

Research Report

SPLITTING THE BEAM: Distribution of Attention Over Noncontiguous Regions of the Visual Field

Arthur F. Kramer and Sowon Hahn

Beckman Institute and Department of Psychology, University of Illinois

Abstract—*In an effort to examine the flexibility with which attention can be allocated in visual space, we investigated whether subjects could selectively attend to multiple noncontiguous locations in the visual field. We examined this issue by precluding two separate areas of the visual field and requiring subjects to decide whether the letters that appeared in these locations matched or mismatched while distractors that primed either the match or mismatch response were presented between the cued locations. If the distractors had no effect on performance, it would provide evidence that subjects can divide attention over noncontiguous areas of space. Subjects were able to ignore the distractors when the targets and distractors were presented as nononset stimuli (i.e., when premasks were changed into the targets and distractors). In contrast, when the targets and distractors were presented as sudden-onset stimuli, subjects were unable to ignore the distractors. These results begin to define the conditions under which attention can be flexibly deployed to multiple noncontiguous locations in the visual field.*

A number of metaphors have been used to describe how organisms selectively process information in the visual field. Some researchers have suggested that attention operates like a spotlight (Broadbent, 1982; Posner, Snyder, & Davidson, 1980). Information that falls within the spotlight is actively processed while information outside the spotlight either passively decays or is actively inhibited. In general, the spotlight is assumed to be of a fixed diameter, so focusing attention on objects in different regions of the visual field requires a movement of the spotlight. Other theorists have proposed a somewhat more flexible model of visual attention. For example, it has been suggested that attention operates like a zoom lens on a camera (Eriksen & St. James, 1986; Jonides, 1983). The lens can be either tightly focused on a limited area of visual space or expanded to include a larger portion of the visual field. Within this framework, the rate of processing is inversely related to the size of the focus. Thus, it is assumed that attention is a limited resource and that processing rate is proportional to the number of resource units allocated per unit of visual space. Processing of information outside the focus involves either expanding the focus of the lens or reorienting a tightly focused beam. One important assumption shared by both the spotlight and the zoom-lens models is that there is a unitary attentional focus. Thus, within these theoretical frameworks, attention cannot be focused simultaneously in noncontiguous areas of the visual field.

There are, however, several models which do suggest that

attention can be focused on noncontiguous regions of the visual field. For example, LaBerge and Brown (1989) have suggested that multiple attentional gradients can be imposed upon different locations in visual space such that objects may be selectively processed within these areas but not in intervening portions of the visual field. A number of other researchers (Baylis & Driver, 1993; Duncan, 1984; Kahneman, Treisman, & Gibbs, 1992; Kramer & Jacobson, 1991) have suggested that attention is focused on objects or perceptual groups in the visual field. Within these object-based models of visual attention, it is possible to divide attention among noncontiguous regions of space in situations in which objects with similar characteristics are distributed across the visual field. For example, according to these models, it should be possible to selectively process traffic that is moving at a constant velocity and direction, and therefore forms a perceptual group, and ignore stationary objects such as route signs, which may be physically interspersed among the moving automobiles.

Empirical tests of the assumption of noncontiguous attentional selection have produced mixed results. Previous studies have varied the probability that a target will appear at one of several locations in the visual field and have found patterns of response times that have been interpreted as evidence that attention was divided on the basis of this probabilistic information (Shaw & Shaw, 1977). However, other researchers have suggested that this pattern of performance could be the result of shifting attention from location to location across trials rather than dividing attention among locations within a single trial (Posner et al., 1980). Other researchers have cued two locations, one in each hemifield, and have found an inverse relationship between the size of a box cue and response time to detect a luminance increment target (Castiello & Umiltà, 1992). Given the assumption that processing rate is inversely related to the size of the attended region, these findings suggest that attention can be divided among multiple locations, at least in situations in which only a single stimulus appears in the visual field.

However, several other studies have failed to find evidence for the division of attention among noncontiguous locations. For example, in studies in which multiple locations were precued as potential target locations, processing time benefits were found either for only the most probable location (Posner et al., 1980) or for the second most probable location only when it was adjacent to the most probable target location (Eriksen & Yeh, 1985). Other studies have found that subjects are unable to selectively ignore distractor stimuli that are located between two cued target locations (Heinze et al., 1994; Pan & Eriksen, 1993).

These results leave unresolved the question of whether it is possible to divide attention among multiple noncontiguous re-

Address correspondence to Arthur F. Kramer, Beckman Institute, University of Illinois, 405 North Mathews Ave., Urbana, IL 61801.

Splitting the Beam

gions of the visual field. We believe that one important characteristic of the studies that have failed to find evidence consistent with the division of attention is that the targets and distractors have been presented as transient events. There is now a substantial body of literature (Muller & Rabbitt, 1989; Theeuwes, 1992; Yantis, 1993) which suggests that transient or sudden-onset events capture attention automatically. Thus, it is conceivable that in previous paradigms, the onset distractors might have captured attention, thereby rendering it difficult or impossible to selectively focus on the target locations.

We tested this hypothesis by presenting subjects with displays like the ones illustrated in Figure 1. The subjects were instructed to decide whether two target letters matched or mismatched. In one condition (onset condition), the targets and distractors occurred as sudden onsets; in another condition (nononset condition), the targets and distractors were created by removing sections of premasks. If the sudden-onset distractors were capturing attention in previous studies, and thereby making it difficult to maintain attention on the target locations, we would expect an interaction between the type of response (match vs. mismatch) and type of distractor (same or different) with the onset but not with the nononset stimuli.

METHODS

Subjects

Eight undergraduate students (age range: 19–26 years) were paid \$5.00 per hour to participate in the study. Four of the subjects were male. All of the subjects possessed corrected visual acuities (Snellen) of at least 20/20.

Apparatus and Stimuli

Stimulus letters were displayed by a Dell 316SX computer with a VGA graphics adaptor and monitor. Uppercase *E*, *H*, *P*, *S*, and *U* served as the targets and distractors. In the onset condition, the target and distractors were presented simultaneously. In the nononset condition, the targets and distractors were presented by removing two of seven segments of a figure-eight premask. Each letter subtended a visual angle of 0.4° in width and 0.6° in height from a viewing distance of 60 cm. Viewing distance was controlled by using a chin rest. Each of the stimuli (premask, targets, and distractors) was presented on an imaginary circle with a radius of 8.5° of visual angle. A fixation point was located in the center of the imaginary circle. The display of two targets and two distractors could be located either above or below fixation. The two target letters appeared at either the 10:30 and 1:30 o'clock positions or at the 4:30 and 7:30 o'clock positions. The two distractors were located on the imaginary circle between the two targets. The center-to-center distance between the two target letters was 12° of visual angle. The locations of the two target letters were precued with two white square boxes, which were 2° in width and height. The precue boxes, premasks, targets, and distractors appeared as white stimuli on a black background.

Procedure

Prior to the beginning of each trial, the fixation cross was presented in the center of the display. When subjects had fixated the cross, they pressed the space bar, which began the trial. A schematic illustration of the temporal structure of the onset and nononset trials is presented in Figure 1. In the onset condition, two precue boxes appeared, in either the upper or the lower visual field, after subjects pressed the space bar. After 150 ms, two targets and two distractors were presented for 60 ms, at which point the boxes and letters were replaced with a pattern mask. In the nononset condition, pressing the space bar led to the presentation of the two precue boxes along with four figure-eight premasks, two in the target positions (i.e., in the boxes) and two in the distractor positions. After 150 ms, two segments were removed from each of the figure eights to reveal two targets and two distractors. The targets and distractors remained on the display for 60 ms and then were replaced with a pattern mask.

We also included onset and nononset control conditions in which the targets were presented in the absence of the distractors. These conditions were included to assess whether differences in reaction time (RT) or accuracy between onset and

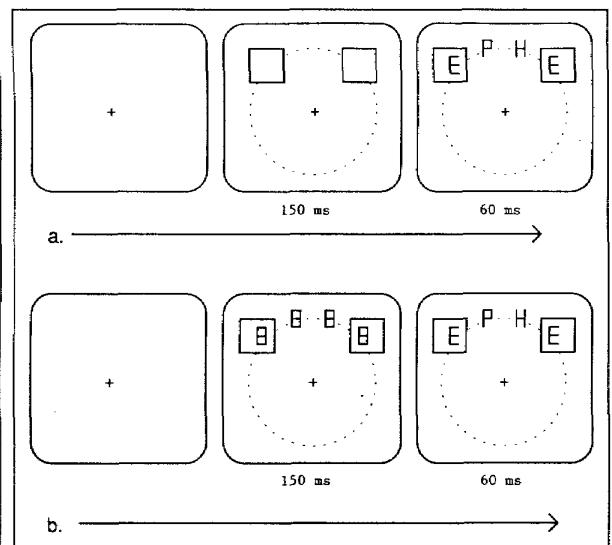


Fig. 1. Graphic illustration of the displays and temporal sequence of events within a trial in Experiment 1. The three panels in (a) represent the temporal sequence of the displays on the onset trials. The fixation cross was presented until the subjects depressed the space bar on the computer keyboard. Next, the precue boxes were presented for 150 ms. Finally, the two targets and two distractors were presented for 60 ms. The three panels in (b) represent the temporal sequence of the displays on the nononset trials. The fixation cross was presented until the subjects depressed the space bar on the computer keyboard. Next, the precue boxes were presented along with the figure-eight premasks for 150 ms. Finally, the two targets and two distractors were displayed by removing segments from the figure-eight premasks, and this display was presented for 60 ms.

nononset conditions might occur independently of the presence of distractors. For example, it is conceivable that onset stimuli, with their associated increase in luminance, might provide more effective temporal cues for the occurrence of the targets than the nononset stimuli. The more effective cuing might, in turn, lead to more rapid and accurate responding on the onset than on the nononset trials. If this were indeed the case, we would expect onset trials to be faster and more accurate than nononset trials whether the distractors were present (experimental conditions) or absent (control conditions).

The subjects' task in both the onset and the nononset conditions was to decide whether the two target letters matched or mismatched. If the two targets matched, subjects depressed one of two keys on the IBM-style keyboard (either the *l* or the *d* key); if the letters mismatched, subjects pressed the other key. Response key was counterbalanced across subjects. Subjects were instructed to respond as quickly as possible while maintaining their accuracy above 85%.

Subjects were also instructed to maintain fixation on the central cross throughout the trial because movement of their eyes to a target location would place the distractors closer to the fovea, thereby making the task more difficult. Subjects were presented with feedback on average speed and accuracy after each block of 60 trials.

Subjects performed two experimental sessions on subsequent days. In each session, subjects performed 15 blocks of 60 trials each. Half of the blocks were onset blocks and the other 50% of the blocks were nononset blocks. Response type (match or mismatch), distractor presence (present or absent), and distractor type (same or different) were within-block variables. In the same-distractor condition, the two distractors were identical. On a same-distractor/match-response trial, the two distractors were also the same as the two target letters. In the different-distractor condition, the two distractor letters were different from each other and also different from the target letters. Fifty percent of the trials were match-response trials, and 50% were mismatch-response trials. Each of the two distractor types occurred 50% of the time on the distractor-present trials. Each session lasted for approximately 1 hr, and subjects were encouraged to take breaks whenever they desired. The first onset and the first nononset trial block were considered practice and were not analyzed.

Design

Four factors were manipulated in the study: display condition (onset or nononset), response type (match or mismatch), distractor presence (present or absent), and distractor type (same or different). Display condition was a blocking factor; distractor type, distractor presence, and response type were randomized within blocks.

RESULTS

The means of single-subject median RTs and accuracy data are presented in Figure 2 for the experimental conditions. The RT data indicate a clear crossover interaction between response

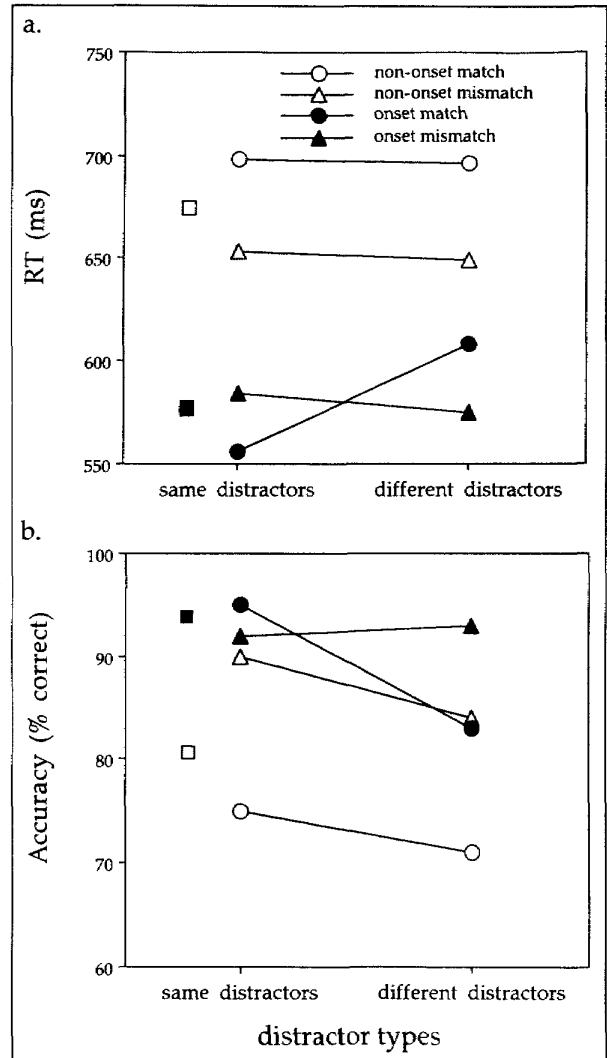


Fig. 2. Mean of the median reaction times (RTs) (a) and accuracy data (b) for the experimental conditions. Results for the control (i.e., nondistractor trials) conditions, averaged across match and mismatch responses, are indicated by solid (onset trials) and unfilled (nononset trials) squares.

type and distractor type for the onset but not the nononset conditions. This interaction also appears to be present in the accuracy data.

The RTs and accuracies were submitted to three-way analyses of variance with display condition (onset vs. nononset), distractor type (prime same vs. different response), and response type (match vs. mismatch) as within-subjects factors. A main effect was obtained for display condition (RT: $F[1, 7] = 17.2, p < .01$; accuracy: $F[1, 7] = 10.1, p < .01$). Subjects were slower and less accurate responding in the nononset than in the onset conditions.

Splitting the Beam

More important, however, with respect to the question of whether subjects are able to divide attention between noncontiguous spatial locations was the significant three-way interaction among display condition, response type, and distractor type (RT: $F[1, 7] = 12.3, p < .01$; accuracy: $F[1, 7] = 8.4, p < .01$). As can be seen in Figure 2, there was a significant interaction between response and distractor type for the onset but not for the nononset trials. Thus, the physically inserted distractors influenced task performance only when both the targets and the distractors were presented as onset stimuli. When the targets and distractors were formed from premasks (i.e., nononset stimuli), subjects' performance was uninfluenced by the nature of the distractor.

There is, however, one aspect of the data that might be interpreted as evidence that target processing was influenced by the distractors on the nononset trials. As can be seen in Figure 2, performance on nononset trials was slower and less accurate than performance on the onset trials. Thus, it might be suggested that these performance differences are consistent with the influence of distractors on target processing on the nononset trials. The data obtained in the control conditions, in which distractors were not present, discount this interpretation. If the distractors were responsible for the RT and accuracy differences between onset and nononset trials, we would not expect onset-versus-nononset performance differences on the control trials. However, if the main effects on RT and accuracy between onset and nononset trials were due to more effective temporal cuing for the appearance of the targets on the onset trials, then we would expect similar onset-nononset performance differences whether the distractors were present (experimental trials) or absent (control trials). The data are consistent with the latter interpretation. There were no significant differences between control and distractor-present trials for onset and nononset conditions for either RT or accuracy measures.

DISCUSSION

The results obtained in our study clearly suggest that attention can be flexibly deployed and maintained on multiple locations in the visual field as long as new objects do not appear among the attended locations. These data are important in that they suggest that the great majority of models of visual attention, with the exception of the gradient model proposed by LaBerge and Brown (1989), require modification if they are to accommodate the flexibility of attentional control demonstrated by young adults. As described previously, space-based models of attention, such as the spotlight (Broadbent, 1982; Posner et al., 1980) and zoom-lens (Eriksen & St. James, 1986; Eriksen & Yeh, 1985) models, suggest that attention is distributed in a unitary area of the visual field. Object-based models of attention (Duncan, 1984; Kahneman et al., 1992) allow for more attentional flexibility because in these models attention can be distributed in noncontiguous locations as long as the information in these areas is perceptually grouped (e.g., a building that is partially occluded by a tree, aircraft flying in formation). However, in the present study, there is no obvious grouping of the target locations. Thus, it appears that attention can be di-

vided between noncontiguous locations even in the absence of some form of grouping between the items in these locations.¹

The fact that sudden-onset distractors (new objects) do disrupt the goal-directed allocation of attention is probably fortunate: From an evolutionary perspective, such events (e.g., a predator, an automobile suddenly approaching a crosswalk) are often significant. It is of interest to note, however, that a number of researchers have found that attentional capture by sudden-onset objects can be overridden under some conditions. Yantis and Jonides (1990) found that if subjects are given a sufficient amount of time to focus their attention on a potential target location, sudden onsets that occur elsewhere in the visual field do not disrupt performance (see also Theeuwes, 1991; Koshino, Warner, & Juola, 1992). In fact, Yantis and Jonides (1990) found that a location precue-target stimulus onset asynchrony (SOA) of 200 ms was sufficient to override the influence of a sudden-onset distractor.

In our study, a precue-target SOA of 150 ms was ineffective in eliminating the influence of the sudden-onset distractors on the processing of the two targets. In an effort to determine whether additional preparatory time would enable subjects to effectively ignore the onset distractors, we conducted another study with equivalent experimental conditions and procedures except for a 600- rather than a 150-ms precue-target SOA. The results of this study with 6 subjects are presented in Figure 3. As in our previous experiment, subjects were unable to ignore the sudden-onset distractors. However, subjects were able to effectively ignore the distractors when they were presented as nononsets. Whether attentional capture by sudden-onset distractors can be overcome with longer precue-target SOAs or additional practice in situations in which subjects must focus on multiple display locations is an important topic for future research.

In our experiment, we used a brief precue-target SOA (150 ms) and a short display duration (60 ms) followed by a pattern mask in an effort to ensure that subjects neither had the time to move their eyes to the target locations nor had sufficient time to shift attention between these locations. Our choice of a 60-ms stimulus presentation duration was based on the results of previous researchers who estimated that it takes from 150 ms to

1. It might be argued that the two target locations in our task were grouped by virtue of the similarity of the precues (i.e., the two boxes). Although we view this as unlikely given the large separation between the two cued locations, we performed a control study in which both the targets and the distractors were precued with boxes. Eight subjects were instructed to compare the targets in the outermost boxes (i.e., the same locations that served as target locations in the main study reported here) and ignore the letters in the distractor (i.e., innermost) boxes. In all other respects, the control study was identical to the main study.

If subjects were grouping the target locations together on the basis of the similarity of the precues in the main study, we would expect that the four boxes used as precues in the control study would result in an interaction between distractor type and response type in both the onset and the nononset trials. If subjects were using the precues to orient attention to specific locations in the main study, we would expect distractor effects only on the onset trials. The data were consistent with this latter possibility. An interaction was obtained between response type and distractor type on the onset but not on the nononset trials in our control study.

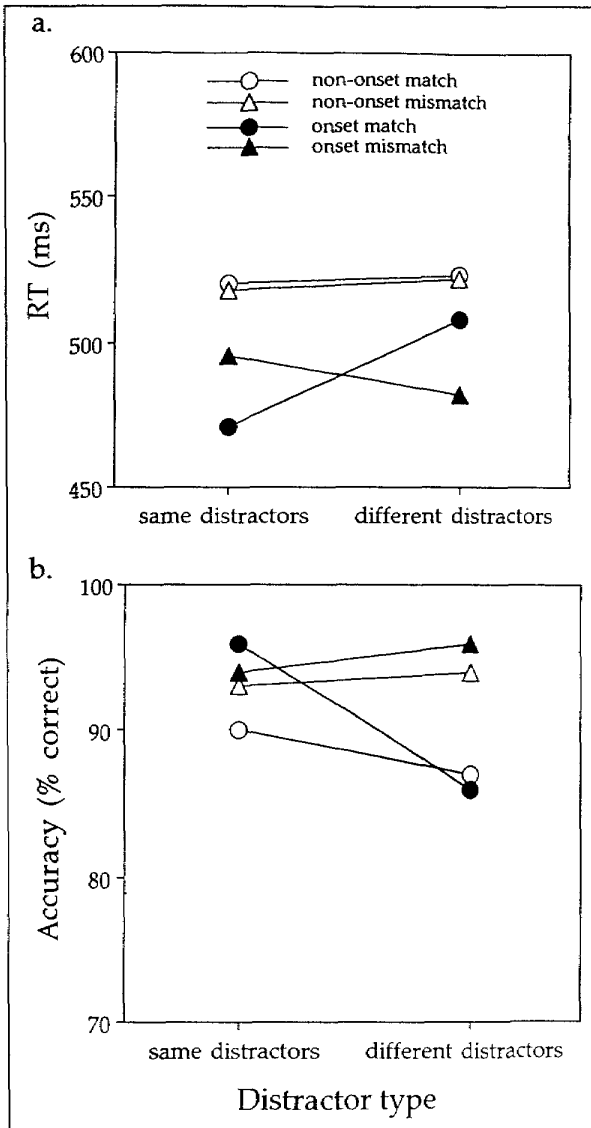


Fig. 3. Mean of the median reaction times (RTs) (a) and accuracy data (b) for each of the conditions when duration of the precue display was 600 ms.

250 ms to identify a stimulus at a precued location and switch attention to a second location (Eriksen & Yeh, 1985; Krose & Julesz, 1989; Madden, 1992; Sperling, 1984). However, research performed in visual search paradigms has found search slopes of less than 30 ms per letter, suggesting that attention may be switched among locations in less than 30 ms, at least with extremely well practiced subjects (Czerwinski, Lightfoot, & Shiffrin, 1992). Although the visual search task appears to lend itself to faster shifts of attention than the matching task, given that in search the target is known in advance and is often

unchanged across many experimental trials, we decided to err on the side of caution and examine the attention-switching hypothesis more directly in an additional study.

To that end, we modified the matching task so that subjects would be required to identify a letter in one of two precued locations and then shift their attention to the second location. In this way, we could estimate directly the speed with which subjects could switch attention in our task. Subjects were instructed that if a particular letter (i.e., either *E*, *H*, *P*, *S*, or *U*) was presented in one of the two precued locations (the specific location in which this letter would appear was defined in the pretrial instructions to the subject), then they should shift their attention to the second precued location, where a second letter was presented, and decide whether the two letters matched or mismatched. If a different letter appeared in the predefined location, subjects were told not to make the match/mismatch response. For example, on a particular trial, a subject might be instructed to complete the matching task only if the letter *E* appeared in the precue box on the left side of the display. Thus, in essence, subjects were required to first identify the letter in one precued location (e.g., the left-most precue box) and then switch their attention to the second precued location (e.g., the right-most precue box).

A critical aspect of the study was the delay between the occurrence of the first target letter (i.e., created by removing segments of a figure-eight premask) and the occurrence of the second target letter. We reasoned that RT would begin to increase at delays that exceeded the amount of time required for subjects to identify the letter in the primary location and shift attention to the second location. Delays between the presentation of the first and second target letters were 0, 14, 28, 42, 56, 70, 84, and 98 ms. Six subjects participated in this study. A significant increase in RT, relative to the 0-ms delay (i.e., the simultaneous presentation of the target letters in the primary and secondary locations), was obtained for the 84- and 98-ms delays. Therefore, it appears reasonable to conclude that subjects did not have a sufficient amount of time to switch attention between locations in our study, but instead our results can be attributed to the division of attention between noncontiguous locations.

Although our results are intriguing in that they suggest a remarkable degree of attentional flexibility—that is, the ability to divide attention among noncontiguous locations in the visual field—there are a number of important unresolved questions. For example, how many locations can be attended simultaneously? Can attention be divided within as well as between hemifields? Is it possible to overcome the influence of sudden onsets on goal-directed attentional allocation? What are the neural mechanisms that underlie the ability to split the attentional beam? We are currently exploring these issues in our laboratory.

Acknowledgments—This research was supported by a grant from the Office of Naval Research (N00014-92-J-1792). We wish to thank Marie Banich and Gordon Logan for helpful comments on a previous version of this manuscript.

REFERENCES

- Baylis, G., & Driver, J. (1993). Visual attention and objects: Evidence for hierarchical coding of location. *Journal of Experimental Psychology: Human Perception and Performance*, 19, 451-470.
- Broadbent, D.E. (1982). Task combination and selective intake of information. *Acta Psychologica*, 50, 253-290.
- Castiello, U., & Umiltà, C. (1992). Splitting focal attention. *Journal of Experimental Psychology: Human Perception and Performance*, 18, 837-848.
- Czerwinski, M., Lightfoot, N., & Shiffrin, R.M. (1992). Automatization and training in visual search. *American Journal of Psychology*, 105, 271-315.
- Duncan, J. (1984). Selective attention and the organization of visual information. *Journal of Experimental Psychology: General*, 113, 501-517.
- Eriksen, C.W., & St. James, J.D. (1986). Visual attention within and around the field of focal attention: A zoom lens model. *Perception & Psychophysics*, 40, 225-240.
- Eriksen, C.W., & Yeh, Y.Y. (1985). Allocation of attention in the visual field. *Journal of Experimental Psychology: Human Perception and Performance*, 11, 583-597.
- Heinze, H., Luck, S.J., Munte, T.F., Goes, A., Mangun, G.R., & Hillyard, S.A. (1994). Attention to adjacent and separate positions in space: An electrophysiological analysis. *Perception & Psychophysics*, 56, 42-52.
- Jonides, J. (1983). Further toward a model of the mind's eye movement. *Bulletin of the Psychonomic Society*, 21, 247-250.
- Kahneman, D., Treisman, A., & Gibbs, B. (1992). The reviewing of object files: Object-specific integration of information. *Cognitive Psychology*, 24, 175-219.
- Koshino, H., Warner, C., & Juola, J. (1992). Relative effectiveness of central, peripheral, and abrupt-onset cues in visual attention. *Quarterly Journal of Experimental Psychology*, 45, 609-631.
- Kramer, A.F., & Jacobson, A. (1991). Perceptual organization and focused attention: The role of objects and proximity in visual processing. *Perception & Psychophysics*, 50, 267-284.
- Kroese, B., & Julesz, B. (1989). The control and speed of shifts in attention. *Vision Research*, 29, 1607-1619.
- LaBerge, D., & Brown, V. (1989). Theory of attentional operation in shape identification. *Psychological Review*, 96, 101-124.
- Madden, D. (1992). Selective attention and visual search: Revision of an allocation model and application to age differences. *Journal of Experimental Psychology: Human Perception and Performance*, 18, 821-836.
- Mueller, H., & Rabbitt, P. (1989). Reflexive and voluntary orienting of attention: Time course of activation and resistance to disruption. *Journal of Experimental Psychology: Human Perception and Performance*, 15, 315-330.
- Pan, K., & Eriksen, C. (1993). Attentional distribution in the visual field during same-different judgments as assessed by response competition. *Perception & Psychophysics*, 53, 134-144.
- Posner, M.I., Snyder, C.R.R., & Davidson, B.J. (1980). Attention and the detection of signals. *Journal of Experimental Psychology: General*, 109, 160-174.
- Shaw, M., & Shaw, P. (1977). Optimal allocation of cognitive resources to spatial location. *Journal of Experimental Psychology: Human Perception and Performance*, 3, 201-211.
- Sperling, G. (1984). A unified theory of attention and signal detection. In R. Parasuraman & D. Davies (Eds.), *Varieties of attention* (pp. 103-182). New York: Academic Press.
- Theeuwes, J. (1991). Exogenous and endogenous control of attention: The effects of visual onsets and offsets. *Perception & Psychophysics*, 49, 83-90.
- Theeuwes, J. (1992). Perceptual selectivity for color and form. *Perception & Psychophysics*, 51, 599-606.
- Yantis, S. (1993). Stimulus-driven attentional capture. *Current Directions in Psychological Science*, 2, 156-161.
- Yantis, S., & Jonides, J. (1990). Abrupt visual onsets and selective attention: Voluntary versus automatic allocation. *Journal of Experimental Psychology: Human Perception and Performance*, 16, 121-134.

(RECEIVED 8/29/94; REVISION ACCEPTED 4/28/95)

This document is a scanned copy of a printed document. No warranty is given about the accuracy of the copy. Users should refer to the original published version of the material.