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1 In press at *Journal of Experimental Psychology: General*

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3 **Moving time: The influence of action on duration perception**

4

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23

24 **Abstract**

25 Perceiving the sensory consequences of action accurately is essential for appropriate  
26 interaction with our physical and social environments. Prediction mechanisms are  
27 considered necessary for fine-tuned sensory control of action, yet paradoxically may  
28 distort perception. Here we examine this paradox by addressing how movement  
29 influences the perceived duration of sensory outcomes congruent with action.  
30 Experiment 1 required participants to make judgments about the duration of vibrations  
31 applied to a moving or stationary finger. In Experiments 2 and 3, participants judged  
32 observed finger movements, congruent or incongruent with their own actions. In all  
33 experiments, target events were perceived as longer when congruent with movement.  
34 Interestingly, this temporal dilation did not differ as a function of stimulus perspective  
35 (first or third person) or spatial location. We propose that this bias may reflect the  
36 operation of an adaptive mechanism for sensorimotor selection and control that pre-  
37 activates anticipated outcomes of action. The bias itself may have surprising  
38 implications both for action control and perception of others - we may be in contact  
39 with grasped objects for less time than we realize and others' reactions to us may be  
40 briefer than we believe.

41

42 Keywords: Motor Processes, Perceptual Motor Coordination, Time Perception, Social  
43 Perception

44

## 45 **Introduction**

46 To interact appropriately with physical and social environments, actors must predict  
47 and evaluate the sensory consequences of their actions. We select actions based on their  
48 predicted outcomes (Greenwald, 1970; Hommel, Müsseler, Aschersleben, & Prinz,  
49 2001), and when the experienced sensory information deviates from our prediction,  
50 corrective adjustments can be made ensuring successful execution. For example, when  
51 picking up a cup of tea, the motor commands generated result in both visual (e.g., sight  
52 of grasping and lifting) and tactile (e.g., pressure on the fingertips) sensory  
53 consequences. If the actual feedback differs from the anticipated sensory outcomes,  
54 rapid corrective actions can be executed to avoid spillage. Similarly, when interacting  
55 with others, rapid response prediction and error correction may facilitate smooth social  
56 interactions (Wolpert, Doya, & Kawato, 2003).

57

58 Successful interaction with the environment requires perception not only of the nature  
59 of our action outcomes (e.g., somatosensation on the fingertips during grasping), but  
60 also crucially, the onset and duration of those outcomes. We are sensitive both to the  
61 ‘what’ and ‘when’ of sensory predictions (Blakemore, Frith, & Wolpert, 1999;  
62 Blakemore, Wolpert, & Frith, 1998; Christensen, Ilg, & Giese, 2011; Fagioli, Hommel, &  
63 Schubotz, 2007; Hommel, 2010; Lee, Young, Reddish, Lough, & Clayton, 1983; Schubotz,  
64 2007). For example, lifting the teacup from a saucer requires an anticipatory response  
65 to maintain postural stability (Diedrichsen, Verstynen, Hon, Lehman, & Ivry, 2003;  
66 Dufossé, Hugon, & Massion, 1985). Similarly, anticipating the duration of the lift phase is  
67 essential for coordinating hand and mouth gestures, and when shaking someone’s hand,  
68 contact must be made for an appropriate length of time, neither too long, nor too short,  
69 to convey the intended social message.

70

71 While the preceding examples underscore the importance of temporal information in  
72 the generation and perception of sensory expectancies, duration perception is  
73 frequently distorted. For example, perceived motion of upright point light walkers is  
74 temporally dilated relative to inverted walkers (Wang & Jiang, 2012; see also Gavazzi,  
75 Bisio, & Pozzo, 2013). The present experiments examine how movement influences the  
76 perceived duration of sensory outcomes of action. Sensory prediction mechanisms  
77 essential for action selection and fine-tuned control may, paradoxically, distort the

78 perceived duration of outcomes, with consequences for action-control and perception in  
79 a variety of contexts. In Experiment 1, participants were required to perform a lifting  
80 movement with either their index or middle finger. A short target vibratory tactile  
81 stimulus was presented to the moving or stationary finger, followed by a second  
82 reference vibration. Participants judged which was of longer duration, allowing us to  
83 determine how action influences duration perception. Experiments 2-4 asked whether  
84 action influences the perceived duration of visual events in a comparable manner.

85

## 86 **Experiment 1**

87 Sixteen right-handed, healthy adults (12 male) with a mean age of 29.3 years (SEM =  
88 2.5) participated in the experiment. Three were replacements for participants who  
89 could not perform the perceptual discrimination (psychometric functions could not be  
90 modeled or their point of subjective equivalence (PSE) fell outside the range of  
91 presented stimuli). All experiments were undertaken in accordance with the 1964  
92 Declaration of Helsinki.

93

94 The experiment was conducted in MATLAB using the Cogent toolbox<sup>i</sup>. Two 5V solenoids,  
95 each driving a metal rod with a blunt conical tip (diameter  $\approx$  1.5 mm, skin indentation  $\approx$   
96 1 mm), were attached to the distal phalange (ventral surface) of the index and middle  
97 fingers on the right hand. Participants held down two keys on the keyboard until an  
98 imperative cue instructed them to lift either their index ('I') or middle ('M') finger. They  
99 were instructed to make large, rapid, single-movement lifts. Their response hand was  
100 visually occluded. Approximately 10 ms after the cued finger was lifted, a target  
101 vibration lasting for one of seven durations (104 – 296 ms, 32 ms steps) was applied to  
102 the moving ('congruent') or stationary ('incongruent') finger (see Fig. 1). After an inter-  
103 stimulus interval (ISI; 300 – 500 ms), a 200 ms reference vibration was applied to the  
104 same finger. Both vibratory stimuli were presented at 62.5 Hz.

105

106 Participants judged whether the target or reference vibration was longer, responding  
107 with a button press made with their left hand. Following this response, they returned  
108 the lifted finger to the start position. The next trial started after 2000 ms. There were  
109 280 trials; 140 in which stimuli were applied to the congruent finger and 140 where

110 they were applied to the incongruent finger. Trial type was randomized and  
111 participants completed 8 practice trials.

112

113 To estimate psychometric functions, the responses for each individual were modeled by  
114 fitting cumulative Gaussians, and associated pDev statistics were calculated to establish  
115 the goodness-of-fit of each function (Palamedes toolbox, Kingdom & Prins, 2010). This  
116 procedure was performed separately for congruent and incongruent response data. In  
117 each condition, bias was inferred from the PSE and precision from the difference  
118 threshold (Fig. 2).

119

120 The participants were more precise in their judgments when the vibratory stimuli were  
121 applied to the congruent relative to incongruent finger ( $t(15) = 2.3, p < 0.05, \eta^2 = 0.26$ ;  
122 Table 1). There was also an effect on PSE: Target events were judged longer when the  
123 stimulus was applied to the congruent relative to incongruent finger ( $t(15) = 2.6, p <$   
124  $0.02, \eta^2 = 0.32$ ; Fig. 2 & Supplementary Fig. 1). In sum, tactile events presented to a  
125 moving effector are perceived to be longer and are judged more consistently than when  
126 that effector is stationary.

127

## 128 **Experiment 2**

129 If prediction mechanisms operate in social contexts, we may predict and evaluate  
130 sensation not only related to our own actions, but also actions produced by interactants  
131 (Wolpert et al., 2003). As such, we should observe comparable action-related predictive  
132 modulation with visual action stimulus events. Additionally, such mechanisms should  
133 operate across perspectives given the range of viewpoints from which others' actions  
134 are observed. Experiment 2 therefore examined duration perception of congruent and  
135 incongruent visual events during action, across stimuli presented from first and third  
136 person perspectives<sup>ii</sup>.

137

138 Sixteen right-handed, healthy adults (12 male) with a mean age of 25.9 years (SEM =  
139 1.9) participated in the experiment. Five were replacements for participants who could  
140 not perform the discrimination. Unless otherwise stated, procedural information  
141 already outlined in Experiment 1 is identical in this, and all subsequent, experiments.  
142 Participants compared the duration of two finger movements simulated visually by

143 gestures of an avatar hand. At the start of the trial, the avatar hand was presented in a  
144 neutral position on the monitor (Fig. 1; screen refresh rate = 85 Hz). An imperative cue  
145 ('1' or '2') was presented between the index and middle fingers. When participants  
146 lifted the cued finger, the neutral hand image was immediately replaced (within the  
147 constraints of the refresh rate) by one depicting the avatar hand performing either an  
148 index or middle finger lift for 120 - 480 ms (7 levels; 60 ms steps). This resulted in  
149 apparent motion of the avatar's finger approximately synchronized with the  
150 participant's action. At the offset of the target event, the avatar hand resumed the  
151 neutral position for an ISI of 300 - 500 ms, followed by a second image of the same lifted  
152 finger for a reference duration of 300 ms, and then the neutral image again (300 ms).  
153 Participants judged which lift lasted longer. The range of durations was chosen to match  
154 discrimination performance in Experiment 1.

155

156 There were four block types. In spatially aligned first person perspective (1PP) blocks,  
157 participants viewed a right avatar hand with fingers aligned in the horizontal plane with  
158 their own right hand (Fig. 1). In spatially aligned third person perspective (3PP) blocks,  
159 the avatar hand was rotated about the horizontal axis (therefore presenting a left hand).  
160 The remaining blocks consisted of these stimuli flipped on a vertical axis, such that  
161 corresponding finger movements did not match in spatial location (necessitating left  
162 hand for 1PP and right hand for 3PP). These blocks thereby controlled for the spatial  
163 location of finger movement (Press, Gherri, Heyes, & Eimer, 2010). The four blocks each  
164 comprised 140 trials and were completed in a counterbalanced order.

165

166 The precision and PSE distributions were analyzed using separate three-way ANOVAs,  
167 with factors of movement congruency (avatar and participant moved the  
168 congruent/incongruent finger), location (avatar and participant finger movements  
169 made in aligned/misaligned horizontal locations), and perspective (1PP or 3PP). No  
170 precision effects were observed (all  $F$ s < 2.1, all  $p$ s > 0.17; Table 1). However, as in  
171 Experiment 1, target events were perceived as longer when the avatar and participant  
172 moved the same finger ( $F(1,15) = 5.3, p < 0.04, \eta^2 = 0.26$ ). There were no other main  
173 effects or interactions (all  $F$ s < 2.5, all  $p$ s > 0.14). These results indicate a bias to judge  
174 target events as longer when observed actions are congruent with self-generated  
175 actions, regardless of whether stimuli are observed from first or third person



176 perspectives. Notably these effects reflect congruency between effectors (same finger)  
177 rather than spatial location.

178

### 179 **Experiment 3**

180 Experiment 2 suggests that action performance influences the perceived duration of  
181 effector-congruent visual events. However, it is possible that, despite informing  
182 participants that the reference event was always presented for the same length of time,  
183 participants' actions might have influenced the perceived duration of the reference  
184 rather than the target event. To control for this possibility, the reference event was  
185 modified in Experiment 3. Rather than define the reference duration by a second avatar  
186 movement, this interval was defined by the stimulus duration of a rectangle, a neutral  
187 stimulus selected because it exhibited no congruency relationship with the fingers.

188

189 Sixteen right-handed, healthy adults (11 male) with a mean age of 28.3 years (SEM =  
190 2.2) participated in the experiment. Three were replacements for participants who  
191 could not perform the discrimination. The imperative cue ('I' or 'M') was presented  
192 between the index and middle fingers of the observed hand. When participants lifted  
193 the cued finger, the neutral hand image was immediately replaced by an image of a hand  
194 with a lifted index or middle finger for 150 – 900 ms (7 levels; 125 ms steps). Following  
195 an ISI of 300 - 500 ms, a rectangle was presented for a reference interval of 525 ms. The  
196 color, luminance, and aspect ratio of the rectangle were identical to that of the avatar  
197 hand. The test stimulus range was selected based on piloting to yield comparable  
198 performance to that observed in Experiments 1 and 2. Participants again judged which  
199 of the two intervals was longer. Given that spatial location had no impact on the effect in  
200 Experiment 2, only aligned blocks were included. Participants completed 1PP and 3PP  
201 blocks, each comprising 140 trials, in a counterbalanced order.

202

203 The precision analysis yielded no main effects or interactions (all  $F_s < 1.4$ , all  $p_s > 0.25$ ;  
204 see Table 1). However, the PSE phenomenon observed in Experiments 1 and 2 was  
205 replicated: Target events were perceived as longer when the observed event was  
206 congruent with the participant's action ( $F(1,15) = 6.5, p < 0.03, \eta^2 = 0.30$ ; see Fig. 2). As  
207 in Experiment 2, this effect did not interact with perspective ( $F(1,15) = 0.05, p = 0.8, \eta^2$

208 = 0.02). These findings demonstrate that action biases perception of the temporally  
209 contiguous target event, rather than reference events presented after a delay.

210

#### 211 **Experiment 4**

212 It is possible that the temporal dilation effects in Experiments 2 and 3 result from  
213 attentional orienting towards the location of the congruent effector. Increased attention  
214 may influence the perceived duration of events at this location irrespective of action-  
215 stimulus congruency. A final experiment was conducted to test this possibility, identical  
216 to Experiment 3, except that target durations were defined by the presentation of a  
217 rectangle over the fingertip of the index or middle finger rather than by a finger  
218 movement (see Fig. 1). If attentional orienting generates temporal dilation effects  
219 irrespective of the nature of the target event, similar influences of congruency will be  
220 observed.

221

222 Sixteen right-handed, healthy adults (7 male) with a mean age of 27.7 years (SEM = 1.7)  
223 participated in the experiment. One was a replacement for a participant who could not  
224 perform the discrimination. The precision analysis yielded no main effects or  
225 interactions (all  $F$ s < 0.7, all  $p$ s > 0.41; see Table 1). Unlike Experiments 1-3, there was  
226 also no PSE effect ( $F(1,15) = 0.7, p = 0.42$ , see Fig. 2). ANOVA conducted on the PSE data  
227 from both Experiments 3 and 4, with experiment as a between-subjects factor, revealed  
228 a congruency  $\times$  experiment interaction ( $F(1,30) = 6.8, p < 0.02, \eta^2 = 0.2$ ). These results  
229 argue against this attentional orienting account of the congruency-induced temporal  
230 dilation.

231

#### 232 **Discussion**

233 The present findings demonstrate a bias to judge sensory events as longer when  
234 congruent with a concurrently performed action. This effect was found when  
235 participants judged the duration of tactile vibrations applied to a moving finger, as well  
236 as when assessing the duration of observed finger movements<sup>iii</sup>. These results indicate  
237 that subjective action-time can be subject to temporal dilation: Events effector-  
238 congruent with performed actions are perceived as longer than events incongruent with  
239 those actions.

240

241 These effects may be a consequence of pre-activated action expectancies during  
242 selection and preparation (Greenwald, 1970; Hommel et al., 2001), whereby congruent  
243 sensory events are perceived to begin before action onset. Imperfect distinctions  
244 between anticipated and actual sensory consequences would cause congruent sensation  
245 to be perceived as longer. In contrast, when action consequences are unexpected, pre-  
246 activated outcomes differ from the actual sensory consequences and can thus be  
247 discriminated. The hypothesis that duration biases result from imperfect distinctions  
248 between predicted and stimulus-driven percepts is consistent with the finding that  
249 imagined and perceived visual events activate common occipital representations  
250 (Kosslyn et al., 1993; Albers, Kok, Toni, Dijkerman, & de Lange, 2013; see also Buetti &  
251 Macaluso, 2010), and that action preparation activates representations of the  
252 anticipated effects (Müsseler & Hommel, 1997; Kühn, Keizer, Rombouts, & Hommel,  
253 2011). Furthermore, the idea that the perceived onset of anticipated events is shifted in  
254 time is consistent with a number of temporal distortions in the action control literature.  
255 For example, it has long been recognized that, when tapping to a metronome,  
256 movements show a phase lead to the pacing signals (Dunlap, 1910; Bartlett & Bartlett,  
257 1959). Moreover, effects resulting from action but at delay are perceived to occur  
258 earlier than in reality (Haggard, 2005).

259

260 Temporal biases resulting from the prediction of congruent action consequences might  
261 be expected to detract from effective action control. However, illusory biases often  
262 result from the operation of adaptive mechanisms. For example, visual aftereffects,  
263 defined by significant sensory distortion, are believed to be the products of ongoing  
264 perceptual recalibration to ambient sensory inputs (Clifford & Rhodes, 2005; Thompson  
265 & Burr, 2009; see also Yarrow, Haggard, Heal, Brown, & Rothwell, 2001). Similarly,  
266 stimulus-general temporal dilation during action planning may maximize information  
267 acquisition prior to movement (Hagura, Kanai, Orgs, & Haggard, 2012). Following this  
268 line of reasoning, we suggest that the dilation of subjective action-time observed for  
269 anticipated sensory outcomes may be indicative of an adaptive mechanism optimized  
270 for online action control. Anticipation of the sensory consequences of action is essential  
271 for action selection and subsequent error correction. Imperfect distinction between  
272 anticipated and actual sensory outcomes may reflect exploitation of mechanisms  
273 adapted for perception during action planning. While these mechanisms broadly benefit

274 actors, there may be surprising consequences for tightly time-locked action control and  
275 social perception. For example, we may be in contact with grasped objects for less time  
276 than we realize and handshakes may be briefer than we believe.

277

278 Equivalent effects when observing sensory events from first and third person  
279 perspectives suggests that common mechanisms anticipate the consequences of our  
280 own actions as well as the imitative reactions of others. Wolpert et al. (2003) proposed  
281 that sensory prediction mechanisms for action control may also operate when  
282 interacting with others, but this possibility has received little empirical investigation.  
283 The present study provides support for this hypothesis, suggesting that we  
284 overestimate the duration, not only of our own actions, but also others' imitative  
285 reactions. Future investigations must establish whether these effects are found when  
286 other individuals react in a non-imitative, but predictable, manner; for example, when  
287 dominant body postures result in complementary submissive postures of an interactant  
288 (Tiedens & Fragale, 2003).

289

290 Neuropsychological and neuroimaging studies have implicated motor structures in  
291 duration perception, even when action is not required. For example, the cerebellum and  
292 basal ganglia are thought to play key roles in a range of temporal judgments  
293 (Harrington, Haaland, & Hermanowicz, 1998; Ivry & Keele, 1989; Ivry, Spencer,  
294 Zelaznik, & Diedrichsen, 2002; Koch et al., 2007). Additionally, greater activation has  
295 been observed in cortical motor areas, including the supplementary motor area (SMA)  
296 and dorsal premotor cortex, when judging the duration of visual events (Coull, Nazarian,  
297 & Vidal, 2008; Ferrandez et al., 2003), than when making intensity or color judgments  
298 about the same stimuli. These duration judgments may recruit the motor system to  
299 exploit mechanisms adapted, either phylogenetically or ontogenetically (Heyes, 2003),  
300 for action control.

301

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307

308 **References**

309

310 Albers, A.M., Kok, P., Toni, I., Dijkerman, H.C., & de Lange, F.P. (2013). Shared representations for  
311 working memory and mental imagery in early visual cortex. *Current Biology*, 23, 1-5. doi:  
312 10.1016/j.cub.2013.05.06

313 Bartlett, N.R., & Bartlett, S.C. (1959). Synchronization of a motor response with an anticipated  
314 sensory event. *Psychological Review* 66(4), 203-218.

315 Blakemore, S. J., Frith, C. D., & Wolpert, D. M. (1999). Spatio-temporal prediction modulates the  
316 perception of self-produced stimuli. *Journal of Cognitive Neuroscience*, 11(5), 551-559.

317 Blakemore, S. J., Wolpert, D. M., & Frith, C. D. (1998). Central cancellation of self-produced tickle  
318 sensation. *Nature Neuroscience*, 1(7), 635-640. doi:10.1038/2870

319 Bueti, D., & Macaluso, E. (2010). Auditory temporal expectations modulate activity in visual  
320 cortex. *NeuroImage*, 51(3), 1168-1183. doi: 10.1016/j.neuroimage.2010.03.023

321 Christensen, A., Ilg, W., & Giese, M. A. (2011). Spatiotemporal Tuning of the Facilitation of  
322 Biological Motion Perception by Concurrent Motor Execution. *Journal of Neuroscience*,  
323 31(9), 3493-3499. doi:10.1523/JNEUROSCI.4277-10.2011

324 Clifford, C. W. G., & Rhodes, G. (2005). *Fitting the mind to the world: Adaptation and after-effects*  
325 *in high-level vision*. Oxford University Press.

326 Coull, J. T., Nazarian, B., & Vidal, F. (2008). Timing, storage, and comparison of stimulus duration  
327 engage discrete anatomical components of a perceptual timing network. *Journal of Cognitive*  
328 *Neuroscience*, 20(12), 2185-2197. doi:10.1162/jocn.2008.20153

329 Diedrichsen, J., Verstynen, T., Hon, A., Lehman, S. L., & Ivry, R. B. (2003). Anticipatory  
330 adjustments in the unloading task: Is an efference copy necessary for learning?  
331 *Experimental Brain Research*, 148(2), 272-276. doi:10.1007/s00221-002-1318-z

332 Dufossé, M., Hugon, M., & Massion, J. (1985). Postural forearm changes induced by predictable in  
333 time or voluntary triggered unloading in man. *Experimental Brain Research*, 60(2), 330-334.  
334 doi:10.1007/BF00235928

335 Dunlap, K. (1910). Reaction to rhythmic stimuli with attempt to synchronize. *Psychological*  
336 *Review*, 17(6), 399-416.

337 Fagioli, S., Hommel, B., & Schubotz, R.I. (2007). Intentional control of attention: Action planning  
338 primes action-related stimulus dimensions. *Psychological Research*, 71, 22-29. doi:  
339 10.1007/s00426-005-0033-3.

340 Ferrandez, A. ., Hugueville, L., Lehericy, S., Poline, J. ., Marsault, C., & Pouthas, V. (2003). Basal  
341 ganglia and supplementary motor area subtend duration perception: an fMRI study.  
342 *NeuroImage*, 19(4), 1532-1544. doi:10.1016/S1053-8119(03)00159-9

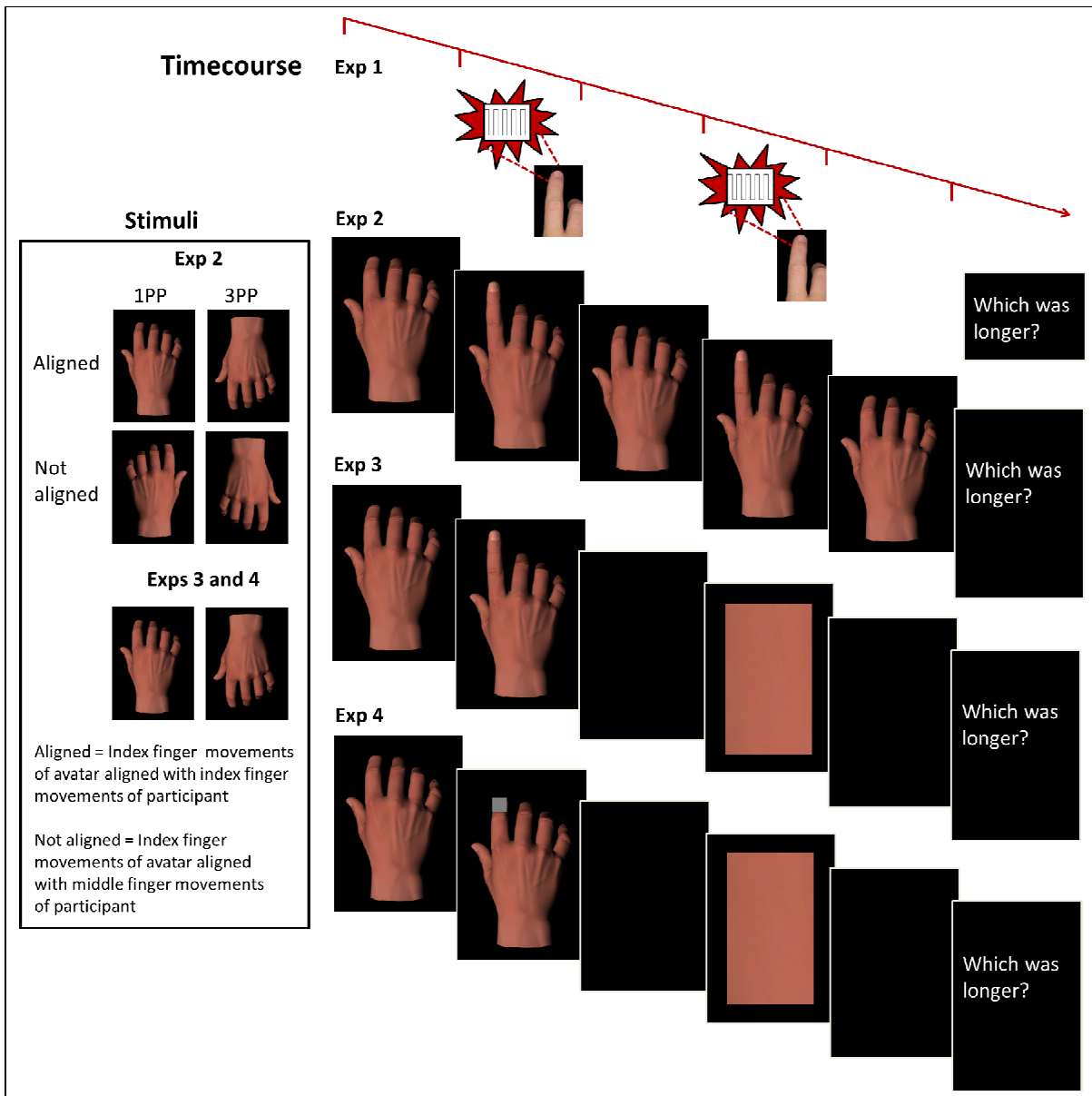
343 Gavazzi, G., Bisio, A., & Pozzo, T. (2013). Time perception of visual motion is tuned by the motor  
344 representation of human actions. *Scientific Reports*, 3, 1168. doi: 10.1038/srep01168

345 Greenwald, A. G. (1970). Sensory feedback mechanisms in performance control: With special  
346 reference to the ideo-motor mechanism. *Psychological Review*, 77(2), 73-99.  
347 doi:10.1037/h0028689

- 348 Haggard, P. (2005). Conscious intention and motor cognition. *Trends in Cognitive Sciences*, 9,  
349 290-295. doi: 10.1016/j.tics.2005.04.012
- 350 Hagura, N., Kanai, R., Orgs, G., & Haggard, P. (2012) Ready steady slow: action preparation slows  
351 the subjective passage of time. *Proceedings of the Royal Society of London, B*. 279(1746),  
352 4399-4406. doi: 10.1098/rspb.2012.1339
- 353 Harrington, D. L., Haaland, K. Y., & Hermanowicz, N. (1998). Temporal processing in the basal  
354 ganglia. *Neuropsychology*, 12(1), 3-12.
- 355 Heyes, C. (2003). Four routes of cognitive evolution. *Psychological Review*, 110(4), 713-727.  
356 doi:10.1037/0033-295X.110.4.713
- 357 Hommel, B., Müsseler, J., Aschersleben, G., & Prinz, W. (2001). The Theory of Event Coding  
358 (TEC): A framework for perception and action planning. *Behavioral and Brain Sciences*,  
359 24(05), 849-878. doi:10.1017/S0140525X01000103
- 360 Hommel, B. (2010). Grounding attention in action control: The intentional control of selection.  
361 In B.J Bruya (ed.), *Effortless attention: A new perspective in the cognitive science of attention*  
362 *and action* (pp.121-140). Boston, MA: MIT Press.
- 363 Ivry, R. B., & Keele, S. W. (1989). Timing functions of the cerebellum. *Journal of Cognitive*  
364 *Neuroscience*, 1(2), 136-152. doi:10.1162/jocn.1989.1.2.136
- 365 Ivry, R. B., Spencer, R. M., Zelaznik, H. N., & Diedrichsen, J. (2002). The cerebellum and event  
366 timing. *Annals of the New York Academy of Sciences*, 978(1), 302-317. doi:10.1111/j.1749-  
367 6632.2002.tb07576.x
- 368 Kingdom, F.A.A., & Prins, N. (2010). *Psychophysics: A practical introduction*. London, UK:  
369 Elsevier.
- 370 Koch, G., Oliveri, M., Torriero, S., Salerno, S., Lo Gerfo, E., & Caltagirone, C. (2007). Repetitive TMS  
371 of cerebellum interferes with millisecond time processing. *Experimental Brain Research*,  
372 179(2), 291-299. doi:10.1007/s00221-006-0791-1
- 373 Kosslyn, S.M., Alpert, N.M., Thompson, W.L., Maljkovic, V., Weise, S.B., Chabris, C.F., et al. (1993).  
374 Visual mental imagery activates topographically organized visual cortex: PET investigations.  
375 *Journal of Cognitive Neuroscience*, 5(3), 263-287. doi:10.1162/jocn.1993.5.3.263
- 376 Kühn, S., Keizer, A., Rombouts, S.A.R.B., & Hommel, B. (2011). The functional and neural  
377 mechanism of action preparation: Roles of EBA and FFA in voluntary action control. *Journal*  
378 *of Cognitive Neuroscience*, 23, 214-220. doi: 10.1162/jocn.2010.21418
- 379 Lee, D. N., Young, D. S., Reddish, P. E., Lough, S., & Clayton, T. M. H. (1983). Visual timing in  
380 hitting an accelerating ball. *The Quarterly Journal of Experimental Psychology Section A*,  
381 35(2), 333-346. doi:10.1080/14640748308402138
- 382 Müsseler, J., & Hommel, B. (1997). Blindness to response-compatible stimuli. *Journal of*  
383 *Experimental Psychology: Human Perception and Performance*, 23, 861-872. doi:  
384 10.1037/0096-1523.23.3.861
- 385 Press, C., Gherri, E., Heyes, C., & Eimer, M. (2010). Action preparation helps and hinders  
386 perception of action. *Journal of Cognitive Neuroscience*, 22(10), 2198-2211.  
387 doi:10.1162/jocn.2009.21409

- 388 Schubotz, R.I. (2007). Prediction of external events with our motor system: towards a new  
389 framework. *Trends in Cognitive Sciences*, 11(5), 211-218. doi: 10.1016/j.tics.2007.02.006
- 390 Thompson, P., & Burr, D. (2009). Visual aftereffects. *Current Biology*, 19(1), R11-R14.  
391 doi:10.1016/j.cub.2008.10.014
- 392 Tiedens, L.Z., & Fragale, A.R. (2003). Power moves: Complementarity in dominant and  
393 submissive nonverbal behaviour. *Journal of Personality and Social Psychology*, 84(3),  
394 558-568. doi:10.1037/0022-3514.84.3.558
- 395 Wang, L. & Jiang, Y. (2012). Life motion signals lengthen perceived temporal duration.  
396 *Proceedings of the National Academy of Sciences*, 109(11), 673-677. doi:  
397 10.1073/pnas.1115515109
- 398 Wolpert, D. M., Doya, K., & Kawato, M. (2003). A unifying computational framework for motor  
399 control and social interaction. *Philosophical Transactions of the Royal Society of London.*  
400 *Series B: Biological Sciences*, 358(1431), 593-602. doi:10.1098/rstb.2002.1238
- 401 Yarrow, K., Haggard, P., Heal, R., Brown, P., & Rothwell, J. C. (2001). Illusory perceptions of space  
402 and time preserve cross-saccadic perceptual continuity. *Nature*, 414(6861), 302-305.  
403 doi:10.1038/35104551
- 404

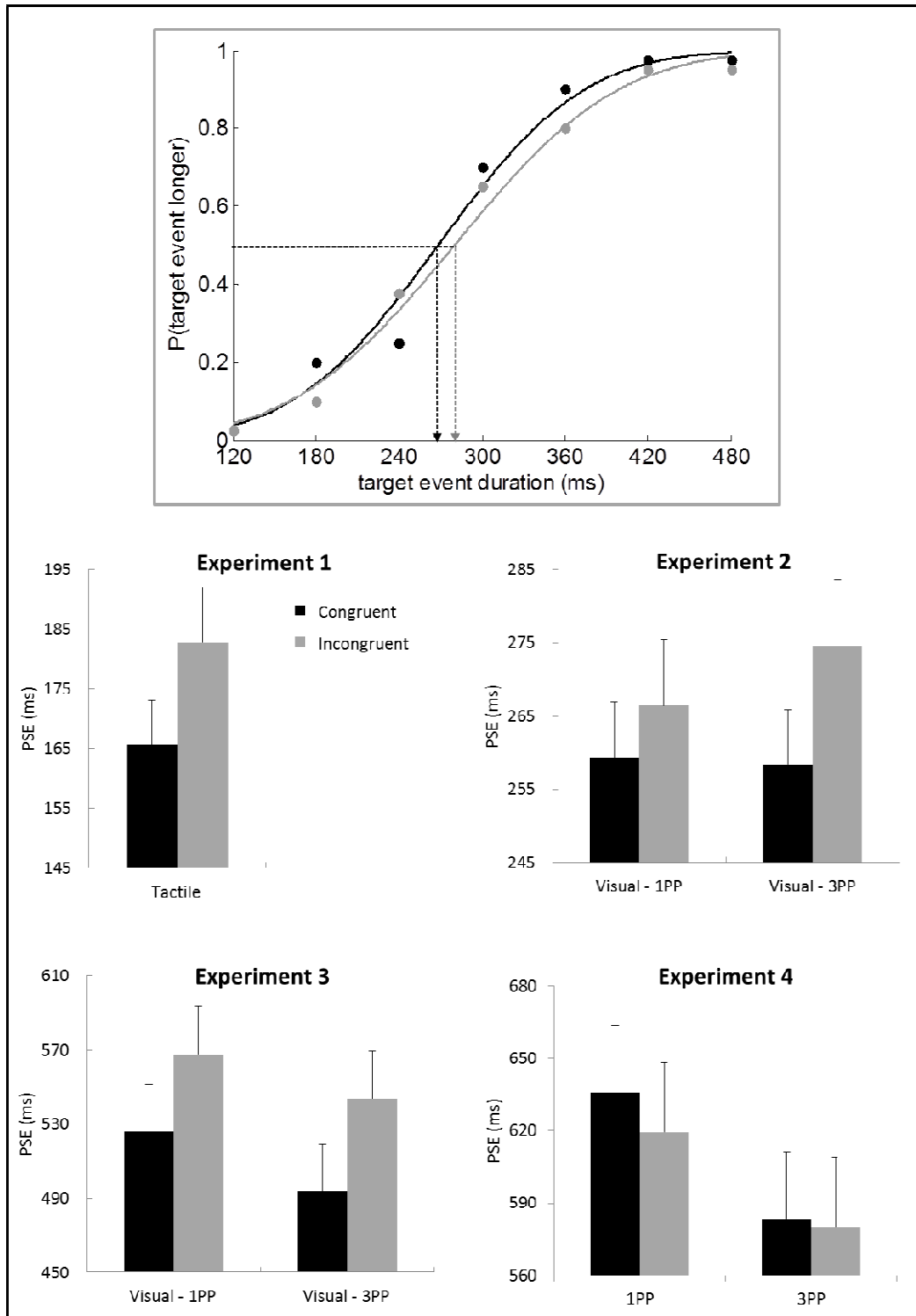




405

406 **Figure 1. The visual stimuli (created using Smith Micro Software's Poser 7.0) and timecourse for the action-**  
 407 **related events in each of the four experiments. Timecourse stimuli depict the avatar hand in first person**  
 408 **perspective.**

409



410

411 **Figure 2. Top panel: Demonstration of how the PSE was calculated with psychometric functions for an**  
 412 **example participant, with stimuli congruent and incongruent with moving fingers. The PSE describes the**  
 413 **point where participants judge the target and reference events as having equal duration. Judgment precision**  
 414 **was inferred from the standard deviation of the Gaussian distribution which best fits the data; it pertains to**  
 415 **the inverse of the slope, and lower thresholds reflect more consistent categorizations, thereby indicating**  
 416 **better performance. Other panels: Mean PSEs for stimuli congruent and incongruent with moving fingers,**  
 417 **for all experiments and perspectives. 1PP = first person perspective, 3PP = third person perspective. Error bars**  
 418 **represent the standard error of the mean.**  
 419

	Congruent	Incongruent
Experiment 1: Tactile	107.5 (52.3)	129.4 (68.5)
Experiment 2: Visual – 1PP	100.6 (13.5)	100.2 (12.1)
Experiment 2: Visual – 3PP	100.5 (14.5)	111.3 (14.8)
Experiment 3: Visual – 1PP	379.9 (67.1)	346.8 (48.2)
Experiment 3: Visual – 3PP	330.3 (40.6)	287.3 (35.2)
Experiment 4: 1PP	294.2 (21.9)	283.1 (28.6)
Experiment 4: 3PP	318.1 (36.5)	319.5 (37.7)

421 **Table 1. Mean precision estimates for stimuli congruent and incongruent with moving fingers, shown**  
422 **separately for each experiment and perspective. Standard error of the mean is displayed in brackets in each**  
423 **condition. 1PP = first person perspective, 3PP = third person perspective.**

424

425

426 **FOOTNOTES**

427 <sup>i</sup> Developed by the Cogent 2000 team at the FIL and the ICN and Cogent Graphics  
428 developed by John Romaya at the LON at the Wellcome Trust Centre for Neuroimaging.

429 <sup>ii</sup> Examining congruency-induced temporal dilation in the visual modality also permits  
430 better isolation of perceptual effects from the direct effects of action performance.  
431 Visually-defined congruency is eliminated when the hands are occluded. It is not  
432 possible to eliminate tactile-defined congruency without some form of sensory  
433 deafferentation.

434 <sup>iii</sup> This similarity was observed across experiments despite changes in the range of  
435 durations presented. It is worth noting that piloting indicated these shifts in duration to  
436 be necessary for two reasons. First, the apparent motion in Experiments 2 and 3 did not  
437 appear natural with short durations. Second, the duration judgments became more  
438 difficult across experiments, moving from punctate touch to apparent motion in vision  
439 in Experiment 2, and changing the nature of the reference relative to the target in  
440 Experiment 3. Given these changes to the durations presented in Experiments 1-3, it is  
441 difficult to draw conclusions concerning the presence of a precision effect in Experiment  
442 1 and its absence in Experiments 2 and 3.

443