

1 Mistaking opposition for autonomy:
2 psychophysical studies on detecting
3 choice bias

4 Ashild Kummen¹, Patrick Haggard¹, Gwydion Williams¹, Lucie Charles¹

5

6 ¹ Institute of Cognitive Neuroscience, University College London, Alexandra House, 17 Queen
7 Square, London WC1N 3AZ, UK

8

9 Corresponding author:

10 Lucie Charles

11 Institute of Cognitive Neuroscience, University College London, Alexandra House, 17 Queen
12 Square, London WC1N 3AZ, UK;

13 Email: l.charles@ucl.ac.uk

14

15 **ABSTRACT (200 WORDS LIMIT)**

16 Do people know when they act freely and autonomously versus when their actions are
17 influenced? While the human aspiration to freedom is widespread, little research has
18 investigated how people perceive whether their choices are biased. Here, we explored how
19 actions congruent or incongruent with suggestions are perceived as influenced or free. Across
20 three experiments, participants saw directional stimuli cueing left or right manual responses.
21 They were instructed to follow the cue's suggestion, oppose it, or ignore it entirely to make a
22 'free' choice. We found that we could bias participants' 'free responses' towards adherence or
23 opposition, by making one instruction more frequent than the other. Strikingly, participants
24 consistently reported feeling less influenced by cues to which they responded incongruently,
25 even when response habits effectively biased them towards such opposition behaviour. This
26 effect was so compelling that cues that were frequently presented with the *Oppose* instruction
27 became systematically judged as having less influence on behaviour, artificially increasing the
28 sense of freedom of choice. Taken together, these findings demonstrate that acting contrarian
29 distorts the perception of autonomy. Crucially, we demonstrate the existence of a novel illusion
30 of freedom evoked by trained opposition. Our results have important implications for
31 understanding mechanisms of persuasion.

32

33

34

35 Keywords: Decision-making, Metacognition, Freedom of choice, Cognitive Control,
36 Psychophysics

37 1 INTRODUCTION (621)

38 How much do we know about the true reasons for our choices, and about the causes of our
39 decisions? Can we truly estimate if our choices are influenced by external factors?

40 Research in psychology has long demonstrated the limits of our voluntary control over our
41 decisions: choices can be swayed by factors beyond our control, and beyond our awareness.
42 For example, we may struggle to ignore sensory inputs even when we know these are
43 irrelevant for the decision, and try to ignore them. This is true when visual attention is captured
44 by distractors that present similarities with the target information like in the Eriksen Flanker
45 Task, [1,2] and its variants [3,4] or when a target stimulus is preceded by a non-conscious
46 prime, affecting response latency and choice [5].
47

48 Decisions are not only influenced by current information, but also by previous stimulation and
49 previous responses. Learned stimulus-response associations influence future performance
50 [6,7], and simple response repetition also biases current choices [8,9]. This influence of past
51 behaviour continues even when it becomes irrelevant or unrewarding, and can only be
52 overcome through additional cognitive control resources [10,11]. This is illustrated in task
53 switching paradigms in which participants frequently stick with the response they performed
54 on the previous trial rather than use the response mapping required for the present trial [12].

55 While the mechanisms of unwanted influence on choice behaviour have been well documented
56 [13] and formally modelled [14], it remains unclear whether people are able to detect when
57 their choices are biased by such influences. Research in social psychology suggests that
58 people are mostly unaware of the underlying drivers of their decisions [15,16]. Even when
59 participants are conscious of using biased strategies in their decisions, feelings of objectivity
60 prevail [17]. Equally, one might feel that extraneous influences are present even when they
61 are not – as during “thought insertion” in psychosis [18]. In other words, introspective access
62 to our own decision-making process may be limited.

63 Although “decision priming” has attracted wide scientific interest, few studies have investigated
64 the subjective experience of external influence on decisions. When participants were shown
65 ambiguous random dot kinematograms (RDK) and asked to respond ‘freely’, their actions were
66 unsurprisingly influenced by small fluctuations in motion energy [19]. Strikingly, however, they
67 reported stronger experiences of autonomy when they opposed the stimulus, than when they
68 followed it [22].

69 In the present study, we have systematically investigated the link between opposition and
70 subjective freedom. Across three experiments, we presented participants with random-dot
71 kinematogram stimuli in which the majority of dots moved either leftward or rightward. Each
72 display was accompanied by one of three instructions: respond congruently to the direction of
73 the motion (Adhere condition), respond incongruently to it (Oppose condition), or make a free
74 choice independent of the visual display (Detach condition). Thus, we could measure both the
75 extent to which participants successfully ignored stimulus information (by quantifying the
76 objective statistical relation between stimuli and response), and also their ability to introspect
77 this stimulus-independence (by obtaining ratings of subjective freedom of choice).

78 We then used a structured series of experimental manipulations to investigate the cognitive
79 basis underlying the sense of acting freely. In experiment 1, we presented participants more
80 frequently with one instruction (Adhere or Oppose) than the other, to test if a learned response
81 strategy would bias free choices. In experiment 2, we more specifically associated one
82 direction of motion (left or right RDK stimuli) with one instruction (Adhere or Oppose) to test
83 how using repeatedly one stimulus-response association would affect the perceived influence
84 of that stimulus on behaviour. Finally in experiment 3, we increased the proportions of one
85 direction of motion to test how the repetition of one visual cue would affect how participants
86 perceived their freedom of choice when responding to it.

87 2 METHOD (2151)

88 2.1 EXPERIMENT 1

89 2.1.1 Participants

90 62 participants with normal or corrected-to-normal vision and no history of psychiatric of
91 neurological illness were recruited. Sample size was based on a previous study [19] to detect
92 an influence of signal fluctuations on free choice with 80% power at $\alpha = .05$ (two-tailed t-test
93 for $d = 0.74$; required sample of at least 30 participants per group). The study was approved
94 by the UCL Research Ethics Committee and participants were compensated £12. Two
95 participants were excluded: Testing was interrupted for one participant as they failed to detect
96 motion direction during training. One participant used one end of the freedom of choice scale
97 in more than 98% of the trials and was therefore excluded from analysis. This left 60
98 participants (43 female; mean age 22.63 ± 3.17 years ranging from 20 to 30), randomly
99 assigned to 30 in Adhere Group and 30 in Oppose Group.

100 2.1.2 Stimuli

101 Participants were presented with a series of random dot kinematograms (RDK) moving either
102 to the left or to the right (see Stimuli in Supplementary Methods). The proportion of the dots
103 moving in the same direction representing the motion coherence was adjusted by a Staircase
104 Procedure to reach 71% coherence (see Staircase Procedure in Supplementary Methods). At
105 a random time during stimulus presentation, the dots changed colour (red, green, or blue;
106 matched for luminance and contrast) providing the instruction on how to respond in that
107 particular trial. On (1) *Adhere* trials, participants were to respond congruently with the direction
108 of the motion they perceived in the stimulus; on (2) *Oppose* trials, participants were to respond
109 with the response key on the opposite side to the perceived motion direction; on (3) *Detach*
110 trials, participants were to choose freely which response key to press, while trying to ensure
111 that their choice was not influenced by the direction of dot motion. In those trials, participants
112 were told not to use a set response strategy, such as responding always with the same key
113 press or alternating between key presses, but try to act randomly.

114 2.1.3 Design & Procedure

115 In this experiment, one group of participants received the instruction to adhere (i.e. respond
116 congruently to the motion direction) in most trials (Adhere group) while a second group
117 received the instruction to oppose (i.e. respond incongruently) in most trials (Oppose group).
118 This design allowed us to test whether a habitual response strategy would influence: (1) the
119 objective ability to detach from the stimuli when given a free choice (objective FoC), and (2)
120 the subjective feeling of acting freely (subjective FoC).

121 **Protocol.** The experiment followed a mixed design with group as the between-subjects
122 variable (Adhere Group, Oppose Group), and instruction condition (Adhere, Oppose, Detach)
123 as the within-subject variable. The main experiment consisted of 7 blocks of 57 trials, totalling
124 to 378 trials of interest (excludes high coherence trials). For the majority Adhere Group, this

125 was split so that 50% were Adhere trials, 16.7% Oppose, and 33.3% Detach. For the majority
126 Oppose Group: 50% of trials were Oppose trials, 16.7% Adhere, and 33.3% Detach (see
127 Figure 1). The experimental session lasted 80-90 minutes and was divided in two parts; the
128 staircase procedure, and the main experiment.

129 **Trial procedure.** In the main experiment, participants started each trial with a fixation cross,
130 before the stimulus was presented for 2500ms (see Figure 1A & B). At a random time-point
131 between 100ms and 2388ms, the dots changed colour to indicate the instruction for this trial
132 (see Figure 1C). Association between colours and instructions were counterbalanced across
133 participants. After the stimulus disappeared, participants had 1500ms to respond with a left-
134 arrow and right-arrow keypress using their right hand (see Figure 1C).

135 Following the response, participants were required to estimate how much their response
136 choice was guided by the dot-motion, reporting their subjective experience of being more or
137 less influenced by the stimulus (Figure 1E). To do so, a scale was presented for 3000ms and
138 participants could move the slider with the '<' and '>' keys. The starting position of the slider
139 was set at a random position on each trial. One end of the scale (counterbalanced across
140 participants) indicated absolute independence of the stimulus ('I decided what to do myself,
141 completely independently of what I saw on the screen'), while the other indicated complete
142 dependent on the stimulus ("My response was determined entirely by what I saw on the
143 screen"). Participants were instructed to use the entire range of the scale rather than only the
144 extreme values and that their response should be guided by how they came up with their
145 response on that given trials rather than the instruction.

146 **2.1.4 Data Analysis**

147 Trials in which no response was given were excluded (Adhere: $M = 4.20\%$ of trials, $SD = 5.65\%$;
148 Oppose: $M = 5.08\%$, $SD = 5.16\%$; Detach: $M = 6.60\%$, $SD = 6.88\%$). Each response was
149 classified as congruent or incongruent with the stimulus direction (response mode factor). The
150 mean proportion of motion congruent responses was calculated for each participant and each
151 instruction. A repeated measure ANOVA was used to estimate the effects of group (between-
152 participant factor) and instruction (within-participant factor) on the proportion of motion-
153 congruent responses for instructed trials (i.e., Adhere and Oppose trials). A separate ANOVA
154 was run for the Detach trials to estimate the effect of group (between-participant factor) on the
155 proportion of motion congruent responses. Mean subjective FoC ratings were calculated for
156 each combination of participant, instruction, and response mode. A repeated measure ANOVA
157 was used to estimate the effect of group (between-participant factor), instruction and response
158 mode (within-participant factor) on the reported freedom of choice in instructed trials. (Adhere
159 and Oppose trials). Follow-up t-tests (two-tailed, unless specified otherwise) were used to
160 compare ratings in error and correct trials for each instruction. A separate repeated measure
161 ANOVA was run on the subjective FoC ratings in the Detach condition to estimate the effect
162 of group (between-participant factor) and response mode (within-participant factor) on the
163 reported freedom of choice.

164

165 **2.2 EXPERIMENT 2**

166 **2.2.1 Participants**

167 39 participants were recruited. Sample size was determined by a-priori power analysis based
168 on the subjective FoC observed in experiment 3 (conducted prior to experiment 2), an effect
169 size of $d = 0.29$ requiring $N=37$ to achieve 80% power, with alpha 0.05 (two-tailed t-test). One
170 participant was excluded as they confused instructions. This left 38 participants (28 females;
171 mean age 22.24 ± 2.94 years ranging from 18 to 30; all right-handed). Participants were
172 randomly assigned to a group (20 for majority left motor responses; 18 for majority right motor
173 responses).

174 **2.2.2 Design, Stimuli, & Procedure**

175 Experiment 2 used a similar design than experiment 1 but that each stimulus direction was
176 now associated more frequently with one instruction. Importantly, this also meant that
177 participant used more frequently one motor response than the other (Figure 1). We therefore
178 also included a second type of detach trial where the stimulus did not have any obvious motion
179 direction (motion-neutral') to estimate the effect of purely motoric response repetition on the
180 ability to detach.

181 The experiment followed a mixed design with group (majority left responses, majority right
182 responses) as the between-subjects variable and instruction (Adhere, Oppose, Detach) as the
183 within- subject variables. The stimuli and procedure for experiment 2 were identical to that of
184 experiment 1, except for the modifications in the proportion of trials: In 469 trials, 59.7% of
185 trials were instructed (half Adhere, half Oppose). Among instructed trials, the proportion of
186 leftward and rightward stimuli differed between instructions. On Adhere trials, 75% of stimuli
187 were in a direction congruent to the most frequent response of the group (e.g. rightward for
188 the "majority right responses" group), whilst the remaining 25% were in the other direction. On
189 Oppose trials, 75% of stimuli were in the incongruent direction to the most frequent response
190 of the group (e.g. leftward for the "majority right responses" group), and 25% in the congruent
191 direction (see Figure 1).

192 The remaining trials (40.3%) corresponded to the Detach instruction. Among the Detach trials,
193 a third were moving leftward, a third moving rightward, and a third were motion-neutral where
194 the motion coherence was set to 0% and *all* dots moved at random. These trials were added
195 to investigate whether the repeated motor response would influence free choices in the
196 absence of visual stimulation.

197 At the end of the experiment, participants were asked to estimate the overall proportion of right
198 responses they had made (see Bias Awareness in Supplementary Results).

199 **2.2.3 Data Analysis**

200 Data analysis followed the same methods as for experiment 1. We separated trials according
201 to whether the stimuli were frequently adhered, or frequently opposed. Trials in which no
202 response were given were excluded (Adhere: $M = 3.36\%$, $SD = 3.14\%$; Oppose: $M = 3.20\%$,

203 $SD = 3.70\%$; motion-present Detach: $M = 5.33\%$, $SD = 4.05\%$; motion-neutral Detach: $M =$
204 4.64% , $SD = 3.84\%$). Mean proportions of motion congruent responses and mean subjective
205 freedom of choice ratings were calculated per participant, split by instruction, stimulus-
206 response association, and response mode and analysed using repeated measures ANOVAs
207 (group as a between subject variable, and instruction, stimulus-response association, and
208 response mode as within subject variables), similarly to Experiment 1. On motion-neutral
209 detach trials, we tested whether small fluctuations in motion energy would influence response
210 choice using reverse correlation (see Reverse Correlation Analysis in Supplementary
211 Methods). Responses were classified by whether they were congruent or incongruent to the
212 motion direction fluctuation and analysed following the same procedure as motion-present
213 trials.

214 **2.3 EXPERIMENT 3**

215 **2.3.1 Participants**

216 42 participants were tested using similar procedures to experiment 1 and 2 (sample size
217 indicated to achieve power of 80% for two-tailed t-test with effect size $d = 0.55$ from experiment
218 1). Five participants were excluded. Two failed to see the dot motion direction during training,
219 one used one end of the scale in more than 98% of trials or two had ceiling accuracy in
220 instructed trials for rare stimulus direction, making it impossible to compare accuracy across
221 response modes. This left 37 participants (24 females; mean age 24.35 ± 2.94 ranging from
222 18 to 30, all but five right-handed) allocated randomly to each group (20 participants: majority
223 leftward stimuli; 17 majority rightward stimuli).

224 **2.3.2 Design, Stimuli, & Procedure**

225 To understand how repetition vs salience of stimuli may influence the ability to detach, the
226 previous design was modified so that now one motion direction (leftward or rightward) was
227 seen more frequently than the other. This allowed us to compare trials where a frequent
228 stimulus was presented and trials where a rare stimulus was presented.

229 The experiment followed a mixed design with group (majority left-direction stimuli, majority
230 right-direction stimuli) as the between-subjects factor, and instruction condition (Adhere,
231 Oppose, Detach) and stimulus frequency (frequent stimulus, rare stimulus) as the within-
232 subjects factors.

233 Stimuli and procedure were identical to those of experiment 1 and 2. The main modification
234 was that 75% of all trials (except motion-neutral trials) had stimuli moving in the frequent
235 direction (determined by participant's group), whilst the other 25% moved in the other direction
236 (rare stimulus). A total of seven blocks with 60 trials of interest (excluding high coherence trials)
237 was presented, giving a total of 420 trials of interest. In each block, 36 trials (60%) were
238 instructed (half Adhere, half Oppose). The remaining 24 (40%) were detach trials; of which six
239 (25%) were motion-neutral and 18 (75%) motion-present.

240 At the end of the experiment, participants were asked to estimate the proportion of leftward
241 versus rightward stimuli (see Bias Awareness in Supplementary Results, for the findings).

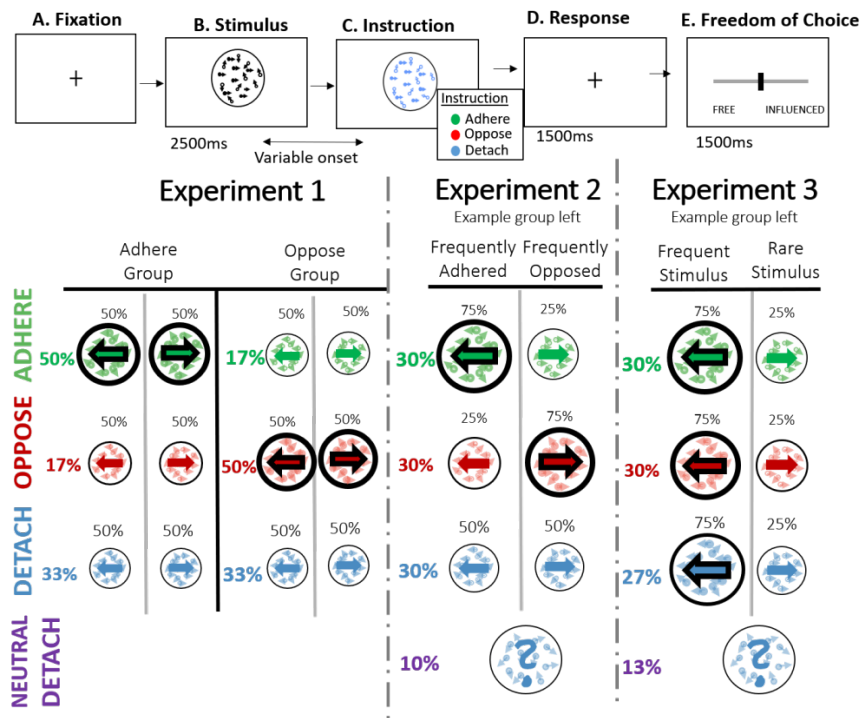
242 2.3.3 Data Analysis

243 Trials with no response were excluded from the analysis (Adhere: $M = 8.34\%$, $SD = 10.38\%$;
244 Oppose: $M = 8.71\%$, $SD = 10.91\%$; motion-present Detach: $M = 10.56\%$, $SD = 14.35\%$;
245 motion-neutral Detach: $M = 10.94\%$, $SD = 16.45\%$). Mean proportion of motion congruent
246 responses and mean subjective freedom of choice ratings were calculated per participant, split
247 by instruction, stimulus frequency, and response mode and analysed using repeated measures
248 ANOVAs (group as a between subject variable, and instruction, stimulus frequency, and
249 response mode as within subject variables), similarly to Experiment 1&2. Motion-neutral
250 detach trials were analysed following the same procedure as Experiment 2.

251 2.3.4 Data Analysis across experiments

252 In an exploratory analysis, we tested whether participants who more strongly used one
253 response strategy in detach trial (for instance adhering more to the stimulus direction) would
254 also report lower subjective freedom when using that same strategy in detach trials. To do so,
255 we computed the difference in proportion of each participant's congruent and incongruent
256 responses in the detach condition and tested whether a linear relationship was present with
257 the difference in subjective freedom ratings between pro and incongruent in a regression
258 model. To maximize power, we pooled the data across all experiments. We also performed
259 the same regression in the specific parts of our design that most encouraged opposition during
260 detachment (Oppose Group of experiment 1, frequently opposed stimuli condition of
261 experiment 2, and frequent stimuli condition of experiment 3).

262

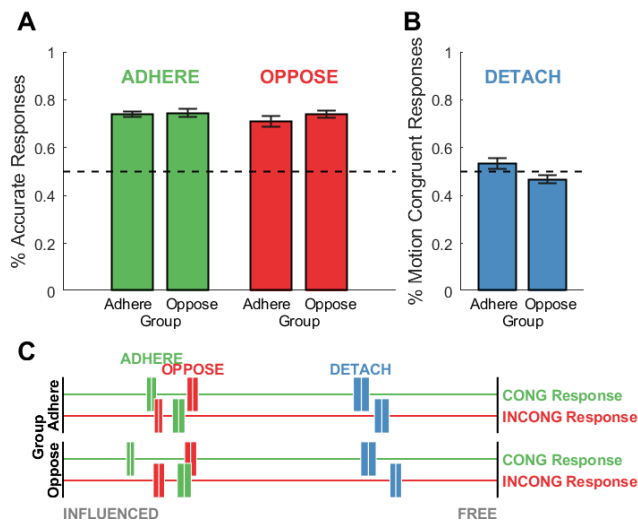
263 **3 RESULTS**264 **3.1 IMPACT OF DOMINANT INSTRUCTION ON FREEDOM OF CHOICE (EXPERIMENT**
265 **1)**

266

267 **Figure 1.** Experimental procedure (top) and design (bottom) of experiments 1-3. (Top)
 268 Following a fixation cross (A), the stimulus was presented for 2500ms (B). At a random
 269 timepoint during stimulation, the dots changed colour to indicate the instruction for that trial
 270 (Adhere, Oppose, or Detach; C). After giving their response (D), participants were asked to
 271 report their subjective feeling of freedom (E). (Bottom) Experiment 1 manipulated the
 272 frequency of instructions between groups so that one group were required to adhere more than
 273 to oppose (example highlighted by bold outline), while the other were required to oppose more
 274 than to adhere. Experiment 2 manipulated the total frequency of one response (example of
 275 Left motor-response group highlighted by bold-outline) by arranging for one stimulus direction
 276 to be preferentially associated with Adhere trials (frequently adhered) and the other stimulus
 277 direction with Oppose trials (frequently opposed). Experiment 3 manipulated the stimulus
 278 frequency so that each group saw more of one stimulus direction than the other (example of
 279 Left stimulus group highlighted by bold-outline).

280 Response accuracy was not affected by instruction frequency (Figure 2A) [$F(1, 58) = 2.77, p$
 281 $= .102, \eta_p^2 = .046$] or by group [$F(1, 58) = 0.59, p = .447, \eta_p^2 = .010$]. However, participants
 282 responded faster on Adhere than Oppose trials [$F(1, 58) = 7.32, p = .009, \eta_p^2 = .112$], as well
 283 as when they responded correctly compared to when they made an error [$F(1, 58) = 66.25, p$

284 < .001, $\eta_p^2 = .533$]. When they were instructed to detach from the stimulus (Figure 2B),
 285 participants in the Adhere group made more congruent responses on detach trials than
 286 participants in the Oppose group [$t(58) = 2.44$, $p = .018$, $d = 0.63$, two-tailed]. Considering each
 287 instruction separately, we observed non-significant trends for participants in the Oppose Group
 288 to make more incongruent responses ($M = 46.74\%$, $SD = 9.06\%$; $t(29) = -1.97$, $p = .06$, $d = -$
 289 0.36 , two-tailed) while participants in the Adhere Group tended to respond congruently in 53.49%
 290 of detach trials ($M = 53.49\%$, $SD = 12.12\%$, $t(29) = 1.58$, $p = .13$, $d = 0.29$, two-tailed). Taken
 291 together, these results suggest that the dominant instruction affected detach trials and habitual
 292 response strategies continued to prevail even when attempting to detach.



293

294 **Figure 2.** Results of Experiment 1. **A.** Mean proportion of correct responses on instructed trials
 295 (Adhere, green; Oppose, red). **B.** Proportion of motion congruent responses on Detach trials
 296 (blue) split by group (Adhere Group, Oppose Group). Error bars show standard error of the
 297 mean. **C.** Mean subjective freedom of choice ratings split by group (Adhere Group, Oppose
 298 Group), instruction (Adhere, Oppose, Detach), and response (congruent vs incongruent
 299 response; green and red lines respectively), for each stimulus motion direction. Ratings range
 300 from feeling completely influenced (left) to completely free (right). For instructed trials (Adhere:
 301 vertical green bars; Oppose: vertical red bars) matching bar and line colours (Adhere trials:
 302 green bars on green lines; Oppose trials: red bars on red lines) correspond to correct
 303 responses while bars on opposite colour lines (Adhere trials: green bars on red lines; Oppose
 304 trials: red bars on green lines) correspond to errors. Box width reflect +/- 1 standard error
 305 across participants..

306 We next considered how participants in each group rated their freedom of choice in instructed
 307 and free-choice trials. On instructed trials (Adhere and Oppose), participants reported feeling
 308 freer on trials where they were instructed to oppose the motion of the stimulus than when
 309 instructed to adhere to it [Figure 2C; main effect of instruction; $F(1, 58) = 11.43$, $p = .001$, η_p^2
 310 $= .165$]. Moreover, they also reported feeling freer when making an error (e.g. incongruent
 311 responses for Adhere trials depicted by green bars on red lines on Figure 2C and congruent
 312 responses for Oppose trials depicted by red bars on green lines on Figure 2C) than when

313 making an accurate response [interaction between instruction and response mode; $F(1, 58) =$
 314 $88.42, p < .001, \eta_p^2 = .604$; Adhere instruction, Correct vs Error : $t(59) = -7.09, p < .001, d = -$
 315 0.92 ; Oppose instruction, Correct vs Error : $t(59) = -8.06, p < .001, d = -1.04$]. Participants
 316 may have paid less attention to the stimulus when they made an error than when responding
 317 correctly. An unseen or unattended stimulus cannot strongly influence behaviour. Thus,
 318 stronger sense of freedom suggests that participants used the subjective scale appropriately.

319 Turning to detach trials, we found that when making a response incongruent with the mean
 320 stimulus-motion direction, participants felt significantly freer (i.e., more detached from the
 321 stimulus) than when making a response congruent with the motion direction [Figure 2C; main
 322 effect of response mode; $F(1, 58) = 18.12, p < .001, \eta_p^2 = .238$]. Importantly, this was the case
 323 in the Adhere Group [$t(29) = 2.68, p = .012, d = 0.49$] but also in the Oppose Group [$t(29) =$
 324 $3.32, p = .002, d = 0.61$], and the effect did not differ significantly between the groups [$F(1, 58)$
 325 $= 0.37, p = .54$]. Since the Oppose group made incongruent responses more frequently than
 326 the Adhere, this suggests that opposition was associated with increased sense of freedom,
 327 irrespective of whether opposition was the most frequent behaviour or not. In other words,
 328 acting contrarian was always associated with an increased sense of freedom, even when
 329 opposition was the dominant response tendency.

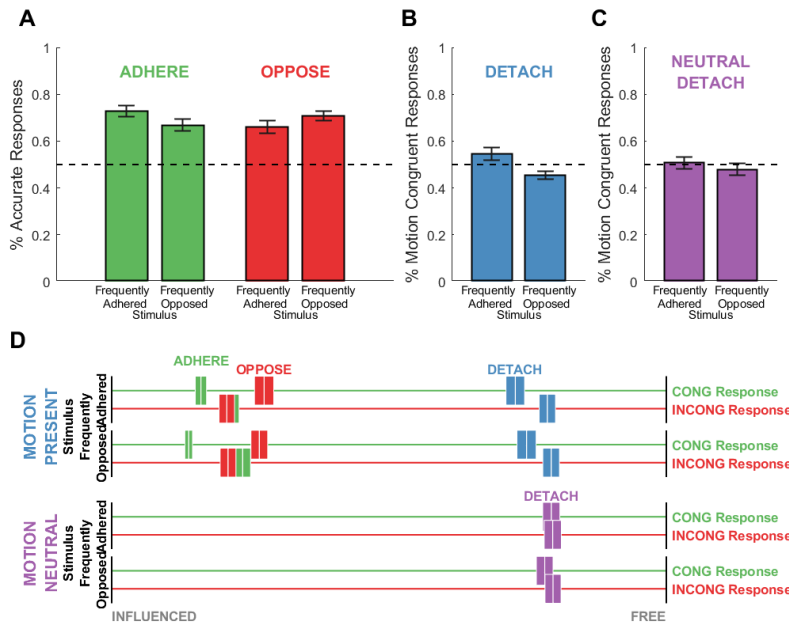
330

331 **3.2 IMPACT OF LEARNED STIMULUS-RESPONSE ASSOCIATION ON FREEDOM OF** 332 **CHOICE (EXPERIMENT 2)**

333 While we showed that participants' objective ability to detach was partially compromised by
 334 following repeatedly one instruction over the other, we wondered whether this persistence
 335 effect would also be observed when one specific stimulus-response association was repeated,
 336 as opposed to one instruction. This was assessed in Experiment 2.

337 In instructed trials, we found that participants were more accurate in the condition
 338 corresponding to the trained stimulus-response association (Figure 3A). As in the first
 339 experiment, participants were also faster on Adhere than on Oppose trials [$F(1, 36) = 4.38, p$
 340 $= .044, \eta_p^2 = .108$], and faster for correct than incorrect responses [$F(1, 36) = 39.53, p < .001,$
 341 $\eta_p^2 = .523$]. When required to detach, (Figure 3B), we found that participants responded
 342 differently to stimuli that they had been trained to adhere and those they had been trained to
 343 oppose [$t(37) = 2.56, p = .015, d = 0.42$, two-tailed]: Participants tended to respond more
 344 congruently to frequently adhered stimuli [$M = 54.56\%, SD = 10.68\%, t(37) = 1.69, p = .050, d$
 345 $= 0.27$, one-tailed] and to respond more incongruently to stimuli they were trained to oppose
 346 [$M = 45.57\%, SD = 10.36\%, t(37) = 2.64, p = .006, d = 0.43$, one-tailed]. This result therefore
 347 replicated and extended those of Experiment 1, showing that training with one particular
 348 stimulus-response association continued to influence response strategy when required to
 349 detach. No effect of motor bias was observed in the motion-neutral trials [proportion of frequent
 350 responses compared to chance level on motion-neutral detach trials; $t(37) = 0.66, p = .510, d$
 351 $= 0.11$] and no further differences were observed when considering if the motion energy
 352 fluctuations favoured the frequently adhered or frequently opposed stimulus direction [Figure

353 3C; $t(37) = 0.65, p = .520, d = 0.10$], suggesting that purely motor repetition in itself did not
 354 affect the ability to detach.



355

356 **Figure 3. Results of Experiment 2.** Accuracy for instructed trials (**A**) and percentage of
 357 motion congruent responses for motion-present Detach trials (**B**) and motion-neutral Detach
 358 trials (**C**) split by stimulus-response association (frequently adhered, frequently opposed). (**D**)
 359 Mean ratings of subjective ratings of freedom of choice split by stimulus-response association
 360 (frequently adhered, frequently opposed). Same legend as Figure 2.

361 Considering the ratings of freedom of choice (Figure 3D), we found that participants reported
 362 greater detachment on Oppose than on Adhere trials [$F(1, 36) = 5.13, p = .030, \eta_p^2 = .125$], and
 363 when making an error (e.g. incongruent responses for Adhere trials depicted by green bars on
 364 red lines on Figure 3C or a congruent response for Oppose trials depicted by red bars on green
 365 lines on Figure 3C; interaction between response mode and instruction: $F(1, 36) = 33.83, p$
 366 $< .001, \eta_p^2 = .484$; Adhere instruction, Correct vs Error $t(37) = -4.26, p < .001, d = -0.69$; Oppose
 367 instruction, Correct vs Error : $t(37) = -2.9, p = .006, d = -0.47$) replicating the results of our
 368 previous experiment. In detach trials (Figure 3D, blue bars), participants again reported they
 369 felt freer making a motion incongruent than a motion congruent response [$F(1, 36) = 12.55, p$
 370 $= .001, \eta_p^2 = .258$], confirming that opposition was always associated with an increased sense
 371 of freedom. Interestingly however, we also found an unexpected effect of previous opposition
 372 in detach trials: participants reported they felt more free when responding to a frequently-
 373 opposed stimulus compared to a frequently adhered stimulus [Main effect of stimulus-response
 374 association; $F(1, 36) = 4.34, p = .044, \eta_p^2 = .108$]. This was true regardless of the response
 375 made on the current trial [$F < 1$]. Hence, when attempting to detach, frequently-opposed stimuli
 376 became associated in themselves with an increased sense of autonomy, irrespective of
 377 whether participants actually opposed them on a given trial or not.

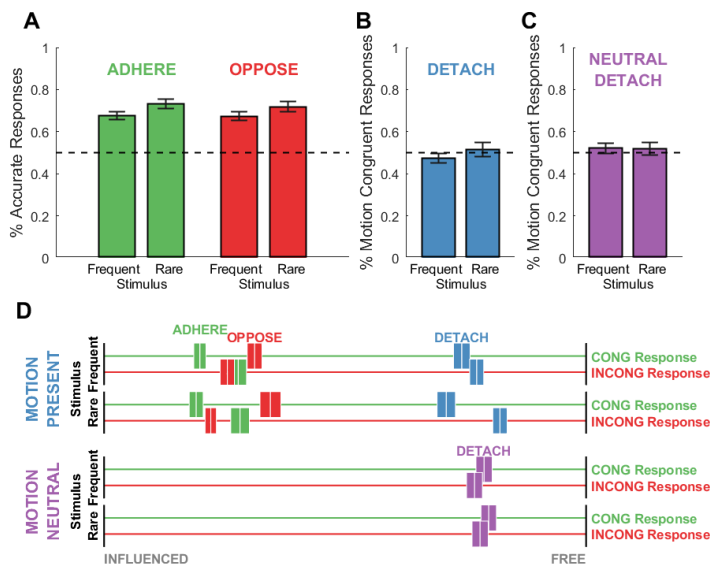
378 Interestingly, no effect of stimulus-response association or motor bias was observed on ratings
 379 of freedom in the motion-neutral detach trials (Figure 3D, purple, all $F < 1$) suggesting that
 380 motor novelty vs. motor repetition did not itself influence subjective freedom.

381 3.3 IMPACT OF STIMULUS FREQUENCY ON FREEDOM OF CHOICE (EXPERIMENT 3)

382 We observed that participants reported feeling freer when presented with a stimulus they had
 383 frequently opposed in the past, compared to one frequently adhered to, regardless of their
 384 response to it. We then tested whether stimulus frequency itself impacted freedom of choice.

385 Stimulus frequency did not significantly affect response accuracy in instructed trials Figure 4A,
 386 although a trend was observed towards participants performing better for frequent stimuli [$F(1,$
 387 $35) = 3.25, p = .08, \eta_p^2 = .085$]. As previously observed, participants responded more rapidly
 388 on Adhere trials compared to Oppose trials [$F(1, 35) = 6.88, p = .013, \eta_p^2 = .164$]. Accurate
 389 responses were again associated with faster response times than inaccurate responses [$F(1,$
 390 $35) = 41.81, p < .001, \eta_p^2 = .544$].

391 In motion-present detach trials (Figure 4B, blue bars), no effect of stimulus frequency was
 392 observed on the proportion of motion congruent responses [frequent stimulus vs rare: $t(36) =$
 393 $1.08, p = .290, d = 0.18$; frequent stimulus vs chance level: $t(36) = 1.18, p = .250, d = 0.19$;
 394 rare stimulus vs chance level: $t(36) = 0.48, p = .640, d = 0.08$], suggesting participants were
 395 equally likely to oppose dot-motion as they were to adhere to it, irrespective of whether the
 396 motion direction had been seen frequently or not. No further effects were observed on motion-
 397 neutral trials (Figure 4C), indicating participants had no response bias in the absence of a clear
 398 visual cue.



399

400 **Figure 4. Results of Experiment 3.** Accuracy for instructed trials (A) and percentage of
 401 motion congruent responses for motion-present Detach trials (B) and motion-neutral Detach

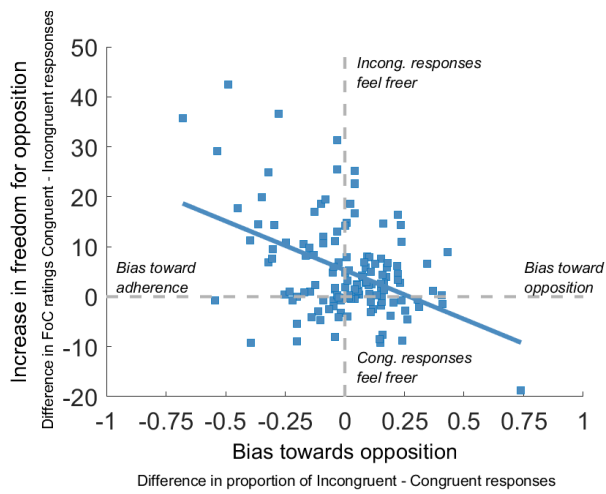
402 trials (C) split by stimulus frequency (frequent, rare). (D) Mean ratings of subjective ratings of
 403 freedom of choice split by by stimulus frequency (frequent, rare). Same legend as Figure 2.

404 As in our two previous experiments, we observed that freedom was rated higher on Oppose
 405 than Adhere trials [Figure 4D; $F(1, 35) = 10.29, p = .003, \eta_p^2 = .227$] and when making an error
 406 than when responding correctly [$F(1, 35) = 33.88, p < .001, \eta_p^2 = .492$; Adhere instruction,
 407 Correct vs Error $t(36) = -4.6, p < .001, d = -0.76, d = -0.69$; Oppose instruction, Correct vs
 408 Error : $t(36) = -3.6, p < .001, d = -0.59$]. Further, in detach trials, we replicated the findings of
 409 experiment 1 and 2 that participants reported feeling more free when making a response
 410 incongruent to the stimulus direction (Figure 4D, blue bars), compared to a congruent response
 411 [$F(1, 35) = 14.35, p < .001, \eta_p^2 = .291$]. Interestingly, we observed an interaction between
 412 stimulus frequency and response mode [$F(1,35) = 16.85, p < .001, \eta_p^2 = .325$], showing that
 413 participants reported feeling more free when making an incongruent response to a rare
 414 stimulus, compared to a frequent stimulus [$t(36) = 3.01, p = .005, d = 0.50$]. This suggested
 415 that the sense of freedom was increased when opposing a rare occurrence. No further effects
 416 were observed in motion-neutral detach trials.

417 3.4 EXPLORING CORRELATIONS BETWEEN SUBJECTIVE AND OBJECTIVE 418 FREEDOM OF CHOICE

419 Finally, we tested whether participants who more strongly tended to adhere to the stimulus
 420 would also experience higher subjective freedom when opposing. Pooling data across all
 421 experiments, we found that this was indeed the case (Figure 5, regression slope = -19.62, beta
 422 = 5.31, $R^2 = .19, t(133) = 5.53, p < .001$; see Table S3 in Supplementary Materials).
 423 Interestingly, the intercept of this regression relation showed that, even when participants
 424 objectively detached, tending neither to oppose nor adhere, there remained a strong and
 425 significant tendency to rate incongruent responses as subjectively freer (beta = 5.32, $t(133) =$
 426 $7.06, p < .001, d = 1.22$).

427 Performing the same regression in the specific parts of our design that most encouraged
 428 opposition during detachment (Oppose Group of experiment 1, frequently opposed stimuli
 429 condition of experiment 2, and frequent stimuli condition of experiment) however, revealed no
 430 significant correlation between the degree of detachment and the subjective sense of freedom
 431 for congruent versus incongruent responses (dataset with majority Oppose behaviour: p
 432 = .130). This suggests that this effect might be more strongly associated with adherence
 433 strategy. However, the intercept of the regressions remained significantly different from 0 ($b =$
 434 $5.41, t(102) = 4.43, p < .001, d = 0.87$), suggesting that even when participants showed an
 435 objective tendency to oppose the stimulus, they still experienced greater freedom when
 436 opposing. Taken together, these results show that both habitual and contrarian behaviour
 437 modulated the sense of freedom of choice, independently of each other.



438

439 **Figure 5.** Difference in subjective freedom of choice (FoC) to incongruent minus congruent
 440 responses on motion-present detach trials according to the difference in proportion of
 441 incongruent minus congruent responses on motion-present detach trials. Each data-point
 442 represent one participant, pooling together all participants from all three experiments.

443

444 **4 DISCUSSION (1358)**

445 How do we know whether we are acting freely, or are influenced in our choices? And
446 is our introspection of influence accurately reflecting our learned biases? In the present study,
447 we tested how the tendency to respond congruently or incongruently to suggestion affected
448 the sense of acting autonomously. To do so, we showed participants stimuli that primed a left
449 or a right response and asked them to make a manual choice by either adhering to the direction
450 suggested, opposing it, or detaching entirely from the visual stimulation to make a choice free
451 from external influence. Crucially, we manipulated the stimulus and response distributions so
452 that participants 1/ had to follow one set of instructions more frequently (experiment 1), 2/ had
453 to respond to one stimulus with one response more frequently (experiment 2), or 3/ were
454 exposed to one stimulus direction more frequently (experiment 3). This systematic set of
455 designs allowed us to test how repeated association between a stimulus and a response could
456 bias free decisions, and also how it could alter the subjective experience of acting freely.

457 Experiment 1 and 2 revealed that participants' ability to make free choices in the detach
458 condition, was strongly compromised by implicitly learned stimulus-response associations. In
459 Experiment 1, we observed that when participants were required to detach from the stimulus
460 and attempted not to be influenced by it, they failed to do so. Instead, they tended to act in
461 accordance with the instruction that they had been trained with the most frequently. This meant
462 that stimulus-response mappings that were formed through repetition persisted in free choice.
463 In experiment 2, participants were predominantly trained to make stimulus-congruent
464 responses to some stimuli, while predominantly opposing others. Participants' free choices
465 tended to perseverate in using the trained stimulus-response mappings. The ability to act
466 autonomously and independently of a stimulus was therefore reduced if one had previously
467 learned to make a particular response to that stimulus.

468 Such findings are in broad agreement with the classical concept of conditioned or
469 habitual responding [21], as well as literature on task-switch costs. Only a few studies
470 investigated how free choices themselves are impacted by previous experience of SR
471 associations [22]. Our study shows that task history profoundly constrains the capacity for truly
472 autonomous action. We show here that even when people try to ignore current stimulus
473 content, their previous experiences strongly shape their behaviour. Importantly, persistence of
474 SR mappings has also been found in subliminal priming paradigms [23], suggesting that it
475 might operate independently of conscious processes of action selection [24]. Thus, the
476 'positive priming effect', in which participants tend to respond congruently to a subliminal prime
477 [25–27] or be positively entrained by a non-conscious stimulus [28] could be reversed if the
478 participants had been trained with the incongruent stimulus-response mapping [23]. Similarly,
479 it has been observed that increasing the proportion of instructed trials strengthens the positive
480 priming effect on free choices [29,30].

481 Our findings extend those results to free choices in the presence of supraliminal stimuli
482 [31–34]. To act freely in the present study meant detaching from stimulus information.
483 Participants however showed only a limited ability to dissociate from previously trained
484 strategies. Crucially, we show that these are not merely effects of visual repetition alone, nor
485 of motor priming alone. Thus, future free choices were influenced by the repeated association
486 of one stimulus with a response, but not by patterns of stimulation alone, or by patterns of

487 responding alone [35]. This finding suggests that volition and habit represent two dissociable,
488 competing and mutually-exclusive routes to action [36] and that true detachment may be
489 difficult to achieve once task habits are present, requiring effortful cognitive control to
490 overcome habit-induced response conflicts [11]. Further research will be needed to understand
491 the mechanisms of such effect and whether it is caused by occasional lapses in voluntary
492 control, like the lapses of attention that cause errors in flanker tasks [11] or whether it
493 corresponds to a more sustained “response priming” effect [23].

494 Importantly, our results bring new lights on whether participants are able to introspect
495 the factors that influenced their choices. Crucially, we found that across all three experiments
496 responding incongruently to the stimulus was associated with an increased subjective sense
497 of acting freely and independently from the stimulus, compared to responses that followed the
498 stimulus. Interestingly, this association between opposition and perceived autonomy was
499 present irrespective of how much the stimulus in fact influenced participants’ choices. In
500 particular, whether participants more frequently followed the cued response, or more frequently
501 opposed it, they always rated their choice as more autonomous when opposing the action
502 suggested by the current circumstances. Indeed, even those participants who tended
503 statistically to oppose the stimulus direction when required to detach nevertheless felt freer
504 when they made incongruent responses on detach trials than when they made a congruent
505 response (Experiment 1). This result was confirmed further by combining the conditions in
506 each of the three experiments in which participants tended to make more opposing responses.

507 Inhibitory control therefore distorts the introspection of one’s own choices: opposition
508 increases the feeling of acting freely. A pooled regression further showed that this relation
509 between opposition and subjective freedom was present even when controlling for participants’
510 dominant response tendency, and for the time they took to make a choice. These findings
511 replicate and extend previous results on the sense of autonomy and freedom [19]. Importantly,
512 we demonstrate here that while free choices can be biased by experimental manipulation, the
513 subjective bias caused by inhibitory control persists despite those manipulations. Thus, the
514 subjective sense of autonomy seems to be driven largely by signals monitoring local conflict
515 related to the external world, and processes that overcome those conflicts. Indeed, participants
516 already rated being less influenced by the stimulus when they followed the Oppose instruction
517 than when they were instructed to Adhere. The effect may recall the feeling of resisting external
518 social influence, as when dissenting from the majority opinion [37] or disobeying norms such
519 as traffic laws [43] [44,45]. The sense of thrill that accompanies rebellion or resistance to
520 external influence, might be due to the metacognitive bias that makes acting contrarian feel
521 like autonomy and freedom. As such, our findings might provide empirical evidence and
522 potential cognitive mechanisms underlying social psychology phenomenon such as reactance
523 [39,40]. Our results suggest that contrarian behaviours that aim to restore or boost subjective
524 autonomy might actually stem from an underlying metacognitive bias in introspecting one’s
525 own decisions.

526 One interesting, and to our knowledge novel, result of this study concerns how
527 participants linked an external stimulus to the feeling of acting freely. When participants
528 learned by repeated experience to make stimulus-incongruent responses, they appeared to
529 start associating that stimulus with an increased sense of freedom. Importantly, this effect was
530 present irrespective of whether participants indeed acted independently from that stimulus or
531 not on a specific trial (experiment 2). This finding shows that the sense of freedom that arose
Page 18 of 33

532 from opposing the influence of an external stimulus stuck to that stimulus, so that the stimulus
533 itself becomes associated with acting freely. This result is striking because the experience of
534 *not* being influenced was paradoxically evoked by the mere exposure to a visual cue. Such
535 findings resonate with the literature on instrumental conditioning where a stimulus can become
536 associated with a particular emotional [21] or cognitive response [41]. Our result could be
537 interpreted as a novel illusion of autonomy, arising from people's limited ability to introspect
538 the reasons for their choices [19]. This result raises the interesting possibility that the sense of
539 autonomy can be increased artificially by training, since an illusory feeling of freedom could be
540 induced by exposure to things that we have been told to reject. Such findings could shed new
541 light on social effects such as irrational belief in conspiracy theories or extreme political
542 radicalization. Indeed, a cognitive bias which conflates opposition and freedom could explain
543 why people might feel autonomous when they are led into rebellions or manipulated into acting
544 contrarian. More research will be needed to determine how our findings can be generalised to
545 contexts of autonomous choice outside the laboratory, and how artificially generated feeling of
546 freedom might influence behaviour.

547

548

REFERENCES

- 549 1. Eriksen BA, Eriksen CW. 1974 Effects of noise letters upon the identification of a target
550 letter in a nonsearch task. *Perception & Psychophysics* (doi:10.3758/BF03203267)
- 551 2. Stroop JR. 1935 Studies of interference in serial verbal reactions. *Journal of*
552 *Experimental Psychology* (doi:10.1037/h0054651)
- 553 3. Williams JMG, Mathews A, MacLeod C. 1996 The Emotional Stroop Task and
554 Psychopathology. *Psychological Bulletin* **122**, 3–24. (doi:10.1037/0033-2909.120.1.3)
- 555 4. Henik A, Tzelgov J. 1982 Is three greater than five: The relation between physical and
556 semantic size in comparison tasks. *Memory & Cognition* **10**, 389–395.
557 (doi:10.3758/BF03202431)
- 558 5. Schlaghecken F, Eimer M. 2004 Masked prime stimuli can bias quot;freequot; choices
559 between response alternatives. *Psychonomic Bulletin and Review* **11**, 463–468.
560 (doi:10.3758/BF03196596)
- 561 6. Talmi D, Seymour B, Dayan P, Dolan RJ. 2008 Human pavlovian-instrumental transfer.
562 *Journal of Neuroscience* **28**, 360–368. (doi:10.1523/JNEUROSCI.4028-07.2008)
- 563 7. Rushworth MFS, Passingham RE, Nobre AC. 2002 Components of Switching
564 Intentional Set. *Journal of Cognitive Neuroscience* **14**, 1139–1150.
565 (doi:10.1162/089892902760807159)
- 566 8. Lages M, Boyle SC, Jaworska K. 2013 Flipping a coin in your head without monitoring
567 outcomes? Comments on predicting free choices and a demo program. *Frontiers in*
568 *Psychology* **4**, 925. (doi:10.3389/fpsyg.2013.00925)
- 569 9. Lages M, Jaworska K. 2012 How Predictable are “Spontaneous Decisions” and “Hidden
570 Intentions”? Comparing Classification Results Based on Previous Responses with
571 Multivariate Pattern Analysis of fMRI BOLD Signals. *Frontiers in Psychology* **3**, 56.
572 (doi:10.3389/fpsyg.2012.00056)
- 573 10. Goschke T. 2003 Voluntary action and cognitive control from a cognitive neuroscience
574 perspective. In *Voluntary action: Brains, minds, and sociality.*, pp. 49–85.
- 575 11. Botvinick M, Braver TS, Barch DDM, Carter CS, Cohen JD. 2001 Conflict monitoring
576 and cognitive control. *Psychological Review* **108**, 624–652. (doi:10.1037/0033-
577 295X.108.3.624)
- 578 12. Arrington CM, Logan GD. 2004 The cost of a voluntary task switch. *Psychological*
579 *Science* **15**, 610–615. (doi:10.1111/j.0956-7976.2004.00728.x)

- 580 13. Newell BR, Shanks DR. 2014 Unconscious influences on decision making: A critical
581 review. *Behavioral & Brain Sciences* **37**, 1–19. (doi:10.1017/S0140525X12003214)
- 582 14. Lavie N. 2010 Attention , Distraction , and Cognitive Control Under Load.
583 (doi:10.1177/0963721410370295)
- 584 15. Pronin E, Lin DY, Ross L. 2002 The bias blind spot: Perceptions of bias in self versus
585 others. *Personality and Social Psychology Bulletin* **28**, 369–381.
586 (doi:10.1177/0146167202286008)
- 587 16. Epstein R, Robertson RE. 2015 The search engine manipulation effect (SEME) and its
588 possible impact on the outcomes of elections. *Proceedings of the National Academy of
589 Sciences* **112**, E4512–E4521. (doi:10.1073/pnas.1419828112)
- 590 17. Hansen K, Gerbasi M, Todorov A, Kruse E, Pronin E. 2014 People Claim Objectivity
591 After Knowingly Using Biased Strategies. *Personality and Social Psychology Bulletin* **40**,
592 691–699. (doi:10.1177/0146167214523476)
- 593 18. Frith CD. 2005 The neural basis of hallucinations and delusions. *Comptes Rendus -
594 Biologies* **328**, 169–175. (doi:10.1016/j.crv.2004.10.012)
- 595 19. Charles L, Haggard P. 2019 Feeling free: External influences on endogenous behaviour.
596 *Quarterly Journal of Experimental Psychology* , 174702181988806.
597 (doi:10.1177/1747021819888066)
- 598 20. Filevich E, Vanneste P, Brass M, Fias W, Haggard P, Kühn S. 2013 Brain correlates of
599 subjective freedom of choice. *Consciousness and Cognition* **22**, 1271–1284.
600 (doi:10.1016/j.concog.2013.08.011)
- 601 21. Maren S. 2001 NEUROBIOLOGY OF PAVLOVIAN FEAR CONDITIONING. *Annual
602 Review of Neuroscience* **24**, 897–931. (doi:10.1146/annurev.psych.55.090902.141409)
- 603 22. Molina MT, Davis G. 2018 Unconscious priming dissociates ‘free choice’ from
604 ‘spontaneous urge’ responses. *Consciousness and Cognition* **60**, 72–85.
605 (doi:10.1016/j.concog.2018.02.003)
- 606 23. O’Connor PA, Neill WT. 2011 Does subliminal priming of free response choices depend
607 on task set or automatic response activation? *Consciousness and Cognition* **20**, 280–
608 287. (doi:10.1016/j.concog.2010.08.007)
- 609 24. Ocampo B. 2015 Unconscious manipulation of free choice by novel primes.
610 *Consciousness and Cognition* **34**, 4–9. (doi:10.1016/j.concog.2015.03.007)
- 611 25. Le Bars S, Hsu YF, Waszak F. 2016 The impact of subliminal effect images in voluntary
612 vs. stimulus-driven actions. *Cognition* **156**, 6–15. (doi:10.1016/j.cognition.2016.07.005)

- 613 26. Wenke D, Fleming SM, Haggard P. 2010 Subliminal priming of actions influences sense
614 of control over effects of action. *Cognition* **115**, 26–38.
615 (doi:10.1016/j.cognition.2009.10.016)
- 616 27. Schlaghecken F, Eimer M. 2004 Masked prime stimuli can bias choices
617 between response alternatives. *Psychonomic Bulletin and Review* **11**, 463–468.
618 (doi:10.3758/BF03196596)
- 619 28. Schurger A, Faivre N, Cammoun L, Trovó B, Blanke O. 2017 Entrainment of Voluntary
620 Movement to Undetected Auditory Regularities. *Scientific Reports* **7**:1 **7**, 1–7.
621 (doi:10.1038/s41598-017-15126-w)
- 622 29. Bodner GE, Mulji R. 2010 Prime Proportion Affects Masked Priming of Fixed and Free-
623 Choice Responses. *Experimental Psychology* **57**, 360–366. (doi:10.1027/1618-
624 3169/a000043)
- 625 30. Fröber K, Dreisbach G. 2017 Keep flexible – Keep switching! The influence of forced
626 task switching on voluntary task switching. *Cognition* **162**, 48–53.
627 (doi:10.1016/j.cognition.2017.01.024)
- 628 31. Francken JC, Gaal S van, de Lange FP. 2011 Immediate and long-term priming effects
629 are independent of prime awareness. *Consciousness and Cognition* **20**, 1793–1800.
630 (doi:10.1016/j.concog.2011.04.005)
- 631 32. Kunde W, Reuss H, Kiesel A. 2012 Consciousness and cognitive control. *Advances in*
632 *Cognitive Psychology*. **8**, 9–18. (doi:10.2478/v10053-008-0097-x)
- 633 33. Desender K, Van Lierde E, Van den Bussche E. 2013 Comparing Conscious and
634 Unconscious Conflict Adaptation. *PLoS ONE* **8**, e55976.
635 (doi:10.1371/journal.pone.0055976)
- 636 34. van Gaal S, de Lange FP, Cohen MX. 2012 The role of consciousness in cognitive
637 control and decision making. *Frontiers in Human Neuroscience*. **6**, 121.
638 (doi:10.3389/fnhum.2012.00121)
- 639 35. Maren S. 2001 Neurobiology of Pavlovian fear conditioning. *Annual Review of*
640 *Neuroscience*. **24**, 897–931. (doi:10.1146/annurev.neuro.24.1.897)
- 641 36. Haggard P. 2019 The Neurocognitive Bases of Human Volition. , 1–20.
- 642 37. Imhoff R, Erb H-P. 2009 What motivates nonconformity? Uniqueness seeking blocks
643 majority influence. *Pers Soc Psychol Bull* **35**, 309–20. (doi:10.1177/0146167208328166)
- 644 38. Scott-Parker B, King MJ, Watson B. 2015 The psychosocial purpose of driving and its
645 relationship with the risky driving behaviour of young novice drivers. *Transportation*
646 *Research Part F: Traffic Psychology and Behaviour* **33**, 16–26.
647 (doi:10.1016/j.trf.2015.06.004)

- 648 39. Steindl C, Jonas E, Sittenthaler S, Traut-Mattausch E, Greenberg J. 2015
649 Understanding psychological reactance: New developments and findings. *Zeitschrift fur*
650 *Psychologie / Journal of Psychology* **223**, 205–214. (doi:10.1027/2151-2604/a000222)
- 651 40. Miron AM, Brehm JW. 2006 Reactance Theory - 40 Years Later. *Zeitschrift fur*
652 *Sozialpsychologie* **37**, 9–18. (doi:10.1024/0044-3514.37.1.9)
- 653 41. Kirsch I, Lynn SJ, Vigorito M, Miller RR. 2004 The Role of Cognition in Classical and
654 Operant Conditioning. *Journal of Clinical Psychology* **60**, 369–392.
655 (doi:10.1002/jclp.10251)
- 656 42. Roitman JD, Shadlen MN. 2002 Response of neurons in the lateral intraparietal area
657 during a combined visual discrimination reaction time task. *The Journal of Neuroscience*
658 **22**, 9475–9489. (doi:10.1016/S0377-2217(02)00363-6)
- 659 43. Kiani R, Hanks TD, Shadlen MN. 2008 Bounded Integration in Parietal Cortex Underlies
660 Decisions Even When Viewing Duration Is Dictated by the Environment. *Journal of*
661 *Neuroscience* **28**, 3017–3029. (doi:10.1523/JNEUROSCI.4761-07.2008)
- 662 44. Adelson EH, Bergen JR. 1985 Spatiotemporal energy models for the perception of
663 motion. *Journal of the Optical Society of America A* **2**, 284.
664 (doi:10.1364/JOSAA.2.000284)
- 665
- 666

667 **ACKNOWLEDGEMENT**

668 We thank Ian Yeung for his help with the testing of participants.

669 **FUNDING**

670 This work was supported by a British Academy Post-doctoral Fellowship to LC.

671

672 **AUTHORS CONTRIBUTIONS**

673 A.K. contributed to designing the experiment, testing participants, analyzing data and writing
674 the manuscript. P.H. and GW contributed to designing the experiment, interpreting data and
675 writing the manuscript. L.C. contributed to designing the experiment, testing participants,
676 analyzing data and writing the manuscript.

677

678 **COMPETING INTERESTS STATEMENT**

679 The authors declare there is no competing interests.

680

681 Data is available at :

682 [https://liveuclac-
683 my.sharepoint.com/:f/g/personal/ucjulch_ucl_ac_uk/EtMvbGTX8tFEqhgAtAPR-
684 BcBx_vHiNPfuMsYr_DZoVB_Dg?e=XecnFr](https://liveuclac-my.sharepoint.com/:f/g/personal/ucjulch_ucl_ac_uk/EtMvbGTX8tFEqhgAtAPR-BcBx_vHiNPfuMsYr_DZoVB_Dg?e=XecnFr)

685 Password: FoCContext

686

688 5 SUPPLEMENTARY MATERIALS

689 5.1 SUPPLEMENTARY METHOD

690 5.1.1 Stimuli

691 The dots were shown for a single frame before being replotted three frames later. When
692 replotted, a subset of the dots were offset from their previous location to produce apparent
693 coherent motion in one direction (to the right or to the left), whilst others were offset and
694 plotted randomly (see [42]).

695 Each block of trials also included three high coherence trials (coherence set to 65%, one for
696 each instruction type) to ensure that participants could follow instructions accurately in Adhere,
697 Oppose, and Detach trials according to the colour change.

698 5.1.2 Staircase Procedure

699 To account for individual differences in sensitivity to dot motion, the motion coherence of the
700 RDK stimuli was predetermined using a 2-down-1-up staircase procedure (Levitt, 1971). To
701 do so, participants started the experiment with a block of 150 trials in which they were
702 instructed to disregard all colour changes and to place their focus on detecting the direction of
703 dot movement. The coherence level was adjusted starting at a motion coherence of 40% and
704 after 12 initial trials, the motion coherence changed by a set amount in such a way that two
705 consecutive correct guesses lowered the coherence, and a single error increased the
706 coherence. The step size started at 6% and halved every time the participant made two
707 consecutive errors (until the minimum step-size of 1.5%). The estimate of the appropriate
708 motion coherence was obtained by averaging the coherence over the last 40 trials. This
709 coherence level was used in the rest of the experiment.

710 5.1.3 Block Design

711 **Experiment 1.** The combination of orientation (left and right) and instruction type (Adhere,
712 Oppose, Detach) generated 6 possible outcomes (i.e. orientation left to adhere; orientation
713 right to adhere, etc) as an experimental trial in each group. Due to the between-subjects design,
714 each group was assigned more Adhere trials than Oppose trials, or more Oppose trials than
715 Adhere trials, whilst the proportion of Detach trials remained the same. Hence, Adhere Group
716 had 50% Adhere trials and 17% Oppose trials (33% Detach trials), whilst Oppose Group had
717 17% Adhere trials, 50% Oppose trials (and 33% Detach trials). Each block consisted of 54
718 trials consisting of the proportions mentioned above. This left a total of 378 trials of interest
719 over 7 blocks. Additionally, each block of trials included 3 high coherence trials.

720 **Experiment 2.** The mixed experimental design was so that each group was assigned to either
721 group majority left responses, or group majority right responses. Hence, the combination of
722 orientation (left and right) and instruction type (Adhere, Oppose, Detach) was altered so that
723 there was more Adhere for one direction, and more Oppose for the other direction. Using group
724 left as an example, the Adhere trials (30% of all trials) consisted of 75% left-direction stimuli,
725 and 25% right-direction stimuli. On Oppose trials (30% of all trials), 75% was right-direction

726 stimuli and 25% left-direction stimuli. The percentage of Detach trials was 40%, of which 75%
727 were motion-present at 25% motion-neutral. For each group, there was a total of 67 randomly
728 arranged experimental trials of interest per block (and a total of 469 trials of interest over 7
729 blocks) following the proportions mentioned above. Additionally, each block of trials included
730 3 high coherence trials.

731 **Experiment 3.** Again, the mixed experimental design was so that participants were assigned
732 to see either left-ward stimulus frequently, or right-ward. Hence, the combination of orientation
733 (left, right) and instruction type (Adhere, Oppose, Detach) was so that there was more of one
734 direction than the other across all three instruction types. Overall, 75% of all trials was one
735 direction (e.g. leftward for group majority left), and 25% the other (e.g. rightward for group
736 majority right. 30% of trials were Adhere, 30% were Oppose, 27% Detach (two thirds motion-
737 present at one third motion-neutral). For each group, there was a total of 60 randomly arranged
738 experimental trials of interest per block (and a total of 420 trials of interest over 7 blocks)
739 following the proportions mentioned above. Additionally, each block of trials included 3 high
740 coherence trials.

741 **5.1.4 Reverse correlation analysis on motion-neutral detach trials**

742 On motion-neutral detach trials, small fluctuations in motion energy could still momentarily
743 favour one direction over the other, even though overall motion energy was balanced across
744 directions. The time-course of the motion energy for each trial was retrieved by applying
745 spatiotemporal motion filters to dot position over time so to compute a time-course of motion
746 direction for each trial [19,43,44]. The net motion energy value was calculated by subtracting
747 the amount of rightward motion from leftward motion on each timeframe of stimulus
748 presentation. These values were then standardized against the mean motion energy and
749 standard deviation of all other motion-neutral detach trials to give a zero mean and unit of
750 standard deviation. Thus, motion energy values now reflected fluctuations in motion direction
751 around the mean coherence level. We then reverse correlated the normalized values so that
752 positive values related to fluctuations favouring the frequently adhered direction (Experiment
753 2) or the frequent direction (Experiment 3), whilst negative values reflected fluctuations
754 favouring the frequently opposed direction (Experiment 2) or the rare direction (Experiment 3).
755 The sum across time (area under curve, AUC) was computed. We then divided trials according
756 to whether responses were classified by whether they were congruent or incongruent to the
757 direction of this fluctuation.

758 **5.2 SUPPLEMENTARY RESULTS**

759 **5.2.1 Bias awareness**

760 **Experiment 2.** All participants were asked to estimate the proportion of right responses
761 across all trials of the experiment on a scale from 0 (only left responses) to 100 (only right
762 responses). An estimate of 50 would reflect 50/50 left and right responses. A one-sample t-
763 test was performed for each group, comparing the estimated perceptual bias to chance level
764 (50%). Both descriptive statistics and t-test comparisons are reported in Table S1. The null
765 effect for both groups meant that participants did not feel they responded more of one button
766 than the other, and hence were not aware of our response bias manipulation.

Table S1

Bias Awareness: Means and Standard Deviation for Estimated Proportion of Right Responses (by group), and T-test Comparison to Chance Level (50%)

Group	M	SD	t	df	p	d
Group Majority Left Responses	48.80%	17.21%	0.31	19	.760	0.07
Group Majority Right Responses	52.33%	19.39%	0.51	17	.620	0.12

767

768 **Experiment 3.** All participants were asked to estimate the proportion of right-direction
 769 trials on a scale from 0 (all left-direction trials) to 100 (all right-direction trials). A one-sample t-
 770 test was performed for each group, comparing the estimated perceptual bias to chance level
 771 (50%). Both descriptive statistics and t-test comparisons are reported in Table S2. The null
 772 effect for group majority leftward stimuli suggest that participants did not feel they saw either
 773 direction more than the other, and hence were not aware of our stimulus frequency
 774 manipulation. However, group majority rightward stimuli reported significantly more right-
 775 direction trials than left-direction trials, suggesting an awareness of the stimulus frequency
 776 manipulation

Table S2

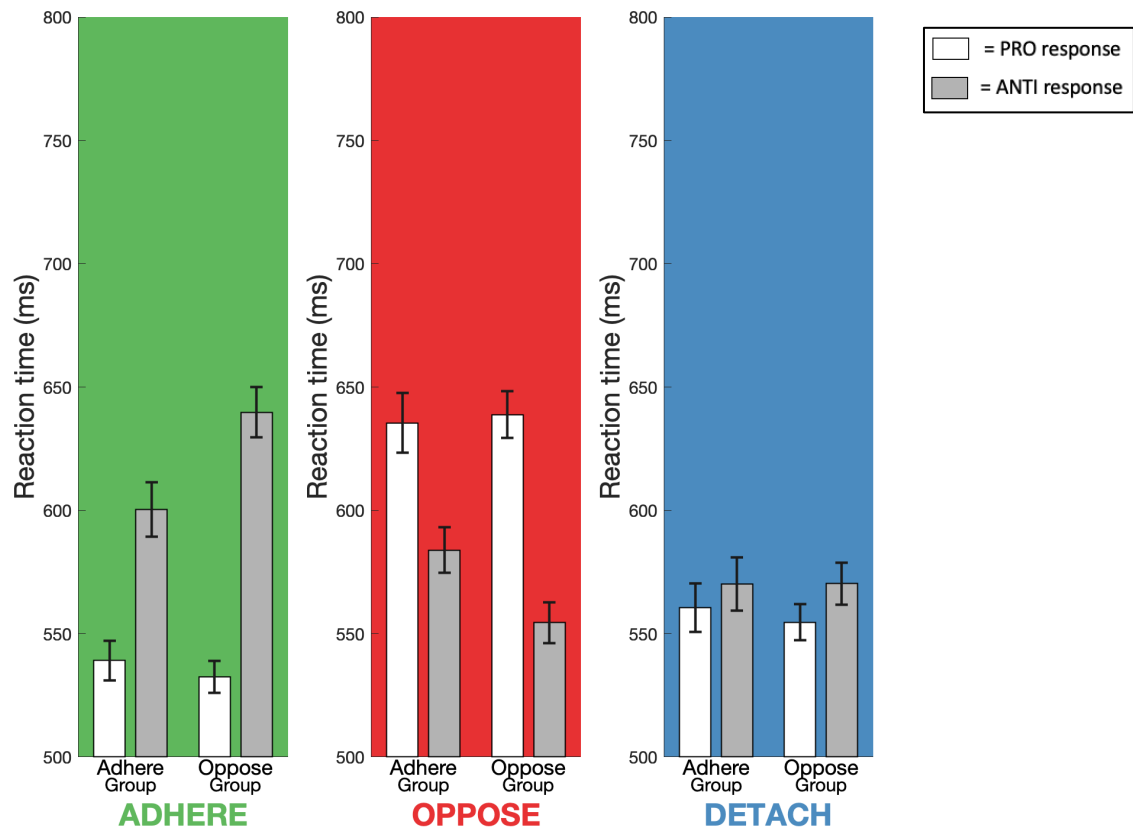
Bias Awareness: Means and Standard Deviation for Estimated Proportion of Right-direction Trials (by group), and T-test Comparison to Chance Level (50%)

Group	M	SD	t	df	p	d
Group Majority Leftward Stimuli	48.70%	19.99%	0.291	19	.770	0.07
Group Majority Rightward Stimuli	59.29%	13.95%	2.75	16	.014	0.67

777

778 5.2.2 Reaction Time on Detach Trials

779 **Experiment 1.** There was no main effect of response mode for reaction time on detach
 780 trials responses [$F(1, 58) = 2.21, p = .142, \eta_p^2 = .000$], reflecting that a congruent response was
 781 no easier made than an incongruent response. There was also no main effect of group [$F(1,$
 782 $58) = 0.01, p = .906, \eta_p^2 = .000$], or an interaction between group and response mode [$F(1, 58)$
 783 $= 0.12, p = .730$], suggesting that being trained for one strategy (Adhere Group vs Oppose
 784 Group) does not influence the ease at which participants go with or against that strategy on
 785 detach trials as measured in reaction time.



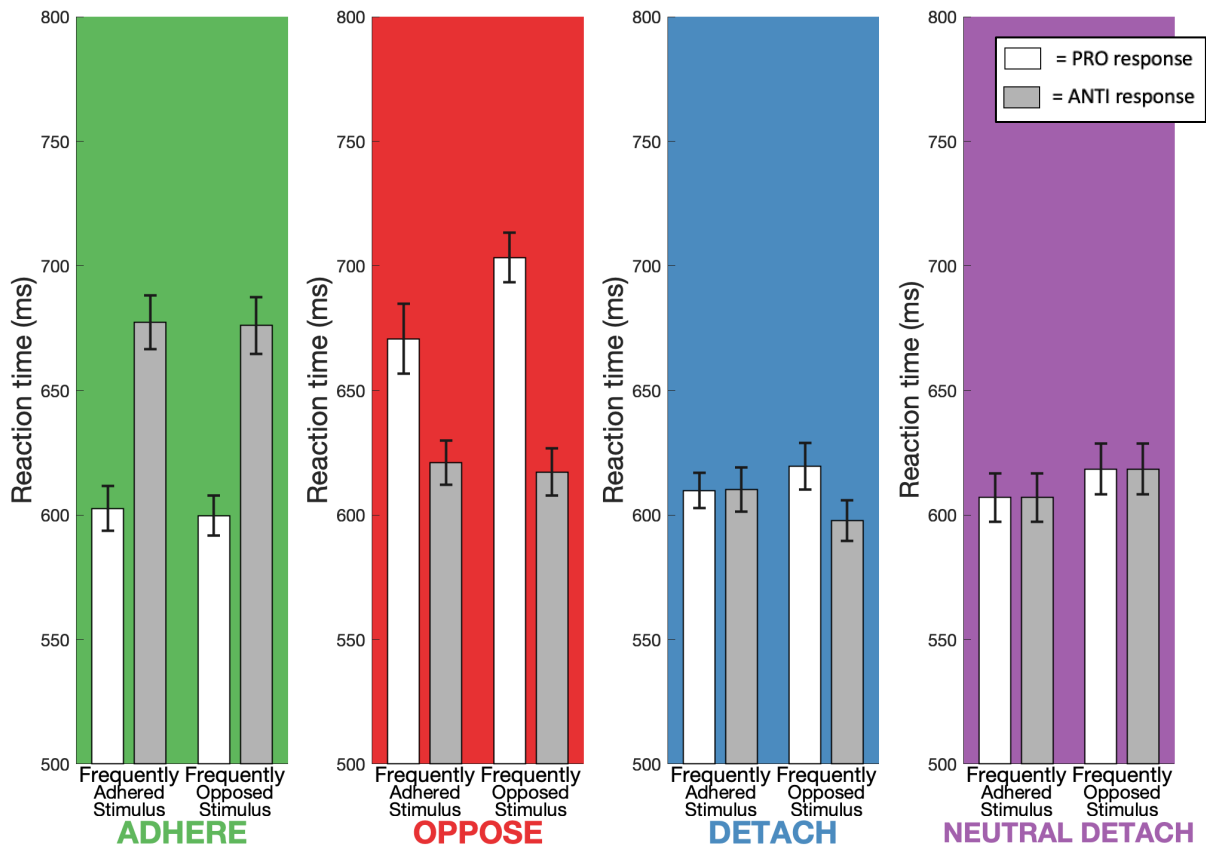
786

787 **Figure S1.** Mean reaction time in milliseconds on Experiment 1 split by instruction (Adhere,
 788 Oppose, Detach), group (Adhere Group, Oppose Group), and response mode (congruent
 789 responses, white bars; incongruent responses, grey bars).

790

791 **Experiment 2.** On motion-present detach trials, there was no difference in the reaction time
 792 between a congruent and an incongruent response [main effect response mode; $F(1, 36) =$
 793 $1.70, p = .201, \eta_p^2 = .045$]. There was also no main effect of stimulus-response association [$F(1,$
 794 $36) = 0.04, p = .846, \eta_p^2 = .001$], or an interaction between stimulus-response association and
 795 response mode [$F(1, 36) = 1.28, p = .266, \eta_p^2 = .034$]. This suggests that being trained for one
 796 strategy at one particular stimulus does not influence the ease at which participants go with or
 797 against that strategy on detach trials as measured in reaction time.

798 The results on motion-neutral detach trials were similar to that of motion-present detach trials,
 799 with no main effect of response mode [$F(1, 36) = 0.73, p = .398, \eta_p^2 = .019$] stimulus-response
 800 association [$F(1, 36) = 0.23, p = .634, \eta_p^2 = .006$], or interaction between response mode and
 801 stimulus-response association [$F(1, 36) = 2.50, p = .122, \eta_p^2 = .063$]. This suggests that
 802 regardless of fluctuations in motion energy favouring one direction over the other, or whether
 803 a motor response is one made frequently or rarely over past trials, there is no difference in the
 804 ease at which participants make such a response on detach trials.



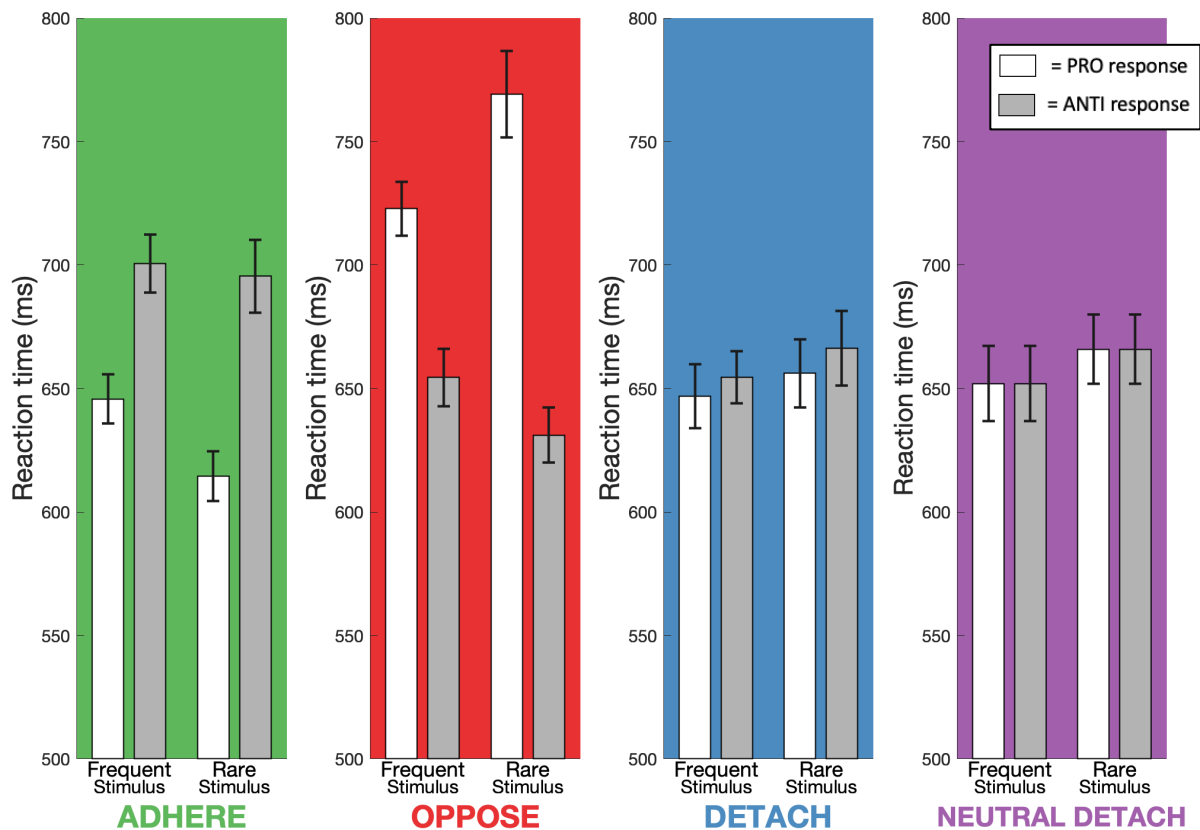
805

806 **Figure S2.** Mean reaction time in milliseconds on Experiment 2 split by instruction (Adhere,
 807 Oppose, Detach), stimulus-response association (frequently adhered, frequently opposed),
 808 and response mode (congruent responses, white bars; incongruent responses, grey bars).

809

810 **Experiment 3.** Again, there was no difference in the reaction time on detach trials between a
 811 congruent and an incongruent response [main effect response mode; $F(1, 35) = 0.39, p = .538,$
 812 $\eta_p^2 = .011$]. There was also no main effect of stimulus frequency [$F(1, 35) = 0.81, p = .373, \eta_p^2$
 813 $= .023$]. This suggests that seeing one stimulus direction more often than the other does not
 814 influence the ease at which one can detach from that stimulus as reflected in reaction times.
 815 There was, however, a 3-way interaction with group, response mode, and stimulus frequency,
 816 which is reported and discussed in Supplementary Results.

817 Motion-neutral detach trials also had no difference in reaction time between response modes
 818 [$F(1, 35) = 0.07, p = .788, \eta_p^2 = .002$] or stimulus frequency [$F(1, 36) = 0.16, p = .691, \eta_p^2 = .005$].
 819 Hence, all detach trials suggested that the frequency at which a cue is presented does not
 820 influence the ability to detach from that cue as measured in reaction time.



821
822 **Figure S3.** Mean reaction time in milliseconds split by instruction (Adhere, Oppose, Detach),
823 stimulus frequency (frequent, rare), and response mode (congruent responses, white bars;
824 incongruent responses, grey bars).

825

826 5.3 GROUP DIFFERENCES ON EXPERIMENT 2 AND 3

827 **Experiment 2.** The same was true for experiment 2, where participants were assigned to press
828 either more of the right button than the left, or more of the left button than the right. Whether
829 making more of a right or left button press influenced the ability to detach or subjective rating
830 of detachment was not of interest to our hypotheses. We did however observe group-
831 dependent findings for accuracy on instructed trials [interaction group and stimulus-response
832 association: $F(1, 36) = 7.91, p = .008$], as well as proportion of motion congruent responses
833 on detach trials [interaction group and motion coherence: $F(1, 36) = 4.16, p = .049$].

834 The group differences observed were perhaps due to a population-based preference for
835 making one motor response over the other, which interacted with our response frequency
836 manipulations.

837 **Experiment 3.** For experiment 3, participants were assigned to either see mostly leftward
838 stimuli, or mostly rightward stimuli. We were interested in the ability to detach and subjective

839 introspection of detachment from the stimulus, and how it differed between a frequent and a
 840 rare stimulus. Group differences in our statistical analyses, reflecting that seeing a lot of right-
 841 ward motion compared to left-ward motion influenced subjective and objective FoC, were not
 842 of interest in regard to our hypotheses. However, we observed group difference in on instructed
 843 trials for subjective rating of freedom [interaction group, instruction, and stimulus frequency:
 844 $F(1, 36) = 5.03, p = .031$], as well as accuracy [interaction group and stimulus frequency: $F(1,$
 845 $36) = 7.29, p = .011$].

846 The group differences observed were likely due to increased or decreased sensitivity to one
 847 motion over the other. Previous research has indicated that subjects that have English or
 848 another left-to right reading language have a higher sensitivity for leftward motion (Morikawa
 849 & McBeath, 1991), but more research is needed to determine the cause of our group effects.

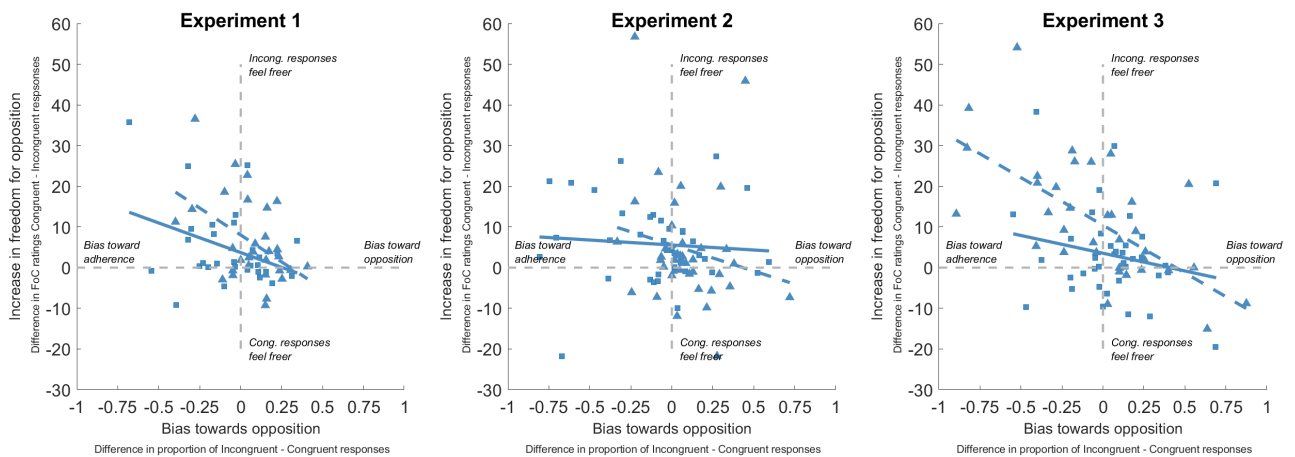
850 **5.4 REGRESSION**

851 **Table S3**

852 Regressing difference in subjective freedom of choice from incongruent to pro responses with
 853 motion consistency and difference in reaction time (RT) as predictors. Datapoints are
 854 combined across experiments.

	b	t	df	p
Adhere Group, frequently adhered stimuli, rare stimuli combined				
Intercept	6.57	6.17	102	.000
Motion Consistency*	-9.87	2.73	102	.008
RT difference pro	0.02	2.25	102	.026
Oppose Group, frequently opposed stimuli, frequent stimuli combined				
Intercept	5.41	4.28	102	.000
Motion Consistency*	-13.36	2.54	102	.130
RT difference incong-cong	-0.01	0.81	102	.420

855 *Note.* Motion consistency measured as proportion of incongruent responses minus the
 856 proportion of congruent responses on detach trials.



857

858

859 **Figure S4.** Difference in subjective freedom of choice (FoC) to incongruent minus congruent
 860 responses on motion-present detach trials correlated with difference in proportion of
 861 incongruent minus congruent responses on motion-present detach trials. This is split by each
 862 experiment, and the experiment-specific main manipulation (Adhere Group vs Oppose Group
 863 on Experiment 1; frequently adhered vs frequently opposed stimuli on Experiment 2; frequent
 864 vs rare stimuli on Experiment 3)

865

866 5.5 SUPPLEMENTARY REFERENCES

867 Morikawa, K., & McBeath, M. K. (1992). Lateral motion bias associated with reading direction.
 868 *Vision Research*, 32(6), 1137–1141. [https://doi.org/10.1016/0042-6989\(92\)90014-A](https://doi.org/10.1016/0042-6989(92)90014-A)

869 Levitt, H. (1971). Transformed Up-Down Methods in Psychoacoustics. *The Journal of the*
 870 *Acoustical Society of America*, 49(2B), 467–477. <https://doi.org/10.1121/1.1912375>