


Please cite the Published Version

Romano-Smith, S, Roberts, JW, Wood, G , Coyles, G and Wakefield, CJ (2022) Simultaneous and alternate combinations of action-observation and motor imagery involve a common lower-level sensorimotor process. *Psychology of Sport and Exercise*, 63. p. 102275. ISSN 1469-0292

DOI: <https://doi.org/10.1016/j.psychsport.2022.102275>

Publisher: Elsevier BV

Version: Accepted Version

Downloaded from: <https://e-space.mmu.ac.uk/630325/>

Usage rights:  [Creative Commons: Attribution-Noncommercial-No Derivative Works 4.0](https://creativecommons.org/licenses/by-nc-nd/4.0/)

Additional Information: This is an Accepted Manuscript of an article which appeared in *Psychology of Sport and Exercise*, published by Elsevier

Enquiries:

If you have questions about this document, contact openresearch@mmu.ac.uk. Please include the URL of the record in e-space. If you believe that your, or a third party's rights have been compromised through this document please see our Take Down policy (available from <https://www.mmu.ac.uk/library/using-the-library/policies-and-guidelines>)

1
2
3
4
5
6
7
8
9
10
11
12
13
14
15
16
17
18
19
20
21
22
23
24
25

Highlights

- Examined the sensorimotor processes underlying motor performance by AOMI
- Simultaneous and alternate AOMI equally enhance motor performance
- Incongruent AO within simultaneous and alternate AOMI attenuates motor performance
- AO and MI may utilise a common lower-level sensorimotor process
- Fixation during AO disrupts motor improvements, which suggests a role for eye movements

26
27
28
29
30
31
32
33
34
35
36
37
38
39
40
41
42
43
44
45
46
47
48
49
50

Abstract

Combining the motor simulation techniques of action observation and motor imagery (AOMI) is known to enhance motor performance more than when these techniques are presented in isolation. The present study examined the involvement of lower-level sensorimotor processes for the improvement in a dart-throwing task using AOMI. Novice participants (n = 70) were assessed on their dart-throwing both before and after a six-week AOMI training intervention that was contingent upon the random allocation of groups. Participants were randomly allocated into groups involving AOMI, where they observed either a congruent action, incongruent action or fixation cross (control), while simultaneously or alternately imagining the dart-throwing task. Dart-throwing performance was significantly more improved for the simultaneous- and alternate-congruent groups compared to the simultaneous-fixation and control groups. There was no indication of improvement by any of the other groups. This improvement appeared to coincide with lower EMG activity at the agonist and antagonist muscles, which would indicate greater movement efficiency. The findings suggest that AOMI involves a common lower-level sensorimotor process, which can lead to motor facilitation or interference, dependent upon whether the simulation techniques are congruent or incongruent with each other, respectively. What's more, this feature does not appear to differ as a function of the structure of delivery (i.e., simultaneous vs. alternate).

Key words: motor performance, motor interference, EMG, eye movements

51 **Simultaneous and alternate combinations of action-observation and motor imagery**
52 **involve a common lower-level sensorimotor process**

53

54 Romano-Smith, S,^{a†} Roberts, J. W.,^{a*} Wood, G.,^b Coyles, G.,^a Wakefield, C. J.^a

55

56 ^a: School of Health Sciences, Liverpool Hope University, Taggart Avenue, Liverpool, L16

57

9JD, UK

58 ^b: Research Centre for Musculoskeletal Science and Sports Medicine, Department of Sport

59

and Exercise Science, Manchester Metropolitan University, UK

60

61 **Corresponding author:**

62 Dr Stephanie Romano Smith

63 School of Life Sciences and Education,

64 Staffordshire University,

65 Leek Road, Stoke-on- Trent,

66 ST4 2DF, UK

67 Tel:0178229495

68 Email: stephanie.romanosmith@staffs.ac.uk

69

70 [†]Affiliation of author SRS has now changed to School of Life Sciences and Education,

71 Department of Sport and Exercise, Staffordshire University, Leek Road, Stoke-on- Trent,

72 ST4 2DF, UK

73 Affiliation of author JWR has now changed to Liverpool John Moores University, Brain and

74 Behaviour Laboratory, Research Institute of Sport and Exercise Sciences (RISES), Liverpool

75 John Moores University, Tom Reilly Building, Byrom Street, Liverpool, L3 5AF

76 **Introduction**

77 Motor simulation techniques have been shown to aid motor learning (O’Shea &
78 Moran, 2017). Among these techniques are motor imagery (MI), which is defined as the
79 internal generation and mental rehearsal of action without overt physical output (Eaves,
80 Riach et al., 2016; Jeannerod, 2001); and action observation, which involves the deliberate
81 and structured viewing of a model demonstration (Hodges et al., 2007; Neuman and Gray,
82 2013; Ste-Marie et al., 2012). Broadly speaking, the benefits served by each of these training
83 or learning interventions may not be mutually exclusive because they each involve the
84 activation of a neural network that is also responsible for motor execution (Jeannerod, 2001).

85 While the benefits of AO and MI interventions on their own are somewhat well
86 established (Agosti & Sirico, 2020; Gatti et al., 2013), more recent evidence indicates that
87 there is an even greater benefit on motor performance and learning when combining AO and
88 MI (AOMI) compared to AO or MI alone (Marshall et al., 2019; Romano-Smith, Wood,
89 Wright, et al., 2018; Romano-Smith, Wood, Coyles, et al., 2019; Scott et al., 2018; Sun et al.,
90 2016; Taub et al., 2014; for a recent review, see Scott et al., 2021). This benefit coincides
91 with the enhanced engagement of the motor system as is indicated by the increased
92 corticospinal excitability (Sakamoto et al., 2009; Wright, et al., 2014; Wright et al., 2016),
93 and more widespread neural activity (Berends et al., 2013; Macuga & Frey, 2012; Villiger et
94 al., 2013) when undertaking AOMI compared to AO or MI alone. In an attempt to explain
95 these findings, the *dual-action simulation hypothesis* (DAS; Bruton et al., 2020; Eaves et al.,
96 2016) suggests there are two parallel processing streams that can simultaneously represent
97 observed and imaged actions, which can then either merge or compete with each other based
98 on their content and relevance (Eaves et al., 2012; Eaves et al., 2016). Alternatively, the
99 *visual guidance hypothesis* (VG; Meers et al., 2020; see also, Vogt et al., 2013) suggests that

100 AO may provide a visual guide in order to facilitate MI, which can alone sufficiently engage
101 the motor system.

102 More recently, our research group has additionally explored the influence of the
103 structure of AOMI interventions; that is, the presentation of AO and MI in simultaneous (S-
104 AOMI) and alternate (A-AOMI) fashion. Initial findings in stroke patients indicated that
105 upper-limb function may improve more following S-AOMI compared to A-AOMI (Sun et al.,
106 2016; see also, Kim et al., 2018). However, it has been shown that the learning of a
107 Taekwondo move can be more greatly advanced by A-AOMI compared to AO or MI alone
108 (Kim et al., 2020). Meanwhile, our recent study in young healthy adults attempting a dart-
109 throwing task indicated that performance outcome scores could be equally improved by both
110 S-AOMI and A-AOMI (Romano-Smith, Wood, Wright, et al., 2018). In an attempt to further
111 detail the underlying neuromuscular function using surface electromyography (EMG), we
112 also found that S-AOMI and A-AOMI generated a similarly lower magnitude of activity at
113 the triceps brachii muscle (agonist) during the latter throwing phase (Romano-Smith et al.,
114 2019); potentially highlighting the increased movement efficiency (Lohse et al., 2010) that is
115 a hallmark of highly-skilled, expert-like movement control (Gatti et al., 2013; Duchateau et
116 al., 2006; Lohse et al., 2010).

117 At this juncture, we may prematurely assume that both S-AOMI and A-AOMI involve
118 similar or the same sensorimotor processes. However, it is possible that while both
119 interventions have obtained similar behavioural outcomes, they may manifest from slightly
120 different processes. This possibility resembles that of previous models that indicate separate,
121 but not mutually exclusive, processes may underpin motor learning (e.g., early vs. late
122 mediation: Vogt & Thomascke, 2007; effector-dependent vs. -independent: Bird et al., 2005;
123 Heyes & Foster, 2002; Gruetzmacher et al., 2011; bottom-up vs. top-down: Hayes et al.,
124 2014; Roberts et al., 2014). To elucidate, the S-AOMI may primarily engage lower-level

125 motor processes, where the neural network that is responsible for execution may be more
126 greatly activated by AOMI (Macuga & Frey, 2012; Wright et al., 2018). This possibility is
127 consistent with previous suggestions that AOMI involves neural processes that unfold at the
128 very same time (e.g., DAS (Bruton et al., 2020; Eaves et al., 2016), VG (Meers et al., 2020)).
129 Meanwhile, the A-AOMI may additionally comprise higher-level cognitive processes that
130 accommodate for some degree of deliberation or comparison between each of the AO and MI
131 trials. Indeed, research from the motor learning literature consistently highlights a benefit of
132 having the opportunity to deliberate or contemplate previous trial attempts (Guadagnoli &
133 Kohl, 2001; Swinnen et al., 1990), or switching between different routines or variations of a
134 set task (Hall et al., 1994; Shea & Morgan, 1979). Moreover, the potential for motor learning
135 through AO and/or MI is additionally accompanied by the recruitment of the lateral
136 prefrontal neural regions, which are synonymous with attentional processes including action
137 monitoring and movement reorganization (Buccino et al., 2004; Higuchi et al., 2012; see also,
138 Eaves, Behmer, et al., 2016). Thus, it is further possible to capture the learning or adaptation
139 from AO and/or MI by also recognizing the importance of events that take place in between
140 each of their presentations. These particular learning processes may lend themselves more to
141 the A-AOMI intervention because of its separated and delayed presentations of AO and MI.

142 The aim of this experiment was to further expand upon our previous findings using
143 dart-throwing (Romano-Smith et al., 2019), and more specifically, disentangle the
144 sensorimotor processes that underpin motor learning by S-AOMI and A-AOMI. In this
145 regard, we adapted a secondary task paradigm, where participants would either
146 simultaneously or alternately observe congruent (dart-throwing; elbow extension along the
147 midsagittal axis) or incongruent (internal shoulder rotation along the frontal axis) movements
148 with respect to the imagined dart-throwing task. This manipulation has often been used to
149 highlight the involvement of lower-level motor processes, where if AO and MI were to

150 similarly engage the neural network that is responsible for execution, then it would facilitate
151 (congruent) and/or interfere (incongruent) with any subsequent movement or motor learning
152 (Kilner et al., 2003; Mattar & Gribble, 2005; Piedimonte et al., 2018; Ramsey et al., 2010).
153 Of interest, a similar principle was adapted for recent investigations on the corticospinal
154 excitability and eye movements generated during AOMI when the MI component was either
155 congruent or incongruent with the AO component (Bruton et al., 2020; Meers et al., 2020).
156 Specifically, these studies demonstrated that excitability was a product solely of the MI
157 component (Meers et al., 2020), or excitability and eye movements reflected a combination of
158 both the AO (i.e., index finger abduction-adduction) and MI (i.e., little finger abduction-
159 adduction) components (Bruton et al., 2020). However, it is important to note that the present
160 study differed in two ways. Firstly, we manipulated AO while retaining MI because the
161 former can be conveniently modified and set-up prior to commencing the study, and the latter
162 can simply remain specific to the learning objective (i.e., dart-throwing) for the learners to
163 control. Secondly, the AOMI could be uniquely delivered in alternate fashion meaning there
164 was potentially less opportunity for the previously stated dual or simultaneous processes to
165 take place.

166 In line with the suggestions of dual or simultaneous processes (Bruton et al., 2020;
167 Eaves et al., 2016; Meers et al., 2020), it is predicted that if the S-AOMI involves AO and MI
168 utilising a common neural network including a similar set of eye movements, then learning
169 will be attenuated having simultaneously observed an incongruent movement because it
170 would interfere with any benefit served by MI. At the same time, if the A-AOMI involves a
171 different set of neural and eye movement processes, then learning should continue to unfold
172 having alternately observed an incongruent movement because any benefit served by MI
173 would still be upheld as if it were delivered on its own. Moreover, we incorporated a
174 condition where participants simultaneously or alternately observed a fixation cross to act as

175 a control. That is, the typical AO (congruent/incongruent) was replaced by the requirement to
176 fixate on a cross, which would presumably limit any interference and continue to enable
177 learning by MI (aside from prohibiting the potential contribution of eye movements;
178 Wakefield, et al., 2020).

179

180 **Method**

181 *Participants*

182 Because the present study was heavily adapted from previous studies that were
183 conducted within our lab (Romano-Smith et al., 2018; 2019), the current number of
184 participants within each group were intended to remain as close as possible to the original
185 target sample. As a result, there were 70 participants (35 males, 35 females; *mean age* = 28.1
186 *years*, *SD* = 5.96) that agreed to take part in the study; all with limited darts experience
187 (including no targeted or deliberate practice, no competitive experience, >3 years from when
188 they recalled having last possibly attempted a dart throw within a recreational setting) and no
189 previous MI training. Participants were equally distributed to 1 out of 7 groups; therefore,
190 there were 10 participants per group that were age- and gender-matched. Participants had
191 normal or corrected-to-normal vision and reported being right-hand dominant (Oldfield,
192 1971). The experiment was ethically approved by the department research ethics board.

193

194 *Measures*

195 *Movement Imagery Questionnaire-Revised (MIQ-R)*

196 The MIQ-R (Hall & Martin, 1997) is an 8-item inventory that assesses the ability to
197 perform visual and kinesthetic imagery. Participants had to imagine a series of movements
198 using an internal kinesthetic or visual modality and rate the degree of ease or difficulty that
199 was experienced by using a 7-point Likert scale (1 = very hard to feel/see; 7 = very easy to

200 feel/see). The validity and consistency of the MIQ-R has been demonstrated by Gregg et al.
201 (2010), and has been used previously in imagery studies involving far-aiming tasks (e.g.,
202 Romano-Smith et al., 2018; Romano-Smith et al., 2019; Smith et al., 2008)

203

204 ***Imagery Diary***

205 Participants were provided with an imagery diary, which they could complete after
206 each MI session throughout the intervention period. Participants noted any difficulties,
207 feelings or concerns that were experienced when engaging in MI (for further guidelines, see
208 Goginsky & Collins, 1996). Engagement within the session was measured using a frequency
209 count of the number of sessions completed from a possibility of 18 sessions. The vividness
210 and controllability of the imagery were also rated on a 7-point Likert scale (1=not at all vivid
211 / controllable;7=very vivid / controllable).

212

213 ***Performance***

214 The performance measure was based on the points scored from dart-throwing toward
215 a dartboard. In line with the American Darts Organization regulations, the dartboard was
216 mounted 172.72 cm from the ground and located at a distance of 236.86 cm from the
217 throwing line. The dartboard was 45.72 cm in diameter and featured 10 concentric circles (2
218 cm wide) with the centre assuming the highest score (10 points) and outer edge assuming the
219 lowest score (1 point). Darts landing outside the board were scored as 0 points (see Fig. 1)

220

221 [Insert Fig.1 about here]

222 ***Electromyography (EMG)***

223 Muscle activation patterns were measured using electromyography (EMG) during the
224 pre-test and post-test procedures, by placing pairs of 10-mm Trigno EMG electrodes (Delsys

225 Inc.) sampling at 1500 Hz with a 20-mm separation on the belly of the biceps brachii, triceps
226 brachii, flexor carpi radialis and extensor carpi radialis (for similar regions, see Lohse et al.,
227 2010; Mousavi et al., 2019). EMG signals were initially processed using a Hamming
228 bandpass filter of 20-350 Hz and converted using a root mean square (RMS) calculation with
229 a 100-ms time-window (Fukuda et al., 2010; for examples in dart-throwing, see Romano
230 Smith et al., 2019; Zachry et al., 2005). Data from each individual trial were then normalised
231 with respect to the peak activation for that corresponding trial.

232 A digital recording of the dart-throwing movement was made via a secure webcam
233 (30 Hz), which was located perpendicular to the throwing direction and adjacent to the non-
234 throwing limb. The EMG and digital recordings were synchronised using Noraxon MR3.10
235 analysis software (Scottsdale, AZ, USA). The points of maximum elbow flexion (throw
236 preparation) and extension (throw completion) were marked as the key events of the
237 complete throwing action courtesy of a frame-by-frame analysis. Mean muscle activation
238 between each of these events was calculated and expressed as a percentage of the peak
239 activity.

240

241 ***Procedure***

242 The study followed a mixed-measures design including a pre- and post-test within
243 seven groups of participants that each undertook different forms of an AOMI intervention.
244 Participants were initially introduced to the task by the experimenter and completed the MIQ-
245 R (Hall & Martin, 1997). Participants physically attempted the dart-throwing task during a
246 pre- and post-test in order to assess their baseline dart-throwing ability and any subsequent
247 improvement, respectively. They were instructed to aim as close as possible to the centre of
248 the dartboard and obtain the highest possible score. In between each of these tests,
249 participants received a training intervention that was assigned according to randomly

250 allocated groups: simultaneous-congruent, alternate-congruent, simultaneous-incongruent,
251 alternate-incongruent, simultaneous-fixation, alternate-fixation, and control (Fig. 2).

252 The experimental groups could be discriminated by their unique combination and
253 structure of MI and AO. For the MI component, each of the experimental groups imagined
254 dart throws from a first-person perspective with the sound of the dart hitting the dartboard to
255 ensure imagery unfolded within real-time; that is, at the same rate of the AO component. In
256 order to facilitate MI and promote a high-fidelity to physical execution (Lang, 1977; 1979),
257 participants were initially prompted by the reading of an individualised imagery script that
258 could be progressively adapted or revised by the participants themselves across the course of
259 their training. Based on the stimulus and response training from Lang et al. (1980), this
260 process involved appropriately capturing the environmental surroundings (stimulus
261 proposition), physical experiences (e.g., muscle contraction, heart rate, etc) (response
262 proposition), and stimulus-response relationships (meaning proposition) (for a similar
263 procedure, see Romano-Smith et al., 2018; Romano-Smith et al., 2019). Following the
264 completion of each training session with MI, participants had to update their own imagery
265 diary.

266 For the AO component, participants viewed either a congruent action, incongruent
267 action, or fixation cross (Fig. 3). The congruent action consisted of a model dart throw taken
268 from the first-person perspective. The incongruent action involved upward internal rotation of
269 the shoulder taken from the first-person perspective. The cross featured two intersecting
270 white lines at the centre of a black background, which participants had to fixate on.
271 Meanwhile, the control group observed one continuous block of video interviews with a
272 professional dart player, which did not provide any technical insights on dart-throwing and
273 roughly equated to the time of the other experimental group interventions.

274 The structure for each of the AO and MI components was also manipulated by
275 presenting them in simultaneous or alternate fashion. The simultaneous structure involved
276 closely observing the visual stimuli (congruent, incongruent, fixation), while also imagining
277 the dart throw at the very same time. The alternate structure involved closely observing the
278 visual stimuli followed by the independent or separate imagining of the dart throw whilst in
279 in view of the dartboard. Each of the training sessions featured 6 blocks of 5 trials for all of
280 the experimental groups (30 trials; see Fig. 2) (Smith & Holmes., 2004; Romano-Smith et al.,
281 2019). A single trial for the simultaneous groups involved AO and MI taking place in one
282 instance, while a single trial for the alternate groups involved AO followed MI with each
283 component taking place in separate instances – thus making the duration of the intervention
284 slightly shorter in time for the former compared to the latter. There were 3 sessions per week
285 for 6 weeks. All participants were instructed to separate each session by a minimum of 48
286 hours of rest to avoid any fatigue and/or boredom (Romano-Smith et al., 2019). The pre- and
287 post-test consisted of 5 initial familiarisation trials followed by 30 performance trials.

288

289 [Insert Fig.2 and Fig, 3 about here]

290

291 ***Data analysis***

292 To evaluate any potential differences between the allocated groups in their inherent
293 imagery ability, the ratings from the kinesthetic and visual sub-scales of the MIQ-R that were
294 measured near the start of testing were analysed using separate one-way Analysis of Variance
295 (ANOVA). To indicate whether participants actively engaged in the required MI, the reported
296 accounts of the sessions from each participant imagery diary were accumulated and analysed
297 using a one-way ANOVA.

298 The points from each individual dart throw were accumulated across all trials within
299 the pre- and post-test to generate a performance score out of 300 (10 max. points x 30 trials).
300 To obtain the most consistent or representative muscle activation patterns, and to avoid any
301 on-off transient phenomena including muscular exertion (Ahmadi et al., 2007; Lohse et al.,
302 2010; Merletti et al., 1985), participant mean EMG data were taken only from trials 2-4
303 within the blocks 2-4.

304 Performance and EMG measures were analysed using Analysis of Covariance
305 (ANCOVA) with pre-test scores as the covariate and group as the fixed factor. The
306 assumption of homogeneity of regression slopes was evaluated courtesy of the interaction
307 term (group x pre-test).¹ Effect sizes were indicated by partial eta-squared (η_p^2). Significant
308 main effects were decomposed using the Fisher Least Significant Difference (LSD) post hoc
309 procedure. Statistical significance was declared at $p \leq .05$.

310

311 **Results**

312 *Self-report data*

313 For the ratings from the MIQ-R, there was no significant main effect of group for the
314 kinesthetic, $F(6,69) = 1.65, p = .144, \eta_p^2 = .13$, and visual, $F(6,69) = .87, p = .522, \eta_p^2 = .07$,
315 subscales (Table 1). For the frequency of sessions taken from the imagery diaries, there was
316 no significant main effect of group, $F(5,54) = 1.69, p = .152, \eta_p^2 = .13$. It appeared all
317 participants clearly undertook the MI with reports available on at least 14 out of a possible 18
318 sessions.

319

320

[Insert Table 1 about here]

321

322 **Performance**

323 For performance outcomes, there was a significant main effect of group, $F(6,62) =$
324 $2.26, p = .049, \eta_p^2 = .18$ (Fig. 4). Post hoc analysis indicated a significantly higher score for
325 the simultaneous-congruent compared to the simultaneous-fixation ($p = .039$) and control (p
326 $= .007$) groups. Additionally, there was a significantly higher score for the alternate-
327 congruent compared to the simultaneous-fixation ($p = .026$) and control ($p = .005$) groups.
328 There were no further significant differences for the remaining pairwise comparisons ($ps >$
329 $.05$).

330

331 [Insert Fig. 4 about here]

332

333 **EMG measures**

334 For the biceps brachii, the main effect of group approached conventional levels of
335 significance, $F(6,62) = 2.15, p = .060, \eta_p^2 = .17$, which indicated a trend toward the
336 simultaneous-fixation group generating the highest muscle activity (Table 2).

337 For the triceps brachii, there was a significant main effect of group, $F(6,62) = 3.08, p$
338 $= .011, \eta_p^2 = .23$. Post hoc analysis indicated significantly less activation for the
339 simultaneous-congruent compared to the alternate-incongruent ($p = .047$), simultaneous-
340 fixation ($p = .003$) and control ($p = .001$) groups. In a similar vein, there was significantly
341 less activation for the alternate-congruent compared to the simultaneous-fixation ($p = .014$)
342 and control ($p = .007$) groups. In addition, there was significantly less activation for the
343 alternate-fixation compared to the control group ($p = .032$).

344 Meanwhile, there was no significant main effect of group for either the flexor carpi
345 radialis, $F(6,62) = .91, p = .494, \eta_p^2 = .08$, nor the extensor carpi radialis, $F(6,62) = .71, p =$
346 $.641, \eta_p^2 = .07$.

347

348

[Insert Table 2 about here]

349

350 **Discussion**

351 The aim of this experiment was to further expand on previous motor learning findings
352 by combining AO and MI, and more specifically, disentangle the sensorimotor processes that
353 underpin their benefit. More specifically, we explored the possibility of whether S-AOMI and
354 A-AOMI feature contributions from different processes. Namely, we anticipated that the
355 lower-level sensorimotor processes that may potentially more heavily contribute toward S-
356 AOMI would render an inability to learn when one of the simulation components (i.e., AO)
357 was incongruent with the other (i.e., MI). On the other hand, we anticipated that the
358 potentially higher-level cognitive processes that may additionally contribute toward A-AOMI
359 would render a continued capacity to learn regardless of whether the simulation components
360 were congruent or incongruent with each other.

361 Firstly, in line with our previous findings (Romano-Smith, et al., 2019), we showed an
362 improvement in motor performance following both the simultaneous-congruent and alternate-
363 congruent interventions compared to the simultaneous-fixation and control groups. However,
364 there was no indication of learning for either of the simultaneous-incongruent and alternate-
365 incongruent interventions. These behavioural findings were corroborated by EMG
366 recordings, where the improvements in dart-throwing performance appeared to coincide with
367 a decrease in activity for the key agonist (triceps brachii) and antagonist (biceps brachii)
368 muscles. This pattern of activity may characterise a skilled level of performance because it
369 represents the recruitment of fewer motor units, and thus greater efficiency, which is one of
370 the hallmarks of expert-like performance (Duchateau et al., 2006; Lohse et al., 2010).

371 The benefit that was served by the combined AO and MI interventions for
372 simultaneous-congruent and alternate-congruent groups has been previously explained by
373 learners more greatly recruiting a common neural network that is synonymous with physical
374 execution (Berends et al., 2013; Eaves et al., 2016; Taube et al., 2015). This learning may also
375 be seen as additive in nature compared to the much smaller and independent benefit that is
376 usually served by AO and MI alone. Likewise, the smaller magnitude muscle activation
377 patterns that occurred within each of these groups would seem to suggest a more enhanced
378 neuromuscular function. To elucidate, an enhanced performance combined with lower muscle
379 activity may indicate a more automatized mode of control as opposed to poorer performance
380 manifesting from a larger magnitude and more abrupt muscle activation pattern, which is
381 synonymous with the conscious constraint of action (Wulf et al., 2001; Zachry et al., 2005).

382 In line with this argument, it was shown that learning could no longer take place when
383 participants had to observe a movement (internal shoulder rotation) that was incongruent with
384 their imagery (dart-throwing). While there was not necessarily a statistically significant
385 difference between the congruent and incongruent groups, their differences with respect to
386 the remaining group interventions including the control group would suggest otherwise (for a
387 similar pattern of results, see Brown et al., 2009; Marshall et al., 2019); particularly as it was
388 aligned with key theoretical stances and related empirical findings. Thus, knowing as we do
389 that AOMI (Bruton et al., 2020; Wright et al., 2018), and AO (Fadiga et al., 1995; Iacoboni et
390 al., 1999; Iacoboni & Dapretto, 2006) and MI (Fadiga et al., 1999; Grezes & Decety, 2001;
391 Villiger et al., 2013) alone are capable of engaging neural regions that are associated with
392 physical execution, it is assumed that this interference operates at the lower-level
393 sensorimotor processes (Mattar & Gribble, 2005; Ramsey et al., 2010). To elucidate, the
394 observation of an incongruent human movement may have awakened an internal

395 representation that was specific to the observed movement, while at odds with the
396 representation recruited through motor imagery (see Blakemore & Frith, 2005).

397 The extent of this learning and interference across the simultaneous and alternate
398 groups appeared to be relatively similar or consistent. In a similar vein, the muscle activation
399 patterns within each of these groups appeared to be highly similar, which would suggest a
400 similar mode of control and related neuromuscular function. This was despite the clear
401 differences in being able to continuously match or actively off-set the AO and MI for
402 simultaneous and alternate groups, respectively. Thus, it would appear that both groups
403 equally engaged lower-level sensorimotor processes. That said, it is possible that there are
404 subtle or other psychological differences that remain elusive or beyond the remits of the
405 present study. For example, it is possible that the simultaneous and alternate interventions
406 may be differentiated by their use of attentional resources. While it has been argued that a
407 simultaneous intervention may occupy fewer resources by allowing AO and MI to be
408 combined (Scott et al., 2021), it is also possible that it inversely uses more resources, given
409 that it could also be described as a dual-task intervention (Eaves, Behmer, & Vogt, 2016; see
410 also, Hayes et al., 2014; Mattar & Gribble, 2005). Moreover, while the alternate intervention
411 appears to indicate the involvement of lower-level sensorimotor process, this does not
412 preclude its use of additional explicit or verbalizable knowledge that can also contribute to
413 the learning process (Beilock & Carr, 2001).

414 Perhaps surprisingly, the simultaneous- and alternate-fixation groups that featured a
415 white-on-black cross in observation failed to exhibit any improvement. Indeed, the absence of
416 any incongruent AO, along with the continuation of MI, may anticipate at least some
417 indication of motor learning.² However, a viable explanation for this outcome may relate to
418 the role of eye movements. That is, in addition to a common neural network, it is suggested
419 that AO and MI share similar patterns of eye movements as physical execution (Causer et al.,

420 2013; Heremens et al., 2008; McCormick et al., 2012a,b; Wakefield et al., 2020; see also
421 Flanagan & Johansson, 2003). Thus, in the presence of a static cross in order to fixate on or
422 generally suppress eye movements, the functional correspondence between MI and execution
423 may be limited, which would diminish the benefit for motor learning (Heremens et al., 2011).
424 This issue may not have been as prevalent for the alternate-fixation group because they had
425 MI without the cross being simultaneously present, which may explain how they were
426 slightly better or had lower muscle activation (greater efficiency) than the simultaneous-
427 fixation and control groups.

428

429 ***Limitations***

430 While the present study further examined the sensorimotor processes that underpin
431 learning using different structures of AOMI, it is important to acknowledge the possible
432 limitations. Firstly, while the present study design captures the improvement of a novel or
433 unpractised movement skill to indicate motor learning, it does not feature the delayed
434 retention and transfer tests that would respectively assess the relative permanence and
435 adaptation that also characterise motor learning (Schmidt et al., 2019; for similar approaches,
436 see Marshall et al., 2019; Scott et al., 2018; Taube et al., 2014). That said, we strongly
437 suspect learning to have unfolded in this instance because the superior outcomes for the
438 simultaneous- and alternate-congruent groups coincided with the muscle activation patterns
439 from the EMG data, which indicated a smaller magnitude and more expert-like response.
440 That said, the present study adopted surface electrode EMG, which may not necessarily
441 provide a direct indication of neuromuscular function that could otherwise be detected using
442 fine-wire intramuscular EMG with a decreased risk of cross-talk between muscles (Yue et al.,
443 1995; Felici & Del Vecchio, 2020). Along these lines, there are perhaps further insights to be

444 drawn from the use of TMS as a direct indication of the cortical level processes during AOMI
445 (e.g., Bruton et al., 2021; Meers et al., 2020; Wright et al., 2018).

446 Despite the interference witnessed when being exposed to an incongruent secondary
447 task, it is possible that the incongruent movement comprising AO within the present study
448 could have partially facilitated or been coordinated with the MI. This possibility has been
449 mostly highlighted by some of the empirical findings underpinning the dual-action simulation
450 hypothesis, where it is possible to simultaneously utilise representations for AO and MI that
451 can perhaps merge together depending on their degree of similarity (Bruton et al., 2021; see
452 also Vogt et al., 2013). It is precisely this logic that could explain why the present differences
453 between congruent and incongruent groups were comparatively limited.

454 However, perhaps most importantly, one of the underlying issues that is inherent
455 within this sort of research; specifically, the comparison between S-AOMI and A-AOMI,
456 involves the potentially confounding influence of volume (i.e., number of trials) and time
457 spent within practice (i.e., simulation). That is, because of the differences in structure for
458 each of these combined interventions, it also incurs differences in either the volume or time.
459 For example, the present study features the same number of AO and MI trials for S-AOMI
460 and A-AOMI interventions, although incidentally there are twice as many simulations for the
461 latter compared to the former. In this regard, such comparisons within the literature may be
462 somewhat misguided in terms of their underlying assumptions or inferences surrounding
463 sensorimotor processes. While the findings from our research group to-date tend to indicate a
464 limited difference between S-AOMI and A-AOMI, we still cannot deny the possibility that
465 manipulations targeting volume and time of practice could also have an influence on learning
466 outcomes. Thus, it is useful perhaps to have future studies systematically control the volume
467 and time that is coincident with any manipulation of structure (e.g., x30 trials AO and MI vs.
468 x15 trials AO and MI in simultaneous and alternate structures). Finally, with this in mind,

469 because of the potential for AO to coincide with spontaneous MI (Meers et al., 2020; Vogt et
470 al., 2013), it is relevant to consider ways to control for this possibility including manipulation
471 checks (e.g., Bruton et al., 2021) that ensure AOMI combinations unfold as intended.

472

473 *Applied implications and future recommendations*

474 In summary, the results of the present study provide further insight on the benefits
475 served by AOMI interventions for motor learning. More specifically, we highlight how the
476 benefit of either simultaneous or alternate presentations of AO and MI appear to utilise the
477 same lower-level sensorimotor processes. What's more, we suspect that the suppression of
478 eye movements may prohibit the benefit of AOMI. These differences were also reflected
479 within the muscle activation patterns, where a smaller magnitude of activity coincided with
480 better outcomes, and thus more efficient neuromotor control.

481 These findings lend further support to the benefits served by AOMI interventions for
482 motor learning. What's more, they allude to the importance of the congruency or similarity
483 between the AO and MI components within a combination of AOMI – the closer they are to
484 each other, then the greater the benefit for motor learning. Meanwhile, there appear to be
485 rather limited differences between the benefits served by simultaneous and alternate
486 structures. In this regard, for example, learners may save themselves more time when
487 undertaking S-AOMI, although if there were to be any need to distinguish presentations of
488 AO and MI as in A-AOMI (e.g., instructions, coaching points, etc), then they should be done
489 without any fear of mitigating key lower-level sensorimotor processes. That said, more needs
490 to be done to explore the potential differences in attentional processes and/or strategies that
491 may coincide with each of these delivery structures. Finally, future research may further
492 explore the additive benefit of AOMI by investigating the potentially different sources of
493 information that are gleaned from each of the AO and MI components. For example, access

494 to the biological motion trajectory during AO (Grossman et al., 2000; Kilner et al., 2003;
495 Press et al., 2011) may provide a spatiotemporal kinematic referent for updating an already
496 engaged internal representation courtesy of MI (for a similar argument, see Glover & Baran,
497 2017).

498 **References**

- 499 Agosti, V., & Sirico, M. (2020). Motor imagery as a tool for motor learning and improving
500 sports performance: A mini review on the state of the art. *Sport Science, 13*, 13-17.
- 501 Ahmadi S, Sinclair PJ, Foroughi N, Davis GM. Electromyographic activity of the biceps
502 brachii after exercise-induced muscle damage. *J Sports Sci Med.* 2007 (4), 461- 470.
- 503 Bek, J., Poliakoff, E., Marshall, H., Trueman, S., & Gowen, E. (2016). Enhancing voluntary
504 imitation through attention and motor imagery. *Experimental Brain Research, 234*(7),
505 1819-1828.
- 506 Berends, H.I., Wolkorte, R., IJzerman, M. J., & van Putten, M. J. A. M. (2013). Differential
507 cortical activation during observation and observation-and-imagination. *Experimental*
508 *Brain Research, 229*(3), 337-345.
- 509 Brown, L. E., Wilson, E. T., & Gribble, P. L. (2009). Repetitive transcranial magnetic
510 stimulation to the primary motor cortex interferes with motor learning by observing.
511 *Journal of Cognitive Neuroscience, 21*(5), 1013-1022.
- 512 Bird, G., & Heyes, C. (2005). Effector-dependent learning by observation of a finger
513 movement sequence. *Journal of Experimental Psychology: Human Perception and*
514 *Performance, 31*(2), 262-275.
- 515 Beilock, S. L., & Carr, T. H. (2001). On the fragility of skilled performance: What governs
516 choking under pressure? *Journal of Experimental Psychology: General, 130*(4), 701.
- 517 Blakemore, S. J., & Frith, U. (2005). The role of motor contagion in the prediction of action.
518 *Neuropsychologia, 43*(2), 260-267.
- 519 Bruton, A. M., Holmes, P. S., Eaves, D. L., Franklin, Z. C., & Wright, D. J. (2020).
520 Neurophysiological markers discriminate different forms of motor imagery during
521 action observation. *Cortex, 124*, 119-136.

522 Buccino, G., Binkofski, F., & Riggio, L. (2004). The mirror neuron system and action
523 recognition. *Brain and language*, *89*(2), 370-376.

524 Causser, J., McCormick, S. A., & Holmes, P. S. (2013). Congruency of gaze metrics in action,
525 imagery and action observation. *Frontiers in Human Neuroscience*, *7*, 604.

526 Felici, F., & Del Vecchio, A. (2020). Surface Electromyography: What Limits Its Use in
527 Exercise and Sport Physiology? *Frontiers in Neurology*, 1508.

528 Duchateau, J., Semmler, J. G., & Enoka, R. M. (2006). Training adaptations in the behavior
529 of human motor units. *Journal of Applied Physiology*, *101*(6), 1766-1775.

530 Eaves, D. L., Behmer, L. P., & Vogt, S. (2016). EEG and behavioural correlates of different
531 forms of motor imagery during action observation in rhythmical actions. *Brain and*
532 *Cognition*, *106*, 90-103.

533 Eaves, D. L., Riach, M., Holmes, P. S., & Wright, D. J. (2016). Motor imagery during action
534 observation: a brief review of evidence, theory and future research opportunities.
535 *Frontiers in Neuroscience*, *10*, 514.

536 Fadiga, L., Fogassi, L., Pavesi, G., & Rizzolatti, G. (1995). Motor facilitation during action
537 observation: a magnetic stimulation study. *Journal of Neurophysiology*, *73*(6), 2608-
538 2611.

539 Fadiga, L., Buccino, G., Craighero, L., Fogassi, L., Gallese, V., & Pavesi, G. (1998).
540 Corticospinal excitability is specifically modulated by motor imagery: a magnetic
541 stimulation study. *Neuropsychologia*, *37*(2), 147-158.

542 Flanagan, J. R., & Johansson, R. S. (2003). Action plans used in action observation. *Nature*,
543 *424*(6950), 769-771.

544 Fukuda, T. Y., Echeimberg, J. O., Pompeu, J. E., Lucareli, P. R. G., Garbelotti, S., Gimenes,
545 R. O., & Apolinário, A. (2010). Root mean square value of the electromyographic

546 signal in the isometric torque of the quadriceps, hamstrings and brachial biceps
547 muscles in female subjects. *The Journal of Applied Research*, 10(1), 32-39.

548 Gatti, R., Tettamanti, A., Gough, P. M., Riboldi, E., Marinoni, L., & Buccino, G. (2013).
549 Action observation versus motor imagery in learning a complex motor task: a short
550 review of literature and a kinematics study. *Neuroscience Letters*, 540, 37-42.

551 Glover, S., & Baran, M. (2017). The motor-cognitive model of motor imagery: Evidence
552 from timing errors in simulated reaching and grasping. *Journal of Experimental*
553 *Psychology: Human Perception and Performance*, 43(7), 1359.

554 Goginsky, A. M., & Collins, D. (1996). Research design and mental practice. *Journal of*
555 *Sports Sciences*, 14(5), 381-392.

556 Gregg, M., Hall, C., & Butler, A. (2010). The MIQ-RS: a suitable option for examining
557 movement imagery ability. *Evidence-Based Complementary and Alternative*
558 *Medicine*, 7(2), 249-257.

559 Grezes, J., & Decety, J. (2001). Functional anatomy of execution, mental simulation,
560 observation, and verb generation of actions: A meta-analysis. *Human Brain Mapping*,
561 12(1), 1-19.

562 Gruetzmacher, N., Panzer, S., Blandin, Y., & Shea, C. H. (2011). Observation and physical
563 practice: Coding of simple motor sequences. *Quarterly Journal of Experimental*
564 *Psychology*, 64(6), 1111-1123.

565 Grossman, E., Donnelly, M., Price, R., Pickens, D., Morgan, V., Neighbor, G., & Blake, R.
566 (2000). Brain areas involved in perception of biological motion. *Journal of cognitive*
567 *neuroscience*, 12(5), 711-720.

568 Guadagnoli, M. A., & Kohl, R. M. (2001). Knowledge of results for motor learning:
569 relationship between error estimation and knowledge of results frequency. *Journal of*
570 *Motor Behavior*, 33(2), 217-224.

571 Hall, K. G., Domingues, D. A., & Cavazos, R. (1994). Contextual interference effects with
572 skilled baseball players. *Perceptual and Motor Skills*, 78(3), 835-841.

573 Hayes, S. J., Roberts, J. W., Elliott, D., & Bennett, S. J. (2014). Top-down attentional
574 processes modulate the coding of atypical biological motion kinematics in the absence
575 of motor signals. *Journal of Experimental Psychology: Human Perception and*
576 *Performance*, 40(4), 1641.

577 Heremans, E., Smits-Engelsman, B., Caeyenberghs, K., Vercruyssen, S., Nieuwboer, A., Feys,
578 P., & Helsen, W. F. (2011). Keeping an eye on imagery: the role of eye movements
579 during motor imagery training. *Neuroscience*, 195, 37-44.

580 Heremans, E., Helsen, W. F., & Feys, P. (2008). The eyes as a mirror of our thoughts:
581 quantification of motor imagery of goal-directed movements through eye movement
582 registration. *Behavioural Brain Research*, 187(2), 351-360.

583 Heyes, C. M., & Foster, C. L. (2002). Motor learning by observation: evidence from a serial
584 reaction time task. *The Quarterly Journal of Experimental Psychology Section A*,
585 55(2), 593-607.

586 Higuchi, S., Holle, H., Roberts, N., Eickhoff, S. B., & Vogt, S. (2012). Imitation and
587 observational learning of hand actions: prefrontal involvement and connectivity.
588 *Neuroimage*, 59(2), 1668-1683.

589 Hodges, N. J., Williams, A. M., Hayes, S. J., & Breslin, G. (2007). What is modelled during
590 observational learning? *Journal of Sports Sciences*, 25(5), 531-545.

591 Jeannerod, M. (2001). Neural simulation of action: a unifying mechanism for motor
592 cognition. *Neuroimage*, 14(1), S103-S109.

593 Jacobson, E. (1930). Electrical measurements of neuromuscular states during mental
594 activities: I. Imagination of movement involving skeletal muscle. *American Journal of*
595 *Physiology-Legacy Content*, 91(2), 567-608.

596 Iacoboni, M., Woods, R. P., Brass, M., Bekkering, H., Mazziotta, J. C., & Rizzolatti, G.
597 (1999). Cortical mechanisms of human imitation. *Science*, 286(5449), 2526-2528.

598 Iacoboni, M., & Dapretto, M. (2006). The mirror neuron system and the consequences of its
599 dysfunction. *Nature Reviews Neuroscience*, 7(12), 942-951.

600 Kim, T., Frank, C., & Schack, T. (2018). The effect of different training schedules of action
601 observation and motor imagery on the changes in mental representation structure,
602 cognitive and skill performance.

603 Kim, T., Frank, C., Schack, T. (2020). The effect of alternate training of action observation
604 and motor imagery on cognitive and skill performance. *International Journal of Sport
605 Psychology*, 51(2), 101-121.

606 Kilner, J. M., Paulignan, Y., & Blakemore, S. J. (2003). An interference effect of observed
607 biological movement on action. *Current Biology*, 13(6), 522-525.

608 Lang, P. J. (1977). Imagery in therapy: An information processing analysis of fear. *Behavior
609 Therapy*, 8(5), 862-886.

610 Lang, P. J. (1979). A bio-informational theory of emotional imagery. *Psychophysiology*,
611 16(6), 495-512.

612 Lang, P. J., Kozak, M. J., Miller, G. A., Levin, D. N., & McLean Jr, A. (1980). Emotional
613 imagery: Conceptual structure and pattern of somato-visceral response.
614 *Psychophysiology*, 17(2), 179-192.

615 Lohse, K. R., Sherwood, D. E., & Healy, A. F. (2010). How changing the focus of attention
616 affects performance, kinematics, and electromyography in dart-throwing. *Human
617 Movement Science*, 29(4), 542-555.

618 O'Shea, H., & Moran, A. (2017). Does motor simulation theory explain the cognitive
619 mechanisms underlying motor imagery? A critical review. *Frontiers in Human
620 Neuroscience*, 11, 72.

621 Macuga, K. L., & Frey, S. H. (2012). Neural representations involved in observed, imagined,
622 and imitated actions are dissociable and hierarchically organized. *Neuroimage*, *59*(3),
623 2798-2807.

624 Marshall, B., Wright, D. J., Holmes, P. S., & Wood, G. (2019). Combining action observation
625 and motor imagery improves eye–hand coordination during novel visuomotor task
626 performance. *Journal of Motor Behavior*.

627 Mattar, A. A., & Gribble, P. L. (2005). Motor learning by observing. *Neuron*, *46*(1), 153-160.

628 Meers, R., Nuttall, H. E., & Vogt, S. (2020). Motor imagery alone drives corticospinal
629 excitability during concurrent action observation and motor imagery. *Cortex*, *126*,
630 322-333.

631 Merletti R, Knaflitz M, De Luca CJ. (1985) Myoelectric manifestations of fatigue in
632 voluntary and electrically elicited contractions. *Journal of Applied Physiology* *69*(5),
633 1810-1820.

634 McCormick, S. A., Causer, J., & Holmes, P. S. (2012). Eye gaze metrics reflect a shared
635 motor representation for action observation and movement imagery. *Brain and*
636 *Cognition*, *80*(1), 83–88.

637 McCormick, S. A., Marple-Horvat, D., Goodwin, P., & Holmes, P. S. (2012). Eye movement
638 metrics are required to justify the efficacy of imagery and observation in stroke
639 patients. *International Journal of Stroke* *7*, 50-50.

640 Mousavi, S. A., Shahbazi, M., Arabameri, E., & Shirzad, E. (2019). The effect of virtual
641 reality training on learning and kinematics characteristics of dart-throwing.
642 *International Journal of School Health*, *6*(1), 1-7.

643 Neuman, B., & Gray, R. (2013). A direct comparison of the effects of imagery and action
644 observation on hitting performance. *Movement & Sport Sciences-Science & Motricité*,
645 *79*, 11-21.

646 O'Shea, J. B., & Morgan, R. L. (1979). Contextual interference effects on the acquisition,
647 retention, and transfer of a motor skill. *Journal of Experimental Psychology: Human*
648 *Learning and Memory*, 5(2), 179.

649 Oldfield, R. C. (1971). The assessment and analysis of handedness: the Edinburgh inventory.
650 *Neuropsychologia*, 9(1), 97-113.

651 Piedimonte, A., Conson, M., Frolli, A., Bari, S., Della Gatta, F., Rabuffetti, M., ... &
652 Garbarini, F. (2018). Dissociation between executed and imagined bimanual
653 movements in autism spectrum conditions. *Autism Research*, 11(2), 376-384.

654 Press, C., Cook, J., Blakemore, S. J., & Kilner, J. (2011). Dynamic modulation of human
655 motor activity when observing actions. *Journal of Neuroscience*, 31(8), 2792-2800.

656 Ramsey, R., Cumming, J., Eastough, D., & Edwards, M. G. (2010). Incongruent imagery
657 interferes with action initiation. *Brain and Cognition*, 74(3), 249-254.

658 Roberts, J. W., Bennett, S. J., Elliott, D., & Hayes, S. J. (2014). Top-down and bottom-up
659 processes during observation: Implications for motor learning. *European Journal of*
660 *Sport Science*, 14(suppl), S250-S256.

661 Romano Smith, S., Wood, G., Coyles, G., Roberts, J. W., & Wakefield, C. J. (2019). The
662 effect of action observation and motor imagery combinations on upper limb
663 kinematics and EMG during dart-throwing. *Scandinavian Journal of Medicine &*
664 *Science in Sports*, 29(12), 1917-1929.

665 Romano-Smith, S., Wood, G., Wright, D. J., & Wakefield, C. J. (2018). Simultaneous and
666 alternate action observation and motor imagery combinations improve aiming
667 performance. *Psychology of Sport and Exercise*, 38, 100-106.

668 Scott, M., Taylor, S., Chesterton, P., Vogt, S., & Eaves, D. L. (2018). Motor imagery during
669 action observation increases eccentric hamstring force: an acute non-physical
670 intervention. *Disability and Rehabilitation*, 40(12), 1443-1451.

671 Scott, M. W., Wood, G., Holmes, P. S., Williams, J., Marshall, B., & Wright, D. J. (2021).
672 Combined action observation and motor imagery: An intervention to combat the
673 neural and behavioural deficits associated with developmental coordination disorder.
674 *Neuroscience & Biobehavioral Reviews*.

675 Shea, J. B., & Morgan, R. L. (1979). Contextual interference effects on the acquisition,
676 retention, and transfer of a motor skill. *Journal of Experimental Psychology: Human*
677 *Learning and Memory*, 5(2), 179.

678 Sakamoto, M., Muraoka, T., Mizuguchi, N., & Kanosue, K. (2009). Combining observation
679 and imagery of an action enhances human corticospinal excitability. *Neuroscience*
680 *Research*, 65(1), 23-27.

681 Ste-Marie, D. M., Law, B., Rymal, A. M., Jenny, O., Hall, C., & McCullagh, P. (2012).
682 Observation interventions for motor skill learning and performance: an applied model
683 for the use of observation. *International Review of Sport and Exercise Psychology*,
684 5(2), 145-176.

685 Schmidt, R. A., Lee, T. D., Winstein, C. J., Wulf, G., & Zelaznik, H. N. (2019). Motor
686 control and learning (6th edition). Champaign, IL: Human Kinetics.

687 Smith, D., Wright, C. J., & Cantwell, C. (2008). Beating the bunker: The effect of PETTLEP
688 imagery on golf bunker shot performance. *Research Quarterly for Exercise and Sport*,
689 79(3), 385-391.

690 Smith D, Holmes P. (2007) The Effect of Imagery Modality on Golf Putting Performance.
691 *Journal of Sport Exercise Psychology*, 26(3), 385-395.

692 Sun, Y., Wei, W., Luo, Z., Gan, H., & Hu, X. (2016). Improving motor imagery practice with
693 synchronous action observation in stroke patients. *Topics in Stroke Rehabilitation*,
694 23(4), 245-253.

695 Swinnen, S. P., Schmidt, R. A., Nicholson, D. E., & Shapiro, D. C. (1990). Information
696 feedback for skill acquisition: Instantaneous knowledge of results degrades learning.
697 *Journal of Experimental Psychology: Learning, Memory, and Cognition*, *16*(4), 706.

698 Taube, W., Lorch, M., Zeiter, S., & Keller, M. (2014). Non-physical practice improves task
699 performance in an unstable, perturbed environment: motor imagery and observational
700 balance training. *Frontiers in Human Neuroscience*, *8*, 972.

701 Villiger, M., Estévez, N., Hepp-Reymond, M. C., Kiper, D., Kollias, S. S., Eng, K., & Hotz-
702 Boendermaker, S. (2013). Enhanced activation of motor execution networks using
703 action observation combined with imagination of lower limb movements. *PloS One*,
704 *8*(8), e72403.

705 Vogt, S., & Thomaschke, R. (2007). From visuo-motor interactions to imitation learning:
706 behavioural and brain imaging studies. *Journal of Sports Sciences*, *25*(5), 497-517.

707 Vogt, S., Di Rienzo, F., Collet, C., Collins, A., & Guillot, A. (2013). Multiple roles of motor
708 imagery during action observation. *Front Hum Neurosci.*, *7*, 807.doi:
709 10.3389/fnhum.2013.00807

710 Wakefield, C. J., Roberts, J. W., & Wood, G. (2020). Eye-movements support chronometric
711 imagery performance even when the task is occluded. *Visual Cognition*, *28*(4), 285-
712 291.

713 Wright, D. J., McCormick, S. A., Williams, J., & Holmes, P. S. (2016). Viewing instructions
714 accompanying action observation modulate corticospinal excitability. *Frontiers in*
715 *Human Neuroscience*, *10*, 17.

716 Wright, D. J., Williams, J., & Holmes, P. S. (2014). Combined action observation and
717 imagery facilitates corticospinal excitability. *Frontiers in Human Neuroscience*, *8*,
718 951.

719 Wright, D. J., Wood, G., Eaves, D. L., Bruton, A. M., Frank, C., & Franklin, Z. C. (2018).
720 Corticospinal excitability is facilitated by combined action observation and motor
721 imagery of a basketball free throw. *Psychology of Sport and Exercise*, 39, 114-121.

722 Wulf, G., McNevin, N., & Shea, C. H. (2001). The automaticity of complex motor skill
723 learning as a function of attentional focus. *The Quarterly Journal of Experimental*
724 *Psychology Section A*, 54(4), 1143-1154.

725 Yue, G., Fuglevand, A. J., Nordstrom, M. A., & Enoka, R. M. (1995). Limitations of the
726 surface electromyography technique for estimating motor unit synchronization.
727 *Biological cybernetics*, 73(3), 223-233.

728 Zachry, T., Wulf, G., Mercer, J., & Bezodis, N. (2005). Increased movement accuracy and
729 reduced EMG activity as the result of adopting an external focus of attention. *Brain*
730 *Research Bulletin*, 67(4), 304-309

731
732
733
734
735
736
737
738
739
740
741
742
743
744

745 **Footnotes**

- 746 1) There was no significant effect for the interaction term (group x pre-test) in any of the
747 dependent measures (F s range = .49-1.73, p s range = .13-.82), which suggests that the
748 assumption of homogeneity of regression slopes was met.
- 749 2) In light of the learning from each of the congruent groups, but neither of the fixation
750 groups, it could be argued that the availability of congruent AO is the sole feature to
751 enable learning given that all the groups had the same MI. Thus, we compared the
752 present data from the congruent groups (simultaneous-congruent, alternate-congruent)
753 with two of our previous data sets involving only AO (Romano-Smith, Wood, Wright, et
754 al., 2018 (study 1); Romano-Smith, Wood, Coyles, et al., 2019 (study 2). Using
755 ANCOVA with pre-test as the covariate and group as the fixed factor (assuming
756 homogeneity of regression slopes; group x pre-test: $F(3,32) = .65, p = .588$), we found a
757 significant main effect of group, $F(3,35) = 3.46, p = .027, \eta_p^2 = .23$, which indicated
758 higher scores for the congruent groups from the present study compared to AO alone
759 (simultaneous-congruent > AO (study 1) ($p = .040$), AO (study 2) ($p = .083$); alternate-
760 congruent > AO (study 1) ($p = .010$), AO (study 2) ($p = .021$). This outcome refutes the
761 suggestion of learning solely through AO, and demonstrates the advantage of having AO
762 accompanied by MI.

763 **Figures and Tables Captions**

764 **Fig.1** A still shot of the task performed.

765 **Fig. 2.** Representative illustration of the timeline of events across the six weeks of training
766 (*upper panel*) and within a single session of training (*lower panel*)

767 **Fig. 3.** Representative illustration of the timeline of events for the AO and MI (left-to-right)
768 within the experimental AOMI interventions. The present illustration features only four
769 events that are each synonymous with an individual trial (total = 30 trials). *Dotted arrows*
770 comprising the incongruent stimuli indicate the direction of the observed arm movement.

771 **Fig. 4.** Adjusted mean dart-throwing scores (out of 300) as a function of group. Error bars
772 represent the standard error (SE) of the mean. (*) indicates a significant pairwise difference at
773 $p < .05$.

774 **Table 1** Mean (\pm SE) MIQ-R scores (kinesthetic and visual) and frequency of training
775 sessions for each group.

776 **Table 2.** Adjusted means (\pm SE) for the EMG activation across all throws (normalized (%) to
777 peak) at the biceps brachii, triceps brachii, flexor carpi radialis and extensor carpi radialis.

778

779

780

781

782

783

784

785

786

787

788
789
790
791
792
793
794
795
796
797
798
799
800
801
802
803
804
805
806
807

808

809

810

811

812

813

814

815

816

817

818

819

820

821

822

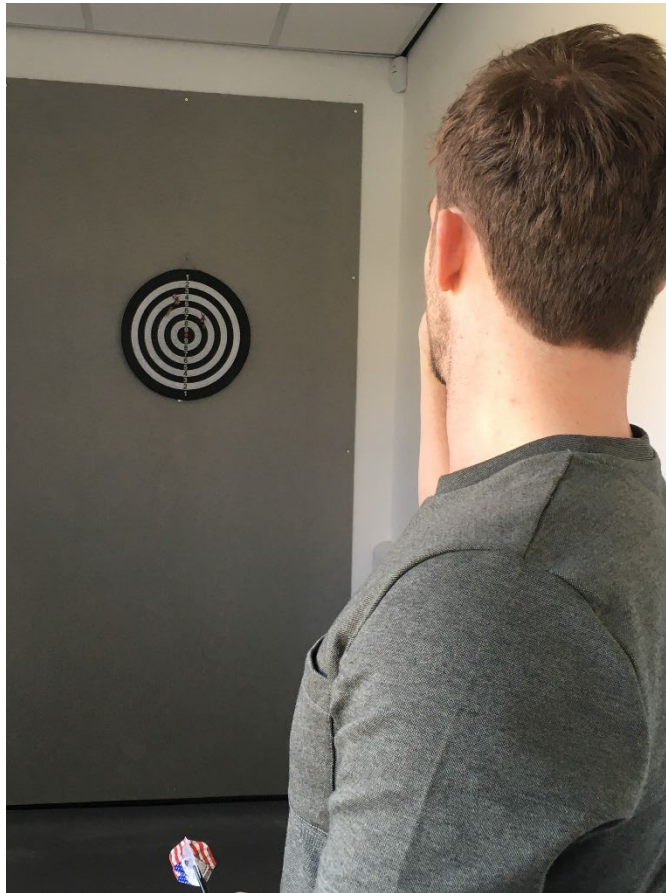
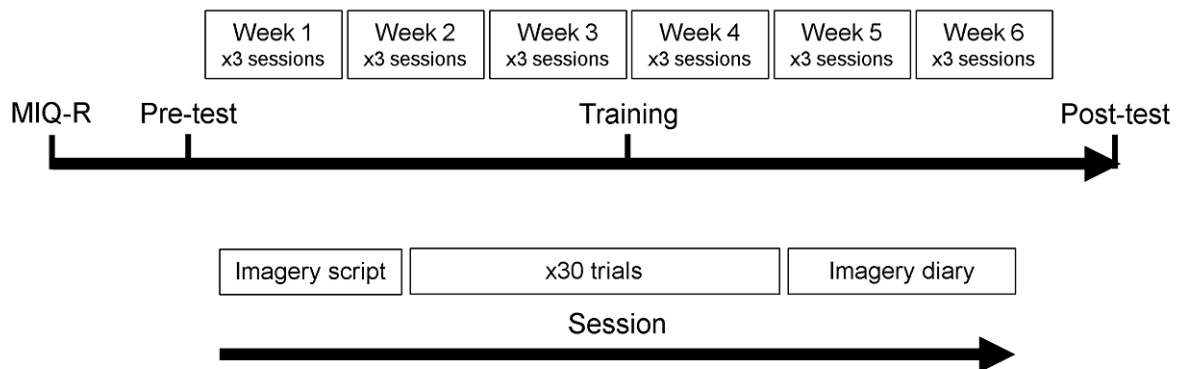


Fig.1 A still shot of the task performed.

823

824



825

826

827 **Fig. 2.** Representative illustration of the timeline of events across the six weeks of training

828 (*upper panel*) and within a single session of training (*lower panel*)

829

830

831

832

833

834

835

836

837

838

839

840

841

842

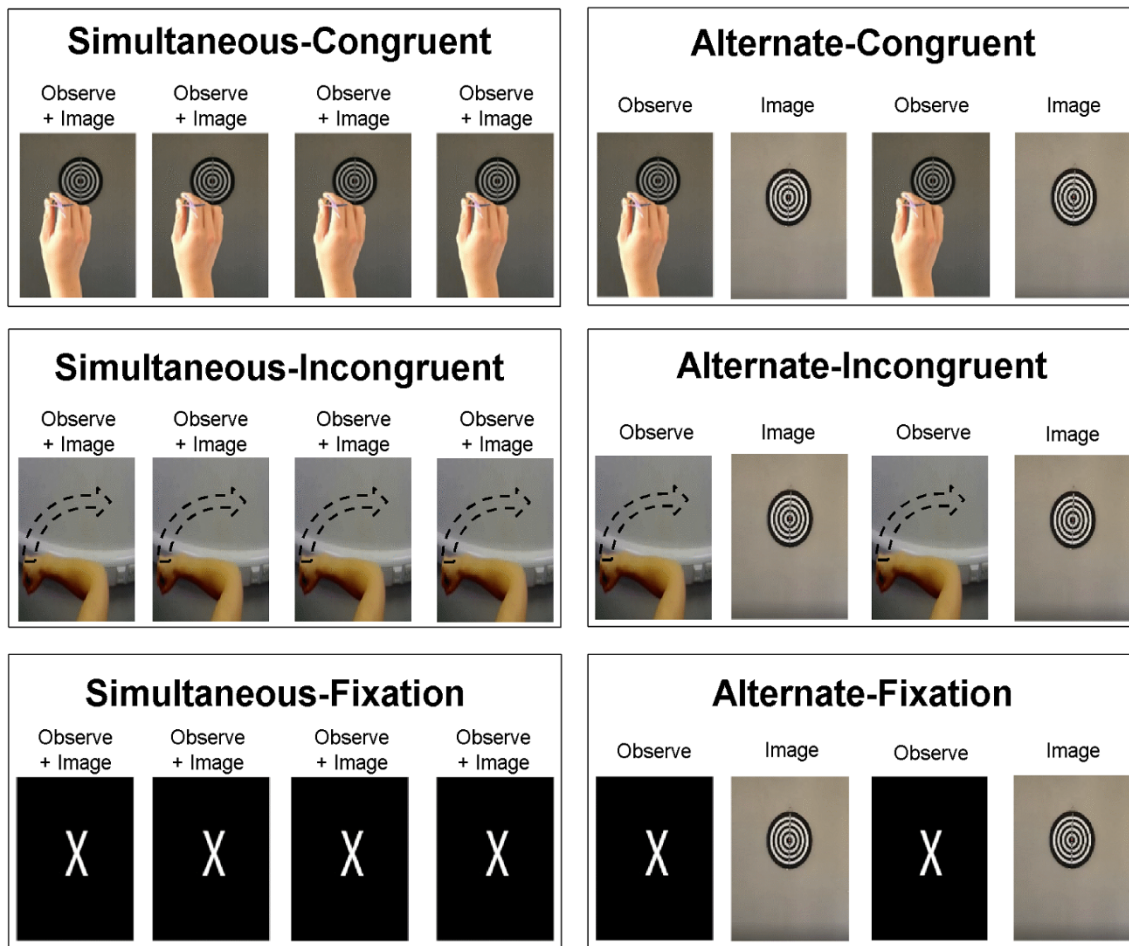
843

844

845

846

847



848

849

850 **Fig. 3.** Representative illustration of the timeline of events for the AO and MI (left-to-right)

851 within the experimental AOMI interventions. The present illustration features only four

852 events that are each synonymous with an individual trial (total = 30 trials). *Dotted arrows*

853 comprising the incongruent stimuli indicate the direction of the observed arm movement.

854

855

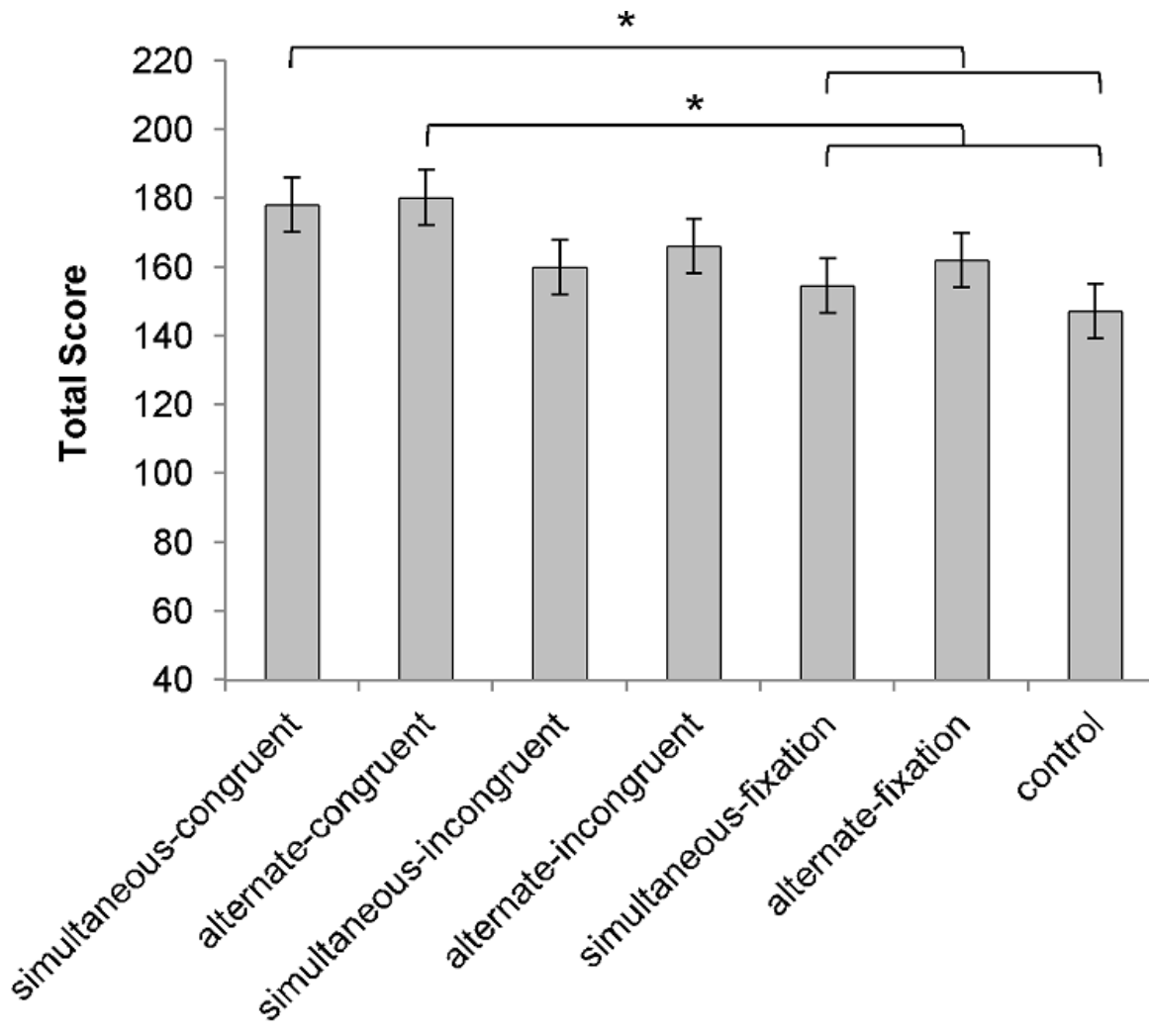
856

857

858

859

860



861

862

863

864

Fig. 4. Adjusted mean dart-throwing scores (out of 300) as a function of group. Error

865 bars represent the standard error (SE) of the mean. (*) indicates a significant pairwise

866 difference at $p < .05$.

867

868

869

870

871
872
873
874
875
876
877
878
879
880
881
882
883
884
885
886
887
888
889

Table 1

Mean (\pm SE) MIQ-R scores (kinaesthetic and visual) and frequency of training sessions for each group.

	MIQ-R Kinaesthetic	MIQ-R Visual	Frequency
simultaneous-congruent	6.30 (0.18)	6.20 (0.26)	17.30 (0.21)
alternate-congruent	6.15 (0.29)	6.31 (0.23)	16.60 (.34)
simultaneous-incongruent	6.35 (0.23)	6.35 (0.29)	16.60 (.40)
alternate-incongruent	6.57 (0.18)	6.50 (0.30)	16.70(.42)
simultaneous-fixation	5.75 (0.29)	5.95 (0.31)	17.70 (.15)
alternate-fixation	6.62 (0.12)	6.60 (0.15)	16.70 (.47)
control	5.90 (0.28)	6.12 (0.20)	-----

890
891
892
893
894
895
896
897
898
899
900
901
902
903
904
905
906
907
908
909

Table 2.

Adjusted means (\pm SE) for the EMG activation across all throws (normalized (%) to peak) at the biceps brachii, triceps brachii, flexor carpi radialis and extensor carpi radialis.

	Biceps	Triceps	Flexor	Extensor
simultaneous-congruent	48.35 (4.76)	49.45 (4.02)	46.74 (4.98)	60.46 (5.72)
alternate-congruent	52.31 (4.89)	52.4 (4.03)	53.17 (4.97)	58.47 (5.70)
simultaneous-incongruent	57.01 (4.78)	59.38 (4.06)	50.32 (5.05)	51.31 (5.70)
alternate-incongruent	40.54 (4.83)	60.90 (4.01)	57.16 (5.02)	58.41 (5.71)
simultaneous-fixation	63.12 (4.83)	66.86 (4.02)	52.38 (5.01)	52.12 (5.75)
alternate-fixation	51.20 (4.76)	55.89 (4.00)	48.11 (5.01)	46.92 (5.70)
control	50.86 (4.79)	68.31 (4.00)	59.88 (5.04)	54.33 (5.73)

910
911
912
913
914
915
916
917
918
919
920
921
922
923
924
925
926
927
928
929
930
931
932
933
934
935

Funding

This research did not receive any specific grant from funding agencies in the public,
commercial, or not for profit sectors

Conflict of interest

None