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1 Images depicting human pain increase exercise-induced pain and impair endurance cycling
2 performance

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23 Abstract

24 Observing others in pain can enhance pain intensity. The current study aimed to investigate
25 whether viewing images of others in pain influences exercise-induced pain (EIP) and
26 endurance cycling performance. Twenty-one recreational cyclists attended five laboratory
27 visits. The first two visits involved the measurement of participants' maximal aerobic
28 capacity and familiarized participants to the fixed power (FP) and 16.1 km cycling time trial
29 (TT) tasks. The FP task required participants to cycle at 70% of their maximal aerobic power
30 for 10-minutes. In the subsequent three visits, participants performed the FP and TT tasks
31 after viewing pleasant, painful or neutral images. Participants rated the subset of painful
32 images as significantly more painful than the pleasant and neutral images; with no difference
33 in the pain ratings of the pleasant and neutral images. In the FP task, ratings of EIP were
34 higher in the painful image condition compared to the pleasant condition, while no
35 differences in EIP were observed between the pleasant and neutral conditions or the neutral
36 and painful conditions. Perceived exertion, heart rate (HR) and blood lactate (B[La]) during
37 the FP task did not differ across conditions. In the TT, performance did not differ between the
38 pleasant and neutral conditions. However, TT performance was reduced after viewing painful
39 images compared to neutral or pleasant images. Despite these performance changes, heart
40 rate HR, B[La], perceived exertion and EIP did not differ between the three conditions. These
41 results suggest that viewing painful images prior to exercise decreases TT performance and
42 increases pain during a fixed intensity exercise task.

43 Key Words: Exercise-induced pain; compassionate hyperalgesia; time trial;
44 performance; empathy.

45		Abbreviations
46	EIP	Exercise Induced Pain
47	FP	Fixed Power
48	TT	Time Trial
49	HR	Heart Rate
50	B[La]	Blood Lactate
51	RPE	Rating of Perceived Exertion
52	IAPS	International Affective Picture System
53	PO	Power Output

54 Images depicting human pain increase exercise-induced pain and impair endurance cycling
55 performance

56 Intense exercise causes a noxious environment in the muscle which typically elicits
57 exercise-induced pain (EIP) (Dannecker & Koltyn, 2014). Tolerance of this sensation has
58 been associated with performance in endurance exercise tasks, with those better able to
59 tolerate EIP producing superior performance (Astokorki & Mauger, 2016). Indeed, the
60 experimental manipulation of EIP has been shown to affect exercise performance. For
61 example, acetaminophen and caffeine have both been shown to reduce EIP and increase
62 endurance cycling performance (Gonglach, Ade, Bemben, Larson, & Black, 2015; Mauger,
63 Jones, & Williams, 2010). Based on this evidence, it is suggested that pain may act as a
64 regulator of work rate during endurance exercise tasks, influencing the athlete's ability to
65 access a physiological reserve (Noakes, 2011; Swart et al., 2009). While these interventions
66 aimed at decreasing EIP have resulted in improvements in performance, interventions that
67 instead increase EIP may provide novel insights into the role of pain as a regulator of
68 endurance exercise performance.

69 Viewing others in pain has been shown to induce the vicarious experience of pain in
70 the observer, termed "*synaesthesia for pain*" (Fitzgibbon, Giummarra, Georgiou-Karistianis,
71 Enticott, & Bradshaw, 2010), and increase one's sensitivity to pain (Godinho et al., 2012;
72 Khatibi, Vachon-Preseu, Schrooten, Vlaeyen, & Rainville, 2014; Loggia, Mogil, &
73 Bushnell, 2008). This psychophysical phenomenon, whereby pain sensitivity is increased
74 when viewing others in pain, is also referred to as *compassional hyperalgesia* and has been
75 observed in both men and women (Godinho et al., 2012). Loggia et al. (2008) reported that
76 when participants observed videos of others in pain, they offered higher pain intensity and
77 pain unpleasantness ratings in response to noxious thermal stimuli. These changes in pain

101 pain manipulation and participant characteristics between the current study and that of
102 Mauger et al. (2010), we conducted a second sample size estimation based on an effect size
103 of 0.34, calculated from the findings of Godinho et al. (2012) who reported hyperalgesic
104 effects of observing images of human pain. This calculation resulted in an estimated
105 minimum sample of 17 participants required to detect an effect with 80% power ($\alpha = 0.05$).

106 Due to the large effect sizes observed in both studies, we sought to recruit a larger
107 sample than the minimum calculated. Therefore, 21 male ($n = 13$) and female ($n = 8$)
108 recreational cyclists (>3 h exercise per week) were recruited for participation (see Table 1).
109 The participation opportunity was advertised using flyers distributed throughout the local
110 community and university. Recruitment also occurred online, through social media platforms.
111 Volunteers were encouraged to contact the primary researcher to register their interest in
112 participating.

113 Participants were given an overview of the study, describing the requirements for their
114 involvement. Specifically, participants were informed that a series of potentially distressing
115 images would be viewed to examine the effects on exercise performance. In order to reduce
116 the possibility of response bias, participants were not informed of the hypothesised effects of
117 the intervention on pain and performance. The participants were aware that all data would be
118 unidentifiable and that they had the right to withdraw from the study at any time. Following
119 this, they were asked to complete the inclusion/exclusion criteria checklist and then asked to
120 sign an informed consent form. Individuals were excluded if they self-reported any of the
121 following: pregnancy; lifetime history of psychological disorders; history of fainting;
122 bleeding disorders (e.g. haemophilia); types I or II diabetes; lifetime history of clinically
123 significant or unstable medical, neuropsychiatric, or chronic pain disorders; history of
124 substance abuse or dependence; history of brain disorders, surgery, tumour or heart disease;

125 intracranial metal implantation; chronic use of medications that affect the central nervous
126 system. Participants were asked to avoid vigorous exercise 24 hours before the laboratory
127 visits, and to refrain from the ingestion of alcohol, caffeine and analgesics 48 h, 8 h and 6 h
128 before any experimental visit. The experimental protocol was approved by the local Ethics
129 Committee.

130 **Procedure**

131 The study followed a within groups, randomised and counterbalanced design,
132 including three experimental conditions (painful, neutral, pleasant). Participants reported to
133 the laboratory on five separate occasions, with each session separated by 2-5 days. The
134 ordering of experimental conditions was randomised by a researcher not otherwise involved
135 in the study. An online tool was used to achieve this randomisation
136 (<https://www.randomizer.org/>).

137 **Session 1.** Participants were first given standard instructions for the use of the
138 numeric pain rating scale (Cook, O'Connor, Eubanks, Smith, & Lee, 1997) and rating of
139 perceived exertion (RPE) scale (Borg, 1998) to be used throughout all physical performance
140 measures. To gain an understanding of participants' aerobic capacity, a cycling-based
141 incremental ramp test was then conducted. After a 5 min warm-up at 75 Watts, the
142 incremental ramp protocol started at 100 Watts and increased by 30 Watts every 2 min until
143 volitional exhaustion or when cadence dropped 5 RPM below the participants' self-selected
144 cadence (Astokorki & Mauger, 2016). Throughout the test and all subsequent cycling tasks,
145 power output (PO) was monitored using a cycle ergometer (Velotron, Racermate, Seattle,
146 WA) and heart rate (HR) was continuously displayed using a Polar Vantage XL HR monitor
147 (Polar Electro Oy, Kempele, Finland). Pain intensity and perceived exertion were recorded 15
148 s before the end of each 2 min stage. Prior to the test, the ergometer was adjusted for each

149 participant and the setting was recorded to allow reproduction at each subsequent visit for
150 both the fixed power (FP) and TT tasks. Expired gases were assessed using an online gas
151 analyser (Cortex Biophysik GmbH, Leipzig, Germany) throughout the test. Following a rest
152 period of 30 min, participants then performed a familiarisation of the FP (see FP Procedure)
153 and TT (see TT Procedure) tests to be used in subsequent sessions. A 5 min rest period
154 separated the FP and TT tasks.

155 **Session 2.** The purpose of the second session was to again familiarise participants to
156 the exercise performance tasks. Specifically, participants attended the laboratory and first
157 completed the FP task. During this familiarisation session, self-selected cadence was
158 monitored to allow for this cadence to be replicated across the subsequent experimental
159 sessions. Following a 5 min rest period, the TT was completed.

160 **Sessions 3-5.** Sessions 3-5 formed the experimental data collection phase of the study.
161 In these sessions, participants first sat quietly for 10 min before viewing 15 either painful,
162 neutral or pleasant images (see Images Procedure), depending on their assigned condition.
163 Immediately after viewing the images, participants were positioned on the cycling ergometer
164 and instructed to complete the FP task. After a 5 min rest period, during which time
165 participants viewed a further 10 painful, neutral or pleasant images, the TT was completed.
166 To reduce the risk of bias, the experimenter involved in the collection of performance and
167 pain data during the FP and TT tasks was not present during the presentation of images. This
168 ensured that they were blinded to the participants' assigned condition.

169 At the completion of the TT in the final session, participants were thanked for their
170 involvement and invited to ask any questions that they had about the study.

171 **Measures**

172 **EIP.** Pain experienced during the cycling tasks was assessed using the scale
173 developed by Cook et al. (1997). This scale required participants to verbally report their
174 perceived pain levels according to a 12-point scale. Standardised instructions (see Cook et al.,
175 2004) were provided to participants to before each cycling task to ensure proper use of the
176 scale. Importantly, participants were asked to rate the feelings of pain and discomfort
177 experienced in the legs and not use the rating as an expression of perceived exertion.

178 **Perceived exertion.** Perceived exertion was assessed using Borg's (1998) 6-20 RPE
179 scale. Prior to each cycling task, participants were asked to report their perceived exertion as
180 the amount of effort required to drive the limbs.

181 **FP procedure.** Prior research has shown that during self-paced exercise tasks (e.g. a
182 TT), participants alter their work rate to maintain a fixed progression in perceptual
183 parameters (Mauger, 2014; Tucker, 2009). Therefore, an FP task was used to examine
184 potential changes in pain, perceived exertion, and physiological parameters of HR and blood
185 lactate concentration (B[La]) when cycling at a fixed PO, across the three experimental
186 conditions. The task required participants to cycle at a fixed power equivalent to 70% of their
187 maximal aerobic power (determined in the incremental ramp task) for 10 min. A fingertip
188 sample of blood was collected at rest, 5 min and 10 min during the FP task for the analysis of
189 B[La]. Pain, perceived exertion and HR were assessed at 2 min intervals throughout the FP
190 task. Scripted verbal encouragement was provided throughout.

191 **TT procedure.** In the TT, participants were instructed to complete a 16.1 km cycling
192 TT on the cycle ergometer (Veltron, Racermate, Seattle, WA), as previously described
193 (Mauger et al. 2010). During the self-paced TT, perceived exertion and pain were assessed
194 every km, using the scales described above. HR was also measured at the end of each km of
195 the TT. Every 4 km, a fingertip sample of blood was taken to assess B[La] concentration. To

196 ensure consistency across sessions and participants, scripted verbal encouragement was
197 offered throughout the TT. At the completion of the 16.1 km, participants completed a 10 min
198 cool-down at a self-selected intensity.

199 **Images procedure.** Seventy-five images were categorised into three subsets (painful,
200 pleasant and neutral). The painful images subset ($n = 25$) included images of athletes in pain
201 (e.g. suffering a severe injury), while the pleasant images subset ($n = 25$) showed athletes
202 enjoying cycling, exercising or in enjoyable situations. The neutral subset of images ($n = 25$)
203 included complex visual stimuli with no overtly emotional content (e.g. a natural scene).
204 Where possible, images (40%) were taken from the International Affective Picture System
205 (IAPS), with IAPS arousal and valence values used to categorise images into the painful,
206 pleasant and neutral subsets (Lang, Bradley, & Cuthbert, 1997) (see Supplementary material
207 for image codes). As a limited number of relevant images (i.e. pain occurring in sporting
208 situations) were present on the IAPS database, the remaining images were obtained from the
209 internet (images available upon request).

210 Images were presented to participants in a PowerPoint presentation, following
211 protocols described elsewhere (Boggio, Zaghi, & Fregni, 2009; Godinho et al., 2012).
212 Briefly, participants viewed a computer screen at a comfortable distance of approximately 60
213 cm. A standardised set of instructions were used to explain the procedure of the study, and
214 participants were informed that a series of images would be viewed. The three subsets of
215 images were presented on separate visits in a counterbalanced and randomised order. Each
216 subset presented a total of 25 images (15 images were viewed before the FP test and 10
217 images before the TT). Each image was viewed for 30 s. After viewing the image for 25 s,
218 participants were asked to provide a rating of their pain affect in response to the question
219 “how do you feel while viewing the image?” (1 = comfortable/no pain, 9 =

220 uncomfortable/pain) (Boggio et al., 2009). The number of images and duration of
221 presentation were selected to produce an overall time-on-task, including an opportunity to
222 provide a pain affect rating, that was approximately consistent with previous research
223 (Boggio et al., 2009). The ordering of the images within each subset was kept consistent
224 across participants.

225 **Statistical Analysis**

226 Prior to statistical analysis, assumptions were checked for each statistical test. Data
227 relating to completion time for the TT violated the assumption of normality. The reciprocal
228 transformation was used to normalise the distribution of TT completion time data, which was
229 then analysed using a repeated measures analysis of variance (ANOVA), with the factor of
230 Condition (painful, neutral, pleasant). Pairwise comparisons with a Bonferroni correction
231 were used to follow up significant differences in TT completion time across conditions. The
232 same analyses were also conducted using non-transformed data, giving the same results.
233 Therefore, to aid in interpretation, results presented here relate to the analysis of non-
234 transformed TT completion time data.

235 In cases where the assumption of sphericity was violated, the Greenhouse-Geisser
236 epsilon was corrected. Mean ratings of pain affect for image subsets were analysed using a
237 repeated measures ANOVA with the factor of Condition (painful, neutral, pleasant), and
238 pairwise comparisons with a Bonferroni correction were used to further investigate
239 significant main effects across the three levels. HR (beats per minute (bpm)), RPE and EIP
240 during the FP task were analysed using 3 (Condition: painful, neutral, pleasant) \times 5 (Time: 2
241 min, 4 min, 6 min, 8 min, 10 min) repeated measures ANOVAs. A 3 (Condition: painful,
242 neutral, pleasant) \times 3 (Time: rest, 5 min, 10 min) ANOVA was used to analyse B[La]
243 measured during the FP task. For the TT task, PO, HR, RPE and EIP were analysed using 3

244 (Condition: painful, neutral, pleasant) \times 16 (Distance: 1km, 2km, 3km, 4km, 5km, 6km, 7km,
245 8km, 9km, 10km, 11km, 12km) repeated measures ANOVAs.. B[La] during the TT task was
246 analysed using a 3 (Condition: painful, neutral, pleasant) \times 4 (Distance: 4km, 8km, 12km,
247 16km) repeated measures ANOVA. Appropriate follow-up pairwise comparisons with
248 Bonferroni corrections were used to further investigate significant main effects on the
249 Condition factor.

250 Statistical analysis was performed using the statistical package SPSS version 22 for
251 Windows programs (SPSS Inc., Chicago, IL, USA). Descriptive data are reported as means \pm
252 SD. Statistical significance was accepted when $p < 0.05$. Cohen's d and partial eta squared
253 (η_p^2) are reported as estimates of the effect size.

254 Results

255 Image Ratings

256 Ratings of pain affect differed across the three experimental conditions, $F(1.105,$
257 $22.094) = 257.87, p = .000, \eta_p^2 = .928$ (Figure 1). Specifically, participants provided
258 significantly higher pain affect ratings for the subset of painful images (6.061 ± 1.301)
259 compared to both the pleasant images ($1.248 \pm 0.303, p = .000, d = 5.095$) and neutral images
260 ($1.328 \pm 0.401, p = .000, d = 4.917$). No significant difference was observed between pain
261 affect ratings of the pleasant and neutral images ($p = .929, d = .225$).

262 FP Task

263 **HR.** Mean HR in the FP task did not differ across the conditions, $F(2, 40) = .360, p$
264 $= .700, \eta_p^2 = .018$. There was a main effect for Time, $F(1.740, 34.798) = 79.521, p = 000, \eta_p^2$
265 $= .799$, but no significant interaction effect during the FP test, $F(8, 160) = .781, p = .620, \eta_p^2$
266 $= .038$. See Table 2 and Figure 2a for data on HR during the FP task.

267 **B[La]**. No significant main effect of Condition was observed for B[La] during the FP
268 task, $F(2, 40) = 1.927, p = .159, \eta_p^2 = .088$. There was a main effect for Time, $F(1.288,$
269 $25.761) = 58.435, p = .000, \eta_p^2 = .745$, but no significant interaction effect was found, $F(4,$
270 $80) = 1.270, p = .289, \eta_p^2 = .060$. See Table 2 and Figure 2b for data on B[La] during the FP
271 task.

272 **Perceived exertion**. No significant main effect of Condition was observed for
273 perceived exertion in the FP task, $F(2, 40) = 2.788, p = .074, \eta_p^2 = .122$. There was a main
274 effect for Time, $F(1.154, 23.079) = 32.688, p = .000, \eta_p^2 = .620$, but no significant
275 interaction effect was found, $F(3.594, 71.874) = .856, p = .485, \eta_p^2 = .041$. See Table 2 and
276 Figure 2c for data on perceived exertion during the FP task.

277 **EIP**. There was a main effect of Condition for EIP, $F(2, 40) = 4.363, p = .019, \eta_p^2 =$
278 $.179$. Pairwise comparisons with a Bonferroni correction showed a significant difference in
279 EIP between the pleasant and painful image conditions ($p = .033, d = .263$). No significant
280 difference between the pleasant and neutral image conditions ($p = 1.00, d = .062$), or between
281 the neutral and painful image conditions was found ($p = .232, d = .206$). There was a
282 significant main effect for Time, $F(1.290, 25.808) = 30.606, p = .000, \eta_p^2 = .605$, but no
283 significant interaction effect for EIP, $F(3.834, 76.674) = .805, p = .521, \eta_p^2 = .039$. See Table
284 2 and Figure 2d for data on EIP during the FP task.

285 **TT Task**

286 **Completion time**. The completion time for the TT differed across conditions, $F(2,$
287 $40) = 9.223, p = 0.001, \eta_p^2 = .316$. Pairwise comparisons revealed that participants performed
288 a significantly faster TT in the pleasant condition (29 min 38 s \pm 4 min 35 s; $p = .005, d =$
289 $.140$) and the neutral condition (29 min 39 s \pm 3 min 34 s; $p = .009, d = .136$) compared to the

290 painful condition (30 min 19 s \pm 5 min 7 s). There was no significant difference in TT
291 completion time between the neutral condition and the pleasant condition ($p = 1.000$, $d =$
292 $.004$).

293 **PO.** Mean PO in the TT differed across the three conditions, $F(2, 40) = 5.536$, $p =$
294 $.008$, $\eta_p^2 = 2.17$) (Figure 3a). Pairwise comparisons employing a Bonferroni correction
295 showed a significantly higher PO in the pleasant condition (209.236 Watts \pm 68.980 Watts; p
296 $= .007$, $d = .131$) and the neutral condition (207.633 Watts \pm 63.956; $p = .024$, $d = .112$)
297 compared to the painful condition (200.218 Watts \pm 68.392 Watts). There was no significant
298 difference between the neutral and pleasant conditions ($p = 1.000$, $d = .024$). There was also a
299 main effect for Distance, $F(3.160, 63.195) = 11.283$, $p = .000$, $\eta_p^2 = .361$, but no interaction
300 effect between Condition and Distance was found, $F(30, 600) = .847$, $p = .702$, $\eta_p^2 = .041$,
301 shown in Figure 3b.

302 **B[La].** The ANOVA revealed a significant main effect of Condition for B[La] during
303 the TT, $F(2, 40) = 5.724$, $p = .007$, $\eta_p^2 = .223$. Pairwise comparisons employing a Bonferroni
304 correction showed no significant difference in mean B[La] between pleasant and painful
305 image conditions ($p = .145$, $d = .556$). There was also no significant difference between the
306 pleasant and neutral image conditions ($p = 1.000$, $d = .194$), or between the neutral and
307 painful image conditions ($p = .113$, $d = .454$). There was a main effect for Distance, $F(1.505,$
308 $30.103) = 20.332$, $p = .000$, $\eta_p^2 = .504$, but no significant interaction effect was found, F
309 $(3.219, 64.374) = 1.961$, $p = .125$, $\eta_p^2 = .089$. See Table 3 and Figure 4a for data on B[La]
310 during the TT.

311 **HR.** A significant difference in the mean HR between conditions during the TT was
312 observed, $F(2, 40) = 4.502$, $p = .017$, $\eta_p^2 = .184$. However, pairwise comparisons employing
313 a Bonferroni correction uncovered no significant difference in HR between the pleasant and

314 neutral conditions ($p = 1.00$, $d = .088$), the pleasant and painful conditions ($p = .095$, $d =$
315 $.408$), nor the painful and neutral conditions ($p = .170$, $d = .292$) . There was a significant
316 main effect for Distance, $F(2.392, 47.849) = 43.410$, $p = .000$, $\eta_p^2 = .685$, but no significant
317 interaction effect was found, $F(30, 600) = .572$, $p = .969$, $\eta_p^2 = .028$ See Table 3 and Figure
318 4b for data on HR during the TT.

319 **Perceived exertion.** No significant differences in RPE were observed across the three
320 conditions, $F(2, 40) = .249$, $p = .781$, $\eta_p^2 = .012$. However, there was a main effect for
321 Distance, $F(1.840, 36.793) = 92.197$, $p = .000$, $\eta_p^2 = .822$, but no significant interaction
322 effect, $F(30, 600) = 1.344$, $p = .106$, $\eta_p^2 = .063$. See Table 3 and Figure 4c for data on
323 perceived exertion during the TT.

324 **EIP.** Pain experienced during the TT did not differ across conditions, $F(2, 40) =$
325 1.865 , $p = .168$, $\eta_p^2 = .085$. Irrespective of condition, pain did change throughout the TT, F
326 $(1.511, 30.220) = 89.387$, $p = .000$, $\eta_p^2 = .817$, but no significant Distance by Condition
327 interaction effect was found, $F(30, 600) = 1.380$, $p = .088$, $\eta_p^2 = .065$. See Table 3 and
328 Figure 4d for data on EIP during the TT.

329 Discussion

330 Pain experienced during exercise is thought to have an impact on endurance exercise
331 performance (Mauger, 2014). In support, factors that attenuate EIP have been shown to
332 enhance exercise performance (Mauger et al., 2010). It is, therefore, possible that endurance
333 exercise performance may be negatively impacted by factors that increase the pain of
334 exercise. Compassional hyperalgesia is a phenomenon whereby the observation of pain in
335 others results in increased pain sensitivity, typically assessed through pain intensity ratings
336 given on a numerical rating scale (Godinho et al., 2012). Therefore, the current study aimed

337 to assess whether viewing images of others in pain impacts on EIP and endurance cycling
338 performance. It was hypothesised that images of others in pain, presented immediately before
339 exercise, would increase perceived pain during exercise and reduce exercise performance.
340 This hypothesis was partially supported, with results indicating that pain experienced during
341 an exercise task, which required participants to cycle at a fixed PO, was elevated after
342 viewing images of other athletes in pain compared with viewing pleasant images. Also, as
343 hypothesised, viewing images of others in pain resulted in longer time-to-completion and
344 lower PO in a cycling TT.

345 The observed change in perceived pain intensity resulting from viewing others in pain
346 aligns with the compassionate hyperalgesic effect (Godinho et al., 2012). Indeed, the
347 hyperalgesic effect of viewing others in pain has been consistently observed across a range of
348 pain modalities, including acute thermally-induced pain (Loggia et al., 2008) and noxious
349 electrical stimulation (Godinho et al., 2012; Godinho et al., 2006; Khatibi et al., 2014;
350 Mailhot, Vachon-Pressseau, Jackson, & Rainville, 2012). However, the current findings are
351 novel as they are the first to describe how the perceived intensity of EIP can also be subject to
352 manipulation by observing others in pain. This is an important advancement on existing
353 knowledge, as it has been suggested that EIP represents a distinct psychophysiological
354 experience to that of pain induced through other experimental methods (Angius, Hopker,
355 Marcora, & Mauger, 2015).

356 The use of the FP task in the current study presents as an important methodological
357 consideration in the assessment of changes in EIP. The FP test was designed to assess
358 whether the intervention resulted in a change in perceptual response (i.e. pain and RPE) for a
359 given exercise intensity. The subsequent TT was then performed to assess whether the
360 intervention would elicit a change in endurance performance. This experimental design was

361 necessary to fully explore the research question, because it has previously been shown that
362 whilst changes in perceptual responses to an intervention can be observed in fixed intensity
363 exercise, in self-paced endurance exercise (such as a TT), participants tend to maintain a
364 fixed progression in perceptual parameters at the expense of changes to work rate (Mauger,
365 2014; Mauger et al., 2010; Tucker, 2009). Thus, in the current study, the TT provided a true
366 measure of self-paced endurance performance, whilst the FP task helped demonstrate that the
367 intervention elicited changes in EIP intensity. Importantly, both tasks induced levels of pain
368 that were consistent with the EIP reported in previous research involving similar maximal
369 and submaximal cycling tasks (Astokorki & Mauger, 2017; Astorino, Cottrell, Talhami
370 Lozano, Aburto-Pratt, & Duhon, 2012; Mauger et al., 2010; Motl, Gliottoni, & Scott, 2007).
371 Future research attempting to explore the role of pain in the regulation of endurance exercise
372 performance should consider implementing a similar methodological approach as that used
373 here. Indeed, in an editorial by Hettinga et al. (2017), it is suggested that the use of both FP
374 and TT tasks may be required to provide a comprehensive understanding of the regulation of
375 endurance performance.

376 In addition to the changes in EIP observed in the FP task, viewing images of others in
377 pain also reduced performance in the 16.1 km cycling TT. These changes in performance
378 occurred without any change in pain experienced during the TT. These findings can be
379 interpreted in the context of the observed increases in performance following the
380 administration of analgesic substances. For example, Mauger et al. (2010) reported increased
381 performance in a cycling TT without changes in perceived pain after the administration of
382 acetaminophen; a finding subsequently replicated in repeated sprint cycling (Foster, Taylor,
383 Christmas, Watkins, & Mauger, 2014), running (Pagotto, Paradisis, Maridaki, Papavassiliou,
384 & Zacharogiannis, 2018) and isometric contractions (Morgan, Bowtell, Vanhatalo, Jones, &

385 Bailey, 2018). Similarly, the analgesic effect of caffeine consumption has been shown to
386 produce performance improvements in a cycling task (Gonglach et al., 2015). Together, these
387 findings provide indirect support for the putative role for pain in the regulation of work-rate
388 during exercise tasks.

389 Whilst it is tempting to attribute the observed increase in EIP during the FP task and
390 subsequent changes in TT performance to compassionate hyperalgesia, alternative
391 explanations should be carefully considered. Research exploring compassionate hyperalgesia
392 has offered greater insight into the phenomenon, suggesting a more complex interpretation of
393 the current findings may be warranted. In particular, the hyperalgesia experienced after
394 viewing others in pain appears to be dependent on an empathic response being elicited in the
395 observer. After experimentally manipulating the degree of empathy that an observer feels for
396 an actor, Loggia et al. (2008) found that those with higher empathy for an actor appearing to
397 be in pain, displayed stronger compassionate hyperalgesia. Similarly, dispositional optimism
398 has also been shown to correlate with compassionate hyperalgesia, with highly empathic
399 individuals showing strong hyperalgesic responses to observing others in pain (Mailhot et al.,
400 2012). In fact, those scoring lowest on dispositional optimism experienced *reduced* pain
401 sensitivity (i.e. analgesia) as a result of viewing pain in others. Without a measure of
402 empathy, we cannot conclude as to whether participants in the current study empathised with
403 those depicted in the painful images. As a consequence, the observed changes in EIP cannot
404 be conclusively attributed to compassionate hyperalgesia.

405 Alternative explanations for the current findings should, therefore, be considered. One
406 potential explanation relates to the likely impact of the painful and pleasant images on affect.
407 Previous research has reported that the induction of a positive affective state decreases pain
408 sensitivity, while negative affect results in increased pain sensitivity (Meagher, Arnau, &

409 Rhudy, 2001; Meng et al., 2012; Zelman, Howland, Nichols, & Cleeland, 1991). These
410 findings support motivational priming theory, which describes how the activation of an
411 appetitive or aversive motivational state can enhance the amplitude of responses to the
412 subsequent presentation of congruent stimuli (Lang, 1995). Therefore, it is possible that the
413 change in pain evoked by the presentation of painful images was due to the induction of
414 negative affect and the activation of an aversive motivational drive. Similarly, it is possible
415 that the pleasant images induced an appetitive motivational state which decreased EIP in the
416 FP task relative to the painful condition. Without an assessment of the valence dimension of
417 affect, it is beyond the scope of the current study to partition the possible effects of
418 motivational priming and compassionate hyperalgesia. This presents as a notable limitation of
419 the current study. We do, therefore, encourage future research to measure changes in affect
420 resulting from experimental manipulations so as to allow for the application of motivational
421 priming theory to exercise performance settings.

422 In addition to the likely influence of the image intervention on affect, the current
423 findings may be explained by a mental fatigue or ego depletion effect. A recent meta-analysis
424 by Giboin and Wolff (2019) reported impaired endurance performance after the induction of
425 a mentally fatigued or ego depleted state. This state is typically achieved through prior mental
426 exertion in a challenging cognitive task (e.g. Stroop test) and is thought to impair subsequent
427 performance by elevating perceived exertion (Marcora, Staiano, & Manning, 2009). In the
428 current study, no such change in perceived exertion was observed in the FP task, suggesting
429 that the observed decrement in TT performance was not due to the induction of mental
430 fatigue or ego depletion. Also, without a measure of mental fatigue or ego depletion, it is
431 unclear whether the images presented in the painful image condition induced such a state.
432 Indeed, issues with operationally defining mental fatigue and ego depletion (Lurquin &

433 Miyake, 2017) and the failure to replicate the phenomena (Hagger et al., 2016), highlight the
434 need for additional research into these constructs.

435 Ratings of perceived exertion during the FP and TT tasks were similar to those
436 reported in previous research (Mauger et al., 2010; Williams et al., 2015). However, it is
437 noteworthy that the intervention resulted in no changes to perceived exertion but a significant
438 change to EIP in the FP task. This provides further evidence that EIP and perceived exertion
439 can be separated, provided participants are given adequate instruction and familiarisation
440 with the two scales (Pageaux, 2016). Of further note, is that despite no apparent effect of the
441 intervention on perceived exertion, performance of the TT was affected by the image
442 intervention. This supports the argument that endurance performance can be moderated
443 without any change in perceived exertion. Such findings question the emphasis placed on
444 perceived exertion as the sole perceptual regulator of work-rate during endurance exercise, as
445 proposed by the psycho-biological model (Marcora, 2008). Indeed, the current findings fail to
446 support Staiano, Bosio, de Morree, Rampinini, and Marcora (2018) and their suggestion that
447 EIP may influence exercise performance indirectly, by altering perceived exertion.

448 It is noteworthy that no differences in EIP and cycling TT performance were observed
449 between the pleasant and neutral image conditions. The lack of a performance improvement
450 in the pleasant condition is inconsistent with previous research reporting increases in cycling
451 performance following the induction of pleasant affective states using IAPS images (Coudrat
452 et al., 2014; Jaafar et al., 2015). However, more recent research by di Fronso et al. (2020)
453 suggests a more complex effect, with some participants showing performance improvements
454 after viewing pleasant images and others displaying improved performance after unpleasant
455 images. Whether these individual differences in responses to the pleasant images were also
456 evident in the findings presented above is beyond the scope of the current study. However,

457 given the likely affective consequences of the images used in the current study, the lack of a
458 measure of affect presents as a potential limitation. As suggested above, future research
459 should extend on the current findings by including measures of affect.

460 Several other limitations should be considered when interpreting the current findings.
461 First, the current study recruited male and female recreational cyclists. It is possible that the
462 findings reported here may not generalise to other populations. For example, athletes and
463 non-athletes have been shown to exhibit differences in their pain responses (Flood,
464 Waddington, Thompson, & Cathcart, 2016; Tesarz, Schuster, Hartmann, Gerhardt, & Eich,
465 2012). Similarly, research has reported differences in the pain responses of contact and non-
466 contact athletes (Ryan & Kovacic, 1966), strength and endurance athletes (Assa, Geva,
467 Zarkh, & Defrin, 2019) and males and females (Greenspan et al., 2007). While it is beyond
468 the scope of the current study to compare the effect of images depicting human pain across
469 these sample populations, the limits to the generalisability of the current findings should be
470 acknowledged and explored in future research. Sex-related differences, in particular, should
471 be addressed given the observed differences in pain responses to the presentation of IAPS
472 images between men and women (Meagher et al., 2001).

473 In the current study, the three experimental conditions were presented in a randomised
474 order. A single blinded design was also used, with the primary researcher unaware of the
475 assigned image condition. To further reduce the potential for bias, researchers used
476 standardised instructions for the presentation of the pain and perceived exertion measures and
477 provided scripted verbal encouragement throughout the FP and TT tasks. However, the nature
478 of the intervention made it impossible to blind the participants to their assigned order of
479 conditions. Therefore, it is possible that participants were biased in their responses. We
480 encourage future research to address this potential limitation through alternative

481 methodological approaches, such as the use of subliminal priming, as used by Godinho et al.
482 (2012).

483 Participants provided higher pain affect ratings in response to the images presented in
484 the painful condition compared to the neutral and pleasant conditions. The measure used to
485 assess responses to the images matched that used by Boggio et al. (2009) to determine the
486 emotional pain and discomfort experienced after viewing images of others in pain. While
487 responses to this measure indicated increased pain affect in the pain condition, alternative
488 measures should be considered in future research. In particular, pain affect is widely assessed
489 using pain unpleasantness numerical rating scales (Rainville, 2002) and multidimensional
490 tools such as the McGill Pain Questionnaire (Melzack, 1987).

491 **Conclusion**

492 In the current study, viewing images of others in pain increased the pain experienced
493 during a cycling task of fixed intensity and decreased exercise performance in a cycling TT.
494 These findings have significant implications for our understanding of the role of pain in
495 exercise performance, indicating that factors that produce hyperalgesic effects, such as
496 viewing pain in others, can be detrimental to performance in fatiguing exercise.

497

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636

637 **Figure Captions**

638 Figure 1. Differences in pain affect while viewing images in the painful, neutral and pleasant
639 conditions. * denotes significant difference between conditions.

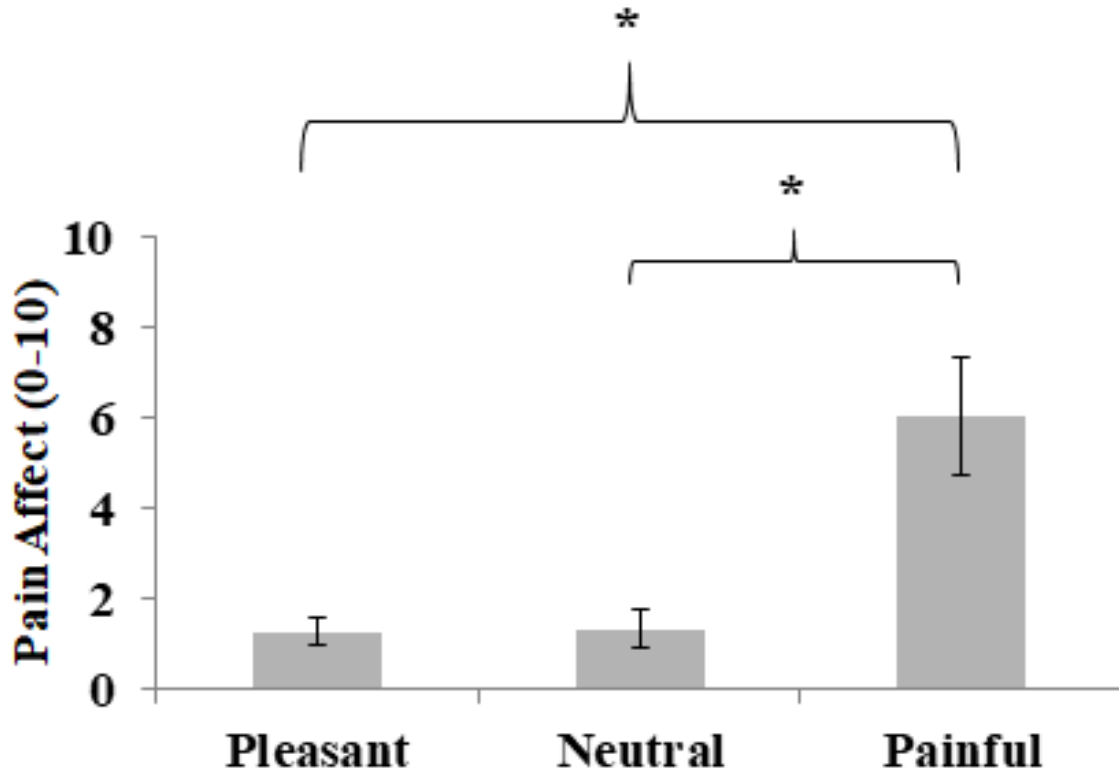
640 Figure 2. Physiological and perceptual measures taken during the fixed power (FP) cycling
641 task. Panel A shows differences in heart rate between conditions over time. Panel B shows
642 differences in blood lactate between conditions over time. Panel C shows differences in
643 ratings of perceived exertion (RPE) between conditions over time. Panel D shows differences
644 in pain between conditions over time. * denotes a significant difference in mean pain across
645 the FP task between the pleasant and painful conditions.

646 Figure 3. Time trial (TT) performance across the painful, neutral and pleasant conditions.
647 Panel A shows differences in power output during the TT between the three conditions. Panel
648 B shows differences in power output between conditions over the distance of the TT. *
649 denotes a significant difference in power output in the TT between the painful and pleasant
650 conditions and the painful and neutral conditions.

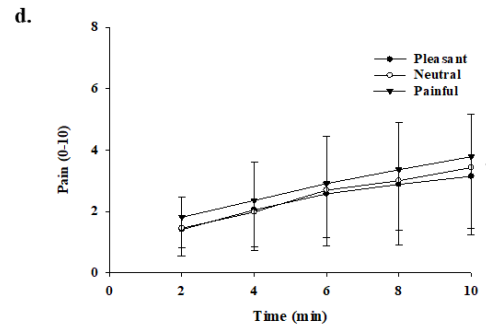
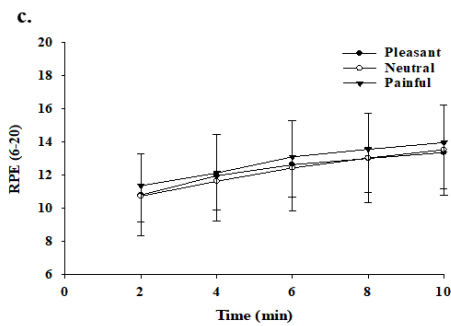
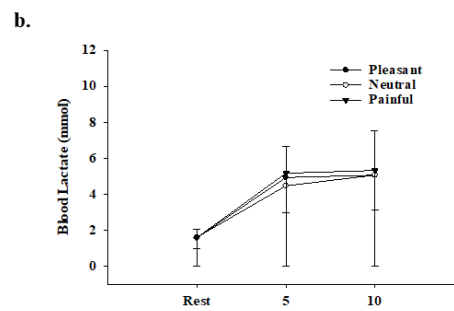
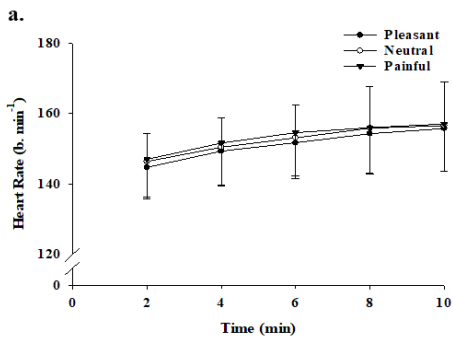
651 Figure 4. Physiological and perceptual measures taken during the cycling TT task. Panel A
652 shows differences in blood lactate between conditions throughout the TT. Panel B shows
653 differences in heart rate between conditions throughout the TT. Panel C shows differences in
654 RPE between conditions throughout the TT. Panel D shows differences in pain between
655 conditions throughout the TT. # denotes a significant main effect of Condition for blood
656 lactate during the TT. * denotes a significant main effect of Condition for heart rate during
657 the TT.

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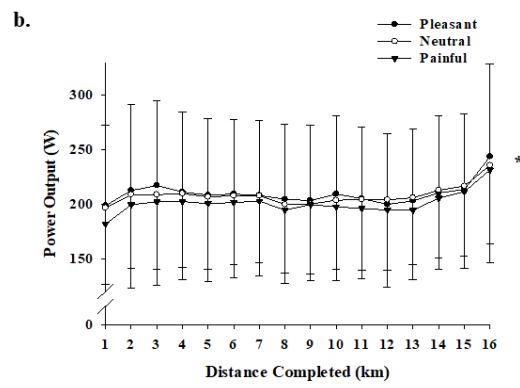
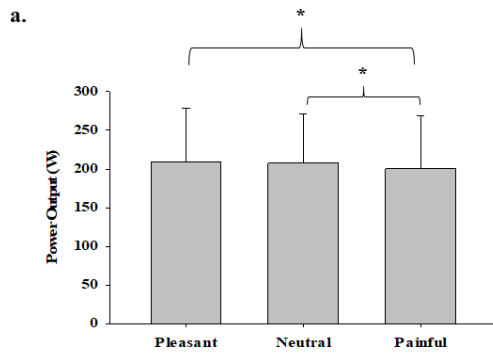
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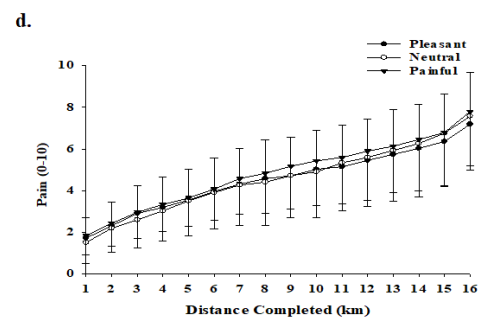
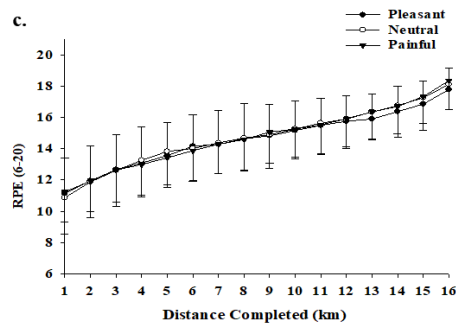
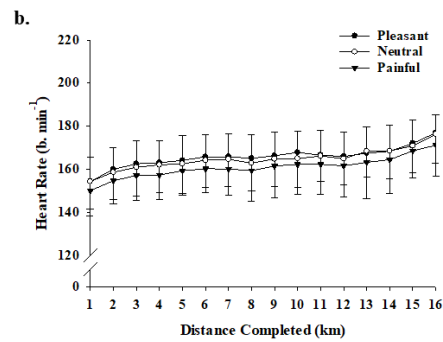
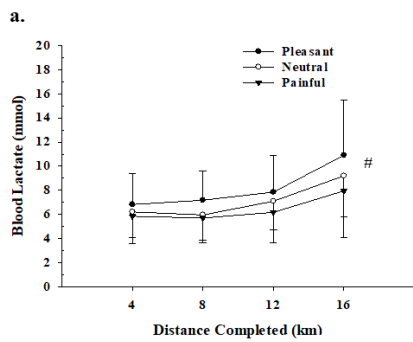
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672 **Table 1.** Participant mean values for anthropometric characteristics and cardiovascular and
 673 performance parameters.

Variable	Male	Female	Total (F & M)
Age (yrs)	31 ± 7	29 ± 8	31 ± 7
Height (cm)	183 ± 9	166 ± 6	176 ± 12
Body mass (kg)	78.5 ± 15.7	59.5 ± 5.9	71.3 ± 15.8
VO _{2max} (mL/kg/min)	56.7 ± 8.9	49.5 ± 10.8	54.0 ± 10.1
Anaerobic Threshold (W)	164.4 ± 53.1	116.5 ± 30.9	146 ± 51
Peak Power Output (W)	336.1 ± 56.5	214.6 ± 51.2	290 ± 81
Ramp end pain	7.9 ± 1.7	5.3 ± 2.6	6.9 ± 2.4
Ramp end RPE	18.0 ± 1.5	17.0 ± 2.6	17.6 ± 2.0
Ramp HR _{max} (beat. min ⁻¹)	181 ± 12	173 ± 18	180 ± 15

675 RPE, rating of perceived exertion; Ramp, incremental ramp test; HR, heart rate; W, watts.

676

677

678 **Table 2.** Means and standard deviations for HR, pain, perceived exertion and B[La] during
 679 the FP task.

Variable	Pleasant	Neutral	Painful
HR (bpm)	151.209 ± 10.981	152.324 ± 11.584	153.295 ± 12.103
EIP	2.410 ± 1.657	2.510 ± 1.589	2.843 ± 1.642
RPE	12.367 ± 2.538	12.286 ± 2.396	12.838 ± 2.282
B[La] (mmol/L)	7.487 ± 2.772	7.019 ± 2.409	7.851 ± 2.900

680 HR, heart rate; bpm, beats per minutes; EIP, exercise induced pain; RPE, rating of perceived
 681 exertion; B[La], blood lactate; mmol/L, millimoles per litre.

682

683 **Table 3.** Means and standard deviations for HR, pain, perceived exertion and B[La] during
 684 the TT task.

Variable	Pleasant	Neutral	Painful
HR (bpm)	165.041 ± 9.391	164.094 ± 11.919	160.545 ± 12.419
EIP	4.408 ± 1.789	4.628 ± 1.698	4.515 ± 1.731
RPE	14.610 ± 1.721	14.732 ± 1.721	14.728 ± 1.655
B[La] (mmol/L)	7.801 ± 2.923	7.316 ± 1.999	6.336 ± 2.311

685 HR, heart rate; bpm, beats per minutes; EIP, exercise induced pain; RPE, rating of perceived
 686 exertion; B[La], blood lactate; mmol/L, millimoles per litre.