

Attentional processes link perception and action

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Behavioural studies on normal and brain-damaged individuals provide convincing evidence that the perception of objects results in the generation of both visual and motor signals in the brain, irrespective of whether or not there is an intention to act upon the object. In this paper we sought to determine the basis of the motor signals generated by visual objects. By examining how the properties of an object affect an observer's reaction time for judging its orientation, we provide evidence to indicate that directed visual attention is responsible for the automatic generation of motor signals associated with the spatial characteristics of perceived objects.

Keywords: vision; attention; action; movement; affordance

1. INTRODUCTION

More than any other sense, vision allows us to navigate our environment and act upon it. The remarkable efficiency with which we are able to do this can be attributed to the fact that some of our actions, such as reaching for an object, are based, in part, on a repertoire of potential movements derived directly from vision (Gibson 1979; Lhermite 1983; Hommel 1995; Hari et al. 1998; Rumiati & Humphreys 1998; Craighero et al. 1999; Grezes & Decety 2002). There is convincing evidence that shifts of attention to the location of visual objects automatically generate some motor response codes (Simon 1969; Nicoletti & Umilta 1989, 1994; Stoffer 1991; Tipper et al. 1992, 1998; Proctor & Lu 1994; Rizzolatti et al. 1994; Rubichi et al. 1997; Sheliga et al. 1997), mimicking the way oculomotor systems are recruited (Fischer & Breitmeyer 1987; Goldberg & Segraves 1987; Rizzolatti et al. 1987; Schneider & Deubel 1995; Corbetta & Shulman 1998). Separate motor response codes are believed to be generated by the object itself, although their basis remains unclear. There is a long-standing view that perceived objects automatically generate motor codes based on the actions most highly associated with them (Craft & Simon 1970; Wallace 1971; Gibson 1979; Michaels 1988; Tucker & Ellis 1998).

It is axiomatic that the selection of movement plans from an existing repertoire of signals is advantageous in terms of both execution time and efficiency. These are crucial factors in making postural adjustments essential for survival; for example, moving your arm to deflect an approaching missile, or regulating your gait while in motion to avoid an obstacle. While the advantage of generating motor response codes with respect to the location of objects is clear, it is not clear why motor codes would be generated automatically by everyday objects on the basis of the actions they afford: speed is not essential when grasping a mug of coffee. With such actions we have the luxury of evaluating the sensory information at hand, allowing the formulation of motor plans on the basis of need. To determine whether the action-related properties of an object (or its elements) are responsible for the generation of motor codes we used, as stimuli, computer-generated images of everyday objects that either afforded a grasping action or not. Our measure was the reaction time for judging the orientation of each stimulus. The rationale for this experimental approach is well documented (Proctor & Reeve 1990; Tucker & Ellis 1998). In brief, if the action-relevant properties of an object are represented automatically during visual perception, then one would expect preferential activation of the hand most suited to perform the action which should, in turn, facilitate speeded responses by that hand. To determine the role of directed visual attention in the generation of motor codes, we used the same experimental paradigm in conjunction with object and non-object patterns that were either symmetrical or asymmetrical about the vertical axis.

2. MATERIAL AND METHODS

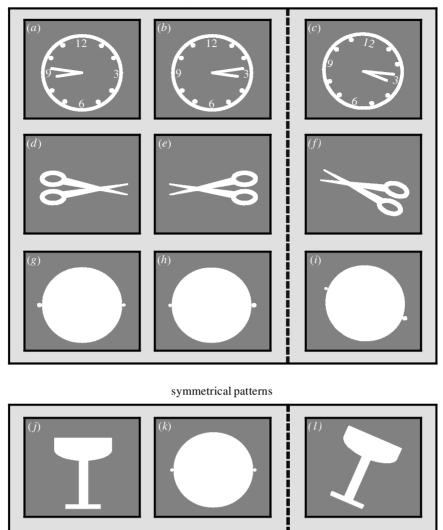
(a) Stimuli

White-on-grey images of object and non-object stimuli were generated using a VSG2/3 graphics board (from Cambridge Research Systems, Rochester, Kent, UK) and displayed on an Eizo T662 monitor. Examples of the stimuli are shown in figure 1. The stimuli were viewed binocularly at either 50 cm or 4 m in dim light, using either central or peripheral vision (centred 7.5° along the horizontal meridian in the left visual field). A fixation target was only employed for eccentric viewing. At 50 cm viewing distance, the object stimuli occupied approximately the same size $(10-14^\circ)$ as they would in life.

(b) *Procedure*

Separate experiments were run for object and non-object stimuli. Within each experiment, a stimulus was selected at random with equal probability and abruptly presented at the centre of the monitor. Our measure was reaction time (ms) for judging whether the stimulus was oriented clockwise or anticlockwise from its 'normal' orientation. The stimulus was presented up to the reaction key press. Observers were shown each stimulus in its 'normal' orientation prior to any measurement trials (see

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asymmetrical patterns

Figure 1. Examples of the stimuli used, as they appeared on the display monitor. Stimuli were either asymmetrical (a-i) or symmetrical (j-l) about the vertical axis. All stimuli were approximately the same size $(10-14^{\circ} \text{ at } 50 \text{ cm viewing distance})$. The panels to the left of the dashed line depict the stimuli in their 'normal' orientation, while those to the right of the line depict how the stimuli would have appeared on the monitor when clockwise-oriented (by 18°). The analogue clock and non-object patterns were deemed not to afford an action whereas both the scissors and wine glass were. The non-object stimuli were termed 'symmetrical' (1k, side patches of equal size), 'left-asymmetrical' (1g, left-side patch larger than right) or 'right-asymmetrical' (1h, right-side patch larger than left).

figure 1). Response buttons were fixed to a table in front of the observer, symmetrically displaced about their midline and separated by 30 cm. Observers were instructed to press the right button with their right hand if the stimulus was clockwise-oriented and the left button with their left hand if it was anticlockwise-oriented. Either the reaction time (ms) for a correct response or a response error was recorded. The next trial followed automatically after a delay of 2 s. Two of the authors (S.J.A. and V.K.) plus a total of 33 naive persons acted as observers: all had normal or corrected-to-normal visual acuity and all were right-handed.

Object stimuli included scissors with handles to the left or right, an analogue clock set at 8.45 or at 3.15 and a wine glass. Each was presented either clockwise- or anticlockwise-oriented by 18°, resulting in a total of 10 different object stimuli. For observers S.J.A. and V.K., five blocks of 100 trials were completed within each experiment, yielding approximately 50 trials per object (minus any response errors made). Reaction time data

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for the naive observers are reported as group means, each observer having completed one block of 100 trials (approximately 10 trials per object per observer). The error rate data for the naive observers are reported as group totals.

Two types of non-object stimuli were used. In one experiment, each stimulus consisted of three white, non-patterned, circular luminance patches: a large centrally viewed patch with a small patch either side of it (examples are shown in figure 1). The stimuli were termed 'symmetrical' (side patches of equal size), 'left-asymmetrical' (left-side patch larger than right) or 'right-asymmetrical' (right-side patch larger than left). Each was presented either clockwise or anticlockwise-oriented, resulting in a total of six different non-object stimuli. In another experiment, the asymmetrical patterns were constructed using a single side patch only. The same symmetrical pattern was used. Again, the stimuli were termed 'symmetrical' (side patches of equal size), 'left-asymmetrical' (left-side patch only) or 'right-asymmetrical' (right-side patch only). In both experiments with non-object

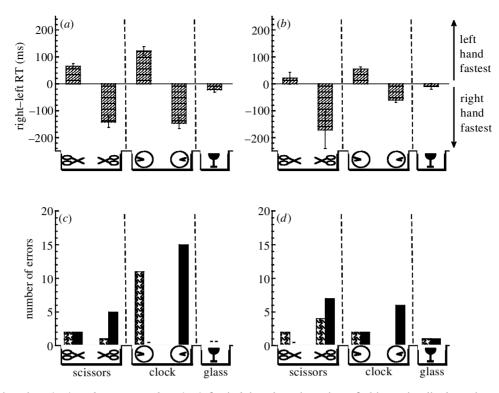


Figure 2. Reaction time (a,b) and error rate data (c,d) for judging the orientation of object stimuli when viewed centrally at a distance of 50 cm, for two experienced observers (S.J.A. (a,c) and V.K. (b,d)). Left-hand reaction times (RT) were subtracted from right-hand reaction times and plotted along the ordinate for each stimulus type (the small icons depict scissors with handles to the left or right of fixation, a clock with hands to the left or right of fixation, and a symmetrical wine glass). Positive values indicate that the left hand was fastest and negative values, that the right hand was fastest. The error bars show \pm one standard error. The total number of response errors is plotted for each stimulus type. Note that for some object patterns, no error was recorded (indicated by a small horizontal dash near the abscissa). Black bars in (c,d) indicate right-hand errors; cross-hatched bars indicate left-hand errors.

patterns the diameter of the large central patch was 10° , matching that of the clock face (figure 1*a*). For S.J.A. and V.K., three blocks of 120 trials were completed within each experiment, yielding approximately 60 trials per non-object stimulus (minus any response errors made). Reaction time data for the naive observers are reported as group means, each observer having completed one block of 60 trials (approximately 10 trials per stimulus per observer). The error rate data for the naive observers are reported as group totals. Note that different sets of naive observers were employed in the experiments with object and non-object stimuli.

3. RESULTS AND DISCUSSION

(a) Object stimuli

(i) Experiment 1

The results for two experienced observers (S.J.A and V.K., the authors) are shown in figure 2 for object patterns viewed centrally at a distance of 50 cm. Observers reported that the hands of the clock, the handles of the scissors and the bowl of the wine glass were salient object features, and that these features were used to judge object orientation on each presentation. The pattern of results for both the clock and scissors was near identical: left-hand responses were fastest when the salient features of the object were to the left of fixation and *vice versa* (figure 2a,b). Response errors usually coincided with the asymmetry of the object. For example, false right-hand responses were to the right of fixation (figure 2c,d). A small

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hand-response difference was also evident with the symmetrical wine glass (figure 2): each individual responded fastest with their right hand (figure 2). We assume this reflects hand dominance, as each individual was righthanded.

Discussion

The reaction time and error rate data for the scissors are in agreement with the hypothesis that perceived objects automatically potentiate components of the actions they afford. This is so because they were within the reaching space of the observer (50 cm), and because it is reasonable to suppose that any grasping action afforded by the scissors would be directed towards the handles and not the open, pointed ends. However, this model cannot explain the results for the clock. Although one could grasp the clock when positioned at 50 cm, it does not afford any left or right bias for so doing. Our results for the clock show that any advantage one hand had over the other in making a response was dependent on the time depicted (figure 2). We further observed that the pattern of results for both the scissors and clock was unaffected by viewing distance (50 cm versus 4 m). This also runs contrary to the affordance hypothesis that states that visual objects should elicit automatic motor codes only if the objects are within reaching distance (Tucker & Ellis 1998).

(ii) Experiment 2

This first experiment was repeated using a group of 21 naive observers, principally to determine whether practice

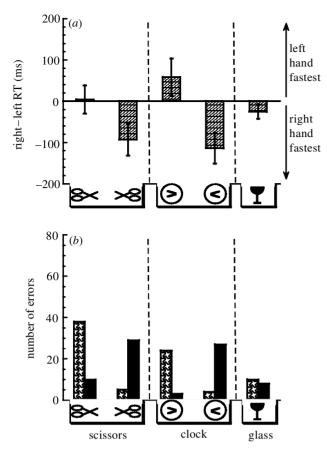


Figure 3. Reported salient feature: handle of scissors. Reaction time (a) and error rate data (b) for judging the orientation of object stimuli when viewed centrally at a distance of 50 cm, for naive observers who reported that the location of the handles of the scissors was used to judge orientation. Reaction time data are reported as group means, and the error rate data are reported as group totals. Data are plotted using the same format as in figure 2. Bar shading as in figure 2.

and/or familiarity with the stimuli affected the authors' results. Following completion of the experiment, each naive observer reported that the hands of the clock, and the bowl of the wine glass, were conspicuous on each presentation, in agreement with the authors. However, only 10 naive observers agreed with the authors that the handles of the scissors were their most salient feature. Ten other observers reported that the open, pointed ends of the scissors attracted their attention on each presentation and were used to judge orientation, while one observer reported that the 'long axis of the scissors' was used to judge orientation. The pooled data for the naive observers showed clear hand-response differences for the clock, in agreement with the authors, but no hand-response differences for the scissors. However, dividing the entire dataset according to which feature of the scissors the observers reported as most conspicuous revealed differences for both the clock and scissors. Figure 3 shows the dataset for those naive observers who reported that the handles of the scissors were used to judge orientation; figure 4 shows the results for those who reported that the pointed ends of the scissors were used to judge orientation. (The data for the naive observer who based their judgement on 'the long axis of the scissors' were necessarily excluded from this

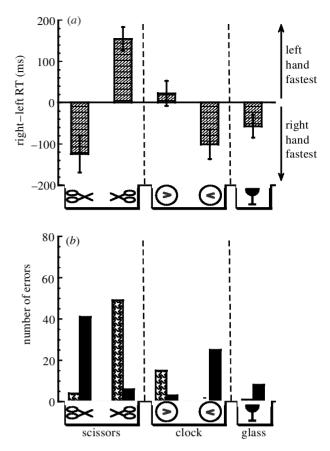


Figure 4. Reported salient feature: points of scissors. Reaction time (a) and error rate data (b) for judging the orientation of object stimuli when viewed centrally at a distance of 50 cm, for naive observers who reported that the location of the pointed ends of the scissors was used to judge orientation. Reaction time data are reported as group means, and the error rate data are reported as group totals. Data are plotted using the same format as in figure 2. Bar shading as in figure 2.

analysis.) The pattern of responses for both the clock and wine glass was similar for each dataset (compare figures 3 and 4), and in general agreement with the authors' data (figure 2). However, the two datasets for the scissors were virtually a mirror image of one another. For example, the number of response errors was greatest for the right hand when either the handles of the scissors were conspicuous and to the right of fixation (figure 3b), or when the pointed ends were conspicuous and to the right of fixation (figure 4b).

Discussion

Given these results, it appears unlikely that the handresponse differences for the scissors were based on behaviourally relevant actions afforded by them. The critical factor was the location of the scissors' conspicuous feature, as reported by each observer, which we assume induced a degree of attentional bias within the object.

(iii) Experiment 3

In this experiment the object stimuli were positioned entirely within the left visual field, centred at 7.5° along the horizontal meridian and viewed at a distance of 50 cm. The results are shown in figure 5 for experienced

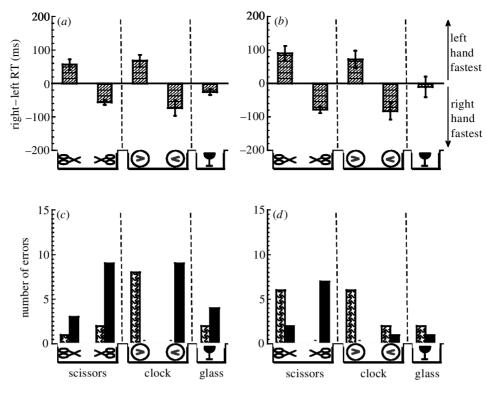


Figure 5. Reaction time (a,b) and error rate data (c,d) for judging the orientation of object stimuli when viewed eccentrically (centred 7.5° along the horizontal meridian in the left visual field), for two experienced observers (S.J.A. (a,c) and V.K. (b,d)). The viewing distance was 50 cm. Data are plotted using the same format as in figure 2. Bar shading as in figure 2.

observers S.J.A and V.K. As with central viewing, both observers reported that the hands of the clock, the handles of the scissors and the bowl of the wine glass were used to judge object orientation on each presentation. Note that the pattern of results was near identical to that for central viewing (compare figures 2 and 5). For example, despite the object being located entirely within the left visual field, right-hand responses remained faster and more error prone than left-hand responses when the object's salient features were to the right.

Discussion

The results of this experiment support the assumption that it was attentional bias within the object and not hemifield bias that was responsible for between-hand motor response differences.

(b) Non-object stimuli

(i) Experiment 1

Each stimulus consisted of a large centrally viewed patch and two small side patches. The side patches were either the same size (symmetrical pattern, figure 1k) or not (asymmetrical pattern, figure $1g_{,h}$), and were displaced from the horizontal meridian such that the stimulus appeared clockwise- (figure 1i) or anticlockwise-oriented. Reaction time for judging orientation was measured using the same protocol employed with object stimuli. The results are shown in figure 6 for one experienced observer (S.J.A) and a group of 12 naive observers. For observer S.J.A (figure 6a,c), left-hand responses were fastest (and more error prone) with left-asymmetrical patterns, and right-hand responses were fastest (and more error prone) with right-asymmetrical patterns. With symmetrical patterns, the error rate was similar for each hand but reaction time was fastest for the right hand. The same pattern of results was evident for the group of naive observers (figure 6b,d). These results are in general agreement with those obtained using object patterns (compare figures 2 and 6).

Discussion

The degree of asymmetry in the 'left-asymmetrical' and 'right-asymmetrical' patterns was subtle, one side patch being only marginally larger than the other (see figure 1g,h). The task of judging stimulus orientation could have been completed by locating the position of either side patch. Nonetheless, a clear hand-response difference was evident, the hand yielding the fastest and most error prone response coinciding with the position (left or right) of the largest side patch. These results are consistent with verbal reports by observers that the largest side patch was used to judge the orientation of asymmetrical patterns.

Because of the abstract nature of the non-object patterns, it is unlikely that the hand-response differences shown in figure 6 could be explained by the affordance hypothesis. The pattern of results leads us to assume that the observer's attention was biased towards the position of the largest side patch in the asymmetrical patterns. Accepting this to be the case, we propose that a motor response advantage for one hand or the other is a consequence of there being a correspondence between the direction of the attentional shift and the hand needed to produce a correct response.

(ii) Experiment 2

The stimuli used in the previous experiment were modified such that the asymmetrical patterns contained a single side patch only. The position (left or right) of the side patch defined whether the pattern was left- or right-asym-

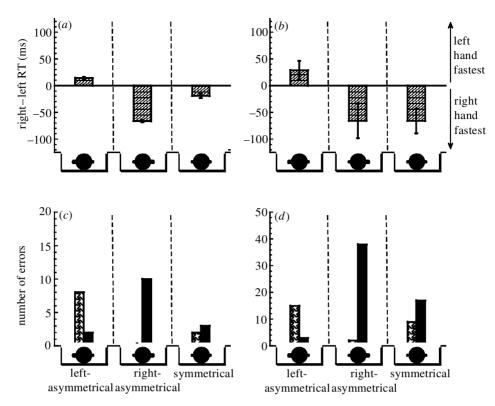


Figure 6. Reaction time (a,b) and error rate data (c,d) for judging the orientation of non-object stimuli when viewed centrally at a distance of 50 cm, for one experienced observer (S.J.A. (a,c)) and 12 naive observers (b,d). For naive observers, the reaction time data are reported as group means, and the error rate data are reported as group totals. Data are plotted using the same format as in figure 2. The small icons depict left-asymmetrical, right-asymmetrical and symmetrical patterns. Note that the asymmetrical patterns were constructed using side patches of unequal size, examples of which are shown in figure 1. Bar shading as in figure 2.

metrical (see small icons in figure 7). The same symmetrical pattern was used. The results for observers S.J.A. and V.K. were similar: clear hand-response differences were evident with asymmetrical patterns but not with symmetrical patterns (figure 7). The fastest and most error prone hand always coincided with the position of the side patch in the asymmetrical patterns.

Discussion

Unlike the previous experiment, the degree of asymmetry in the left-asymmetrical and right-asymmetrical patterns was marked, there being only one side patch. The task of judging stimulus orientation could only have been completed by shifting attention to the side patch. The similarity of results across all experiments leads us to conclude that directed visual attention is responsible for the automatic generation of motor signals associated with the spatial characteristics of perceived object- and nonobject patterns.

4. GENERAL DISCUSSION

Our results support the hypothesis that some of our actions are based, in part, on a repertoire of potential movements derived directly from vision. However, they do not support the idea that the automatic generation of motor signals by visual objects is based on the actions they afford. Previous studies claiming evidence in support of the affordance hypothesis generally used visually asymmetrical targets (e.g. Tucker & Ellis 1998; Grezes &

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Decety 2002). We show that a motor response advantage for one hand or the other accompanies the perception of asymmetrical targets, be they object or non-object patterns. The hand with the advantage coincided with the direction of attentional bias within the target, irrespective of the target's location in the visual field or the functional properties of its elements. This indicates that the attentional shift may stem from a coding strategy whereby the features of the target are labelled as a single entity. Such a strategy may be crucial for the control of motor actions but, assuming it takes place in the visuomotor stream (Milner & Goodale 1995; Sakata et al. 1997), would not necessarily lead to object recognition. We suggest that cooperative activity between the visuomotor and perceptual streams, as has been postulated for efficient and coordinated actions within our environment (Milner & Goodale 1995), may not be necessary if the encoded features of a target were flagged as a single entity within the visuomotor stream. Such a representation would help explain the puzzle of why patients with dorsal stream lesions manifest symptoms of 'object-based' neglect, whereby the contralesional side of an individual object is ignored regardless of its position in the visual field (Rizzolatti et al. 1994; Walker 1995).

We conclude that visually asymmetrical targets induce an attentional bias that in turn leads to the generation of a basis set of motor signals for potential arm or hand movements. Depending on the nature of the target and the needs of the observer, the motor signals so generated may or may not be utilized. If action is required, and the

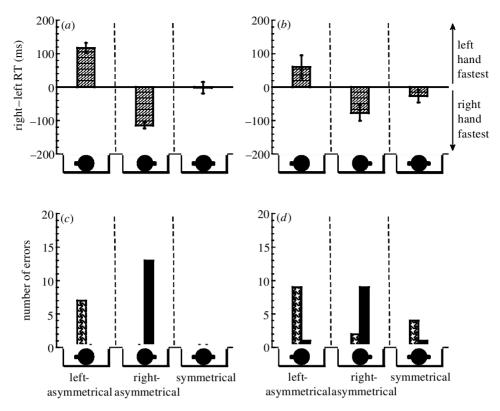


Figure 7. Reaction time (a,b) and error rate data (c,d) for judging the orientation of non-object stimuli when viewed centrally at a distance of 50 cm, for experienced observers S.J.A. (a,c) and V.K. (b,d). Data are plotted using the same format as in figure 2. The small icons depict left-asymmetrical, right-asymmetrical and symmetrical patterns. Note that the asymmetrical patterns were constructed using a single side patch. Bar shading as in figure 2.

direction of the attentional shift is also that which effects the correct motor response, the action will be facilitated. However, incongruent motor signals could be generated if the direction of the attentional shift is opposed to that of the appropriate motor response. Where simultaneous, contradictory motor signals exist, the frontal cortex may be important for selecting which signals are needed for goal-directed movements (Goldberg & Segraves 1987).

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REFERENCES

- Corbetta, M. & Shulman, G. L. 1998 Human cortical mechanisms of visual attention during orienting and search. *Phil. Trans. R. Soc. Lond. B* 353, 1353–1362. (DOI 10.1098/ rstb.1998.0289.)
- Craft, J. L. & Simon, R. 1970 Processing symbolic information from a visual display: interference from an irrelevant directional cue. *J. Exp. Psychol.* 83, 415–420.
- Craighero, L., Fadiga, L., Rizzolatti, G. & Umilta, C. 1999 Action for perception: a motor-visual attentional effect. J. Exp. Psychol. Hum. Percept. Perform. 25, 1673–1692.
- Fischer, B. & Breitmeyer, B. 1987 Mechanisms of visual attention revealed by saccadic eye movements. *Neuropsychologia* 25, 73–83.
- Gibson, J. J. 1979 *The ecological approach to visual perception*. Boston: Houghton-Mifflin.
- Goldberg, M. E. & Segraves, M. A. 1987 Visuospatial and motor attention in the monkey. *Neuropsychologia* 25, 107– 118.

Grezes, J. & Decety, J. 2002 Does visual perception of object

afford action? Evidence from a neuroimaging study. *Neuro-psychologia* **40**, 212–222.

- Hari, R., Forss, N., Avikainen, S., Kirveskari, E., Salenius, S. & Rizzolatti, G. 1998 Activation of human primary motor cortex during action observation: a neuromagnetic study. *Proc. Natl Acad. Sci. USA* 95, 15 061–15 065.
- Hommel, B. 1995 Stimulus-response compatibility and the Simon effect: toward an empirical clarification. J. Exp. Psychol. Hum. Percept. Perform. 21, 764–775.
- Lhermite, F. 1983 Utilisation behavior and its relation to lesions of the frontal lobes. *Brain* **106**, 237–255.
- Michaels, C. F. 1988 S–R compatibility between response position and destination of apparent motion: evidence of the detection of affordances. *J. Exp. Psychol. Hum. Percept. Perform.* 14, 231–240.
- Milner, A. D. & Goodale, M. A. 1995 *The visual brain in action*. Oxford University Press.
- Nicoletti, R. & Umilta, C. 1989 Splitting visual space with attention. *J. Exp. Psychol. Hum. Percept. Perform.* 15, 164–169.
- Nicoletti, R. & Umilta, C. 1994 Attention shifts produce stimulus spatial codes. *Psychol. Res.* 56, 144–150.
- Proctor, R. W. & Lu, C.-H. 1994 Referential coding and attention-shifting accounts of the Simon effect. *Psychol. Res.* 56, 185–195.
- Proctor, R. W. & Reeve, T. G. 1990 Research on stimulusresponse compatibility: toward a comprehensive account. In *Stimulus-response compatibility* (ed. R. W. Proctor & T. G. Reeve), pp. 483–494. Amsterdam: Elsevier.
- Rizzolatti, G., Riggio, L., Dascola, I. & Umilta, C. 1987 Reorienting attention across the horizontal and vertical meridians: evidence in favor of a premotor theory of attention. *Neuropsychologia* 25, 31–40.
- Rizzolatti, G., Riggio, L. & Sheliga, B. M. 1994 Space and

selective attention. In *Attention and performance*, vol. XV (ed. C. Umilta & M. Moscovitch), pp. 231–265. Cambridge, MA: MIT Press.

- Rubichi, S., Nicoletti, R., Iani, C. & Umilta, C. 1997 The Simon effect occurs relative to the direction of an attention shift. *J. Exp. Psychol. Hum. Percpt. Perform.* 23, 1353–1364.
- Rumiati, R. I. & Humphreys, G. W. 1998 Recognition by action: dissociating visual and semantic routes to action in normal observers. *J. Exp. Psychol. Hum. Percept. Perform.* 24, 631–647.
- Sakata, H., Taira, M., Kusunoki, M., Murata, A. & Tanaka, Y. 1997 The parietal association cortex in depth perception and visual control of hand action. *Trends Neurosci.* 20, 350–357.
- Schneider, W. X. & Deubel, H. 1995 Visual attention and saccadic eye movements: evidence for obligatory and selective spatial coupling. In *Eye movement research* (ed. J. M. Findlay, R. Walker & R. W. Kentridge), pp. 317–324. Amsterdam: Elsevier.
- Sheliga, B. M., Craighero, L., Riggio, L. & Rizzolatti, G. 1997 Effects of spatial attention on directional manual and ocular responses. *Exp. Brain Res.* 114, 339–351.

- Simon, J. R. 1969 Reactions toward the source of stimulation. *J. Exp. Psychol.* **81**, 174–176.
- Stoffer, T. H. 1991 Attentional focussing and spatial stimulusresponse compatibility. *Psychol. Res.* 53, 127–135.
- Tipper, S. P., Lortie, C. & Baylis, G. C. 1992 Selective reaching: evidence for action-centered attention. *J. Exp. Psychol. Hum. Percept. Perform.* 18, 891–905.
- Tipper, S. P., Howard, L. A. & Houghton, G. 1998 Actionbased mechanisms of attention. *Phil. Trans. R. Soc. Lond.* B 353, 1385–1393. (DOI 10.1098/rstb.1998.0292.)
- Tucker, M. & Ellis, R. 1998 On the relations between seen objects and components of potential actions. J. Exp. Psychol. Hum. Percept. Perform. 24, 830–846.
- Walker, R. 1995 Spatial and object-based neglect. *Neurocase* 1, 371–383.
- Wallace, R. J. 1971 S-R compatibility and the idea of a response code. *J. Exp. Psychol.* **88**, 354–360.

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