1	CARDIORESPIRATORY FITNESS PREDICTS HIGHER INHIBITORY					
2	CONTROL IN PATIENTS WITH SUBSTANCE USE DISORDER					
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24 Abstract

Background: Impaired inhibitory control has been shown in individuals with substance 25 26 use disorder (SUD). Cardiorespiratory fitness has been described as a potential factor to improve inhibitory control, however, the benefits in individual with SUD are unclear. 27 Aims: To investigate the relationship between cardiorespiratory fitness with general and 28 drug-specific inhibitory control in individuals with SUD. Methods: Sixty-two male 29 participants under treatment for SUD performed a general and drug-specific inhibitory 30 31 control test (Go/NoGo) and a cardiorespiratory fitness test. Results: Cardiorespiratory fitness, age and years of drug use were inversely associated with reaction time for both 32 general and drug-specific inhibitory control. In addition, regression models show that 33 34 cardiorespiratory fitness predicts general and drug-specific inhibitory control adjusted 35 for age and time of drug use. However, cardiorespiratory fitness predicts equally both general and drug-specific inhibitory control. Conclusions: These findings suggest that 36 37 increasing cardiorespiratory fitness could provide benefits in inhibitory function of individuals with SUD. 38

Keywords: Drug addiction, aerobic exercise, alternative therapies, inhibitory control,cognition.

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1. Introduction

Chronic drug misuse is a worldwide public health problem. The Diagnostic and 46 Statistical Manual of Mental Disorders of the American Psychiatric Association (DSM-47 V 2013) classifies drug addiction as Substance Use Disorders (SUD) and Addictive 48 Disorders. Several harmful consequences of SUD have been shown, including 49 biological, mental, social, and financial problems (Beck & Heinz, 2013; Blanco-Gandía 50 et al., 2015). In addition, SUD is also associated with metabolic and cardiovascular 51 dysfunctions (Vongpatanasin, Taylor, & Victor, 2004; Whitman et al., 2017), increased 52 risk of death (Fischbach, 2017) and harmful consequences to others including 53 54 harassment, vandalism, physical aggression, and family problems (Nayak, Patterson, 55 Wilsnack, Karriker-Jaffe, & Greenfield, 2019). The main neural mechanism associated with the persistent use of psychoactive substances is the release of dopamine 56 57 neurotransmitter from the ventral tegmental area throughout the others areas in the reward system (i.e., nucleus accumbens, prefrontal cortex, striatum, and hippocampus) 58 that leads to increased feelings of pleasure (Leshner, 1997). Repetitive drug use results 59 in a decreased concentration of dopamine receptors (D2) in the reward system areas, 60 leading to greater drug tolerance, abstinence, negative affect and craving feelings while 61 62 provoking the urge to take higher doses (N. D. Volkow, Fowler, Wang, Baler, & 63 Telang, 2009; Zorrilla & Koob, 2019). Chronically, this drug misuse also yields lower metabolic activity in the prefrontal cortex (PFC), while impairing higher order cognitive 64 65 processes (i.e. executive functions) (Badre & Nee, 2018) and loss of control over drug seeking and behaviors (Goldstein & Volkow, 2011). 66

67 Inhibitory cognitive control is one component of executive functioning and can 68 be defined as the ability to inhibit habitual impulses or behaviors according to 69 advantageous future consequences (Feil et al., 2010). When it comes to addictive 70 behaviors, poorer performance in inhibitory control is related to impulsiveness, and may lead to further harmful consequences (Brewer & and Potenza, 2008). On the other hand, 71 72 greater performance on the inhibitory control may favor reduced impulsiveness (Bechara, 2005a) and improve decision making related to drug-seeking behaviors 73 (Bechara, 2005b; Shenoy & Yu, 2011). Furthermore, poorer response inhibition in 74 75 individuals with SUD is associated with difficulties in resisting the consumption of drug 76 substances especially when exposed to higher salient substance-related cues (Robinson & Berridge, 2008; Strickland et al., 2018; Weafer & Fillmore, 2012) increasing drug-77 78 seeking and drug-taking behaviors (Fillmore & Rush, 2002; Fu et al., 2008; Luijten, Littel, & Franken, 2011; Rubio et al., 2008; Smith, Mattick, Jamadar, & Iredale, 2014; 79 Volkow, Koob, & McLellan, 2016). Inhibitory control training have been suggested as 80 81 a method to improve inhibitory control towards addictive behaviors (Bos et al., 2019), 82 however, its efficacy is still under investigation.

83 In fact, different strategies have been proposed to treat drug addiction, such as pharmacotherapy, cognitive behavioral therapy and social support groups (Volkow & 84 Li, 2005). However, it has been described a rate of 60% of relapse chances after 85 86 treatment (Maisto, Pollock, Cornelius, Lynch, & Martin, 2003). Thus, new strategies are necessary to help the treatment of individuals with SUD. Regularly-performed physical 87 exercise (defined as planned and structured activity to cardiorespiratory fitness) 88 (Caspersen, Powell, & Christenson, 1985) has been shown to induce several benefits on 89 the human body and has been considered an important complementary tool to treat 90 91 different pathologies, including those associated with the neural functioning (Pedersen & Saltin, 2015; Sallis, 2009). The practice of physical exercise may improve 92 cardiorespiratory fitness (Garber et al., 2011), which is associated with reduced all 93 causes mortality risk (Lee et al., 2011; Blair, Kohl III, Barlow, 1995) and lower risk of 94

poorer health development (Blair, Cheng, & Holder, 2001). Physical exercise have already been suggested to benefit the treatment of patients with SUD by decreasing drug related behaviors, such as abstinence, consumption and craving (Buchowski et al., 2011; Wang, Wang, Wang, Li, & Zhou, 2014). However, few studies have investigated the benefits of physical exercise in cognitive functions using neurobiological markers in individuals with SUD. Crucially, this is important to develop physical activity treatment strategies aiming to improve cognitive function (Costa, Cabral, Hohl, & Fontes, 2019).

102 For instance, previous research have demonstrated that acute exercise can 103 decrease craving levels and abstinence feelings while improving inhibitory control (Wang, Zhou, & Chang, 2015; Wang, Zhou, Zhao, Wu, & Chang, 2016). Corroborating 104 with this idea, we have also demonstrated that one single session of cycling exercise 105 106 decreased drug craving feelings and increased PFC oxygenation in individuals with SUD, which was associated to higher inhibitory control performance (Grandjean da 107 108 Costa et al., 2017). Recently, we have showed that greater cardiorespiratory fitness predicted better cardiac autonomic activity in response to an induced stressful situation 109 in individuals with SUD (Cabral et al., 2019) while a 3-month running exercise program 110 111 (3 times/week) improved PFC oxygenation, cardiac autonomic regulation, and 112 inhibitory control in an alcoholic patient under treatment (Cabral et al., 2017). However, all of these studies have used general inhibitory cognitive measurements without drug 113 114 Color-Matching Stroop task), which might cues (e.g., not trigger the psychophysiological responses related to drug cue-reactivity (i.e., increased heart rate 115 116 and blood pressure, elevated cortisol and dopamine levels) (Papachristou, Nederkoorn, Havermans, Van Der Horst, & Jansen, 2012). To our knowledge, only two studies 117 showed the benefits of aerobic exercise on an inhibitory cognitive task using drug-118 specific pictures (i.e., drug-specific inhibitory control) (Wang et al., 2015; Wang, Zhu, 119

120 Zhou, & Chang, 2017a). However, these studies did not measure cardiorespiratory 121 fitness, which make it difficult to infer any associations between the chronic exercise 122 adaptations and cognitive response to drug-specific inhibition. Thus, we believe that, by 123 improving the understanding of the link between cardiorespiratory fitness and drug-124 specific inhibitory control, we may bring new insights regarding the treatment of 125 individuals with SUD.

Here we investigate the relationship between cardiorespiratory fitness and cognitive performance on a general and drug-specific inhibitory control task in individuals with SUD. We hypothesize that cardiorespiratory fitness would be associated with enhanced inhibitory control in individuals with SUD. We further predict that this association will be higher with drug-specific inhibitory control.

131 Methods

132 2.1 Participants

The study initially composed of 76 male adults under treatment for substance 133 use disorder at five different rehabilitation community settings that are free of 134 135 medications on their routine rehabilitation practice. To be eligible in this study, participants had to score the minimum of 24 points on the MMSE (Batista, Klauss, 136 137 Fregni, Nitsche, & Nakamura-Palacios, 2015) and be approved on the physical activity screening questionnaire (PAR-Q) (Roy J, 1988) by answering "No" to all questions. 138 There were no exclusion criteria for a specific substance. Sixty-two volunteers were 139 140 used in the final sample. 14 individuals were excluded since they did not reached minimum score of 24 points on Mini-Mental State Examination (MMSE) and/or were 141 142 not approved on the cardiac risk screening questionnaire (PAR-Q) (Roy J, 1988). The preferred substances of each participant (alcohol, nicotine, marijuana, cocaine/crack, 143

LSD, amphetamines, hypnotic sedatives, and ecstasy) were defined by applying the 144 145 Alcohol, Smoking and Substance Involvement Screening Test (ASSIST) (Henrique, 146 Iara Ferraz Silva; Micheli, De Lacerda, De Lacerda, & Formigoni, 2008). Among the participants, 33 were multiple drug users (25 alcohol/crack, 4 crack/marijuana, 2 147 Alcohol/Marijuana, 2 Crack/Marijuana/Alcohol) and 15 were crack/cocaine users, 3 148 were marijuana users, 11 were alcohol users. All participants met at least two criteria for 149 150 Substance Use Disorder from the DSM-V (Hasin et al., 2013). The study followed the Declaration of Helsinki standards and all participants signed the informed consent 151 approved by the local ethics committee 152

153 2.2 Experimental Design

154 This cross-sectional study was performed in three visits at different rehabilitation community settings with a minimum interval of 48 hours between each 155 156 visit. On the first visit, patients attended a lecture describing the harmful effects of 157 drugs on the brain and the benefits of physical exercise. At the end, the study purposes 158 and procedures were presented. On the second visit, the patients that agreed to participate in our study completed questionnaires for psychosocial state assessment, 159 160 physical activity readiness (PARQ) and drug-specific risk status. On the third visit, the participants completed the general and drug-specific inhibitory task followed by the 161 162 cardiorespiratory fitness test.

163 2.3 Measurements

164 2.3.1 Psychosocial Questionnaire

Psychosocial questionnaire for stress, anxiety and depression (DASS-21) was administered to the participants (Ribeiro, Honrado, & Leal, 2004). The DASS-21 consists of three scales of seven items each (total of 21 items). Each item consists of a 168 phrase or statement referring to negative emotional symptoms. For each sentence there 169 is a choice of four possible responses, presented on a Likert scale (e.g., 0 Did not apply 170 to me at all; 1 Applied to me to some degree, or some of the time; 2 Applied to me to a considerable degree or a good part of time; 3 Applied to me very much or most of the 171 172 time. The participants carried out the test in a quiet environment and in a reserved room. 173 The DASS-21 score was quantified through three scales: depression, anxiety and stress, 174 deriving scores for depression – dysphoria (two items); Discouragement, (two items); Life devaluation (two items); Auto depreciation (two items); Lack of interest or 175 176 involvement (two items); Anhedonia (two items); Inertia (two items). Anxiety excitation of the autonomous system (five items); Skeletal muscle effects (two items); 177 Situational anxiety (three items); Subjective experiences of anxiety (four items). Stress 178 - Difficulty in relaxing (three items); Nervous excitation (two items); Easily 179 180 agitated/upset (three items); Irritable/exaggerated reaction (three items); Impatience (three items). 181

182 2.3.2 ASSIST Questionnaire

This questionnaire, developed by the World Health Organization (Group, 2002), 183 184 assesses the risks and problems related to the use of alcohol, marijuana, cocaine/crack, LSD, sedatives, hallucinogens, heroin, Inhalants, opioids, and other drugs. The 185 186 questionnaire consists of seven questions that include a score and classifies the individual as being without the need for intervention (< 3 pts), needing a brief 187 intervention (> 4 pts), or a need of immediate intervention (> 27 pts) according to the 188 189 preferred drug. In this study, the ASSIST was used to identify the drug of preference, 190 since all volunteers were on a regimented treatment.

191 2.3.3 Cardiorespiratory Fitness Test

192 The participants initially had their weight and height assessed followed by the 193 multistage 20-m shuttle run (a progressive effort test) proposed by Leger (1988). This 194 test has been demonstrated to indirectly predict the VO₂max. A Meta-analysis showed that this test has been validated and has good reliability predicting VO₂max (Mayorga-195 Vega, Aguilar-Soto, & Viciana, 2015 (Léger, Mercier, Gadoury, & Lambert, 1988). In 196 197 this test, the participants are asked to run from one cone to another cone with a fixed 198 distance of 20m between them, and reversing the direction at each cone, thus returning to the opposite one. The running pace should occur according to the sound signals 199 200 emitted by an audio recording specifically for this test. Initial speed was 8.5 km/h, with 201 an increasing of 0.5 km/h each minute. An exception was given at the first minute, with 202 an increase of 1 km/h. As the test speed increases, the interval between the sound 203 signals decreases. The test was finished when the participant interrupted its 204 displacement by voluntary exhaustion (ratings of perceived exertion = 10 on CR-10 Borg Scale) (Borg, G., Linderholm, 1970) or had not been at least 2m apart from the 205 206 cone at the sound signal for two times, not necessarily consecutive times. The estimative of maximum oxygen consumption (VO₂max) was calculated (VO₂ =(Y) = -207 (4.4) + [(6.0*X)] (Y = ml/kg/min; X = velocity in km/h at the stage reached).208

209 2.3.4 Go/no-go inhibitory control task

An adapted go/no-go inhibitory control task was developed based on previous 210 study with food related images (Price, Lee, & Higgs, 2016). For the drug-specific trials, 211 212 images of marijuana, crack, cocaine and alcoholic beverages were used as no-go images. For the neutral trials, sports images were used as no-go images. In both cases, 213 214 the go images were bathroom objects. All drug specific images were taken from the 215 database "addiction pics" experimental for researchers 216 (https://pixabay.com/pt/photos/addiction/). Images with other objects were taken from 217 the BOSS normative database of photographs of objects (Brodeur, Dionne-Dostie, 218 Montreuil, & Lepage, 2010). Images were randomly presented on a computer screen 219 with a ratio of 20% (No Go) and 80% (Go). The presentation order of the drug-specific task and neutral task was counterbalanced. The individuals were told to press the space 220 221 key button as fast as possible whenever they see a bathroom object (Go) and to not 222 press the space key when the image was drug-cue or sports (neutral) images as relevant 223 (No Go). Two hundred images were presented in total. Each image was shown on the screen for 750ms. Intercalating each image, a blank screen (500ms) and another screen 224 with a "+" (fixing point for 500ms) were presented (Figure 1). Between each drug 225 image and sports images, bathroom images were randomly inserted (3, 4 or 5 between 226 227 every cue picture). The total test lasted approximately 7 minutes. Inhibitory control was 228 evaluated by the number of times the space bar was pressed incorrectly in no-go trials 229 (commission errors) and by reaction times (ms) on Go trials. Instructions were standardized and comprehension and willingness of the participants were assured by a 230 231 short preceding practice trial. Drugs images were selected according to the drug 232 preference of each subject. For example, if an individual was considered as a cocaine 233 and alcohol user (multiple drug user), images of both drugs would be used during the, 234 No Go trials.



Figure 1. General and drug-specific cognitive inhibitory task.





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244 2.4 Statistical analysis

245 The Shapiro-wilk test was used to verify data normality and Levene's test was performed to check data homogeneity. Parametric data are described as mean \pm standard 246 deviation and non-parametric variables as median (confidence interval) (see table 1). 247 248 Initial Spearman correlations were conducted for the variables of interest (commission 249 errors and reaction time of go no/go task, VO2max). Potential covariables (age, time of drug use, body mass index (BMI), days in abstinence, and DASS-21 scores) were also 250 included on the correlation analysis. Significant covariables were included in the main 251 252 regression analysis. Multiple linear regression analyses were applied to investigate the 253 independent contribution of VO₂max to the variance in inhibitory control parameters. Assumptions of equality of variance, independence, linearity and normality were 254 255 plotted, inspected, and verified using Studentized residuals. Multicollinearity was not 256 observed among any of the independent variables. Statistical significance was set at p<0.05 and was used bootstrapping power analysis values are described in CI 257 (Banjanovic & Osborne, 2016). We used the software SPSS® 22.0 for Windows (SPSS, 258 259 Inc., Chicago, IL).

260

261 **2. Results**

Bivariate correlations (2-tailed) showed that cardiorespiratory fitness (VO₂max) was inversely associated with age (r = -.46, p<.001), time of drug use (r = -.52, p<.001) and reaction time for general (r = -.43, p<.001), and drug-specific inhibitory control (r = -.47, p<.001). No associations were found between VO₂max with BMI (r = -.04, p<.709), days in abstinence (r = -.02, p<.851), and DASS-21 scores (stress: r = -.02,

267	p<.832; anxiety: $r =11$, p<.360; depression: $r =13$, p<.286). In addition, no
268	association was found between commission errors and cardiorespiratory fitness (general
269	inhibitory control errors (r =16, p<.189), and drug-specific inhibitory control errors (r
270	=07, p<.580). Therefore, two multiple regression analyses were run for predicting
271	reaction time on the general and drug-specific trials of the go/no-go task (see Table 2).
272	Results show that cardiorespiratory fitness (VO2max) predicts the reaction time for
273	general [F (_{6; 3}); (t = -3.0; β =41; p = .003] and drug-specific inhibitory control [F (_{8;}
274	3); t = -3.2; β =42; p < .002] when we adjusted the analyses for age and time of drug
275	use.

Sample general characteristics				
	Median (CI) (n=62)			
Age	34 (31.9 - 36.4)			
BMI (kg/m ²) [#]	24.6 ± 2.9			
VO ² _{max} (ml/kg/min)	39 (38.4 – 42)			
Time of drug use (yrs)	13 (12.9 – 18.3)			
DSM-V (pts)	5.7 (5.2 – 6.2)			
Abstinence use (days)	105 (103 – 173.6)			
Anxiety (a.u)	28.5 (24.7 – 37.2)			
Depression (a.u)	23.8 (22.3 - 33.8)			
Stress (a.u)	23.8 (23.8 – 34.6)			
Commission errors (general)	2 (1.7 - 2.8)			
Commission errors (specific)	1 (1.2 - 2.1)			
RT (general) (ms) [#]	482.5 ± 56.3			
RT (specific) (ms) [#]	498.9 ± 61.5			

Drug preferences	N (%)
Crack/Cocaine	15 (24.1)
Marijuana	3 (4.8)
Alcohol	11 (17.7)
Crack/Alcohol	25 (40.3)
Crack/Marijuana	4 (6.4)
Alcohol/Marijuana	2 (3.2)
Crack/Marijuana/Alcohol	2 (3.2)

Table I. *Describes the sample general characteristics and cognitive performance on the*

277 *go/no-go task.*

278

279 Legend. RT (Reaction time) #Mean and standard deviation; BMI: body mass index; DSM-V:280 Substance Use Disorders.

Table 2. *Regression analyses between cardiorespiratory fitness and reaction time of*

282 general and specific inhibitory control adjusted by age and time of drug use.

Pradictors	Reaction time (General)		Reaction time (Drug-specific)	
riediciois	β (CI)	ΔR^2	β (CI)	ΔR^2
Model 1		.13		.16
Age	.34* (09 – 54)		.43** (15 - 60)	
Time of Drug use	.03 (04 – 49)		.00 (03 – 54)	
Model 2		.25		.27
Age	.23 (09 – 54)		.32 (15 – 60)	
Time of Drug use	10 (04 – 49)		13 (03 – 54)	
VO ₂ max	41** (- 62 19)		42** (- 66 23)	

283 *p<0.05; **p<0.01

284

285 **Discussion**

286 The present study sought to investigate whether cardiorespiratory fitness was associated with performance on a general and drug-specific inhibitory control task in 287 288 individuals with SUD. We found that higher cardiorespiratory fitness predicted better performance on the cognitive test when controlling for age and length of time using the 289 290 drug. However, cardiorespiratory fitness was not associated with stress, depression or 291 anxiety scores and we could not find differences between general and specific inhibitory 292 control performance predicted by cardiorespiratory fitness. Nevertheless, our results indicate the importance of cardiorespiratory fitness in cognitive control deficits in 293 individuals with SUD. 294

Physical exercise is fundamentally important in the evolutionary history of 295 human beings (Bramble and Lieberman, 2004), aiding survival in hunter-gatherer 296 297 societies. Findings have demonstrated that levels of cardiorespiratory fitness during human evolution are correlated with increases in brain size, with the PFC being the 298 299 most developed neural area when compared to any other primates (Raichlen & Polk, 2012). The cognitive functions of the PFC have been suggested to play role in exercise 300 tolerance and performance (Robertson & Marino, 2016), with a cohort study revealing a 301 302 preventive effect of exercise on drug use. In fact, McElrath and colleagues showed that 303 individuals that performed physical exercise regularly were less likely to consume alcohol, cigarettes, and marijuana (McElrath, O'Malley, & Johnston, 2011). Thus, we 304 305 speculate that our results showing that individuals with higher cardiorespiratory fitness have better inhibitory control may be related to these preventive effects on drug use. 306 Individuals with SUD have been shown to display impairments of inhibitory control 307 and PFC function during exercise (Grandjean da Costa et al., 2017), which could 308 hamper exercise adherence and, consequently, the development of cardiorespiratory 309 fitness. Previous research has also shown that exercise practice can be an effective 310

complementary treatment for SUD rehabilitation (Weinstock, Farney, Elrod, 311 312 Henderson, & Weiss, 2018). To date, most of the research that has analyzed the effects 313 of physical exercise on behavioral outcomes in drug addiction, have not attempted to explore the mechanism of change (Wang et al., 2015; Wang, Zhu, Zhou, & Chang, 314 2017a). We have shown that cardiorespiratory fitness is related to inhibitory control in 315 individuals with SUD, suggesting that the benefits of physical exercise in improving 316 317 cardiorespiratory fitness may be related to improvements in inhibitory control. However, due to the cross-sectional nature of this study we cannot establish causality 318 and longitudinal research is needed to confirm this assumption. 319

Typically, the commission errors during Go/No go tasks are the main parameter 320 321 for inhibitory control performance. In our study, few commission errors were made, and 322 may have induced a floor effect whereas the recorded data was unable to discriminate among the participants' cognitive performance (Catts, Petscher, Schatschneider, 323 324 Bridges, & Mendoza, 2009). Thus, we believe that this may help to explain why we did not find associations between cardiorespiratory fitness and commission errors. 325 However, we did find an association between cardiorespiratory fitness and reaction 326 327 times, which has also been indicated as a parameter of cognitive performance 328 (Papachristou et al., 2012). In go/no-go tasks, slower reaction time in the go trials can indicate increased difficulty in inhibiting the response on the no go trials via a speed-329 330 accuracy trade-off (Bogacz, Wagenmakers, Forstmann, & Nieuwenhuis, 2010). In fact, previous studies used mean reaction time on the go condition as measure of inhibitory 331 332 control performance (Smith et al., 2011; Wang, Zhu, Zhou, & Chang, 2017b). Thus, reaction time can also be used as an efficiency index of cognitive performance (Hirose et 333 al., 2012), and we infer that better inhibitory control in individuals with higher 334

cardiorespiratory fitness, as they performed the same amount of correct answers whilehaving faster reaction time.

337 Few studies have investigated the relationship between physical exercise with general and drug-specific inhibitory control in individuals with SUD. Our results found 338 that cardiorespiratory fitness independently predicts reaction time for general and drug-339 340 specific inhibitory control, which indicates the possible benefits of having higher 341 cardiorespiratory fitness on cognition of individuals with SUD. Several studies have 342 demonstrated the benefits of cardiorespiratory fitness on cognition in different 343 populations (Colcumbe et al., 2006; Hillman, Erickson, & Kramer, 2008; Kramer, 2009). These benefits might also be transposed to SUD individuals as shown in our 344 study. These findings have been discussed in terms of the effects of exercise on 345 neuroplasticity in the prefrontal cortex, which may enhance executive functions (Maass 346 et al., 2016). Therefore, we highlight the importance of investigating the effects of 347 348 exercise on cognition in individuals with SUD to provide further understanding of the rehabilitation alternative methods. 349

However, we could not find differences between the associations of 350 351 cardiorespiratory fitness and performances on the general and drug-specific inhibitory control. One possible explanation for this finding could be that the VO₂ max 352 353 measurement in our study is an indirect measure and may not be a reliable indicator of 354 cardiovascular fitness in individuals SUD. It may be the case that the drug-specific 355 inhibitory control test is only sensitive to laboratory-based tests of VO₂max. Moreover, 356 despite we have used specific images of drug cues for the drug preference of each individual, we speculate that these cues did not produce a physiological cue-reactivity 357 response in order to difficult the inhibition process. Further studies measuring 358 359 physiological markers (i.e. Heart rate variability, skin conductance, electrocortical

360 activity) could help to evaluate such responses. For instance, studies have shown that cue-reactivity responses induce feelings of cravings that activate brain frontal areas and 361 362 predict relapse in individuals with SUD (Wilson, Sayette, & Fiez, 2004). On the other hand, randomized control trials have failed to translate inhibitory control training to 363 changes in addictive behaviors interventions (Bos et al., 2019; Jones et al., 2018). Thus, 364 further studies are necessary to test if there is a difference in physiological and 365 366 behaviors responses between drug-specific inhibitory and general inhibitory control in the drug-specific go/no-go task proposed in our study. 367

368 We do acknowledge that our study is potentially limited by the heterogeneity of the sample in terms of preferred drug and the small sample for each subgroup of drug 369 use (cocaine/crack, alcohol, marijuana). This may have affected the cue-reactive 370 371 responses during the drug-specific inhibitory control due to the different action mechanisms promoted by the drugs. However, studies have shown that all of these 372 373 drugs impair PFC function and inhibitory control performance (Herbsleb et al., 2013; Luijten et al., 2014; Nora D. Volkow et al., 2016), which was the investigative focus of the 374 present study. In this preliminary research study, we also did not have a healthy control 375 376 group, which could have helped to further compare the benefits of cardiorespiratory 377 fitness in inhibitory control. However, we believe that our findings are useful to guide future research in this area. We also did not use any neurobiological measurement (e.g., 378 379 MRI, EEG), which could have masked some cerebral differences between the two inhibitory tasks. Future research should try to replicate our study but using some neural 380 381 instrument. Based on our findings, further longitudinal studies with specific samples of SUD are necessary to test the efficacy of the drug-specific go/no-go task used in this 382 study as an indicator of cognitive changes associated to improvements in 383 cardiorespiratory fitness. 384

385 Clinical Implications

Our results suggest that improved cardiorespiratory fitness might be beneficial to 386 387 inhibitory control in individuals with SUD. However, to date, there is no specific exercise prescription for patients with SUD. Thus, we suggest that professionals from 388 389 therapeutic community settings apply the prescription based on the American College of 390 Sports Medicine (ACSM) guidelines (Garber et al., 2011) which includes light to 391 moderate intensity exercise for 150 minutes or more per week. Moreover, we believe 392 that the PFC impairments in individuals with SUD might have a disadvantage on 393 internal exercise regulatory process that may impact the adherence to exercise programs (Grandjean da Costa et al, 2019). We suggest that activities that promote higher 394 distraction from internal cues, such as adding music, outdoors environment and group 395 training might be important strategies to increase affective feelings while exercising. As 396 a complement to the training program, future research could use the drug-specific 397 398 cognitive task to understand the changes induced by long term exposure to chronic 399 exercise programs.

400

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403

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