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2

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 toolboxⁱ. 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 perspectivesⁱⁱ.

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 movementsⁱⁱⁱ. 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.

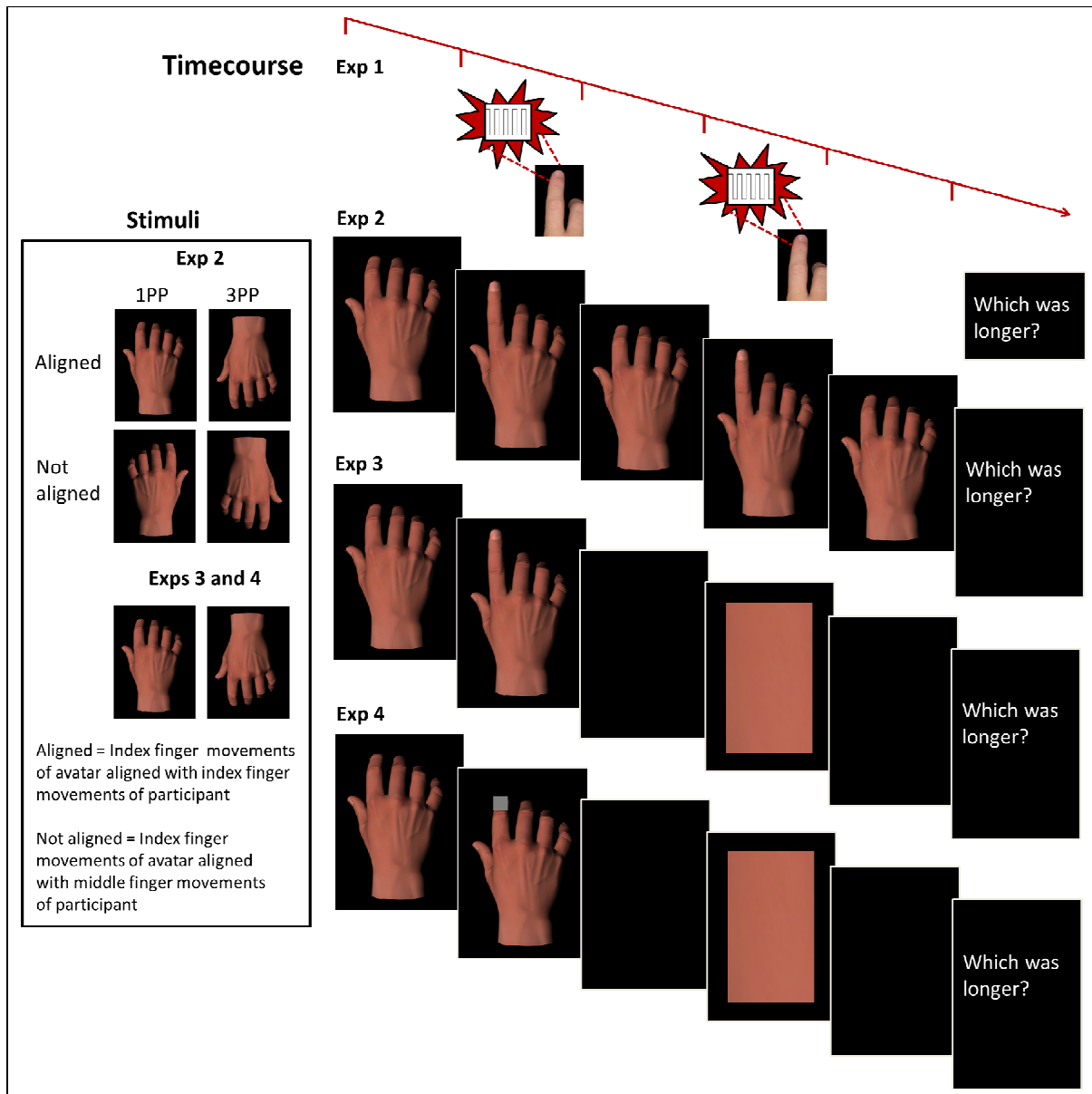
340 Ferrandez, A. ., Hugueville, L., Lehericy, S., Poline, J. ., Marsault, C., & Pouthas, V. (2003). Basal
341 ganglia and supplementary motor area subtend duration perception: an fMRI study.
342 *NeuroImage*, 19(4), 1532-1544. doi:10.1016/S1053-8119(03)00159-9

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 Hagara, 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

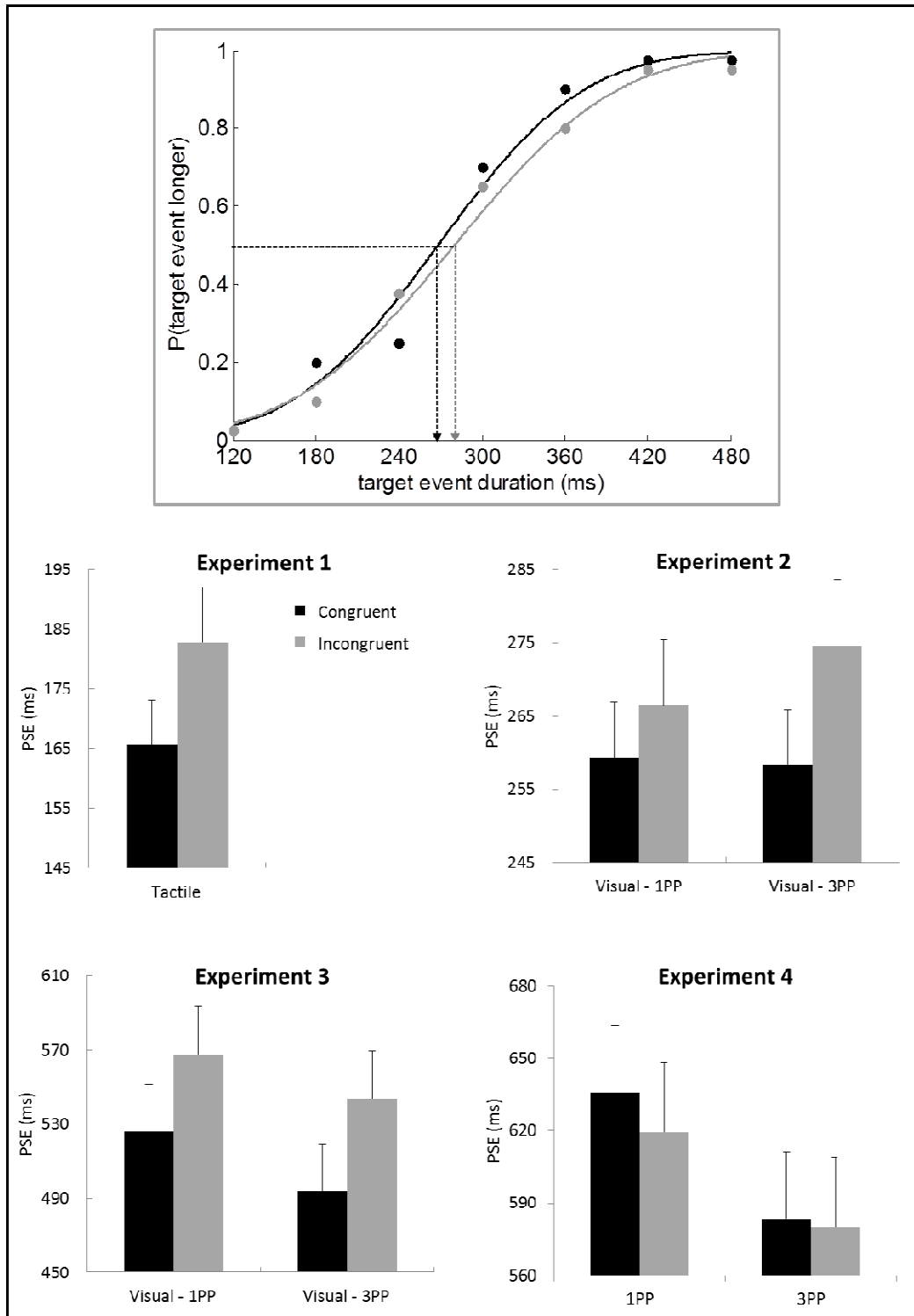
- 388 Schubotz, R.I. (2007). Prediction of external events with our motor system: towards a new
389 framework. *Trends in Cognitive Sciences*, 11(5), 211-218. doi: 10.1016/j.tics.2007.02.006
- 390 Thompson, P., & Burr, D. (2009). Visual aftereffects. *Current Biology*, 19(1), R11-R14.
391 doi:10.1016/j.cub.2008.10.014
- 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



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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.**

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425

426 **FOOTNOTES**

427 ⁱ 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 ⁱⁱ 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 ⁱⁱⁱ 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