants identified a target and then detected a probe within a stream of distractors, responding when prompted at the conclusion of the stream. The target was a red letter, the probe was a white letter X, and distractors were other white letters. The auditory attentional blink (AAB) task was similar, save targets were complex tones, the probe was a high-pitched pure tone, and distractors were other pure tones. SOA = stimulus onset asynchrony.

For the VAB, stimuli consisted of upper-case letters presented in Courier New Font on a dark gray background (Figure 1). Targets were red letters D or F, whereas the probe was a white letter X. Distractors were white letters save D, F, X, I, L, O, and Q. Targets and probes were embedded within a rapid serial presentation stream, with no items repeated during each trial. The stimulus onset asynchrony (SOA) between successive items in the 16-stimulus stream was 117 ms (no gap). A target appeared during every trial at serial position 3, 4, or 5, whereas the probe appeared during 75% of trials. When present, the probe appeared at lags 1, 2, 3, 5, 7, or 9 relative

195

196

197

198

199

200

201

202

203

204

205

206

207

208

209

210

211

212

213

214

215

216

block contained both.

to the target, with the same number of trials per lag condition in each block. The first three lags were expected to be within the AB window, whereas the last three were expected to be outside it. In a small percentage of trials (14%), an irrelevant surprise stimulus (randomly selected from a set of 24 grayscale male faces and 24 colorful objects) was presented at lags 2 or 6 relative to the target. These surprise stimuli are not relevant to this current study; trials containing surprises were excluded from the following analyses and are not reported further. Each 6.25-second trial began with the presentation of a white fixation cross (0.8° x 0.8°) for 500 ms, which became larger (1.0° x 1.0°) and turned yellow to signify that the rapid serial visual presentation (RSVP) stream would begin in 750 ms. Participants searched this RSVP stream (1867 ms) for the target and probe, which they then indicated by button box press after a blank gap (233 ms) at the trial's conclusion. A maximum of 2.9 s was given for participants to respond to both the target and probe response prompts. After this period, no further responses were recorded and the white fixation cross returned until the start of the next trial. Failure to respond was rare: No session had more than 0.16% no-target response trials or 0.64% no-probe response trials averaged across participants, and no participant had more than 3.57% (target) or 4.17% (probe) no-response trials in any given session. The timing between each trial was optimized for functional Magnetic Resonance Imaging (fMRI). As such, it followed an exponential distribution with a range of 1.25-10 s and mean of 3.75 s. Each session contained six blocks of 28 trials each, with the trials presented in a pseudorandom order. The 168 trials took approximately 40 minutes to complete, including breaks. Before the main experimental blocks in each session, each participant completed 3 practice blocks of 8 trials each. The first block contained targets but no probes; the second block contained probes but no targets; and the third

218

219

220

221

222

223

224

225

226

227

228

229

230

231

232

233

234

235

236

237

238

For the AAB, targets were low-pitched or high-pitched complex tones comprised of five log-related frequencies (794 to 1260 Hz, or 1349 to 2142 Hz), whereas the probe was a 4000 Hz pure tone. Distractors were 19 pure tones of log-related frequencies ranging from 697 to 2911 Hz. Sound stimuli were adjusted to have equal mean absolute amplitudes, after which the probe and distractor intensities were set to 45% and 30% of the target intensity (~70 dB). These values were based on performance in Obana, Lim, & Asplund (under review) and additional pilot tests. As in the visual task, a small percentage of auditory trials (14%) contained an irrelevant surprise stimulus, randomly selected from a set of 24 sounds (including an alarm, a cough, and spoken letters), presented at lags 2 or 6 relative to the target. Again, surprise trials were not analyzed here. With the exception of a change in stimuli, trial structure was identical to the VAB. Each 6.25-second trial began with a reminder of the target and probe sounds, which were played for 110 ms each with a gap of 85 ms. After 750 ms, the rapid auditory stream (RAP) began, through which participants searched for the target and probe. Similar to the visual task, the identity of the target and the presence of the probe were indicated by keypress after a blank gap (233 ms). Response prompts for the target and probe then appeared, with a maximum allowed time of 2.9 s for both responses. The timing between each trial was fixed at 0.75 s. Except for the blank gap and the response prompts, a white fixation cross (0.8° x 0.8°) was shown on the screen throughout the block. Each session contained three blocks of 56 trials each, with the trials presented in a pseudorandom order. The 168 trials took approximately 25 minutes to complete, including breaks. Before the main experimental blocks in each session, each participant completed 3 practice blocks of 8 trials each. The first block contained targets but no probes; the second block contained probes but no targets; and the third block contained both.

240

241

242

243

244

245

246

247

248

249

250

251

252

253

254

255

256

257

258

259

260

261

each session.

Before beginning each practice and task block of the auditory task, participants could play the target and probe sounds as many times as desired. Sustained attention. To better understand VAB performance in relation to other forms of attention and their neural underpinnings, we also ran three sustained attention tasks. These paradigms included the visual and auditory versions of the Sustained Attention to Response Task (SART; Robertson, Manly, Andrade, Baddeley, & Yiend, 1997), which we adapted from Seli, Cheyne, Barton, & Smilek (2012), and the Gradual-Onset Continuous Performance Task (GradCPT; Esterman, Noonan, Rosenberg, & DeGutis, 2013; Rosenberg, Noonan, DeGutis, & Esterman, 2013). The SART has been frequently used to examine moment-to-moment fluctuations of sustained attention, requiring participants to make continuous responses to most stimuli but withhold responses to a few. However, due to its trial-based structure, which may provide a short 'break' between trials and not tax attention sufficiently, the GradCPT was later designed to present images that gradually transition from one to the next using a linear pixel-bypixel interpolation. The GradCPT has been shown to show reliable and large interindividual variability amongst high-functioning young adults, such as those in our sample (Rosenberg, Finn, et al., 2016; Rosenberg et al., 2013). In the VSART, participants were presented single digits, one after another, in the center of the display screen. They were asked to press the spacebar if they saw a number from 1 to 9, but withhold their response if they saw the number 3. Each digit was presented for 250 ms, followed by an encircled "x" mask for 900 ms. Digits were presented in Symbol font in white, against a black background, at sizes 0.57°, 1.03°, 1.43°, 1.89°, 2.35° of visual angle. The order of the digits and their sizes were randomized. Participants completed 675 trials (~ 13.5 mins) in

263

264

265

266

267

268

269

270

271

272

273

274

275

276

277

278

279

280

281

282

283

284

For the ASART, stimuli consisted of spoken single numbers. As in the visual task, participants were asked to press the spacebar when they heard the numbers 1 to 9, but to withhold their response if they heard the number 3. The numbers were presented in random order, each for 250 ms, following by a pink noise mask of 900 ms. Throughout the experiment, participants maintained fixation on a cross in the middle of the display. Each session consisted of 675 trials (~13.5 mins). The GradCPT (Rosenberg et al., 2013) consisted of images that gradually transitioned from one to the next using a linear pixel-by-pixel interpolation (ISI = 800 ms). Images consisted of 10 mountain and 10 city scenes, randomly presented with 10% and 90% probability, respectively, without repeats in consecutive images. Participants were instructed to press the spacebar when a city scene was presented, but to withhold their response when a mountain scene was presented. To tax sustained attention sufficiently, the GradCPT was performed in a single block over a relatively long duration (15 min). Selective attention. To understand the AB's relationship to other selective attention tasks and their neural underpinnings, we also employed the Attentional Network Task (ANT), by Fan, McCandliss, Sommer, Raz, & Posner (2002). This paradigm was designed to test three separable components of selective attention: alerting, orienting and executive control (Posner & Petersen, 1990) within a single experimental session. Stimuli consisted of 5 black lines (some with arrowheads) arranged horizontally in a row, against a grey background. The target, always an arrow in the center, was flanked on each side by (i) two arrows pointing in the same direction as the target (congruent condition), (ii) two arrows pointing in the opposite direction from the target (incongruent condition), or (iii) two black lines without arrowheads (neutral condition). Each line or arrow measured 0.55° horizontally, and the space between two adjacent objects measured

0.06°. To trigger attention orienting, all stimuli were presented either 1.06° above or below a central fixation cross. Participants were asked to keep their eyes fixated on the fixation cross and respond whether the target was pointing right or left.

Each trial started with a central fixation cross (400 – 1600 ms), followed by a warning cue (100 ms), a second central fixation cross (400 ms), and finally, the stimuli consisting of target and flankers presented either above or below a central fixation cross. Target and flankers were presented for 1700 ms, or until a response was made, whichever was shorter. A fixation cross was then presented until the end of trial (4000 ms after the first fixation period). For the warning cue, four types of cues were presented: (i) no cue (a central fixation cross, similar to that presented during the first and second fixation period, was presented), (ii) a center cue (an asterisk was presented in the center, thus alerting the participant to the impending stimuli presentation), (iii) double cue (two asterisks were presented above and below a central fixation cross, at both possible locations of the target), and (iv) a spatial cue (an asterisk was presented at the impending location of the target). Participants completed 3 blocks of 96 trials (4 cue conditions x 2 target locations x 2 target directions x 2 repetitions), with trials presented in random order. The entire task lasted about 30 min.

Fluid intelligence. As a final comparison domain for understanding VAB performance and the associated neural underpinnings, we measured fluid intelligence. Participants completed a shortened, nine-item version (Form A; Bilker et al., 2012) of the original 60-item Raven's Standard Progressive Matrices (Raven, Raven, & Court, 1998). The task was completed on a laboratory computer, and there were no response time limits. The task consists of pattern matching questions designed to measure abstract reasoning skills, and it has been typically used in clinical settings as a non-verbal test of fluid intelligence.

Statistical analyses for behavioral data

308

309

310

311

312

313

314

315

316

317

318

319

320

321

322

323

324

325

326

327

328

329

330

Participants had to meet a minimum target discrimination (letter D or F?) score of 75% for the VAB and 60% for the AAB to be included in the final sample. As a result, 73 sets of data were available for all tasks, with the exception of the AAB task (n=71). For the AAB, two additional participants were excluded because their target discrimination performance did not meet the minimum threshold in each session. For all behavioral comparisons, p-values were based on two-tailed comparisons. We also did not correct any behavioral comparisons for multiple corrections, as we wanted to find any normality violations and used the task correlations to identify any relationships that might affect our CPM results. The computation of each behavioral measure is detailed in the following sections, again organized by task domain. To obtain stable behavioral metrics, we computed a 'best score' for each task metric. When data was available across two different sessions, the final score was averaged across both sessions. (When data was available only from a single session, the final score set to that session's.) We assessed whether the distribution of 'best scores' for each metric departed from normality using Jarque-Bera tests. As many normality violations were found, we used Spearman correlations of the 'best scores' to compare each pair of tasks. For tasks with two sessions, we also calculated test-retest reliability. Due to the aforementioned normality violations, Spearman correlations were again used. Attentional blink. The VAB and AAB deficits were calculated in the same way. For each participant, we first computed the mean probe detection accuracy for each lag condition, contingent upon correct identification of the target. Probe detection accuracy scores were then averaged across short-lag (lags 1, 2, or 3) and long-lag (lags 5, 7, 9) conditions. The former condition was expected to be within the attentional blink window, whereas the latter condition

332

333

334

335

336

337

338

339

340

341

342

343

344

345

346

347

348

349

350

351

352

353

was expected to be outside it. The AB deficit was computed by regressing out the long-lag scores from the short-lag scores of each participant (short-lag scores ~ long-lag scores), and then saving the residuals (MacLean & Arnell, 2012). Larger and positive values indicated smaller attentional deficits, and thus better task performance. Note that simply subtracting the short-lag from the long-lag scores yielded highly similar VAB deficit scores (r(71) = .955). Visual AB scores (VABresid) were obtained from sessions 1 and 5, and the auditory AB scores (AABresid) were obtained from sessions 2 and 4. Sustained attention. D-prime values were computed for the VSART, ASART and GradCPT, with larger values indicative of better performance in sustained attention. VSART scores (VSARTdprime) were obtained from sessions 1 and 3, ASART scores (ASARTdprime) were obtained from sessions 2 and 4, and GradCPT scores (GradCPTdprime) were obtained from sessions 2 and 4. Selective attention. To measure overall task performance in the ANT task, we computed the mean error rate (ANTerr) across all trials and the intra-individual variability of RTs (ANTrtvar) for each participant. Intra-individual RT variability was computed as the standard deviation divided by mean of correct-trial RTs. Arguably, this metric is a more sensitive measure of task performance than mean error rate (Rosenberg et al., 2018; Wojtowicz, Berrigan, & Fisk, 2012), with higher RT variability being linked to lower accuracy in ANT tasks (Adolfsdottir, Sorensen, & Lundervold, 2008; Lundervold et al., 2011). We also calculated metrics for each attentional network in the ANT, which putatively reflect their efficiencies. To do so, we compared the RTs between different trial conditions for each participant (Fan et al., 2002; Rosenberg et al., 2018). For the alerting network, efficiency was computed by subtracting the mean RT of double-cue condition from the mean RT of the no-

cue condition (ANTaert). Larger values would indicate faster responses due to the cue and thus better task performance. For the orienting network, efficiency was computed by subtracting the mean RT of spatial-cue condition from the mean RT of the center-cue condition (ANToert), thus larger positive values would also imply faster responses due to the cue and better task performance. For the executive control network, efficiency was computed by subtracting the mean RT of the congruent condition from the mean RT of the incongruent condition (ANTcert). In this case, smaller values would imply less interference by the flanker arrows and better task performance. For all computations of efficiency with RTs, only trials with correct responses were included.

For easier comparisons with the other behavioral measures, we re-coded the raw values of mean error (ANTerr), RT variability (ANTrtvar), and executive control efficiency (ANTcert), multiplying them by -1 such that larger values would also imply better task performance. ANT scores were obtained from sessions 1 and 3.

Fluid intelligence. Accuracy scores for the nine-item Raven's test (RavensAcc) were computed for each participant, with higher scores implying better performance.

Table 2. Summary of behavioral metrics, including their task domain and a description of their calculation. Raw values for ANTerr, ANTrtvar and ANTcert were re-coded such that larger values indicated better task performance.

Task domain	Metric	Description
Attentional Blink	VABresid	Residuals from regressing out long-lag scores from short-lag scores.
Attentional Billik	AABresid	Residuals from regressing out long-lag scores from short-lag scores.
Sustained Attention	VSARTdprime	Dprime scores.

	ASARTdprime				
	GradCPTdprime				
	ANTerr	Mean error across all trials.			
	ANTrtvar	SD/M of correct-trial RTs.			
Selective Attention	ANTaert	Alerting network efficiency. No-cue RT minus double-cue RT.			
	ANToert	Orienting network efficiency. Center-cue RT minus spatial-cue RT.			
	ANTcert	Executive control network efficiency. Incongruent RT minus			
	711 VI COIT	congruent RT.			
Fluid Intelligence	RavensAcc	Mean accuracy.			

MRI data collection and initial processing

Acquisition. MRI data were acquired at the Clinical Imaging Research Centre (Singapore) on a Siemens 3T MAGNETOM Prisma MRI scanner (Siemens, Erlangen, Germany) with a 32-channel head coil. Scanning parameters were adapted from the Human Connectome Project (HCP), and were chosen to ensure that full-brain coverage, including the cerebellum, was achieved for each participant. MRI data were acquired in Sessions 1 and 5 of the study, with the first session's data analyzed here. Each fMRI session started with a 5-min anatomical localizer scan, followed by four 8-min resting-state scans, six 5.25-min task-based (VAB task) scans, and a 5-min anatomical scan. During the resting-state runs, participants were asked to maintain fixation at a cross displayed in the center of the screen.

Imaging parameters. A 3D high-resolution (1 mm x 1 mm x 1 mm) T1-weighted MPRAGE pulse sequence was used to obtain whole-brain anatomical images for each

participant, allowing for subsequent normalization to standard space. For each participant, 128 1-

389

390

391

392

393

394

395

396

397

398

399

400

401

402

403

404

405

406

407

408

409

410

mm thick contiguous sagittal slices (0.5 mm skip; 1 x 1 mm in-plane resolution) were acquired. Other scanning parameters included a repetition time (TR) of 2300 ms, an effective echo time (TE) of 2.22 ms, a flip angle of 8° and 260 mm field of view. Functional MRI data were acquired with a multiband echoplanar imaging (MB-EPI; CMRR release R2015; Feinberg et al., 2010; Moeller et al., 2010; Xu et al., 2013) sequence with a MB acceleration factor of 8. 768 whole-brain images were obtained for each resting-state run while 504 images were acquired for each task-based run. T2*-weighted images were acquired using a TR of 625 ms, a TE of 33.2 ms and FA of 50° . Interleaved slices (imaging matrix = 64 x64) were collected using a 220 mm field of view, with slice thickness at 2.50 mm (no gap). The effective voxel size was thus 2.5 x 2.5 x 2.5 mm³. *Preprocessing.* Both task and localizer data were pre-processed using a previously published pipeline for functional connectivity analyses (Fong et al., 2019; Kong et al., 2019) publicly available at https://github.com/ThomasYeoLab/CBIG/tree/master/stable_projects/preprocessing/CBIG_fMRI Preproc2016. Pre-processing steps across resting-state and task-based runs were the same, except when mentioned otherwise. First, the initial four frames from each run were removed to aid with BOLD signal stabilisation. Motion correction using FSL's MCFLIRT was then applied such that runs with more than 50% of the frames exceeding a motion threshold were discarded to ameliorate any contributions of head motion. The motion threshold was defined as frame displacement (FD) > 75 and frame-to-frame intensity (DVARS) > 0.2 (Power, Barnes, Snyder, Schlaggar, & Petersen, 2012). From this step, one task-based run (and no resting-state runs) was removed from our data. FSL's bbregister function was used for intrasubject registration of the T1 anatomical images to the T2*-weighted images. The best run for each subject was used as the

412

413

414

415

416

417

418

419

420

421

422

423

424

425

426

427

428

429

430

431

432

433

registration file across all functional runs. Subsequently, motion parameters and their derivatives, the global whole brain signal, the white matter signal, the cerebral spinal fluid signal and linear trends were regressed out. Additionally, for task-based runs, we regressed out the haemodynamic response signal aligned to trial onset times. Frames with excessive motion, identified earlier on, were interpolated over (Power et al., 2014), and a temporal filter was applied to retain frequencies between 0.009 and 0.08 Hz. The resulting BOLD signal was projected to fsaverage6 surface space and spatially smoothed with an isotropic Gaussian kernel of 6 mm (FWHM). Functional connectivity. Functional connectivity was evaluated in fsaverage6 surface space for 400 cortical regions (Schaefer et al., 2017) and in MNI152 volumetric space for 19 subcortical regions (including the brain stem, and the left and right hemispheres of the accumbens area, amygdala, caudate, cerebellum, ventral diencephalon, hippocampus, pallidum, putamen, and thalamus), with a total of 419 parcellations. For each run, the mean time course of all the parcellations were correlated using Pearson's product moment correlation, resulting in a 419 (rows) x 419 (columns) correlation matrix, with $(419 \times 419 - 419) / 2 = 87.571$ unique values. Each cell in the correlation matrix represents a functional connection (edge) between a pair of parcellations. Fisher r-to-z transformation was applied to increase normality (Van Dijk et al., 2010). For each participant, Fisher-transformed matrices for all four resting-state scans and all six task-based scans were averaged separately, forming two functional connectivity (FC) matrices: resting-state FC (RSFC) and Visual Attentional Blink task FC (VABFC). Motion control. To reduce the effects of motion on functional connectivity, we adapted motion control procedures from Rosenberg et al. (2018) to remove edges that were correlated with motion. For each participant, we measured the following five motion parameters from their resting-state and task-based scans: (i) maximum displacement, (ii) maximum rotation, (iii) mean

frame-to-frame displacement, (iv) mean frame displacement (FD), and (v) mean frame-to-frame intensity (DVARS). Spearman's rank correlation was computed across participants between each edge in the FC matrix and each motion parameter. To be comparable with Rosenberg et al. (2018), in which 72.7% of edges remained after controlling for motion, we removed edges where r > .275 (two-tailed p < .02), leaving 69,617 edges, or 79.50% of the initial 87,571 edges. Edges were removed from both VABFC and RSFC if they met the criteria for removal in either FC matrix. **General approach for Connectome-based Predictive Models (CPM)**We adapted the Connectome-based Predictive Model (CPM) approach (Finn et al., 2015; Rosenberg, Finn, et al., 2016; Rosenberg et al., 2018; Shen et al., 2017) to predict individual differences in behavior from functional connectivity information (Figure 2). In the following section, we first describe our CPM procedure and then summarize our different models. Data from all 73 participants were included in these analyses (71 for models involving the AAB score, AABresid).

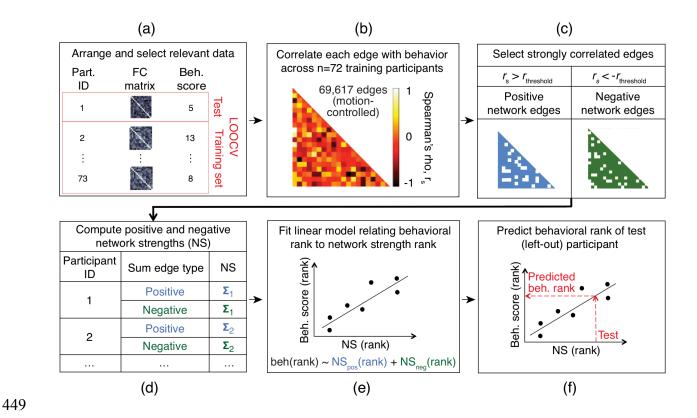


Figure 2. Procedure for Connectome-based Predictive Model (CPM) construction in the current study (adapted from Shen et al., 2017). CPMs predict individual differences in behavior from functional connectivity information. (a) Functional connectivity matrices and a behavioral score of interest for each participant were calculated. One pair was held out of model construction for each round of leave-one-out cross-validation (LOOCV) (Webb et al., 2011). (b) Functional connectivity edges were correlated with behavior across participants. (c) Edges that correlated most strongly, either positively or negatively, were selected. (d) Values from selected edges were summed separately for positive and negative network edges, yielding two network strengths for each participant. (e) A linear regression model relating (rank) network strengths to (rank) behavioral scores was computed. (f) The model was tested on a novel, out-of-sample participant (the individual left out in step (a)). After repeating steps a-f for each participant, the model was evaluated by correlating the predicted behavioral scores with the actual scores.

463

464

465

466

467

468

469

470

471

472

473

474

475

476

477

478

479

480

481

482

483

Model training. We first selected an FC matrix (i.e., VABFC or RSFC) and a behavioral score (e.g., VABresid). As part of the leave-one-out cross-validation procedure (Figure 2a), we then set aside the data for one participant as test data and proceeded to train the model on the remaining data (n = 72). To identify edges that most strongly correlated with behavior, we computed Spearman's rank correlations between each unique edge in the FC matrix and the behavioral score across 72 participants (Figure 2b), yielding 69,617 Spearman's rho values (r_s) . Edges positively related to behavior (positive network edges) were identified as those whose r_s was greater than a pre-defined threshold, $r_{\text{threshold}}$, and edges negatively related to behavior (negative network edges) were defined as those whose r_s was less than $-r_{\text{threshold}}$ (Figure 2c). Next, we computed network strengths (NS) for each participant by summing up values in their individual FC matrices across all positively and negatively correlated edges (Figure 2d), resulting in 72 sets of positive NS and negative NS values. Subsequently, we converted the positive NS, negative NS, and behavioral scores to rank space by ordering them according to participants' values. Finally, we formulated a multiple linear regression model (Figure 2e) with positive NS rank and negative NS rank as independent variables and behavioral rank as the dependent variable i.e., Behavior (rank) $\sim NS_{positive}$ (rank) + $NS_{negative}$ (rank). *Model test.* We proceeded to predict the behavioral score (in rank space) of the test participant (Figure 2f) by applying the training model to his/her FC matrix. Using the same positive and negatively correlated edges identified from training, positive NS and negative NS were first computed by summing up values in the FC matrix across the respective edges. Following this step, positive and negative NS values of the test participant were ranked against

the relevant NS values of the training participants, and entered into the multiple linear regression model to predict a behavioral rank.

Model evaluation. As part of the leave-one-out cross-validation procedure, the above training and prediction steps were repeated on all participants (N = 73) such that each participant was left out of training once, resulting in 73 sets of predicted behavioral ranks. To measure the predictive power of the model, we obtained correlations between the predicted and observed values, controlling for motion (Rosenberg et al., 2018). To this end, partial Spearman's rank correlation was computed between predicted and observed behavioral ranks, with the motion parameters (see section on Motion control) included as covariates. Where FC matrices were different for training and test, motion parameters from both the resting-state and task-based scans were included (i.e., ten covariates); where the same FC matrix was used, only motion parameters from the relevant scan were included (i.e., five covariates). P-values for model evaluation were left uncorrected, as most of the comparisons represented planned replications of previous work, and the pattern of predictions across tasks was more informative than single model predictions. For other neuroimaging-based statistical tests, we corrected for multiple comparisons, as detailed in the respective Results sections (Functional connectivity and Network overlap).

CPM model specification for data from the current study

For each of the eleven behavioral scores (e.g., VABresid, AABresid, VSARTdprime, etc.), we repeated the CPM procedure to train and test four different types of models. Two model types (vabCPM model variants) were trained using FC information and behavioral responses acquired from the main VAB task (i.e., VABFC and VABresid), and used to predict behavioral performance of novel participants using either their VAB *task* (VABFC; Model type A), or their *rest* (RSFC; Model type B) FC information. With these models, we sought to identify attentional

508

509

510

511

512

513

514

515

516

517

518

519

520

521

522

523

524

525

526

527

528

networks that were predictive of good VAB performance (positive network) and poor VAB performance (negative network), and then to determine whether these networks generalized to make similar predictions about other tasks. Two other model types (task-specific model variants) were trained using task-specific behavioral data and either VAB task (VABFC; Model type C) or rest (RSFC; Model type D) information. Behavioral performance was then predicted from the same type of FC data that was used during training. With these models, we tested whether predictive networks for each task could be predicted from FC data that was unrelated to that task; any such predictive networks would be useful for identifying and comparing the edges that are predictive of performance on a given task. Model type A (train on VABFC and behavioral data, predict with VABFC data). For training, the functional connectivity matrix VABFC and behavioral score VABresid were used to select edges and form the linear model. For test, the training model was applied to the VABFC matrix of the left-out participant. For evaluation, the predicted behavior from the leave-one-out procedure was correlated with the behavioral score from a selected task. Model type B (train on VABFC and behavioral data, predict with RSFC data). As in the previous model, VABFC and VABresid were used for training, but for test, the training model was applied to the RSFC matrix of the left-out participant. For evaluation, the predicted behavior was correlated with the behavioral score from a selected task. Model type C (train and test on VABFC data). During training, VABFC and the behavioral score from a selected task were used to form the model. During test, the training model was applied to VABFC of the left-out participant. For evaluation, the predicted behavior was correlated with the observed score of the selected task.

530

531

532

533

534

535

536

537

538

539

540

541

542

543

544

545

546

547

548

549

550

551

Model type D (train and test on RSFC data). For training, the functional connectivity matrix RSFC and the behavioral score from a selected task were used to form the model. For test, the training model was applied to the RSFC of the left-out participant. During evaluation, the predicted behavior was correlated with the observed score of the selected task. Controls. For the primary analyses in the study, we implemented the CPM procedure using edge selection cutoffs ($r_{\text{threshold}} = .232$, p = .05) previously used in Rosenberg et al. (2018). As an exploratory control, we also investigated whether our predictions were reasonably stable across edge selection cutoff values. To do so, we repeated the leave-one-out cross-validation procedure with $r_{\text{threshold}}$ ranging from .005 to .5, in steps of .005. Thus, in total, the CPM procedure was repeated (11 behavior x 4 models x 101 edge selection thresholds) 4,444 times. As p-values from LOOCV procedures can be biased, we verified the significance of our models using permutation testing (Shen et al., 2017). A null distribution with 1000 iterations was generated for each $r_{\text{threshold}}$. For each iteration, we randomly shuffled participants' behavioral scores and repeated the prediction steps above. P-values (uncorrected for this exploratory analysis) were computed as the proportion of permutation r_s with values greater than the observed r_s . Sustained Attention CPM from Rosenberg et al. (2016) To compare our predictions and networks with another attention-related model, we applied the Sustained Attention CPM (saCPM) (Rosenberg, Finn, et al., 2016) to our FC data. The saCPM was constructed using FC data computed with 268 parcellations (Shen, Tokoglu, Papademetris, & Constable, 2013). As our FC data was computed with 419 parcellations (Schaefer et al., 2017), we transformed the Shen parcellations to Schaefer parcellations in MNI152 space (91x109x91, 2mm voxels). For each Shen parcellation, we located the

corresponding Schaefer parcellation at a corresponding spatial location, excluding those that accounted for less than 10% of the voxels in the Shen parcellation. Next, we mapped the saCPM edges in the following way: if Shen parcellation A mapped to Schaefer parcellations 1, 2, and 3 and Shen parcellation C mapped to Schaefer parcellations 7 and 8, a functional connection (edge) between Shen parcellations A and C would be mapped to Schaefer edges 1-7, 2-7, 3-7, 1-8, 2-8, and 3-8. As before, edges that were removed previously due to motion were also removed in the mapped saCPM edges.

We computed network strengths by taking the dot product between the saCPM edges and our FC matrix (VABFC or RSFC), and entered the result into a linear model: Behavior ~ NS_{positive} + NS_{negative}. Note that these were motion-controlled FCs (69,617 edges), but during evaluation, we did not implement partial correlation with motion parameters as co-variates, following Rosenberg, Finn, et al. (2016). As the saCPM model was trained using FC data and dprime scores acquired during the GradCPT task, predicted scores from the model were also dprime scores. We evaluated the result of applying the saCPM to our FC data by computing Spearman's rank correlation between the predicted scores and the scores from each of our behavioral tasks.

Network overlaps and edge locations

Degree of overlap. To better understand the relationships across CPMs, we calculated the percentage of network overlap for model pairs. First, for each model, we identified edges that were common across all iterations of the leave-one-out procedure; we reasoned that these edges were most representative of the model. For each pair of models, we expressed the number of edges in common percentages of the number of edges in each model, and we then computed the

average of the two percentages. These percentages were calculated separately for positive-positive, negative-negative, positive-negative, and negative-positive network overlaps.

To statistically assess whether the edges for each pair of models significantly overlapped, we used the hypergeometric cumulative density function to determine the probability of drawing up to x out of K possible items with n drawings without replacement, from a population of size M (Rosenberg, Zhang, et al., 2016). The procedure was implemented with the hygecdf function in (MATLAB, 2014), as p = 1 – hygecdf(x, M, K, n), with x as the number of overlapping edges, x0 as the total number of edges, x0 as the number of edges from one model, and x0 as the number of edges from the other model. To control for multiple comparisons, FDR correction was applied across the complete set of tests (Benjamini & Hochberg, 1995).

Anatomical location of networks and overlaps. To determine the anatomical locations of network edges and the overlaps between the vabCPM and saCPM networks, we first grouped the 419 parcellations into network groups. The parcellations were matched to 17 network labels (Yeo et al., 2011), from which they were aggregated into eight cortical groups (Yeo, Tandi, & Chee, 2015) and a subcortical group. For each pair of network groups (e.g. Visual and Salience / Ventral Attention), we computed the number of connections (network edges) between them. We then reported this value as a percentage of the total number of possible connections between that pair of network groups.

593 Results

In the sections below, we first report the behavioral results and the functional connectivity matrices. We then present the behavioral predictions based on models derived from the present dataset, followed by the behavioral predictions based on an external model (the

saCPM; Rosenberg et al., 2016). After comparing the pattern of predictions, we compare the degree of overlap across the predictive networks. Finally, we examine the anatomical locations of the overlaps between the vabCPM and saCPM networks.

Behavioral performance

We found a robust visual attentional blink (VAB) in each session, with substantially impaired probe detection performance for the shorter lags (1, 2, 3) but not the longer lags (5, 7, 9) (Figure 3). For subsequent individual differences analyses, the VAB deficit was defined as short-lag performance when controlling for long-lag performance (see Methods), a definition the results supported. An AAB was also evidenced, though it was both smaller and less robust across sessions.

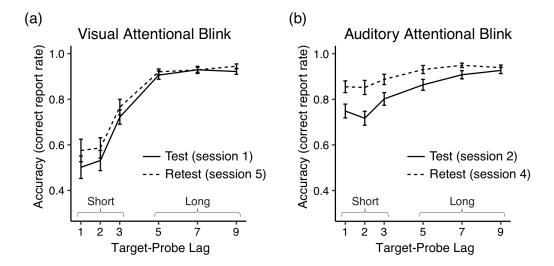


Figure 3. Accuracy scores (probe hit rates) for the VAB and AAB. Note the substantial impairment for the short target-probe lags (1, 2, and 3) in each session, especially for the VAB. Error bars represent standard error of the mean (SEM).

Test-retest reliability was high for most metrics, and their ranges were reasonable (

Table 3). Jarque-Bera tests of normality revealed that several behavioral scores were not normally-distributed (see relevant plots in Figure 4). Hence we adopted non-parametric approaches in our subsequent analyses and models.

Table 3. Summary of behavioral data. Test-retest reliability (Spearman correlations across sessions) was high for most metrics and significant for all (ps < .002). Jarque-Bera tests indicated some significant departures from normality. Metrics are presented before re-coding. Metrics that were subsequently reversed so that larger values would indicate better task performance are marked with an asterisk (*). N = 73 for all metrics except AABresid (N = 71).

Task domain	Metric	Mean	SD	Range	Reliability	Jarque-
rask domain					$(r_{\rm s})$	Bera (p)
Attentional	VABresid	0.003	0.213	[-0.577, 0.356]	0.75	.126
Blink	AABresid	0.001	0.118	[-0.412, 0.274]	0.38	.002
Sustained	VSARTdprime	3.469	0.77	[1.016, 5.256]	0.54	.077
	ASARTdprime	3.187	0.795	[0.364, 5.127]	0.62	.004
Attention	GradCPTdprime	2.844	0.655	[1.607, 4.359]	0.65	.144
	*ANTerr	2.463	2.222	[0, 11.111]	0.66	.001
	(% incorrect)				0.00	.001
Selective	*ANTrtvar (s)	0.147	0.035	[0.088, 0.24]	0.65	.058
Attention	ANTaert (s)	0.074	0.022	[0.02, 0.138]	0.42	.447
	ANToert (s)	0.04	0.017	[0, 0.085]	0.36	.500
	*ANTcert (s)	0.129	0.03	[0.078, 0.207]	0.74	.100
Fluid	RavensAcc	82.5	15.15	[22.22, 100]		001
Intelligence	(% correct)				-	.001

To examine how behavioral performance was related across the various tasks, we computed Spearman's rank correlations between each pair of behavioral scores (Figure 4). Significant correlations were primarily found within a given attentional domain. The three measures of Sustained Attention (VSARTdprime, ASARTdprime and GradCPTdprime) were significantly correlated, as were the primary measures of Selective Attention (ANTerr and ANTrtvar) and some of their component measures. Many of these metrics were significantly correlated, positively and negatively, across the Sustained and Selective Attention task domains as well. The negative correlations with the alerting (ANTaert) and orienting (ANToert) metrics may be due to their reflecting stimulus-driven attentional control, as opposed to the goal-directed control required for many other metrics. Attentional Blink (VABresid and AABresid) and Fluid Intelligence (RavensAcc) metrics generally were not significantly correlated with other metrics.

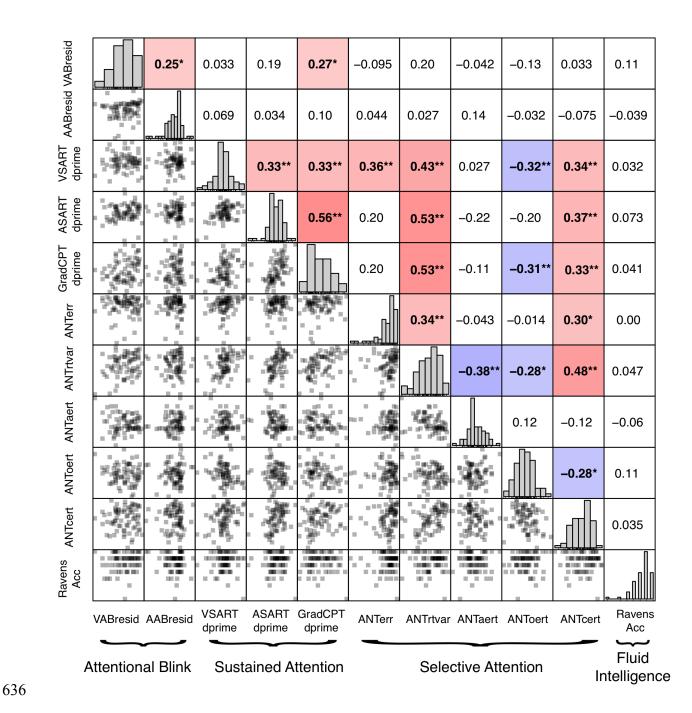
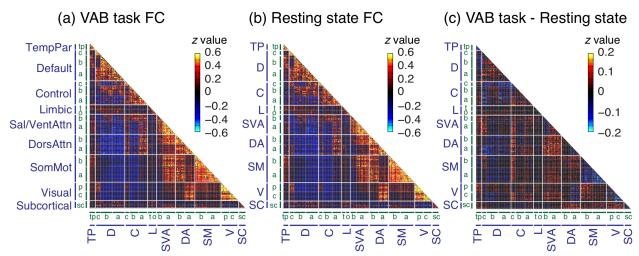


Figure 4. Behavioral score distributions and correlations. (Above diagonal) Spearman correlation coefficients for pairs of behavioral metrics. Most significant correlations were found within task domains, but some metrics correlated across the Sustained and Selective Attention domains. In contrast, Attentional Blink and Fluid Intelligence metrics largely did not correlate with other

metrics. Since the purpose of this analysis was to identify any behavioral relationships that might explain subsequent CPM results, no correction for multiple comparisons was applied. *p < .05, **p < .01. Red and blue shading indicates positive and negative relationships, respectively. (Diagonal) Histograms of behavioral data. The behavioral data had been re-coded so that larger values indicate better task performance for each measure. (Below diagonal) Scatterplots for each pair of behavioral metrics.

Functional connectivity matrices

To assess and compare network connectivity during the VAB task and during rest, we plotted group-averaged functional connectivity (FC) matrices (Figure 5). The 419 parcellations from the FC matrices were matched to 17 network labels (Yeo et al., 2011), from which they were aggregated into eight cortical groups (Yeo, Tandi, et al., 2015) and a subcortical group. FC data from both the VAB task (VABFC; Figure 5a) and resting state (RSFC; Figure 5b) showed similar connectivity patterns, with largely positive within-network correlations and mixed directions for between-network correlations. The network correlation patterns were generally similar to those observed in other data sets (Yeo et al., 2011; Yeo, Tandi, et al., 2015). Using Network-based statistics to correct for multiple comparisons (Zalesky, Fornito, & Bullmore, 2010), we observed small differences between the two FC matrices, with many connections linking the Salience/Ventral attention and the Dorsal attention networks (Figure 5c). Such differences are consistent with the reported neural correlates of the attentional blink, which are in frontal and parietal areas associated with the ventral and dorsal attention networks (Marois, Chun, & Gore, 2000; Marois & Ivanoff, 2005).



665

666

667

668

669

670

671

672

673

674

675

676

677

678

679

Figure 5. Functional connectivity (FC) matrices. Each edge was Fisher-transformed, and the resulting z-scores were averaged across participants. Edges found to correlate with motion were set to zero. (a) VAB task FC matrix (VABFC). (b) Resting-state FC matrix (RSFC). VABFC and RSFC patterns were similar to one another and to other data sets (Yeo et al., 2011; Yeo, Tandi, et al., 2015). (c) Difference FC matrix (VABFC - RSFC), showing edges that were significant at p = .05, corrected for multiple comparisons using network-based statistics. Differences between FC matrices were small, though they notably included connections linking the Salience/Ventral attention and the Dorsal attention networks. The 419 parcellations from the FC matrices were matched to 17 network labels (Yeo et al., 2011) (green labels), from which they were aggregated into eight cortical groups (Yeo, Tandi, et al., 2015) and a subcortical group (blue labels, spelled out in full in (a)). Subcortical regions include the brain stem, accumbens area, amygdala, caudate, cerebellum, ventral diencephalon, hippocampus, pallidum, putamen, and thalamus. For the green labels, letters represent the networks within the corresponding group, e.g., a(Default A), b(Default B), c(Default C), tp(TempPar), t(temporal pole in limbic region), o(orbital frontal cortex in limbic region), p(peripheral visual area), c(central visual area), and sc(subcortical).

Behavioral predictions from CPMs constructed using the present dataset

680

681

682

683

684

685

686

687

688

689

690

691

692

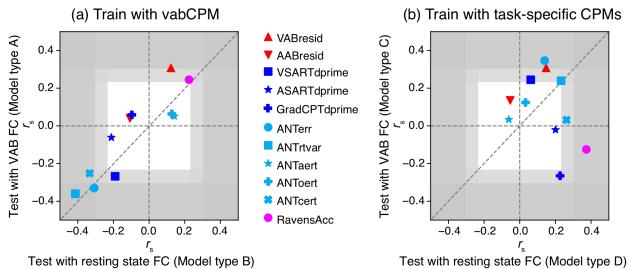
693

694

695

696

Models constructed from VAB functional connectivity and behavioral data (vabCPMs) positively predicted VAB performance from task data (VABFC; Model type A) but not from resting state data (RSFC; Model type B) (Figure 6a; corresponding values are tabulated in Figure 6-1 in the Extended Data). Task predictions were made in relative performance ranks (Spearman correlations), so these predictions readily applied to other rank-order behavioral scores. When so applied, we found that Fluid Intelligence performance could be positively predicted from both task and rest FC data. Critically, the correlations between predicted and actual performance for Sustained Attention metrics were *negative*, as were these correlations for Sustained Attention tasks, albeit less consistently. Such results are counterintuitive, as all behavioral scores were recoded so that larger values indicated better performance. Furthermore, one might expect that individuals whose network data predicted better performance on the VAB and Fluid Intelligence tasks would perform worse on Sustained Attention and Selective Attention tasks. In reality, however, VAB performance correlated insignificantly or weakly positively with other task metrics, whereas Fluid Intelligence performance was not significantly correlated with any other tasks. We return to this intriguing result in the Discussion.



699

700

701

702

703

704

705

706

707

708

709

710

711

712

713

Figure 6. Behavioral predictions from CPMs. Each point represents a pair of Spearman's rank correlation coefficients (r_s) computed between observed and predicted behavioral scores for a given model type. (a) Predictions from vabCPMs, which were trained with VABFC and VABresid. Note the successful positive predictions for the VAB and Fluid Intelligence, but successful *negative* predictions for some Selective and Sustained Attention metrics. (b) Predictions from task-specific models. Many models could successfully predict behavioral performance, though results often varied greatly across the FC source. For both panels, the dark gray region indicates where r_s values are significant at the p = .01 level, and the lighter gray region indicates where r_s values are significant at the p = .05 level (uncorrected, with d.f. = 71). The r_s values and corresponding p-values are tabulated in Figure 6-1 in Extended Data. A standard edge selection threshold ($r_{\text{threshold}} = .232, p = .05$) was used for all models, though results were similar across a wide range of threshold values (Figure 6-2 and Figure 6-3 in Extended Data). Finally, as p-values from LOOCV procedures can be biased, we verified our results for the VAB using permutation testing; significance from this method and parametric approaches was consistent across edge selection thresholds (Figure 6-4 in in Extended Data).

715

716

717

718

719

720

721

722

723

724

725

726

727

728

729

730

731

732

733

734

735

736

Models built from task-specific behavioral data significantly predicted performance for each of the task metrics predicted from the vabCPMs, although the results were less consistent between VABFC-based (Model type C) and RSFC-based (Model type D) predictions (Figure 6b; corresponding values are tabulated in Figure 6-1 in the Extended Data). Although it is possible that the different predictions reflect different information in the two FC data sources (Figure 3), it is unclear whether the differences are stable or simply reflect difficulty in building models from fMRI data for which the behavioral data were collected separately. Indeed, CPMs constructed from fMRI data collected during task performance and that task's behavioral scores tend to be more robust (Rosenberg, Finn, et al., 2016; Rosenberg et al., 2018; Yoo et al., 2017). Regardless, due to the same FC-behavior pair being used during both training and test, significant negative predictions from the vabCPMs became positive, as expected (e.g., VSARTdprime, ANTerr, ANTrtvat and ANTcert). To examine whether model predictions were sensitive to the number of edges selected during CPM training, we explored how r_s changes as a function of edge selection thresholds. R_s values were observed to be reasonably stable across edge selection thresholds, though more variation was observed in task-specific models (Model types C and D) than in vabCPMs (Model types A and B) (Figure 6-2 and Figure 6-3 in Extended Data). Additionally, as p-values from LOOCV procedures can be biased, we verified our results for the VAB using permutation testing; significance from this method and parametric approaches was consistent across edge selection thresholds (Figure 6-4 in in Extended Data). Each of these issues may have contributed to the unexpected negative predictions for the GradCPTdprime metric, an aspect of the results to which we return below. Behavioral predictions from an external CPM (saCPM; Rosenberg et al., 2016).

738

739

740

741

742

743

744

745

746

747

748

749

750

751

752

753

For external validation of our model predictions, we applied the Sustained Attention CPM (saCPM) to our data. The saCPM was trained on fMRI and behavioral data from the GradCPT, a sustained attention task (Rosenberg, Finn, et al., 2016). When applied to our data, the saCPM predictions were virtually mirror images of our vabCPM predictions (Figure 7). Specifically, whereas the vabCPM predicted VAB and Fluid Intelligence performance positively and predicted Sustained Attention and Selective Attention performance negatively, the saCPM predicted VAB and Fluid Intelligence performance *negatively* and predicted Sustained Attention and Selective Attention performance *positively*. With these findings, we also replicated the results from Rosenberg et al. (2018). Specifically, the saCPM was able to predict the error rates (ANTerr), reaction time variability (ANTrtvar), and conflict (ANTcert) metrics for novel individuals in the ANT task. Conversely, we failed to replicate the significant predictions for GradCPTdprime in our data, contrary to expectations. This replication failure is puzzling because the GradCPT behavioral data showed a reasonable spread of scores and good test-retest reliability (Table 3), and the saCPM and vabCPM did make significant predictions on other Sustained Attention metrics in our data set. We are currently exploring our GradCPT task and results further in our laboratory.

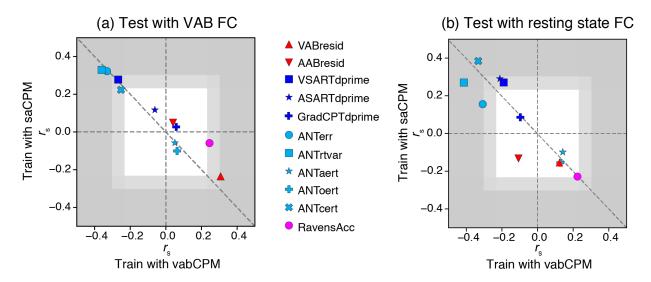


Figure 7. Comparison of predictions from the vabCPM and saCPM. Each point represents a pair of Spearman's rank correlation coefficients (r_s) computed between observed and predicted behavioral scores for a given model type. (a) Predictions from VABFC data. Note the saCPM's successful positive predictions for some Selective and Sustained Attention metrics, with a successful *negative* prediction for the VAB. As noted above, the prediction directions were reversed for the vabCPM; indeed, the prediction points fall close to the diagonal. (b) Predictions from RSFC data. The vabCPM and saCPM predictions went in opposite directions, and were generally similar to the predictions from the VABFC. For both panels, the dark gray region indicates where r_s values are significant at the p = .01 level, and the lighter gray region indicates where r_s values are significant at the p = .05 level (uncorrected, with d.f. = 71). The edge selection threshold ($r_{threshold}$) corresponded to p = .05 for all models. The r_s values and corresponding p-values for the saCPM are tabulated in Figure 7-1 in the Extended Data. (See Figure 6 for additional vabCPM details.)

Degree of overlap between predictive network pairs

771

772

773

774

775

776

777

778

779

780

781

782

783

784

785

786

787

788

789

790

791

To better understand how the underlying functional connectivity networks contributed to the model predictions, we analyzed the extent to which edges were shared between pairs of CPMs. For each pair, we calculated overlaps between positive network edges (those that predicted better behavioral performance for the model's task), between negative network edges, and across positive and negative network edges (Figure 8). The networks derived from the vabCPM (VAB network) and saCPM (SA network) did not significantly overlap at similar network edges (e.g. positive-positive; Figure 8a and 8b). Instead, they significantly overlapped only at *opposing* network edges, the positive edges from one model and the negative edges from the other (Figure 8c and 8d). The overlaps between the VAB network and SA network accorded with their overlaps with networks derived from other task-specific CPMs. VAB network edges tended to overlap more with the opposing network edges of the Sustained and Selective Attention models (Figure 8c and 8d) as compared to their similar network edges (Figure 8a and 8b). In contrast, SA network edges overlapped significantly with similar edges from each of the Sustained and Selective Attention models, but with none of their opposing edges. Finally, although VAB and Fluid Intelligence networks overlapped significantly only at similar network edges, SA network edges did not significantly overlap with either similar or opposing Fluid Intelligence network edges. Taken together, the pattern of edge overlaps accords with the pattern of behavioral predictions. The VAB and SA networks overlapped at opposing edges, and their predictions were also negatively related (Figure 7). Predictions for individual metrics also aligned with model overlaps. In general, when a metric's observed scores positively correlated with scores

predicted from the vabCPM or saCPM, that metric's CPM-derived network tended to overlap with the VAB or SA network at similar network edges.

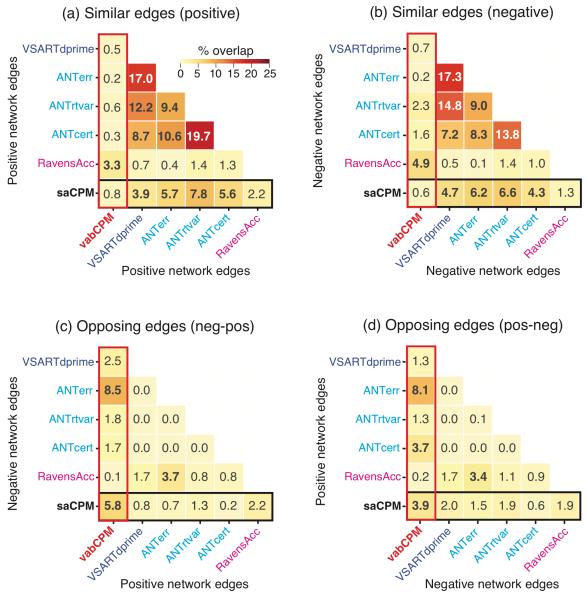


Figure 8. Percentage of edge overlap between networks from selected pairs of CPMs. Each task-specific model was based on VABFC data (Model type C). Model pairs with statistically significant overlap (p < .05, FDR corrected) are indicated in bold. Positive network edges predicted better behavioral performance for their associated metric, whereas negative network

edges predicted worse behavioral performance. Overlap between *similar* network edges, (a) positive-positive and (b) negative-negative. Sustained Attention (including the saCPM) and Selective Attention models overlap primarily on similar edges, as do Fluid Intelligence and the vabCPM. Overlap between *opposing* network edges, (c) negative-positive and (d) positive-negative. The vabCPM network primarily overlaps with Sustained Attention (including the saCPM) and Selective Attention on their opposing edges. Only task metrics that were significantly predicted by vabCPMs using VABFC information are shown. The full set of overlaps for all CPMs can be found in the Extended Data (Figure 8-1, 8-2, 8-3, and 8-4). Note that the highest overlap value for any pairwise comparison, including each task metric predicted from VABFC and RSFC, was 26.5%.

We also observed significant overlaps between similar network edges from Sustained and Selective Attention models (Figure 8a and 8b). Similarly, Rosenberg and colleagues observed substantial overlaps between a high attention (positive) network from the GradCPT task and networks predicting high accuracy and low RT variability (better performance) in the ANT (Rosenberg et al., 2018). Taken together, these results suggest that sustained and selective attention share similar functional networks, at least in part. Such results also justify labels such as the "successful attention network" (Rosenberg et al., 2018). Importantly, these network overlaps are consistent with both the CPM predictions and the behavioral relationships (Figure 4). Significant network overlaps involving the VAB network, however, were not accompanied by significant behavioral relationships. We return to this observation in the Discussion.

Anatomical locations of network edges

823

824

825

826

827

828

829

830

831

832

833

834

835

836

837

838

839

840

841

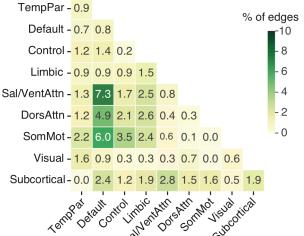
842

843

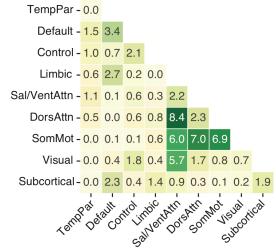
844

To better understand the networks that contribute to the successful behavioral predictions from the vabCPM and the saCPM, we investigated the anatomy of their underlying network edges, Briefly, for each pair of network groups (e.g. Visual and Salience / Ventral Attention), we expressed the number of shared connections (network edges) as a percentage of the number of possible connections between that pair (Figure 9). Edges that positively predicted VAB performance occurred primarily between the Default network and several other networks, including the Salience/Ventral attention, Dorsal attention, and Somatomotor networks (Figure 9a). For negative network edges, the most frequent connections occurred between the Salience/Ventral attention network and several other networks, including the Dorsal attention, Somatomotor, and Visual networks (Figure 9b). Note that such connections were also enhanced in VABFC compared to RSFC (Figure 5c), and that the connections across attention networks are consistent with the neural correlates of the attentional blink (Marois et al., 2000; Marois & Ivanoff, 2005). Negative network edges were also frequently found in Somatomotor network connections to the Dorsal attention network and to itself (within-network connections). The SA network's positive edges primarily included connections that involved the Visual and Subcortical network groups (Figure 9d). This pattern represents the remapping of the connections between the cerebellum and the occipital lobe of the "high attention" network (Rosenberg, Finn, et al., 2016) from the Shen-268 parcellation (Shen et al., 2013) to the Schaefer-419 one (Schaefer et al., 2017). Negative network edges included within-group connections in the TempPar and Subcortical networks (Figure 9c), which accord well with the intra-temporal, intra-cerebellar, and temporo-parietal connections in the "low attention" network (Rosenberg, Finn, et al., 2016). We observed additional negative edges within the Visual network and from the TempPar network to various other networks.

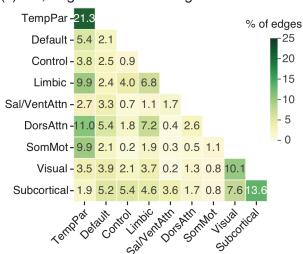
(a) VAB, positive network edge locations



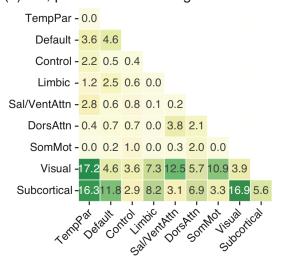
(b) VAB, negative network edge locations



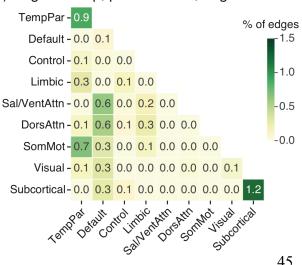
(c) SA, negative network edge locations



(d) SA, positive network edge locations



(e) Edge overlap; positive VAB, negative SA



(f) Edge overlap; negative VAB, positive SA

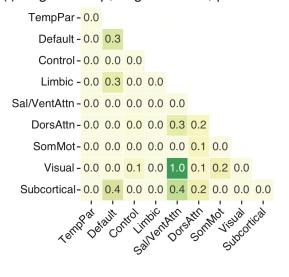


Figure 9. Anatomical locations of predictive attentional network edges. Each cell represents the number of shared connections between a pair of network groups, expressed as a percentage of the number of possible connections between that pair. For details about network groups, see Figure 5. (a) Positive and (b) negative VAB network edges. (c) Negative and (d) positive SA network edges. The panel order has been reversed for easier comparison with the opposing edges from the VAB network. (e) Positive VAB network edges and negative SA network edges (i.e. overlaps between (a) and (c)). (f) Negative VAB network edges and positive SA network edges (i.e. overlaps between (b) and (d)). Overlap percentages were numerically small, but included key network groups, particularly Default and Salience / Ventral Attention (Sal/VentAttn).

Although the VAB and SA networks involved largely dissociable sets of edges, some critical edges appeared to be shared (Figure 9e and 9f). Note that these overlaps were at opposing edges (Figure 8). Although overlap percentages were numerically small, they included key network groups. In particular, multiple identified connections involved the Default, Salience/Ventral attention, or TempPar networks.

862 Discussion

Predictive attentional networks

We used Connectome-based Predictive Modeling (CPM), a machine learning-based technique that associates task performance with functional connectivity measures, to construct a predictive model of Visual Attentional Blink (VAB) performance. Our model (vabCPM) successfully predicted VAB performance in novel individuals from fMRI data. The model's predictions generalized to other task domains, including fluid intelligence (Finn et al., 2015).

870

871

872

873

874

875

876

877

878

879

880

881

882

883

884

885

886

887

888

889

890

Critically, vabCPM predictions for many sustained and selective attention task scores correlated *negatively* with the actual scores. As such, these significant predictions represent both model generalization and an extension of previous CPM results (Rosenberg, Finn, et al., 2016; Rosenberg et al., 2018; Yoo et al., 2017), but with new insights into attentional function owing to the divergent predictions. For external validation of these results, we applied the Sustained Attention CPM (saCPM) (Rosenberg, Finn, et al., 2016), to our data. Previously, the saCPM successfully predicted task performance for sustained attention (GradCPT) (Rosenberg, Finn, et al., 2016) and selective attention (Attention Network Task, ANT) (Rosenberg et al., 2018). Here we broadly replicated these results: The saCPM successfully predicted sustained and selective attention task performance when applied to our participants' data. In the sustained attention domain, however, significant predictions were found only for visual and auditory SART (Sustained Attention to Response Task) scores, but curiously not for GradCPT scores. In results that mirrored the vabCPM predictions, the saCPM predictions for VAB scores were negatively correlated with the observed scores. This pattern of divergent predictions was also reflected in network overlaps. For the networks derived from the vabCPM and the saCPM, opposing network edges (i.e. positive from one, negative from the other) overlapped significantly, whereas similar network edges did not. Models constructed from each behavioral task and our fMRI data corroborated these results. As all behavioral data had been coded such that larger values indicated better performance, these divergent predictions indicate that "good" or "bad" network function was contingent on the task and its cognitive underpinnings. Moreover, the tasks were performed over several days,

suggesting that the individual differences were stable and trait-like, not due to session-specific state effects.

Implications for Cognitive Mechanisms

891

892

893

894

895

896

897

898

899

900

901

902

903

904

905

906

907

908

909

910

911

912

913

Although the observed pattern of CPM predictions may seem counterintuitive, it is consistent with both empirical evidence and theoretical positions. Foremost, our study's attention tasks represent different ways of deploying voluntary attention. The VAB task requires rapid attentional engagement, disengagement, and re-engagement; sustained attention tasks require engagement over a prolonged period of time; the attention network task (ANT) requires the direction of attention to relevant spatial information. Skogsberg et al. (2015) proposed that the VAB and sustained attention tasks lie at opposite ends of a transient-sustained attention continuum. Rensink suggested that the VAB and ANT involve different core attentional functions: In the VAB, 'attentional holding' of one visual object leads to the failure to create a second visual object, whereas in the ANT, 'attentional filtering' selects spatial information (Rensink, 2013, 2015). Empirical findings support the relative uniqueness of the VAB, while also suggesting that the ANT and sustained attention are more closely related. In our data, we found a general lack of behavioral correlations between the VAB and other attention task measures, consistent with previous studies of individual differences (Dale et al., 2013; Skogsberg et al., 2015). Conversely, we found moderately strong correlations between sustained attention tasks and the ANT. These tasks have been found to share similar functional networks (Rosenberg et al., 2018), a result we also replicated. Nevertheless, the conceptual separation of the VAB from other attention tasks does not explain the *opposing* pattern of predictions from the same networks (saCPM and vabCPM), and

915

916

917

918

919

920

921

922

923

924

925

926

927

928

929

930

931

932

933

934

935

936

the significant overlaps between their opposing network edges. Two observations provide important context for this finding. First, the VAB has spawned a variety of theoretical accounts (Dux & Marois, 2009), and its magnitude is sensitive to numerous disparate manipulations, ranging from requiring online responses (Jolicoeur, 1998) to concurrently listening to music (Olivers & Nieuwenhuis, 2005). As such, the VAB may have multiple causes. Second, our VAB predictions were generally moderate (r = .31 for vabCPM and r = .24 for saCPM; 5-10% of the variance), far smaller than the observed degree of stable individual differences (test-retest: r =.71; 50% of the variance). It is possible that the CPMs capture the variance associated with few, or even one, of the factors that affect VAB magnitude. If so, what could that factor be? One plausible explanation is that our predictions reflect an individual's propensity to maintain a more diffuse state of attention. This idea is consistent with the overinvestment hypothesis, in which the VAB results from too much attention being allocated to the first target; consequently, reducing attention on the RSVP stream improves performance (Dale & Arnell, 2010, 2015; Olivers & Nieuwenhuis, 2006). Similarly, task-concurrent mind-wandering, such as listening to music or thinking about a vacation, reduces the VAB deficit (Olivers & Nieuwenhuis, 2005, 2006). Similar effects are found in studies of disposition: Individuals with a greater propensity for mind-wandering tend to perform better in the VAB task (Thomson et al., 2015). Furthermore, mind-wandering has been linked to higher fluid intelligence and better problem-solving abilities (Baird, Smallwood, & Schooler, 2011; Godwin et al., 2017; Unsworth & McMillan, 2014), consistent with our CPM findings. Diffuse attentional states, however, are associated with lower performance on tasks requiring more focused cognition. For example, in sustained attention tasks, individuals more prone to lapses in attention perform more poorly on the SART (Manly, 1999; Robertson et al.,

938

939

940

941

942

943

944

945

946

947

948

949

950

951

952

953

954

955

956

957

958

959

1997; Smilek et al., 2010). Within individuals, distractive thoughts are associated with lower SART accuracy, prolonged and more variable RTs, and poorer response inhibition (Kam & Handy, 2014; Leszczynski et al., 2017; Stawarczyk, Majerus, Maj, Van der Linden, & D'Argembeau, 2011). In selective attention tasks, individuals more prone to mind-wandering performed worse on the ANT task (Gonçalves et al., 2017), and showed impaired exogeneous orienting of attention (Hu et al., 2012). Both mind-wandering states and traits are also reflected in patterns of brain activity. Activity in the Default network and frontoparietal control regions increases during mindwandering (Fox, Spreng, Ellamil, Andrews-Hanna, & Christoff, 2015). Similarly, individuals more prone to mind-wander had increased connectivity both within the Default network and between the Default network and frontoparietal control regions (Godwin et al., 2017). Such results are partially consistent with our findings. Edges between the Default and attentional networks (not the Control network) were related to positive VAB network edges and negative SA network edges. On the weight of the available evidence, we propose that our vabCPM reflects individuals' propensity towards diffuse attentional deployment. That propensity could indicate an individual's ability to diffusely attend, their tendency to be in that state or mode, or both. We also suggest that other CPMs, including the saCPM, could reflect the complementary propensity towards more focused attentional deployment. **Predictions from "Resting State"** Neuroimaging studies using 'resting state' data, in which subjects are scanned while not engaged in any particular task, have become increasingly popular. Initially used to identify functional architecture (Biswal, Yetkin, Haughton, & Hyde, 1995; Schaefer et al., 2017; Yeo et

al., 2011), resting state data have recently been used to test whether functional architecture persists even when an individual is not engaged in a task that requires a given neurocognitive network (Finn et al., 2015; Jangraw et al., 2018; Lin et al., 2018; Rosenberg, Finn, et al., 2016; Yoo et al., 2017). Resting state studies also have many logistical advantages, including relatively easy standardization across multiple test sites and the potential for numerous applications from a single data set. In the current study, resting state CPMs (Model type D) replicated previous studies by successfully predicting sustained attention (Yoo et al., 2017), selective attention (Yoo et al., 2017) and fluid intelligence (Finn et al., 2015) task performance. We also replicated the result that models trained with task-concurrent FC data generally predict task performance better than models trained from resting state data (Yoo et al., 2017). Finally, when applying our trained vabCPM model to novel participants, predictions from task FC data were superior to predictions from resting-state FC data (Finn et al., 2017; Greene, Gao, Scheinost, & Constable, 2018; Rosenberg, Finn, et al., 2016; Rosenberg et al., 2018). This result might be due to amplification of behaviorally relevant individual differences in network patterns while performing a task (Greene et al., 2018).

Methodological Considerations

960

961

962

963

964

965

966

967

968

969

970

971

972

973

974

975

976

977

978

979

980

981

982

Although we followed the guidelines in Shen et al. (2017) when developing our CPMs, there are two notable methodological differences. First, previous studies used the volumetric Shen-268 parcellation (Shen et al., 2013), but we used the surface-based Schaefer-419 parcellation (Schaefer et al., 2017). Second, previous studies have constructed linear models for associating network strength and behavioral scores, with predictions assessed using Spearman correlations (Fountain-Zaragoza et al., 2019; Lin et al., 2018; Rosenberg, Finn, et al., 2016; Rosenberg et al., 2018). We instead formed linear models from the ranks directly. Despite these

differences, we still observed prediction patterns that were largely consistent and comparable to previous studies, providing evidence that CPMs are reasonably robust across such variations.

986 References 987 Adolfsdottir, S., Sorensen, L., & Lundervold, A. J. (2008). The attention network test: A 988 characteristic pattern of deficits in children with ADHD. Behavioral and Brain 989 Functions, 4(1), 9. https://doi.org/10.1186/1744-9081-4-9 990 Asplund, C. L., Todd, J. J., Snyder, A. P., & Marois, R. (2010). A central role for the lateral 991 prefrontal cortex in goal-directed and stimulus-driven attention. *Nature Neuroscience*, 992 13(4), 507–512. https://doi.org/10.1038/nn.2509 993 Baird, B., Smallwood, J., & Schooler, J. W. (2011). Back to the future: Autobiographical 994 planning and the functionality of mind-wandering. Consciousness and Cognition, 20(4), 995 1604–1611. https://doi.org/10.1016/j.concog.2011.08.007 996 Benjamini, Y., & Hochberg, Y. (1995). Controlling the False Discovery Rate: A Practical and 997 Powerful Approach to Multiple Testing. Journal of the Royal Statistical Society. Series B 998 (Methodological), 57(1), 289–300. 999 Bilker, W. B., Hansen, J. A., Brensinger, C. M., Richard, J., Gur, R. E., & Gur, R. C. (2012). 1000 Development of Abbreviated Nine-Item Forms of the Raven's Standard Progressive 1001 Matrices Test. Assessment, 19(3), 354–369. https://doi.org/10.1177/1073191112446655 1002 Biswal, B., Yetkin, F. Z., Haughton, V. M., & Hyde, J. S. (1995). Functional connectivity in the 1003 motor cortex of resting human brain using echo-planar MRI. Magnetic Resonance in 1004 *Medicine*, 34(4), 537–541. 1005 Chun, M. M., Golomb, J. D., & Turk-Browne, N. B. (2011). A Taxonomy of External and 1006 Internal Attention. Annual Review of Psychology, 62(1), 73–101. 1007 https://doi.org/10.1146/annurev.psych.093008.100427

1008 Chun, M. M., & Potter, M. C. (1995). A two-stage model for multiple target detection in rapid 1009 serial visual presentation. Journal of Experimental Psychology. Human Perception and 1010 Performance, 21(1), 109–127. 1011 Corbetta, M., & Shulman, G. L. (2002). Control of goal-directed and stimulus-driven attention in 1012 the brain. Nature Reviews Neuroscience, 3(3), 201–215. https://doi.org/10.1038/nrn755 1013 Dale, G., & Arnell, K. M. (2010). Individual differences in dispositional focus of attention 1014 predict attentional blink magnitude. Attention, Perception, & Psychophysics, 72(3), 602– 1015 606. https://doi.org/10.3758/APP.72.3.602 1016 Dale, G., & Arnell, K. M. (2015). Multiple measures of dispositional global/local bias predict 1017 attentional blink magnitude. Psychological Research, 79(4), 534–547. 1018 https://doi.org/10.1007/s00426-014-0591-3 1019 Dale, G., Dux, P. E., & Arnell, K. M. (2013). Individual differences within and across attentional 1020 blink tasks revisited. Attention, Perception, & Psychophysics, 75(3), 456–467. 1021 https://doi.org/10.3758/s13414-012-0415-8 1022 Desimone, R., & Duncan, J. (1995). Neural Mechanisms of Selective Visual Attention. Annual 1023 *Review of Neuroscience*, 18(1), 193–222. 1024 https://doi.org/10.1146/annurev.ne.18.030195.001205 1025 Di Lollo, V., Kawahara, J., Shahab Ghorashi, S. M., & Enns, J. T. (2005). The attentional blink: Resource depletion or temporary loss of control? Psychological Research, 69(3), 191– 1026 1027 200. https://doi.org/10.1007/s00426-004-0173-x Duncan, J. (2010). The multiple-demand (MD) system of the primate brain: Mental programs for 1028 1029 intelligent behaviour. Trends in Cognitive Sciences, 14(4), 172–179. 1030 https://doi.org/10.1016/j.tics.2010.01.004

1031 Dux, P. E., & Marois, R. (2009). The attentional blink: A review of data and theory. Attention, 1032 Perception & Psychophysics, 71(8), 1683–1700. https://doi.org/10.3758/APP.71.8.1683 1033 Egeth, H. E., & Yantis, S. (1997). Visual attention: Control, representation, and time course. 1034 Annual Review of Psychology, 48(1), 269–297. 1035 Esterman, M., Noonan, S. K., Rosenberg, M., & DeGutis, J. (2013). In the Zone or Zoning Out? 1036 Tracking Behavioral and Neural Fluctuations During Sustained Attention. Cerebral 1037 Cortex, 23(11), 2712–2723. https://doi.org/10.1093/cercor/bhs261 1038 Fan, J., McCandliss, B. D., Sommer, T., Raz, A., & Posner, M. I. (2002). Testing the Efficiency 1039 and Independence of Attentional Networks. Journal of Cognitive Neuroscience, 14(3), 1040 340–347. https://doi.org/10.1162/089892902317361886 1041 Feinberg, D. A., Moeller, S., Smith, S. M., Auerbach, E., Ramanna, S., Glasser, M. F., ... 1042 Yacoub, E. (2010). Multiplexed Echo Planar Imaging for Sub-Second Whole Brain 1043 FMRI and Fast Diffusion Imaging. *PLoS ONE*, 5(12), e15710. 1044 https://doi.org/10.1371/journal.pone.0015710 1045 Finn, E. S., Scheinost, D., Finn, D. M., Shen, X., Papademetris, X., & Constable, R. T. (2017). 1046 Can brain state be manipulated to emphasize individual differences in functional 1047 connectivity? NeuroImage, 160, 140–151. 1048 https://doi.org/10.1016/j.neuroimage.2017.03.064 1049 Finn, E. S., Shen, X., Scheinost, D., Rosenberg, M. D., Huang, J., Chun, M. M., ... Constable, R. T. (2015). Functional connectome fingerprinting: Identifying individuals using patterns 1050 1051 of brain connectivity. *Nature Neuroscience*, 18(11), 1664–1671. 1052 https://doi.org/10.1038/nn.4135

1053 Fong, A. H. C., Yoo, K., Rosenberg, M. D., Zhang, S., Li, C.-S. R., Scheinost, D., ... Chun, M. 1054 M. (2019). Dynamic functional connectivity during task performance and rest predicts 1055 individual differences in attention across studies. *NeuroImage*, 188, 14–25. 1056 https://doi.org/10.1016/j.neuroimage.2018.11.057 1057 Fountain-Zaragoza, S., Samimy, S., Rosenberg, M. D., & Prakash, R. S. (2019). Connectome-1058 based models predict attentional control in aging adults. NeuroImage, 186, 1–13. 1059 https://doi.org/10.1016/j.neuroimage.2018.10.074 Fox, K. C. R., Spreng, R. N., Ellamil, M., Andrews-Hanna, J. R., & Christoff, K. (2015). The 1060 1061 wandering brain: Meta-analysis of functional neuroimaging studies of mind-wandering 1062 and related spontaneous thought processes. *NeuroImage*, 111, 611–621. 1063 https://doi.org/10.1016/j.neuroimage.2015.02.039 1064 Godwin, C. A., Hunter, M. A., Bezdek, M. A., Lieberman, G., Elkin-Frankston, S., Romero, V. 1065 L., ... Schumacher, E. H. (2017). Functional connectivity within and between intrinsic 1066 brain networks correlates with trait mind wandering. *Neuropsychologia*, 103, 140–153. 1067 https://doi.org/10.1016/j.neuropsychologia.2017.07.006 1068 Gonçalves, Ó. F., Rêgo, G., Oliveira-Silva, P., Leite, J., Carvalho, S., Fregni, F., ... Boggio, P. S. 1069 (2017). Mind wandering and the attention network system. Acta Psychologica, 172, 49– 1070 54. https://doi.org/10.1016/j.actpsy.2016.11.008 1071 Greene, A. S., Gao, S., Scheinost, D., & Constable, R. T. (2018). Task-induced brain state 1072 manipulation improves prediction of individual traits. *Nature Communications*, 9(1). 1073 https://doi.org/10.1038/s41467-018-04920-3

1074 Hu, N., He, S., & Xu, B. (2012). Different efficiencies of attentional orienting in different 1075 wandering minds. Consciousness and Cognition, 21(1), 139–148. 1076 https://doi.org/10.1016/j.concog.2011.12.007 1077 Jangraw, D. C., Gonzalez-Castillo, J., Handwerker, D. A., Ghane, M., Rosenberg, M. D., 1078 Panwar, P., & Bandettini, P. A. (2018). A functional connectivity-based neuromarker of 1079 sustained attention generalizes to predict recall in a reading task. NeuroImage, 166, 99– 1080 109. https://doi.org/10.1016/j.neuroimage.2017.10.019 1081 Jolicoeur, P. (1998). Modulation of the attentional blink by on-line response selection: Evidence 1082 from speeded and unspeeded Task1 decisions. Memory & Cognition, 26(5), 1014–1032. 1083 https://doi.org/10.3758/BF03201180 1084 Kam, J. W. Y., & Handy, T. C. (2014). Differential recruitment of executive resources during 1085 mind wandering. Consciousness and Cognition, 26, 51–63. 1086 https://doi.org/10.1016/j.concog.2014.03.002 Kong, R., Li, J., Orban, C., Sabuncu, M. R., Liu, H., Schaefer, A., ... Yeo, B. T. T. (2019). 1087 1088 Spatial Topography of Individual-Specific Cortical Networks Predicts Human Cognition, 1089 Personality, and Emotion. Cerebral Cortex, 29(6), 2533–2551. 1090 https://doi.org/10.1093/cercor/bhy123 1091 Leszczynski, M., Chaieb, L., Reber, T. P., Derner, M., Axmacher, N., & Fell, J. (2017). Mind 1092 wandering simultaneously prolongs reactions and promotes creative incubation. Scientific 1093 Reports, 7(1). https://doi.org/10.1038/s41598-017-10616-3 1094 Lin, Q., Rosenberg, M. D., Yoo, K., Hsu, T. W., O'Connell, T. P., & Chun, M. M. (2018). 1095 Resting-State Functional Connectivity Predicts Cognitive Impairment Related to

1096 Alzheimer's Disease. Frontiers in Aging Neuroscience, 10. 1097 https://doi.org/10.3389/fnagi.2018.00094 1098 Lundervold, A. J., Adolfsdottir, S., Halleland, H., Halmøy, A., Plessen, K., & Haavik, J. (2011). 1099 Attention Network Test in adults with ADHD - the impact of affective fluctuations. 1100 Behavioral and Brain Functions, 7(1), 27. https://doi.org/10.1186/1744-9081-7-27 1101 MacLean, M. H., & Arnell, K. M. (2012). A conceptual and methodological framework for 1102 measuring and modulating the attentional blink. Attention, Perception, & Psychophysics, 1103 74(6), 1080–1097. https://doi.org/10.3758/s13414-012-0338-4 1104 Manly, T. (1999). The absent mind: Further investigations of sustained attention to response. 1105 Neuropsychologia, 37(6), 661–670. https://doi.org/10.1016/S0028-3932(98)00127-4 1106 Marois, R., Chun, M. M., & Gore, J. C. (2000). Neural correlates of the attentional blink. 1107 *Neuron*, 28(1), 299–308. 1108 Marois, R., & Ivanoff, J. (2005). Capacity limits of information processing in the brain. *Trends* 1109 in Cognitive Sciences, 9(6), 296–305. https://doi.org/10.1016/j.tics.2005.04.010 1110 MATLAB. (2014). Natick, MA: The MathWorks, Inc. 1111 Moeller, S., Yacoub, E., Olman, C. A., Auerbach, E., Strupp, J., Harel, N., & Uğurbil, K. (2010). 1112 Multiband multislice GE-EPI at 7 tesla, with 16-fold acceleration using partial parallel 1113 imaging with application to high spatial and temporal whole-brain fMRI. Magnetic 1114 Resonance in Medicine, 63(5), 1144–1153. https://doi.org/10.1002/mrm.22361 Olivers, C. N. L., & Nieuwenhuis, S. (2005). The Beneficial Effect of Concurrent Task-1115 1116 Irrelevant Mental Activity on Temporal Attention. Psychological Science, 16(4), 265– 1117 269. https://doi.org/10.1111/j.0956-7976.2005.01526.x

1118 Olivers, C. N. L., & Nieuwenhuis, S. (2006). The beneficial effects of additional task load, 1119 positive affect, and instruction on the attentional blink. Journal of Experimental 1120 *Psychology: Human Perception and Performance*, 32(2), 364–379. 1121 https://doi.org/10.1037/0096-1523.32.2.364 1122 Peirce, J. W. (2007). Psychopy—Psychophysics software in Python. Journal of Neuroscience 1123 Methods, 162(1-2), 8-13. https://doi.org/10.1016/j.jneumeth.2006.11.017 1124 Posner, M. I., & Petersen, S. E. (1990). The Attention System of the Human Brain. Annual 1125 Review of Neuroscience, 13(1), 25–42. 1126 https://doi.org/10.1146/annurev.ne.13.030190.000325 1127 Power, J. D., Barnes, K. A., Snyder, A. Z., Schlaggar, B. L., & Petersen, S. E. (2012). Spurious 1128 but systematic correlations in functional connectivity MRI networks arise from subject 1129 motion. NeuroImage, 59(3), 2142–2154. 1130 https://doi.org/10.1016/j.neuroimage.2011.10.018 1131 Power, J. D., Mitra, A., Laumann, T. O., Snyder, A. Z., Schlaggar, B. L., & Petersen, S. E. 1132 (2014). Methods to detect, characterize, and remove motion artifact in resting state fMRI. 1133 NeuroImage, 84, 320–341. https://doi.org/10.1016/j.neuroimage.2013.08.048 1134 Raven, J., Raven, J. C., & Court, J. H. (1998). Manual for Raven's progressive matrices and 1135 vocabulary scales. Oxford: Oxford Psychologists. 1136 Raymond, J. E., Shapiro, K. L., & Arnell, K. M. (1992). Temporary suppression of visual 1137 processing in an RSVP task: An attentional blink? *Journal of Experimental Psychology*. 1138 *Human Perception and Performance*, 18(3), 849–860. 1139 Rensink, R. A. (2013). Perception and Attention. In D. Reisberg (Ed.), Oxford Handbook of 1140 Cognitive Psychology (pp. 97–116). New York: Oxford University Press.

1141 Rensink, R. A. (2015). A Function-Centered Taxonomy of Visual Attention. In P. Coates & S. 1142 Coleman (Eds.), Phenomenal Qualities: Sense, Perception, and Consciousness. (pp. 347– 1143 375). Oxford: Oxford University Press. 1144 Robertson, I. H., Manly, T., Andrade, J., Baddeley, B. T., & Yiend, J. (1997). 'Oops!': 1145 Performance correlates of everyday attentional failures in traumatic brain injured and 1146 normal subjects. Neuropsychologia, 35(6), 747–758. https://doi.org/10.1016/S0028-1147 3932(97)00015-8 1148 Rosenberg, M. D., Finn, E. S., Scheinost, D., Papademetris, X., Shen, X., Constable, R. T., & 1149 Chun, M. M. (2016). A neuromarker of sustained attention from whole-brain functional 1150 connectivity. Nature Neuroscience, 19(1), 165–171. https://doi.org/10.1038/nn.4179 1151 Rosenberg, M. D., Hsu, W.-T., Scheinost, D., Constable, T. R., & Chun, M. M. (2018). 1152 Connectome-based Models Predict Separable Components of Attention in Novel 1153 Individuals. *Journal of Cognitive Neuroscience*, 30(2), 160–173. 1154 https://doi.org/10.1162/jocn a 01197 1155 Rosenberg, M. D., Noonan, S., DeGutis, J., & Esterman, M. (2013). Sustaining visual attention 1156 in the face of distraction: A novel gradual-onset continuous performance task. *Attention*, 1157 Perception, & Psychophysics, 75(3), 426–439. https://doi.org/10.3758/s13414-012-0413-1158 X 1159 Rosenberg, M. D., Zhang, S., Hsu, W.-T., Scheinost, D., Finn, E. S., Shen, X., ... Chun, M. M. 1160 (2016). Methylphenidate Modulates Functional Network Connectivity to Enhance 1161 Attention. Journal of Neuroscience, 36(37), 9547–9557. 1162 https://doi.org/10.1523/JNEUROSCI.1746-16.2016

1163 Schaefer, A., Kong, R., Gordon, E. M., Laumann, T. O., Zuo, X.-N., Holmes, A. J., ... Yeo, B. 1164 T. T. (2017). Local-Global Parcellation of the Human Cerebral Cortex from Intrinsic 1165 Functional Connectivity MRI. Cerebral Cortex, 1–20. 1166 https://doi.org/10.1093/cercor/bhx179 Seli, P., Cheyne, J. A., Barton, K. R., & Smilek, D. (2012). Consistency of sustained attention 1167 1168 across modalities: Comparing visual and auditory versions of the SART. Canadian 1169 Journal of Experimental Psychology/Revue Canadienne de Psychologie Expérimentale, 66(1), 44–50. https://doi.org/10.1037/a0025111 1170 1171 Shen, X., Finn, E. S., Scheinost, D., Rosenberg, M. D., Chun, M. M., Papademetris, X., & 1172 Constable, R. T. (2017). Using connectome-based predictive modeling to predict 1173 individual behavior from brain connectivity. *Nature Protocols*, 12(3), 506–518. 1174 https://doi.org/10.1038/nprot.2016.178 1175 Shen, X., Tokoglu, F., Papademetris, X., & Constable, R. T. (2013). Groupwise whole-brain 1176 parcellation from resting-state fMRI data for network node identification. *NeuroImage*, 1177 82, 403–415. https://doi.org/10.1016/j.neuroimage.2013.05.081 1178 Skogsberg, K., Grabowecky, M., Wilt, J., Revelle, W., Iordanescu, L., & Suzuki, S. (2015). A 1179 relational structure of voluntary visual-attention abilities. Journal of Experimental 1180 *Psychology: Human Perception and Performance*, 41(3), 761–789. 1181 https://doi.org/10.1037/a0039000 1182 Smilek, D., Carriere, J. S. A., & Cheyne, J. A. (2010). Failures of sustained attention in life, lab, 1183 and brain: Ecological validity of the SART. *Neuropsychologia*, 48(9), 2564–2570. 1184 https://doi.org/10.1016/j.neuropsychologia.2010.05.002

1185 Stawarczyk, D., Majerus, S., Maj, M., Van der Linden, M., & D'Argembeau, A. (2011). Mind-1186 wandering: Phenomenology and function as assessed with a novel experience sampling 1187 method. Acta Psychologica, 136(3), 370–381. 1188 https://doi.org/10.1016/j.actpsy.2011.01.002 Tamber-Rosenau, B. J., Dux, P. E., Tombu, M. N., Asplund, C. L., & Marois, R. (2013). Amodal 1189 1190 Processing in Human Prefrontal Cortex. Journal of Neuroscience, 33(28), 11573–11587. 1191 https://doi.org/10.1523/JNEUROSCI.4601-12.2013 1192 Thomson, D. R., Ralph, B. C. W., Besner, D., & Smilek, D. (2015). The more your mind 1193 wanders, the smaller your attentional blink: An individual differences study. *Quarterly* 1194 Journal of Experimental Psychology, 68(1), 181–191. 1195 https://doi.org/10.1080/17470218.2014.940985 1196 Unsworth, N., & McMillan, B. D. (2014). Similarities and differences between mind-wandering 1197 and external distraction: A latent variable analysis of lapses of attention and their relation 1198 to cognitive abilities. Acta Psychologica, 150, 14–25. 1199 https://doi.org/10.1016/j.actpsy.2014.04.001 1200 Van Dijk, K. R. A., Hedden, T., Venkataraman, A., Evans, K. C., Lazar, S. W., & Buckner, R. L. 1201 (2010). Intrinsic functional connectivity as a tool for human connectomics: Theory, 1202 properties, and optimization. *Journal of Neurophysiology*, 103(1), 297–321. 1203 https://doi.org/10.1152/jn.00783.2009 1204 Webb, G. I., Sammut, C., Perlich, C., Horváth, T., Wrobel, S., Korb, K. B., ... Raedt, L. D. 1205 (2011). Leave-One-Out Cross-Validation. In C. Sammut & G. I. Webb (Eds.), 1206 Encyclopedia of Machine Learning (pp. 600–601). https://doi.org/10.1007/978-0-387-1207 30164-8 469

1208 Wojtowicz, M., Berrigan, L. I., & Fisk, J. D. (2012). Intra-individual Variability as a Measure of 1209 Information Processing Difficulties in Multiple Sclerosis. *International Journal of MS* 1210 Care, 14(2), 77–83. https://doi.org/10.7224/1537-2073-14.2.77 1211 Xu, J., Moeller, S., Auerbach, E. J., Strupp, J., Smith, S. M., Feinberg, D. A., ... Uğurbil, K. 1212 (2013). Evaluation of slice accelerations using multiband echo planar imaging at 3T. 1213 NeuroImage, 83, 991–1001. https://doi.org/10.1016/j.neuroimage.2013.07.055 1214 Yeo, B. T. T., Krienen, F. M., Eickhoff, S. B., Yaakub, S. N., Fox, P. T., Buckner, R. L., ... 1215 Chee, M. W. L. (2015). Functional Specialization and Flexibility in Human Association 1216 Cortex. Cerebral Cortex, 25(10), 3654–3672. https://doi.org/10.1093/cercor/bhu217 1217 Yeo, B. T. T., Krienen, F. M., Sepulcre, J., Sabuncu, M. R., Lashkari, D., Hollinshead, M., ... 1218 Buckner, R. L. (2011). The organization of the human cerebral cortex estimated by 1219 intrinsic functional connectivity. Journal of Neurophysiology, 106(3), 1125–1165. 1220 https://doi.org/10.1152/jn.00338.2011 1221 Yeo, B. T. T., Tandi, J., & Chee, M. W. L. (2015). Functional connectivity during rested 1222 wakefulness predicts vulnerability to sleep deprivation. NeuroImage, 111, 147–158. 1223 https://doi.org/10.1016/j.neuroimage.2015.02.018 1224 Yoo, K., Rosenberg, M. D., Hsu, W.-T., Zhang, S., Li, C.-S. R., Scheinost, D., ... Chun, M. M. 1225 (2017). Connectome-based predictive modeling of attention: Comparing different functional connectivity features and prediction methods across datasets. NeuroImage. 1226 1227 https://doi.org/10.1016/j.neuroimage.2017.11.010 1228 Zalesky, A., Fornito, A., & Bullmore, E. T. (2010). Network-based statistic: Identifying 1229 differences in brain networks. *NeuroImage*, 53(4), 1197–1207. 1230 https://doi.org/10.1016/j.neuroimage.2010.06.041

1232

1233

1234

1235

1236

1237

1238

1239

1240

1241

1242

1243

1244

1245

1246

1247

1248

1249

1250

1251

1252

1253

Legends Table 1. Schedule of tasks and data used in analysis. Participants were encouraged to take breaks between the tasks to prevent fatigue. With the exception of the Raven's Progressive Matrices test, each task was performed twice on non-successive days. Task domains and tasks were as follows. Attentional Blink: Visual Attentional Blink (VAB) and Auditory Attentional Blink (AAB). Sustained Attention: Visual Sustained Attention to Response Task (VSART), Auditory Sustained Attention to Response Task (ASART), and Gradual-onset Continuous Performance Task (GradCPT). Selective Attention: Attentional Network Task (ANT). Fluid Intelligence: Raven's Progressive Matrices test (Raven's). Figure 1. Experimental paradigm for the visual attentional blink (VAB) task. Participants identified a target and then detected a probe within a stream of distractors, responding when prompted at the conclusion of the stream. The target was a red letter, the probe was a white letter X, and distractors were other white letters. The auditory attentional blink (AAB) task was similar, save targets were complex tones, the probe was a high-pitched pure tone, and distractors were other pure tones. SOA = stimulus onset asynchrony. Table 2. Summary of behavioral metrics, including their task domain and a description of their calculation. Raw values for ANTerr, ANTrtvar and ANTcert were re-coded such that larger values indicated better task performance. Figure 2. Procedure for Connectome-based Predictive Model (CPM) construction in the current study (adapted from Shen et al., 2017). CPMs predict individual differences in behavior from

1255

1256

1257

1258

1259

1260

1261

1262

1263

1264

1265

1266

1267

1268

1269

1270

1271

1272

1273

1274

1275

1276

functional connectivity information. (a) Functional connectivity matrices and a behavioral score of interest for each participant were calculated. One pair was held out of model construction for each round of leave-one-out cross-validation (LOOCV) (Webb et al., 2011), (b) Functional connectivity edges were correlated with behavior across participants. (c) Edges that correlated most strongly, either positively or negatively, were selected. (d) Values from selected edges were summed separately for positive and negative network edges, yielding two network strengths for each participant. (e) A linear regression model relating (rank) network strengths to (rank) behavioral scores was computed. (f) The model was tested on a novel, out-of-sample participant (the individual left out in step (a)). After repeating steps a-f for each participant, the model was evaluated by correlating the predicted behavioral scores with the actual scores. Figure 3. Accuracy scores (probe hit rates) for the VAB and AAB. Note the substantial impairment for the short target-probe lags (1, 2, and 3) in each session, especially for the VAB. Error bars represent standard error of the mean (SEM). Table 3. Summary of behavioral data. Test-retest reliability (Spearman correlations across sessions) was high for most metrics and significant for all (ps < .002). Jarque-Bera tests indicated some significant departures from normality. Metrics are presented before re-coding. Metrics that were subsequently reversed so that larger values would indicate better task performance are marked with an asterisk (*). N = 73 for all metrics except AABresid (N = 71). Figure 4. Behavioral score distributions and correlations. (Above diagonal) Spearman correlation coefficients for pairs of behavioral metrics. Most significant correlations were found within task

1278

1279

1280

1281

1282

1283

1284

1285

1286

1287

1288

1289

1290

1291

1292

1293

1294

1295

1296

1297

1298

domains, but some metrics correlated across the Sustained and Selective Attention domains. In contrast, Attentional Blink and Fluid Intelligence metrics largely did not correlate with other metrics. Since the purpose of this analysis was to identify any behavioral relationships that might explain subsequent CPM results, no correction for multiple comparisons was applied. *p < .05, **p < .01. Red and blue shading indicates positive and negative relationships, respectively. (Diagonal) Histograms of behavioral data. The behavioral data had been re-coded so that larger values indicate better task performance for each measure. (Below diagonal) Scatterplots for each pair of behavioral metrics. Figure 5. Functional connectivity (FC) matrices. Each edge was Fisher-transformed, and the resulting z-scores were averaged across participants. Edges found to correlate with motion were set to zero. (a) VAB task FC matrix (VABFC). (b) Resting-state FC matrix (RSFC). VABFC and RSFC patterns were similar to one another and to other data sets (Yeo et al., 2011; Yeo, Tandi, et al., 2015). (c) Difference FC matrix (VABFC - RSFC), showing edges that were significant at p = .05, corrected for multiple comparisons using network-based statistics. Differences between FC matrices were small, though they notably included connections linking the Salience/Ventral attention and the Dorsal attention networks. The 419 parcellations from the FC matrices were matched to 17 network labels (Yeo et al., 2011) (green labels), from which they were aggregated into eight cortical groups (Yeo, Tandi, et al., 2015) and a subcortical group (blue labels, spelled out in full in (a)). Subcortical regions include the brain stem, accumbens area, amygdala, caudate, cerebellum, ventral diencephalon, hippocampus, pallidum, putamen, and thalamus. For the green labels, letters represent the networks within the corresponding group, e.g., a(Default

1300

1301

1302

1303

1304

1305

1306

1307

1308

1309

1310

1311

1312

1313

1314

1315

1316

1317

1318

1319

1320

1321

A), b(Default B), c(Default C), tp(TempPar), t(temporal pole in limbic region), o(orbital frontal cortex in limbic region), p(peripheral visual area), c(central visual area), and sc(subcortical). Figure 6. Behavioral predictions from CPMs. Each point represents a pair of Spearman's rank correlation coefficients (r_s) computed between observed and predicted behavioral scores for a given model type. (a) Predictions from vabCPMs, which were trained with VABFC and VABresid. Note the successful positive predictions for the VAB and Fluid Intelligence, but successful *negative* predictions for some Selective and Sustained Attention metrics. (b) Predictions from task-specific models. Many models could successfully predict behavioral performance, though results often varied greatly across the FC source. For both panels, the dark gray region indicates where r_s values are significant at the p = .01 level, and the lighter gray region indicates where r_s values are significant at the p = .05 level (uncorrected, with d.f. = 71). The r_s values and corresponding p-values are tabulated in Figure 6-1 in Extended Data. A standard edge selection threshold ($r_{\text{threshold}} = .232$, p = .05) was used for all models, though results were similar across a wide range of threshold values (Figure 6-2 and Figure 6-3 in Extended Data). Finally, as p-values from LOOCV procedures can be biased, we verified our results for the VAB using permutation testing; significance from this method and parametric approaches was consistent across edge selection thresholds (Figure 6-4 in in Extended Data). Figure 7. Comparison of predictions from the vabCPM and saCPM. Each point represents a pair of Spearman's rank correlation coefficients (r_s) computed between observed and predicted behavioral scores for a given model type. (a) Predictions from VABFC data. Note the saCPM's successful positive predictions for some Selective and Sustained Attention metrics, with a

1323

1324

1325

1326

1327

1328

1329

1330

1331

1332

1333

1334

1335

1336

1337

1338

1339

1340

1341

1342

1343

successful negative prediction for the VAB. As noted above, the prediction directions were reversed for the vabCPM; indeed, the prediction points fall close to the diagonal. (b) Predictions from RSFC data. The vabCPM and saCPM predictions went in opposite directions, and were generally similar to the predictions from the VABFC. For both panels, the dark gray region indicates where r_s values are significant at the p = .01 level, and the lighter gray region indicates where r_s values are significant at the p = .05 level (uncorrected, with d.f. = 71). The edge selection threshold ($r_{\text{threshold}}$) corresponded to p = .05 for all models. The r_s values and corresponding p-values for the saCPM are tabulated in Figure 7-1 in the Extended Data. (See Figure 6 for additional vabCPM details.) Figure 8. Percentage of edge overlap between networks from selected pairs of CPMs. Each taskspecific model was based on VABFC data (Model type C). Model pairs with statistically significant overlap (p < .05, FDR corrected) are indicated in bold. Positive network edges predicted better behavioral performance for their associated metric, whereas negative network edges predicted worse behavioral performance. Overlap between *similar* network edges, (a) positive-positive and (b) negative-negative. Sustained Attention (including the saCPM) and Selective Attention models overlap primarily on similar edges, as do Fluid Intelligence and the vabCPM. Overlap between opposing network edges, (c) negative-positive and (d) positivenegative. The vabCPM network primarily overlaps with Sustained Attention (including the saCPM) and Selective Attention on their opposing edges. Only task metrics that were significantly predicted by vabCPMs using VABFC information are shown. The full set of overlaps for all CPMs can be found in the Extended Data (Figure 8-1, 8-2, 8-3, and 8-4). Note

1345

1346

1347

1348

1349

1350

1351

1352

1353

1354

1355

1356

1357

1358

1359

1360

1361

1362

1363

1364

1365

1366

that the highest overlap value for any pairwise comparison, including each task metric predicted from VABFC and RSFC, was 26.5%. Figure 9. Anatomical locations of predictive attentional network edges. Each cell represents the number of shared connections between a pair of network groups, expressed as a percentage of the number of possible connections between that pair. For details about network groups, see Figure 5. (a) Positive and (b) negative VAB network edges. (c) Negative and (d) positive SA network edges. The panel order has been reversed for easier comparison with the opposing edges from the VAB network. (e) Positive VAB network edges and negative SA network edges (i.e. overlaps between (a) and (c)). (f) Negative VAB network edges and positive SA network edges (i.e. overlaps between (b) and (d)). Overlap percentages were numerically small, but included key network groups, particularly Default and Salience / Ventral Attention (Sal/VentAttn). Table 1-1. Detailed protocol from the full study from which the present data derives. Figure 6-1. Behavioral predictions from CPMs. Values indicate r_s and uncorrected, two-tailed pvalues from Spearman's partial rank correlation, computed between predicted and observed behavioral scores, controlled for motion. The p-value corresponding to each r_s was found by transforming the correlation coefficients to Student's t values by the partialcorr.m function in (MATLAB, 2014). Degrees of freedom for these tests were 66 for Models A and C, and 61 for Models B and D. (For AABresid, these d.f. values were 2 lower.) Note that all task scores had been re-oriented so that larger values indicate better task performance.

1368

1369

1370

1371

1372

1373

1374

1375

1376

1377

1378

1379

1380

1381

1382

1383

1384

1385

1386

1387

1388

Figure 6-2. CPM predictions for models trained with VAB functional connectivity and behavioral data (Model types A and B) across edge selection thresholds. Within each subplot, the v-axis represents Spearman's rank correlation values, r_s , computed between predicted and observed task performance, controlled for motion. Horizontal gray lines indicate the corresponding p = .01, .05, .10, .10, .05, .01 (top to bottom) uncorrected levels of significance from standard r-to-p conversions, d.f. = 71 (69 for AAB). The x-axis represents edge selection thresholds, $r_{\text{threshold}}$. The vertical gray line indicates the $r_{\text{threshold}}$ at the p = .05 level of significance, d.f. = 70 (68 for AAB) due to one left-out participant during training. X-axis labels at the top of each plot indicate the average number of edges selected across all leave-one-out iterations at the corresponding $r_{\text{threshold}}$ on the bottom x-axis. Figure 6-3. CPM predictions for models trained from task-specific behavioral data (Model types C and D) across edge selection thresholds. Within each subplot, the y-axis represents Spearman's rank correlation values, r_s , computed between predicted and observed task performance, controlled for motion. Horizontal gray lines indicate the corresponding p = .01, .05, .10, .10, .05,.01 (top to bottom) uncorrected levels of significance from standard r-to-p conversions, d.f. = 71(69 for AAB). The x-axis represents edge selection thresholds, $r_{\text{threshold}}$. The vertical gray line indicates the $r_{\text{threshold}}$ at the p = .05 level of significance, d.f. = 70 (68 for AAB) due to one leftout participant during training. X-axis labels at the top of each plot indicate the average number of edges selected across all leave-one-out iterations at the corresponding $r_{\text{threshold}}$ on the bottom xaxis.

1390

1391

1392

1393

1394

1395

1396

1397

1398

1399

1400

1401

1402

1403

1404

1405

1406

1407

1408

1409

1410

Figure 6-4. Permutation results of predicting VABresid data from VAB FC information, using the vabCPM (Model type A). The black line indicates Spearman's rank correlation values (r_s) computed between predicted and observed task performance, controlled for motion. Horizontal gray lines indicate the corresponding p = .01, .05, .10, .05, .01 (top to bottom) uncorrected levels of significance from standard r-to-p conversions, based on N = 73. Green sections indicate p < .05 level of significance from permutation testing. Magenta sections indicate $p \ge .05$ level of significance from permutation testing. Our analysis demonstrates a high level of consistency in the significance of r_s values between using standard r-to-p conversions (black line) and using permutation testing (green/magenta line) for the majority of edge selection thresholds. Figure 7-1. Results of applying the saCPM model to data in the present study. Values indicate r_s and uncorrected, two-tailed p-values from Spearman's rank correlation, computed between predicted and observed behavioral scores. The p-value corresponding to each r_s was found using standard r-to-p conversions, with d.f. = 71 (69 for AABresid). All task scores had been re-coded so that larger values indicate better task performance. Figure 8-1. Percentage of edge overlap for positive network edges between each pair of models. The value within each cell indicates the percentage of overlap between the pair of models on the corresponding x and y axes. Overlaps between the vabCPM (saCPM) with the other models are illustrated in the column (row) bounded in the red (black) of each plot. Axes labels also reflect the FC-Task pair used for training the task-specific model.

1412

1413

1414

1415

1416

1417

1418

1419

1420

1421

1422

1423

1424

1425

1426

1427

1428

Figure 8-2. Percentage of edge overlap for negative network edges between each pair of models. The value within each cell indicates the percentage of overlap between the pair of models on the corresponding x and y axes. Overlaps between the vabCPM (saCPM) with the other models are illustrated in the column (row) bounded in the red (black) of each plot. Axes labels also reflect the FC-Task pair used for training the task-specific model. Figure 8-3. Percentage of edge overlap between negative network (y-axis) and positive network (x-axis) edges. The value within each cell indicates the percentage of overlap between the pair of models on the corresponding x and y axes. Overlaps between the vabCPM (saCPM) with the other models are illustrated in the column (row) bounded in the red (black) of each plot. Axes labels also reflect the FC-Task pair used for training the task-specific model. Figure 8-4. Percentage of edge overlap between positive network (y-axis) and negative network (x-axis) edges. The value within each cell indicates the percentage of overlap between the pair of models on the corresponding x and y axes. Overlaps between the vabCPM (saCPM) with the other models are illustrated in the column (row) bounded in the red (black) of each plot. Axes labels also reflect the FC-Task pair used for training the task-specific model.

1429 Extended Data

Table 1-1. Detailed protocol from the full study from which the present data derives.

Session	Tasks/ Questionnaires	Session Description	Session	
			Dur (hr)	
1	Visual Attentional Blink (VAB)	Participants were briefed on the study protocol in	1	
	task	the first session and completed a target-probe		
		Attentional Blink (AB) task to obtain a measure of		
		baseline performance. Participants were only		
		eligible to continue with the subsequent sessions if		
		they obtained an average of 75% and above for		
		target discrimination accuracy in the first session.		
		Participants were not informed about their		
		performance until the end of all experimental		
		sessions.		
2	Visual AB and surprise-induced	Participants had to return to the laboratory for the	2.5	
	blindness (SiB) task (in-scanner)	second session 3 to 14 days after the first session.		
	Short-Sleep Questionnaire	The second session consisted of a 1.5 hr long fMRI		
	Visual Sustained Attention to	scan. Task order was fixed across participants.		
	Response task	No time limit was imposed on any of the		
	(12 mins)	questionnaires so participants could complete them		
	Attentional Network Task	at their own pace. All questionnaires were		
	(30 mins)	administered online using Qualtrics (Qualtrics,		
	Theories of Intelligence	2005).		
	Questionnaire			
3	Gradual-onset continuous	The third session took place the day after the	2	
	Performance Task	second session.		
	(15 mins)	Task order for the first 3 tasks was counterbalanced		

	Auditory Sustained Attention to	across participants.	
	Response task		
	(12 mins)		
	Auditory AB and Surprise-		
	induced deafness (SiD) task		
	Visual AB task	-	
	Visual AB task with colour-	•	
	salient training		
4	Visual AB and SiB task	The fourth session took place the day after the third	2
	Visual Sustained Attention to	session.	
	Response task	The Raven's Progressive Matrices task was an	
	(12 mins)	adapted version of the actual task that only	
	Attentional Network Task	consisted of 9 questions with no time limit given.	
	(30 mins)	Task order was fixed across participants.	
	Adult ADHD Self-Report Scale		
	Questionnaire		
	Online Dimensional Change Card		
	Sorting (DCCS) task		
	Raven's Progressive Matrices		
	Loss Aversion task		
5	Gradual-onset continuous	The fifth session took place the day after the fourth	2
	Performance Task	session.	
	(15 mins)	Decision-making tasks included a cups task and an	
	Auditory Sustained Attention to	explore-exploit task.	
	Response task	Task order for the first three computerized tasks	
	(12 mins)	and questionnaires were counterbalanced and	
	Auditory AB and Surprise-	randomized across participants.	
5	Adult ADHD Self-Report Scale Questionnaire Online Dimensional Change Card Sorting (DCCS) task Raven's Progressive Matrices Loss Aversion task Gradual-onset continuous Performance Task (15 mins) Auditory Sustained Attention to Response task (12 mins)	Task order was fixed across participants. The fifth session took place the day after the fourth session. Decision-making tasks included a cups task and an explore-exploit task. Task order for the first three computerized tasks and questionnaires were counterbalanced and	2

	induced deafness (SiD) task	-	
	Media-Multitasking Video Game	-	
	Questionnaire		
	Personality Inventory for DSM-5	-	
	for adults		
	Wender Utah Rating Scale	<u>-</u>	
	Musical background and	-	
	experience questionnaire		
	Decision-making tasks	_	
	Demographic Questionnaire	_	
6	Visual AB and surprise-induced	Session 6 was held between 5 to 14 days from the	1.5
	blindness (SiB) task (in-scanner)	second session and consisted of a 1.5 hr long fMRI	
		scan that used a protocol identical to that of the first	
		in-scanner session so as to investigate test-retest	
		reliability. Participants were encouraged to take	
		breaks between experimental blocks and between	
		tasks to alleviate fatigue.	

Figure 6-1. Behavioral predictions from CPMs. Values indicate r_s and uncorrected, two-tailed p-values from Spearman's partial rank correlation, computed between predicted and observed behavioral scores, controlled for motion. The p-value corresponding to each r_s was found by transforming the correlation coefficients to Student's t values by the partialcorr.m function in (MATLAB, 2014). Degrees of freedom for these tests were 66 for Models A and C, and 61 for Models B and D. (For AABresid, these $d_s f$. values were 2 lower.) Note that all task scores had been re-oriented so that larger values indicate better task performance.

		Mod	lel A	Mo	del B	Mo	del C	Mo	del D
Task	Train	VABFC- VABresid		VABFC- VABresid		VABFC-Task		RSFC-Task	
domain	Test	VABFC-Task		RSFC-Task		VABFC-Task		RSFC-Task	
	Task	$r_{\rm s}$	p	$r_{\rm s}$	p	$r_{\rm s}$	p	$r_{ m s}$	p
Attentional	VABresid	0.308	0.011*	0.123	0.338	0.308	0.011*	0.148	0.228
Blink	AABresid	0.04	0.748	-0.107	0.411	0.135	0.279	-0.053	0.672
G 1	VSARTdprime	-0.268	0.027*	-0.19	0.135	0.245	0.044*	0.062	0.617
Sustained Attention	ASARTdprime	-0.062	0.617	-0.212	0.096	-0.021	0.866	0.201	0.1
Attention	GradCPTdprime	0.058	0.637	-0.097	0.451	-0.265	0.029*	0.226	0.063
	ANTerr	-0.33	0.006*	-0.308	0.014*	0.346	0.004*	0.14	0.256
Selective	ANTrtvar	-0.36	0.003*	-0.414	0.001*	0.24	0.049*	0.232	0.057
Attention	ANTaert	0.051	0.682	0.141	0.269	0.033	0.787	-0.061	0.619
rttention	ANToert	0.063	0.609	0.128	0.318	0.124	0.313	0.032	0.793
	ANTcert	-0.252	0.038*	-0.333	0.008*	0.032	0.798	0.262	0.031*
Fluid Intelligence	RavensAcc	0.245	0.044*	0.224	0.077	-0.125	0.31	0.374	0.002*

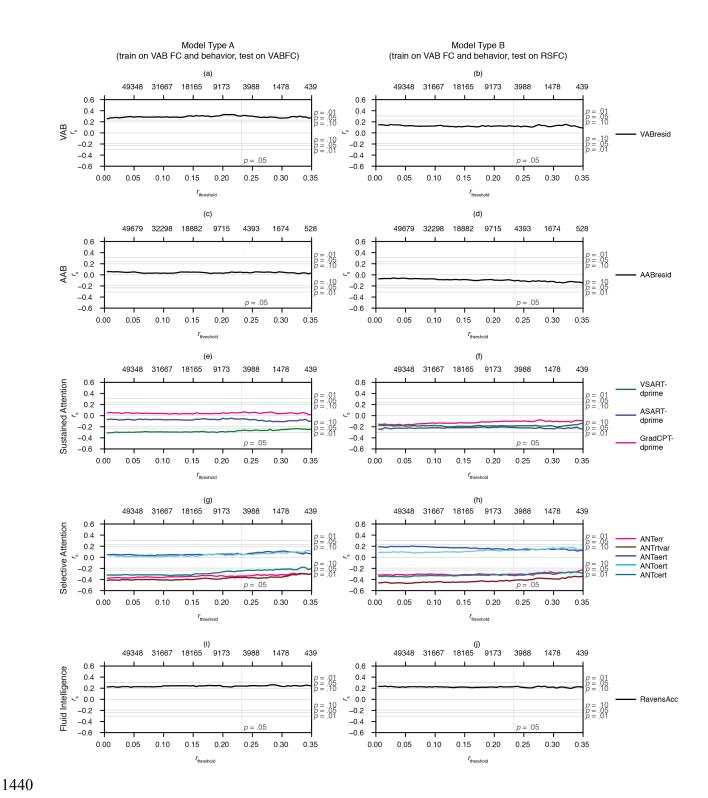


Figure 6-2. CPM predictions for models trained with VAB functional connectivity and behavioral data (Model types A and B) across edge selection thresholds. Within each subplot, the

y-axis represents Spearman's rank correlation values, r_s , computed between predicted and observed task performance, controlled for motion. Horizontal gray lines indicate the corresponding p = .01, .05, .10, .10, .05, .01 (top to bottom) uncorrected levels of significance from standard r-to-p conversions, $d_sf = 71$ (69 for AAB). The x-axis represents edge selection thresholds, $r_{threshold}$. The vertical gray line indicates the $r_{threshold}$ at the p = .05 level of significance, $d_sf = 70$ (68 for AAB) due to one left-out participant during training. X-axis labels at the top of each plot indicate the average number of edges selected across all leave-one-out iterations at the corresponding $r_{threshold}$ on the bottom x-axis.

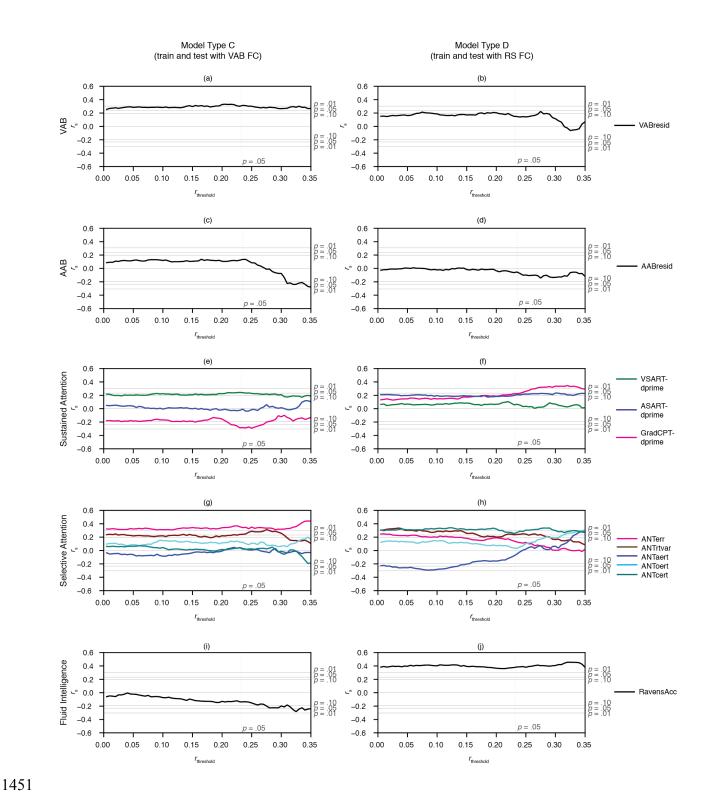


Figure 6-3. CPM predictions for models trained from task-specific behavioral data (Model types C and D) across edge selection thresholds. Within each subplot, the y-axis represents Spearman's

rank correlation values, r_s , computed between predicted and observed task performance, controlled for motion. Horizontal gray lines indicate the corresponding p = .01, .05, .10, .10, .05, .01 (top to bottom) uncorrected levels of significance from standard r-to-p conversions, df = 71 (69 for AAB). The x-axis represents edge selection thresholds, $r_{\text{threshold}}$. The vertical gray line indicates the $r_{\text{threshold}}$ at the p = .05 level of significance, df = 70 (68 for AAB) due to one left-out participant during training. X-axis labels at the top of each plot indicate the average number of edges selected across all leave-one-out iterations at the corresponding $r_{\text{threshold}}$ on the bottom x-axis.

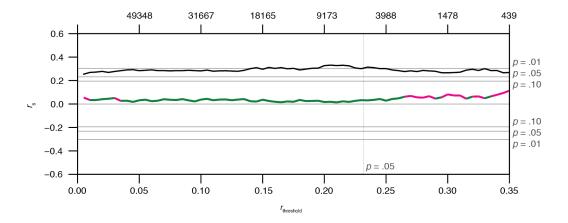


Figure 6-4. Permutation results of predicting VABresid data from VAB FC information, using the vabCPM (Model type A). The black line indicates Spearman's rank correlation values (r_s) computed between predicted and observed task performance, controlled for motion. Horizontal gray lines indicate the corresponding p = .01, .05, .10, .10, .05, .01 (top to bottom) uncorrected levels of significance from standard r-to-p conversions, based on N = 73. Green sections indicate p < .05 level of significance from permutation testing. Magenta sections indicate $p \ge .05$ level of significance from permutation testing. Our analysis demonstrates a high level of consistency

in the significance of r_s values between using standard r-to-p conversions (black line) and using permutation testing (green/magenta line) for the majority of edge selection thresholds.

Figure 7-1. Results of applying the saCPM model to data in the present study. Values indicate r_s and uncorrected, two-tailed p-values from Spearman's rank correlation, computed between predicted and observed behavioral scores. The p-value corresponding to each r_s was found using standard r-to-p conversions, with d_sf . = 71 (69 for AABresid). All task scores had been re-coded so that larger values indicate better task performance.

Task domain	Task	saCPM (VABFC)	saCPM (RSFC)	
		$r_{\rm s}$	p	$r_{\rm s}$	p
Attention Blink	VABresid	-0.235	0.045*	-0.155	0.191
Attention Billik	AABresid	0.049	0.684	-0.132	0.271
	VSARTdprime	0.278	0.018*	0.271	0.021*
Sustained Attention	ASARTdprime	0.117	0.323	0.290	0.013*
	GradCPTdprime	0.027	0.822	0.086	0.469
	ANTerr	0.322	0.005*	0.156	0.189
	ANTrtvar	0.329	0.005*	0.270	0.021*
Selective Attention	ANTaert	-0.057	0.629	-0.099	0.405
	ANToert	-0.101	0.397	-0.155	0.190
	ANTcert	0.224	0.057	0.384	0.001*
Fluid Intelligence	RavensAcc	-0.059	0.620	-0.229	0.052

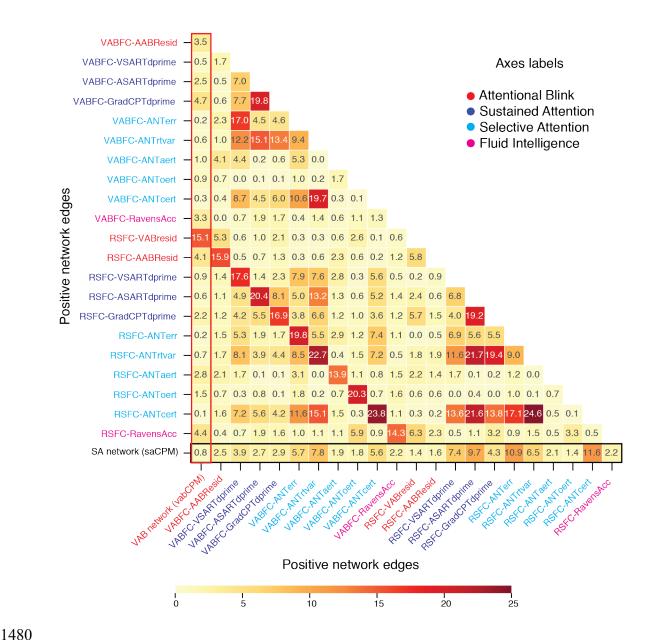


Figure 8-1. Percentage of edge overlap for positive network edges between each pair of models. The value within each cell indicates the percentage of overlap between the pair of models on the corresponding x and y axes. Overlaps between the vabCPM (saCPM) with the other models are illustrated in the column (row) bounded in the red (black) of each plot. Axes labels also reflect the FC-Task pair used for training the task-specific model.

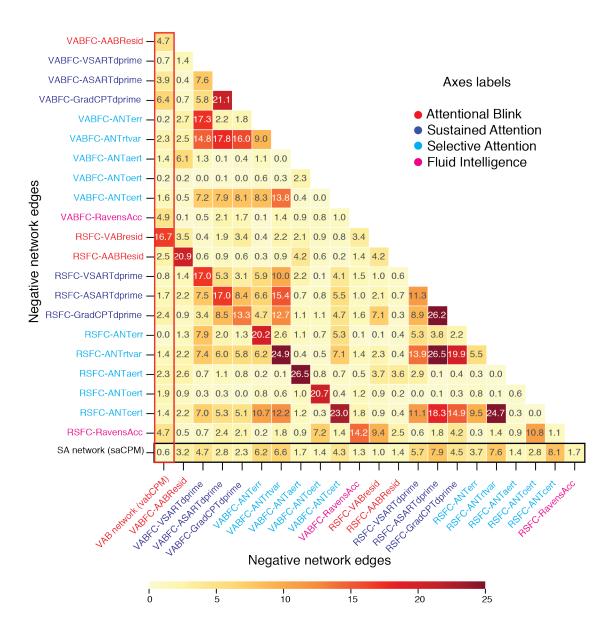


Figure 8-2. Percentage of edge overlap for negative network edges between each pair of models. The value within each cell indicates the percentage of overlap between the pair of models on the corresponding x and y axes. Overlaps between the vabCPM (saCPM) with the other models are illustrated in the column (row) bounded in the red (black) of each plot. Axes labels also reflect the FC-Task pair used for training the task-specific model.

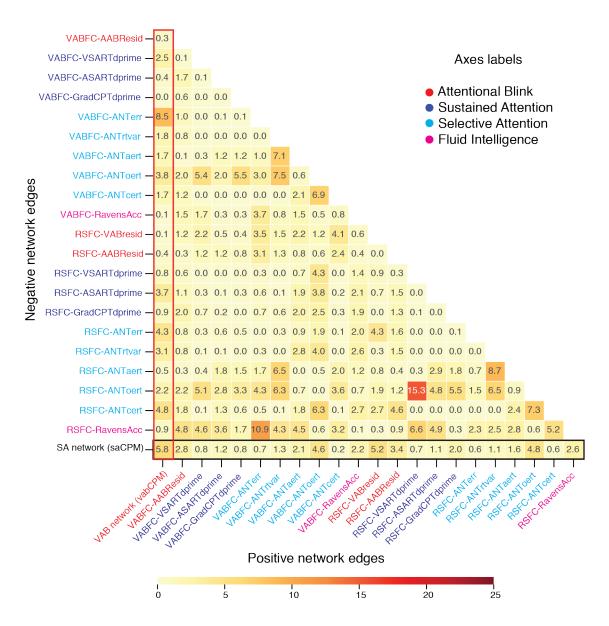


Figure 8-3. Percentage of edge overlap between negative network (y-axis) and positive network (x-axis) edges. The value within each cell indicates the percentage of overlap between the pair of models on the corresponding x and y axes. Overlaps between the vabCPM (saCPM) with the other models are illustrated in the column (row) bounded in the red (black) of each plot. Axes labels also reflect the FC-Task pair used for training the task-specific model.

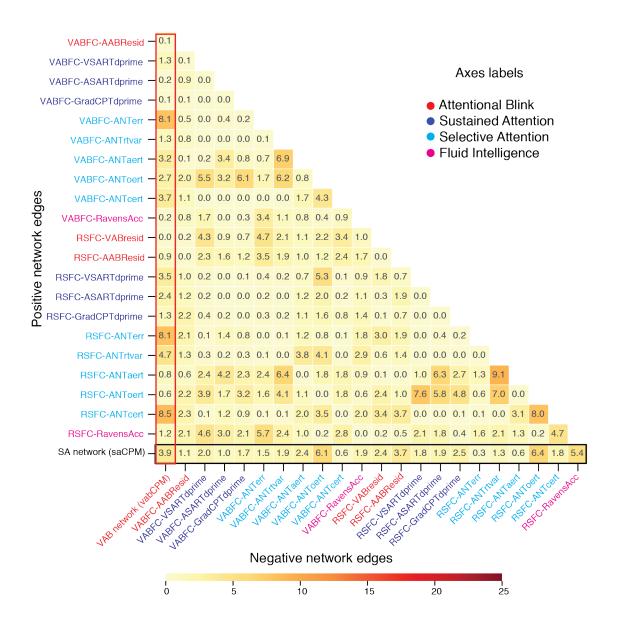


Figure 8-4. Percentage of edge overlap between positive network (y-axis) and negative network (x-axis) edges. The value within each cell indicates the percentage of overlap between the pair of models on the corresponding x and y axes. Overlaps between the vabCPM (saCPM) with the other models are illustrated in the column (row) bounded in the red (black) of each plot. Axes labels also reflect the FC-Task pair used for training the task-specific model.