Relationships between attentional blink magnitude, RSVP target accuracy, and performance on other cognitive tasks

KAREN M. ARNELL and ASHLEY E. HOWE Brock University, St. Catharines, Ontario, Canada

MARC F. JOANISSE University of Western Ontario, London, Ontario, Canada

and

RAYMOND M. KLEIN Dalhousie University, Halifax, Nova Scotia, Canada

When two masked, to-be-attended targets are presented within approximately half a second of each other, performance on the second target (T2) suffers, relative to when the targets are presented further apart in time or when the first target (T1) can be ignored. This pattern of results is known as the attentional blink (AB). Typically, participants differ with respect to the magnitude of their AB and their overall target accuracy. Despite investigations as to what participant characteristics may influence AB performance (e.g., age, brain damage, or mood state), there has been no focused examination of whether individual differences in cognitive performance measures predict the magnitude of the AB or overall rapid serial visual presentation (RSVP) target accuracy. Our university student participants performed single-target and dual-target RSVP tasks, as well as a selection of cognitive tasks that did not use RSVP presentations, with color, letter, digit, and object stimuli. Overall performance on each of the RSVP targets (T1, T2, and single target) was predicted by speeded manual and vocal identification times to isolated stimuli and by performance with other RSVP targets. However, the magnitude of the AB was predicted only by T1 accuracy, not by any other performance measures. The results suggest that individual differences in AB magnitude do not result from differences in effective RSVP target encoding and are not well explained by varied information-processing abilities.

When two masked, to-be-attended targets are presented within approximately half a second of each other, performance on the second target (T2) is impaired, relative to when the targets are presented further apart in time or when the first target (T1) can be ignored (Broadbent & Broadbent, 1987; Raymond, Shapiro, & Arnell, 1992). This pattern of results is known as the *attentional blink* (AB; Raymond et al., 1992). The AB is most often investigated by embedding targets in rapid serial visual presentation (RSVP) streams in which stimuli are presented rapidly one at a time in the same spatial location. Dozens of studies have examined the presentation conditions that modulate the AB. For example, studies have examined the

Note—This article was accepted by the previous editorial team, when Colin M. MacLeod was Editor. importance of masking T1 and T2 in order to produce an AB (e.g., Chun & Potter, 1995; Giesbrecht & Di Lollo, 1998; Grandison, Ghirardelli, & Egeth, 1997; Raymond et al., 1992; Seiffert & Di Lollo, 1997), the AB with targets from different modality combinations (e.g., Arnell & Jolicœur, 1999; Arnell & Larson, 2002; Duncan, Martens, & Ward, 1997; Hillstrom, Shapiro, & Spence, 2002; Mondor, 1998; Potter, Chun, Banks, & Muckenhoupt, 1998; Soto-Faraco & Spence, 2002; Soto-Faraco et al., 2002), how the AB can be modulated by response selection demands (e.g., Arnell & Duncan, 2002; Jolicœur, 1998, 1999; Jolicœur & Dell'Acqua, 1999), and how the AB might vary when the difficulty of extracting the first target's identity is manipulated (McLaughlin, Shore, & Klein, 2001; Shapiro, Raymond, & Arnell, 1994; Shore, McLaughlin, & Klein, 2001; Ward, Duncan, & Shapiro, 1997). Other studies have examined the degree to which T2 was processed even when it could not be reported (e.g., Luck, Vogel, & Shapiro, 1996; Shapiro, Driver, Ward, & Sorensen, 1997; Vogel, Luck, & Shapiro, 1998) or have investigated the nature of targets that can overcome the AB (e.g., Anderson, 2005; Keil & Ihssen, 2004; Shapiro, Caldwell, & Sorensen, 1997).

The research was assisted by a grant from the Canadian Language and Literacy Research Network (CLLRNet) to K.M.A., M.F.J., and R.M.K. and by grants from NSERC, CFI, and OIT to K.M.A. Correspondence concerning this article should be addressed to K. M. Arnell, Department of Psychology, Brock University, St. Catharines, ON, L2S 3A1 Canada (e-mail: karnell@brocku.ca).

In addition to explorations of procedural variables, some investigators have begun to look at group differences in AB magnitude when different participant populations are tested. For example, the AB has been shown to be larger for elderly participants (Lahar, Isaak, & McArthur, 2001; Maciokas & Crognale, 2003), schizophrenics (Cheung, Chen, Chen, Woo, & Yee, 2002; Li et al., 2002), unilateral neglect patients (Husain, Shapiro, Martin, & Kennard, 1997), ADHD patients (Li, Lin, Chang, & Hung, 2004), Alzheimer's patients (Kavcic & Duffy, 2003), and those reporting severe depression symptoms (Rokke, Arnell, Koch, & Andrews, 2002), relative to matched controls. Green and Bavelier (2003) have also reported a reduced AB for action video game players. Typical university student participants also show variability in the magnitude of their AB and their overall target accuracy. It is somewhat surprising, then, that there has been no focused examination of whether individual differences in cognitive performance measures can predict individual differences in RSVP target accuracy or AB magnitude in this population. The question we will address here is the following: What cognitive performance measures might predict the magnitude of an individual's AB and his or her RSVP target accuracy?

Almost all theoretical models of the AB suggest that T2 performance suffers while limited capacity attentional resources are occupied with the processing of T1 (Shapiro, Arnell, & Raymond, 1997). A popular class of these models postulates a bottleneck for conscious stimulus identification and/or consolidation in working memory where conscious identification and consolidation of T2's representation must wait until conscious identification and consolidation of T1 is complete (Arnell, Helion, Hurdelbrink, & Pasieka, 2004; Chun & Potter, 1995; Jolicœur, 1998, 1999; Jolicœur & Dell'Acqua, 1998, 1999). If T2 is masked, its temporary representation will be overwritten by the mask while waiting for T1 to exit the bottleneck. If T1 processing in the bottlenecked stage(s) outlasts T2's representation, consolidation of T2 will fail, and subsequent T2 report accuracy will be reduced, resulting in an AB. Therefore, bottleneck models predict that under dual-task conditions, more efficient consolidation of T1 should reduce the wait at the bottleneck, thereby reducing the AB. Assuming that an individual's T1 accuracy score reflects the effectiveness and efficiency of the individual's T1 processing, at least in part, one might expect a negative relationship between T1 accuracy and AB magnitude, where those with higher T1 accuracy would show smaller ABs.¹ To the extent that those participants who efficiently and effectively encode T1 are the same participants who effectively and efficiently encode single RSVP targets, one might also predict that individuals who were better able to identify a single target in an RSVP stream of information would also show a smaller AB and have better performance on T1 and T2 on dual-task trials. Indeed, if RSVP target performance reflects the efficiency of identification and consolidation more generally, it is possible that faster and better identification of stimuli outside the

AB task (e.g., in speeded stimulus naming) would also predict improved RSVP target accuracy (T1, T2, or single target) and reduced AB magnitude.

Jolicœur's bottleneck model (e.g., Jolicœur, 1998, 1999; Jolicœur & Dell'Acqua, 1998, 1999) also suggests that the processing resources needed for stimulus consolidation in working memory are the same as those required for response selection operations. Several studies have shown that when T1 requires a speeded response, the response selection requirements of the T1 task modulate the magnitude of the AB, so that larger ABs are observed with greater T1 response selection requirements (Arnell & Duncan, 2002; Jolicœur, 1998, 1999). Thus, to the extent that individuals differ in their response selection ability and the AB is sensitive to such abilities, as postulated by Jolicœur (1998), it is reasonable to predict that individuals with longer response times (RTs) on a separate response selection task would produce larger ABs even when a nonspeeded AB paradigm is used. It is also reasonable to predict that individuals with longer RTs on a response selection task will have lower accuracy on T1, T2, or singletarget RSVP performance.

The Present Study

The goal of the present investigation was to examine individual differences in AB magnitude and RSVP target performance in relation to performance on several non-RSVP tasks that require identification of the same RSVP targets. Although previous research has provided a basis for several key predictions, studies to date have yet to directly examine these issues. Indeed, little direct evidence exists concerning the extent to which individual differences in AB magnitude or RSVP performance are associated with performance on other cognitive tasks.

In the present study, we measured the magnitude of the AB for each participant. We also measured each participant's overall accuracy for T1 and T2 in the AB task and his or her single-target RSVP accuracy across four stimulus types: letters, digits, objects, and colors. We predicted that individuals with higher single-target RSVP accuracy would also show higher T1 accuracy, higher T2 accuracy, and smaller AB magnitude, given that more efficient processing of single RSVP targets might predict more efficient processing of T1 and T2 in an AB task should reduce the likelihood that the T2 representation will be overwritten while waiting to be consolidated.

The participants in the present study also performed a variety of other non-RSVP tasks using the same stimulus items. These tasks required cognitive processes that have been implicated in RSVP performance and the AB. Vocal naming times (*vocal RTs*) to lone stimuli were measured, as were vocal naming times to 5×10 grids of stimuli (the rapid automatized naming, or RAN, task; Denckla & Rudel, 1974). The RAN task provided a measure of naming times in the context of competing stimuli. The accuracy with which a participant could correctly identify which stimulus from an array had been briefly presented

at a postcued location (the *location probe* task) was also measured. To the extent that rapid extraction of identity information from a stimulus is critical to RSVP target accuracy and/or AB magnitude, one might expect to see short naming times on vocal RT and RAN tasks, as well as high location probe accuracies associated with higher RSVP target accuracy and smaller ABs.

A manual RT task (pressing one of four keys matching the identity of the stimulus) was included to provide a measure of response selection speed. If response selection and stimulus consolidation do share a common bottlenecked processing stage (e.g., Jolicœur, 1998, 1999) and individuals differ in their response selection abilities, then to the degree that overall manual RTs reflected central processing demands, one would expect to see short response selection times associated with higher RSVP target accuracy and smaller ABs. A delayed RT task (in which the participants did not press the correct key until a tone was sounded after the stimulus) was also used to measure the speed of manual response execution without the identification and response selection components. This task was not expected to correlate with RSVP target accuracy or AB magnitude but was used simply as a motor execution control task. If response selection abilities, not response execution speed, underlie the expected relationships between manual RTs and AB magnitude and manual RTs and RSVP target accuracies, one would expect no relationship between delayed RTs and AB magnitude or RSVP target accuracy. However, if response execution speed underlies the expected relationships between manual RTs and AB magnitude and manual RTs and RSVP target accuracies, one would expect manual RTs and delayed RTs to be equally good predictors of AB magnitude and RSVP target accuracy.

METHOD

Participants

The present experiment draws on data from a previously conducted study (Arnell, Joanisse, Klein, Busseri, & Tannock, 2006), conducted to better understand the relationship between reading performance and performance on the clinical RAN task. Sixty-four undergraduate students from Brock University (n = 34) and the University of Western Ontario (n = 30) received course credit or a small monetary payment for their participation. The participants ranged in age from 19 to 26 years, and all reported learning English before 8 years of age and having normal (or corrected-to-normal) visual acuity. Each participant performed the experiment individually in a single session lasting approximately 2 h. All the participants performed the tasks in the following order: the RAN, manual RT, location probe, single-item vocal naming, delayed RT, single-target RSVP, and dual-target RSVP. The participants performed each task four times (once each with letters, digits, colors, and objects) in the same fixed order.

Apparatus

Stimulus presentation and response collection were controlled using E-Prime software (Schneider, Eschman, & Zuccolotto, 2002) running on a Windows XP-based desktop PC with a 17-in. CRT color monitor. The participants made manual responses with the computer keyboard and vocal responses with a voice key integrated into a serial response box (Psychology Software Tools).

Stimuli and Procedures

RAN task. A 5-row \times 10-column grid of items was presented on the computer screen. Each grid contained colors (red, green, blue, and yellow), digits (2, 4, 6, and 9), letters (g, k, m, and r), or object pictures (dog, hand, book, and chair). Each grid measured approximately 25 cm wide \times 18 cm high, subtending approximately 26.6° of visual angle at an unfixed binocular viewing distance of approximately 50 cm. Each individual element was approximately 1.5 cm high and wide, for a visual angle of approximately 1.7°. The participants pressed a key to display the grid and start the timer. The participants were instructed to accurately name each stimulus item as quickly as possible, beginning immediately after their keypress. The participants were told to name the grid items starting in the upper left element and ending at the lower right element, working their way across the rows from left to right. The participants pressed a key immediately after naming the last item to stop the computer timing. Items were named out loud, and errors were recorded by the experimenter. Each participant performed four trials, one with each stimulus type.

Manual RT task. A single stimulus from the RAN grid (e.g., a blue square) was presented in the center of the computer screen on each trial. The participants were instructed to press the key matching the identity of the stimulus as quickly and accurately as possible, using the index and middle fingers of both hands. Each trial began with a 500-msec presentation of a fixation cross and a 500-msec blank interval; then the stimulus was presented and remained on the screen until a response was made. A 500-msec intertrial interval followed the response. The "z," "x," "n," and "m" computer keyboard keys were labeled with the stimulus names to facilitate stimulus response mappings. Each participant performed one block for each of the four RAN stimulus categories (letters, digits, colors, and objects). Each block contained 48 trials, with each of the four stimulus exemplars presented 12 times each in random order (e.g., each of the four colors was presented 12 times in random order in the color block).

Delayed RT task. The delayed RT task was the same as the manual RT task, with the exception that the participants were told to delay their manual response until a tone sounded. The tone was randomly presented 1,500 or 2,000 msec after the onset of the stimulus element, thereby allowing the participant sufficient time to identify the stimulus and to select and prepare a response prior to the tone. The participants were instructed to prepare their response prior to the tone and then to make a speeded response as soon as the tone sounded. Each participant performed one block of 48 trials for each of the four stimulus categories.

Vocal naming task. The blocks, trials, and stimuli for the vocal naming task were identical to those for the manual RT task. However, in the vocal naming task, the participants reported the identity of the stimulus by vocally naming the stimulus into a microphone, instead of making a manual response. The stimulus remained on the screen until the vocal response had been detected. An experimenter recorded the accuracy and any spoiled trials (e.g., failure of the voice key) during the session. Each participant performed one block for each of the four stimulus categories.

Location probe task. On each trial, the participant viewed all four exemplars from a given stimulus category (e.g., all four colors), presented in random order in a row in the center of the computer screen. Each trial began with the presentation of a fixation cross for 500 msec, followed by a blank interval for 500 msec, and then the row of four stimuli for 125 msec. Immediately after the stimuli were removed from the screen a "^" probe was presented randomly just below one of the four stimulus locations and remained on the screen until a response was made. The participant was asked to make an unspeeded response indicating which of the four exemplars had been presented in that location, guessing if unsure. Responses were made using the same keys and mappings as those used in the manual RT task. Each participant performed one block of 48 trials for each of the four stimulus categories.

Single RSVP task. The participants were instructed to look for a specific target RAN element (e.g., the dog) in an RSVP stream of similar distractors and to report whether the target element was present or absent in the stream. Each trial began with a fixation cross that was presented for 500 msec and a 500-msec blank screen; then an RSVP stream of 16 items was presented one at a time in the center of the computer screen. At the end of each stream, a sentence appeared that asked whether the target was present or absent in the stream. The participants made an unspeeded manual present/absent response. The target was present in the RSVP stream on two thirds of the trials and absent on one third of the trials. When present, the target was always the 6th or the 10th item in the stream. For object and color streams, each RSVP item was presented for 33 msec and was followed by a 17-msec blank interstimulus interval (ISI). For digit and letter streams, each item was presented for 50 msec and was followed by a 17-msec blank ISI.2 Six items were used as RSVP distractors for each stimulus category, but the same distractor was never presented in two successive positions within a stream. All the distractors had the same size, color, shading, and/or font as the targets from the same category. For color trials, distractor colors included the colors purple, orange, pink, brown, olive, and plum. For digit trials, distractors were the numbers 0, 1, 3, 5, 7, and 8. For letters, the distractors were b, c, h, p, x, and y; for objects, the distractor pictures were a teddy bear, a hat, a table, a wheelbarrow, a cup, and a fan. Each participant performed one block for each of the four stimulus categories. Each block contained 48 trials, which were divided into four sections of 12 trials each, so that every 12 trials, the target changed to the next stimulus in the set (e.g., for the first 12 trials in the color block, the participants searched for the red color, in the next 12 they searched for the blue color, etc.). The participants were informed when the target changed by a sentence, which appeared onscreen telling them the identity of the target for the next 12 trials. This sentence remained on the screen until a key was pressed.

Dual RSVP task. RSVP streams used for the dual-task trials were the same as those used for the single-task RSVP trials, with the following exceptions. One of the RSVP items was singled out from the others in the stream by virtue of a unique feature that was meant to attract attention. For the color RSVP trials, one of the colored squares contained an asterisk in the center, whereas all the others did not. The participants were instructed to identify the color of the square that contained the asterisk (blue, red, or yellow) for their first target task. For digit, letter, and object trials, this first target (T1) was colored red, whereas all the other RSVP items remained black. The participants were instructed to identify the red item for their first target task (the red item could be 2, 4, or 6 for digits; g, k, or m for letters; and dog, hand, or chair for objects). After the RSVP stream, the participants were prompted to report the identity of T1 by presentation of a sentence onscreen. The participants identified T1 with an unspeeded buttonpress, using the labeled keys, and were told to guess if unsure. The second task on each trial was to report whether the fourth element on that block (i.e., the green color, the 9, the *r*, or the book, which had not been used for T1) was present or absent in the RSVP stream anytime after the T1. After making their T1 response, the participants were prompted by a sentence on the computer screen to report whether this second target (T2) was present (press "1") or absent (press "0"), using an unspeeded response. Once the first and the second responses had both been entered, the next trial began after a 1-sec blank intertrial interval. T1 was present on all trials as the sixth or eighth item in the RSVP stream. T2 was present on two thirds and absent on one third of all the trials. When present, T2 was presented equally often either two items after T1 or seven items after T1 in the RSVP stream. Each item in the color RSVP stream was presented for 66 msec, with a 17-msec blank ISI. Each item in the digit, letter, and object streams was presented for 83 msec, with a 17-msec blank interval. Each participant performed one block of 48 trials for each of the four stimulus categories.

RESULTS

Mean performance measures for the single- and dual-task RSVP trials are presented in Tables 1 and 2 separately for each task and stimulus category. Single-task RSVP target performance is presented in terms of hit rate (the proportion of target-present trials on which the participant responded "present"), false alarm rate (the proportion of target-absent trials on which the participant responded "present"), and accuracy (the proportion of hits minus the proportion of false alarms). Similarly, dual-task RSVP T2 performance is presented in terms of hit rate, false alarm rate, and accuracy (hits - false alarms) separately for lag 2 and lag 7 and collapsed across lag. AB magnitude was calculated as T2 accuracy (hits - false alarms) at lag 2 subtracted from T2 accuracy (hits - false alarms) at lag 7. T1 accuracy was calculated as the proportion of trials on which the correct identity of T1 was reported. Proportion correct was used as the measure of T1 accuracy, given that a T1 was always present and the task was a three-alternative forced choice decision. Hits minus false alarms was used for T2 in the AB task and for single-target RSVP performance, given that the target task was a present/absent decision. Table 1 shows the accuracy scores used in the subsequent analyses, and Table 2 shows the hits and false alarms used to create the accuracy scores. All the trials were used for calculations of T2 accuracy and AB magnitude (not just T1 correct trials), so that the relationships between T1 and T2 accuracy and between T1 accuracy and the AB could also be examined. Although the following analyses are presented using singletarget and T2 accuracy defined as hits minus false alarms,

Table 1
Mean Accuracy for Dual- and Single-Target Rapid Serial Visual Presentation (RSVP) Tasks
(With Standard Deviations) for Each Stimulus Type and Collapsed Across All Stimulus Types

	(
	Single	RSVP		al T1 oortion		ıl T2 Lags		al T2 g 2		al T2 Ig 7		al T2 agnitude
	0	- FA)	< I	rect)		- FA)		- FA)		- FA)		– Lag 2)
Stimuli	М	SD	М	SD	М	SD	М	SD	М	SD	M	SD
Overall	.573	.115	.953	.043	.521	.183	.367	.184	.675	.221	.308	.179
Colors	.661	.158	.929	.105	.694	.272	.619	.306	.771	.293	.151	.251
Digits	.719	.169	.961	.044	.589	.227	.426	.280	.753	.247	.327	.271
Letters	.590	.165	.958	.046	.426	.243	.252	.267	.600	.291	.348	.273
Objects	.322	.205	.963	.050	.375	.202	.172	.190	.577	.288	.405	.274

Note-H - FA, hits minus false alarms; T1, first target; T2, second target; AB, attentional blink.

			Sin	ngle			Dua	1 T 2			Dua	l T2			Dua	ıl T2
	Sir	ngle	RS	VP	Dua	l T2	Fa	lse	Dua	l T2	Fa	lse	Dua	al T2	Fa	lse
	RS	VP	Fa	lse	Н	its	Ala	rms	Н	its	Ala	rms	Н	its	Ala	rms
	Н	its	Ala	rms	All	Lags	All	Lags	La	g 2	La	g 2	La	g 7	La	g 7
Stimuli	M	SD	M	SD	M	SD	М	SD	M	SD	M	SD	M	SD	M	SD
Overall	.727	.109	.154	.094	.651	.146	.130	.100	.498	.176	.130	.099	.805	.167	.130	.099
Colors	.849	.120	.188	.123	.772	.200	.077	.111	.696	.250	.077	.111	.848	.222	.077	.111
Digits	.798	.135	.079	.098	.692	.184	.103	.127	.528	.278	.102	.127	.856	.167	.102	.127
Letters	.758	.131	.168	.155	.586	.188	.160	.166	.412	.256	.160	.166	.760	.206	.160	.166
Objects	.503	.209	.181	.174	.556	.195	.181	.163	.354	.233	.181	.163	.759	.244	.181	.163

 Table 2

 Mean Hits and False Alarms Used to Calculate Single-Target and Second Target (T2) Accuracies in Table 1

the same pattern of results was also observed when each participant's single-target and T2 RSVP performance was calculated in terms of d' or hit rate.

A significant AB was observed for each of the four stimulus types (all ps < .001 for lag 2 vs. lag 7 T2 accuracy; see Table 1 for percentage of AB magnitude for each stimulus type). For most of the subsequent analyses, scores on all the RSVP measures were averaged across the four stimulus categories (colors, digits, letters, and objects) to create a composite score for that measure (e.g., an average T2 accuracy score). As is evidenced by the results of principal components analyses, there were consistent individual differences in single-target RSVP accuracy, T1 accuracy, T2 accuracy, and AB magnitude across the stimulus types, with the exception of single-target color RSVP trials, which showed somewhat less variability. For example, with AB magnitude, a single component accounted for 45% of the variance among the four AB estimates (one for each stimulus type), and each AB measure loaded strongly on that component (loadings were .57-.76). Furthermore, the mean correlation across stimulus types (i.e., the average of the correlations for all stimulus type pairs) was .26 for AB magnitude. For T2 accuracy, the mean interstimulus correlation was .48, and a single component accounted for 61% of the variance in T2 accuracy, with strong loadings on that component for each stimulus type (.65-.86). For T1 accuracy, the mean interstimulus correlation was .34; a single component accounted for 53% of the variance in T1 accuracy, and loadings on that component ranged from .41 to .90. For single-target RSVP accuracy, the mean interstimulus correlation was .24; a single component accounted for 45% of the variance in single-target accuracy, and loadings on that component varied from .37 to .81.³ Age and gender did not predict performance on any of the task measures (all rs < .17, all ps > .18).

Associations Among RSVP Measures

Table 3 shows the pattern of correlations between singletask RSVP accuracy, T1 accuracy, overall T2 accuracy, and AB magnitude for dual-task RSVP trials, averaged across the four stimulus types. As was predicted, the participants with higher accuracy in the single-target RSVP task also showed higher T2 accuracy in the dual task. However, in contrast to the predictions, neither singletarget accuracy nor T2 accuracy predicted AB magnitude significantly. AB magnitude was predicted by dual-task T1 accuracy, where the participants who had higher T1 accuracy showed a larger AB.

A simultaneous regression was performed in which single-target accuracy, T1 accuracy, and AB magnitude were entered as predictors of T2 accuracy across lags. Combined, the three predictors explained 50% of the variability in overall T2 accuracy (R = .71, p < .001). Both single-target accuracy and T1 accuracy emerged as significant unique predictors of T2 accuracy (standardized regression coefficient $\beta = .48$, semipartial r = .41, p < .001, and $\beta = .31$, semipartial r = .25, p < .01, respectively). However, AB magnitude did not explain a significant amount of unique variability in overall T2 accuracy ($\beta = .03$, semipartial r = .02, p > .79).

A simultaneous regression was then performed in which AB magnitude was the criterion and T1 accuracy, singletarget accuracy, and T2 accuracy were entered as predictors. The model explained 14% of the variability in AB magnitude (R = .37, p < .05). Only T1 accuracy emerged as a significant unique predictor of AB magnitude ($\beta =$.36, semipartial r = .28, p < .05). Neither single-target accuracy nor T2 accuracy was a significant unique predictor of AB magnitude ($\beta = -.03$, semipartial r = -.02, p > .85, and $\beta = .05$, semipartial r = .03, p > .79, respectively). Indeed, when T1 accuracy was removed as a

Zero-Order Correlation		ible 3 ial Visual Present	tation (RSVP) Acc	uracy Scores
	T2 Accuracy (Lag 2 + Lag 7)	T1 Accuracy	Single-Task RSVP Accuracy	AB Magnitude (Lag 7 – Lag 2)
T1 accuracy Single-task RSVP accuracy AB magnitude	.58*** .66*** .23	.54*** .37**	 .19	_

Note—T2, second target; T1, first target; AB, attentional blink. **p < .01. ***p < .001.

predictor, less than 5% of the variability in AB magnitude was explained by single-target and T2 accuracies. Overall, the results suggest that the magnitude of the AB is unrelated to target detection ability (either when the target is presented in a single-task situation or when it is presented as T2 in a dual-task situation). In contrast, target accuracy in one situation (i.e., when the target is presented as a single target or as T1 or T2 on dual-target trials) predicts target accuracy in another situation.

To better understand the relationship between T1 accuracy and AB magnitude, AB magnitude was calculated using only trials on which T1 accuracy was correct (i.e., conditionalizing T2 accuracy on correct T1 accuracy, as is often done when AB data are analyzed). The AB magnitude on correct T1 trials was then correlated with T1 accuracy on all the trials. Using the conditionalized data, the correlation between T1 accuracy and AB magnitude was now eliminated (r = .05, p > .65), suggesting that T1 accuracy predicts AB magnitude only due to T1 misses that reduce the potential for an AB (i.e., T2 should not be blinked if attention was never deployed to T1).⁴

Associations Among RSVP Measures and Other Measures

Mean performance measures for the non-RSVP tasks are presented in Table 4 separately for each task and stimulus category. For the manual RT, vocal RT, and delayed RT tasks, RTs are from correct trials only. Accuracy was at least 97% on these tasks. Fewer than 4% of the RTs were removed using the Van Selst and Jolicœur (1994) outlier elimination procedure.

Table 5 shows the pattern of correlations between singletask RSVP accuracy, T1 accuracy, T2 accuracy, AB magnitude, and measures from the non-RSVP tasks. T2, T1, and single-target accuracies were related significantly to manual RTs, vocal RTs, and RAN times. T1 accuracy was also related to probe location scores. In contrast, AB magnitude was not related to performance measures from any of the non-RSVP tasks.

A simultaneous regression was performed where all five non-RSVP measures (RAN times, manual, vocal and delayed RTs, and probe location scores) were entered as predictors of T2 accuracy. The combined predictors explained a significant 40% of the variability in T2 accuracy (R = .64, p < .001). Manual RT was the only predictor

to account for significant unique variability (semipartial r = -.42, p < .001; see Table 6), since much of the variability was shared by more than one predictor. When T1 accuracy, AB magnitude, and single-target accuracy were added in a second step, the variability accounted for increased significantly to 60% (R^2 change = .20, p <.001). Manual RT, single-target accuracy, and T1 accuracy emerged as significant unique predictors (semipartial r =-.28, p < .01, semipartial r = .28, p < .01, and semipartial r = .17, p < .05, respectively; see Table 6). Note that when entered alone as predictors, the non-RSVP tasks accounted for 40% of the variability in T2 accuracy. This approximates the 50% accounted for by the other RSVP targets when they were entered alone.

When a simultaneous regression was performed in which all five non-RSVP measures were entered as predictors of AB magnitude, the combined predictors accounted for only 2% of the variability in AB magnitude (R = .13, p > .96), with no predictor accounting for significant unique variability (all ps > .37; see Table 7). When T1 accuracy, T2 accuracy, and single-target accuracy were added in a second step, the variability increased significantly to 19% (R^2 change = .17, p < .05). T1 accuracy emerged as the only significant unique predictor (semipartial r = .30, p < .05; see Table 7).

Correlations Between Measures for Individual Stimulus Types

The analyses above were performed using data collapsed across the four stimulus types (colors, digits, letters, and objects). The use of the four stimulus types for each task also gives us an opportunity to examine the interrelationships among measures for each stimulus type individually. Despite the fact that there were fewer observations for each stimulus type than across stimulus types, the pattern of relationships was markedly similar for each stimulus type and overall across stimulus types. As was shown above, overall T2 accuracy (across stimulus types) was found to be related to T1 accuracy, singletarget RSVP accuracy, manual RTs, vocal RTs, and RAN times but was unrelated to delayed RTs, probe location accuracy, and AB magnitude. Replicating this pattern, T2 accuracy was also found to be significantly related to T1 accuracy (rs = .23-.53), single-target RSVP accuracy (rs =.21–.59), manual RTs (rs = -.26 to -.42), and vocal RTs

Table 4
Mean Response Times (RTs, in Milliseconds) or Accuracy
(Proportion Correct) With Standard Deviations for Each
Non-RSVP Task and Stimulus Combination

	RAN	Times		nual Ts	Vo R		Dela R	2		obe zation
Stimuli	M	SD	M	SD	M	SD	M	SD	M	SD
Overall	25,251	3,717	642	113	454	56	305	76	.939	.034
Colors	29,731	6,464	607	113	488	71	306	89	.940	.045
Digits	19,223	3,099	544	80	414	60	304	82	.938	.058
Letters	20,074	3,181	716	171	412	52	305	78	.954	.041
Objects	31,974	4,806	700	159	504	66	304	84	.923	.061

Note-RAN, rapid automatized naming.

Weasures and Other Tasks									
Measure	Single-Target Accuracy	T2 Accuracy	T1 Accuracy	AB Magnitude					
RAN times	40***	39**	41**	.01					
Manual RT	46***	56***	31^{*}	01					
Vocal RT	49***	40^{***}	43**	05					
Delayed RT	09	18	11	03					
Location probe accuracy	.10	23	.32*	.12					

Table 5 Zero-Order Correlations Between Rapid Serial Visual Presentation Measures and Other Tasks

Note—T2, second target; T1, first target; AB, attentional blink; RAN, rapid automatized naming. *p < .05. **p < .01. ***p < .001.

(rs = -.23 to -.43) for each of the four stimulus types individually. T2 accuracy was also significantly related to RAN times for two of the four stimuli (letter and object, rs = -.30 and -.51, respectively) and was in the expected direction for digits and colors (rs = -.19 and -.14, respectively). Also, just as with the overall data, T2 accuracy was unrelated to delayed RT and probe location accuracy for any of the four stimulus types (all ps > .05). Thus, in the stimulus-specific analyses, 14 of the 16 possible correlations with T2 accuracy produced the significant effects that were also observed when the scores were collapsed across stimulus type, and 8 of 8 possible correlations with T2 accuracy produced the same null effects as those observed with the collapsed data.

Collapsed across stimulus type, AB magnitude was not related to any performance measures except T1 accuracy. This general pattern was also observed for each stimulus type. For each of the four stimulus types, AB magnitude was unrelated to manual RTs (rs = .04 to .18), delayed RTs (rs = -.17 to .18), vocal RTs (rs = -.19 to .13), or probe location accuracy (rs = .02 to .12). AB magnitude was also found to be unrelated to RAN performance except for color stimuli (r = .27), unrelated to singletarget RSVP accuracy with the exception of object stimuli (r = .36), and unrelated to T2 accuracy except for object stimuli (r = .42). AB magnitude was significantly related to T1 accuracy for color and object stimuli (rs = .35 and .26, respectively), but not for digits and letters (rs = .08and -.01, respectively). Thus, 25 of the 28 opportunities produced the same nonsignificant correlations with AB magnitude as those observed in the overall correlations collapsed across stimulus type, and two of the four stimulusspecific correlations produced the significant effects that were observed in the collapsed data. Overall, the highly similar results for each of the four stimulus types and the overall average suggests that the absence of significant relationships between AB magnitude and the other performance measures did not arise from the creation of a composite AB magnitude score averaged across stimulus types. Furthermore, the consistency in the pattern across stimulus types acts as a sort of within-experiment replication, which strengthens confidence in the overall results.

DISCUSSION

Summary

The predictors of T2 accuracy and AB magnitude in an AB task were investigated. Correlations among RSVP target performance, AB magnitude, and performance on non-RSVP tasks were examined. Our results showed the following: (1) There was no significant relationship between

Table 6 Results of Simultaneous Regression Predicting Overall T2 Accuracy							
Measure	Standardized Regression Coefficients (β)	Semipartial Correlations	n Values				
Wiedsure	Coefficients (p)	Correlations	p Values				
Step 1: Non-RSVP Predictors Onl	у						
RAN times	20	17	.09				
Vocal RT	05	04	.71				
Manual RT	48	42	< .001				
Delayed RT	.02	.02	.86				
Location probe accuracy	.17	.17	.11				
Step 2: All Predictors							
RAN times	09	07	.41				
Vocal RT	.14	.10	.24				
Manual RT	34	28	.002				
Delayed RT	07	06	.46				
Location probe accuracy	.09	.09	.31				
AB magnitude	.06	.06	.52				
T1 accuracy	.24	.17	.05				
Single-target RSVP accuracy	.38	.28	.002				

Note—RSVP, rapid serial visual presentation; RAN, rapid automatized naming; AB, attentional blink; T1, first target.

Measure	Standardized Regression Coefficients (β)	Semipartial Correlations	p Values
Step 1: Non-RSVP Predictors Or	nly		
RAN times	.05	.04	.76
Vocal RT	05	04	.77
Manual RT	.01	.01	.97
Delayed RT	.01	.01	.99
Location probe accuracy	.12	.12	.38
Step 2: All Predictors			
RAN times	.17	.14	.26
Vocal RT	.09	.06	.63
Manual RT	.14	.10	.40
Delayed RT	06	05	.68
Location probe accuracy	01	01	.97
T2 accuracy	.13	.08	.52
T1 accuracy	.41	.30	.02
Single-target RSVP accuracy	.06	.04	.75

 Table 7

 Results of Simultaneous Regression Predicting Attentional Blink Magnitude

Note—RSVP, rapid serial visual presentation; RAN, rapid automatized naming; T2, second target; T1, first target.

AB magnitude and single-target or T2 accuracy, (2) high T1 accuracy was associated with a larger AB magnitude, (3) individual differences in RSVP target accuracy (T1, T2, or single target) were predicted by performance on manual RT, vocal RT, and RAN tasks, (4) non-RSVP tasks accounted for a total of 40% of the variability in T2 accuracy, and (5) individual differences in AB magnitude were not predicted by anything except T1 accuracy, and this relationship appeared to result simply from the fact that T2 was not blinked if T1 was unattended.

The results show that the magnitude of the AB is largely unrelated to individual differences in stimulus naming, response selection, and rapid stimulus identification and consolidation. In contrast, RSVP target performance (single or dual) is related to individual differences in tasks requiring rapid identification/consolidation and response selection. The participants with shorter manual RTs and shorter naming times showed greater RSVP target accuracy, but not larger AB magnitudes. Thus, although one can predict individual RSVP target performance on the basis of these cognitive performance measures, individual AB magnitude cannot be predicted with these same measures, suggesting independence between single-target processing abilities and AB magnitude.

The finding that AB magnitude is not predicted by tasks requiring stimulus naming, response selection, and rapid stimulus identification and consolidation is, of course, a null effect, but it is not one that results from a lack of power. With the present sample size, relationships of about .25 or larger would be considered reliable, yet the correlations between AB magnitude and RAN times, vocal naming RTs, and manual RTs averaged .03, and the strongest of these relationships correlated at only .05. In contrast, the correlation between these three tasks and T2 accuracy averaged .45. The difference in the ability of these tasks to predict T2 accuracy, but not AB magnitude, is especially

provocative in light of the fact that T2 accuracy is the sum of T2 performance at lags 2 and 7, whereas AB magnitude is the difference between these same two performance measures. Thus, whereas the sum of the T2 accuracy measures across both lags is clearly related to the RAN, vocal RT, and manual RT tasks, the difference in these same measures is clearly unrelated. Furthermore, analyses indicated that the AB was a fairly stable individualdifference variable across the four stimulus types, with just under half of the variability in AB magnitudes shared by a single common factor (i.e., variability reflecting the AB per se, separated from stimulus specific variability). This common variability was similar to the levels of common variability that were observed for T1 accuracy and singletarget RSVP accuracy. The latter two variables shared substantial correlations with many performance measures, whereas the AB did not. This suggests that differences in internal consistency of the RSVP variables do not underlie the pattern of relationships across tasks. Indeed, AB magnitude did correlate with a single variable (T1 accuracy), although this relationship simply suggested that no AB was observed when T1 was not attended.

Predicting RSVP Target Accuracy

Fairly strong positive relationships were observed between single-target accuracy, T1 accuracy, and T2 accuracy. These relationships were expected, given that all three tasks required the participants to extract and consolidate identity information from RSVP streams. However, the present results are the first that we know of to show positive associations and reliable individual differences in overall target performance across RSVP tasks.

Just as novel, but more surprising, was the finding that RSVP target performance (single target, T1, and T2) was significantly predicted by vocal naming times and manual RTs. The participants with higher RSVP target accuracy scores made faster manual responses to identify single stimuli and also named stimuli more rapidly. The manual RT and vocal naming tasks are very different from RSVP tasks, because speed (not accuracy) is the dependent variable and the stimuli are presented without distractors. Indeed, performance on these non-RSVP tasks predicted RSVP performance almost as well as did performance on other RSVP tasks.

The fact that vocal RTs to single stimuli predicted RSVP target accuracies as well as or better than the time to name an entire grid of stimuli suggests that the ability to select a target from a set of distractors prior to naming is not critical to the relationship between RAN times and RSVP target performance. The RAN task requires participants to track the stimuli spatiotemporally and to select the required stimulus from among 49 distractors that have recently been named or are soon to be named. This selection among distractors is not required in the vocal naming task, where a lone stimulus is presented for naming on each trial. Both the RAN task and the vocal RT task require participants to extract the identity of the to-benamed stimulus, consolidate it into working memory so that he or she is consciously aware of its identity, and then produce the vocal response. An online vocal response is not made in the RSVP tasks, so it is likely that performance on RSVP targets and naming times are related due to individual differences in the ability to rapidly identify and/or consolidate stimulus identities.

When T2 accuracy was predicted, regression results showed that manual RT was a unique predictor over and above vocal RT and RAN times, although all three were significant predictors when examined individually. Both vocal naming times and manual RT tasks require the participant to extract the identity of the stimulus and consolidate it into working memory. However, the manual RT task also requires participants to map the stimulus identity onto the appropriate key response (response selection) and then to execute the manual RT task suggests that response selection and/or execution abilities can explain additional variability in T2 accuracy scores.

Performance on the delayed RT task was unrelated to RSVP task performance (single target, T1, or T2). The delayed RT task has response execution requirements identical to those for the manual RT task. However, the long delay between the presentations of the stimulus and the tone removes individual differences in the speed of stimulus identification, consolidation, and response selection, since even the slowest participant will have completed these processes before the tone on the vast majority of trials. This makes the delayed RT task a purer measure of response execution. Therefore, the lack of a relationship between delayed RT and RSVP target accuracies suggests that it is not the response execution speed that accounts for the relationship between manual RT and RSVP target performance but, rather, the response selection speed.

Several studies have shown that when T1 requires a speeded response, increasing the response selection requirements of the T1 task increases the magnitude of the AB (Arnell & Duncan, 2002; Jolicœur & Dell'Acqua, 1998, 1999). Jolicœur (1998, 1999) proposed a bottleneck model of the AB in which consolidation of T2 into working memory cannot proceed until consolidation of T1 has been completed. Jolicœur further suggested that the processing resources needed for stimulus consolidation in working memory are the same as those required for response selection operations. Thus, in Jolicœur's model, working memory consolidation and response selection efficiency modulate RSVP target accuracy and the magnitude of the AB. The present finding that individual differences in the ability to rapidly name items and select manual responses outside RSVP tasks predict target RSVP accuracy is consistent with Jolicœur's model. However, if one assumes that longer RTs on the manual RT task reflect longer consolidation and/or response selection times (at least in part), bottleneck models such as Jolicœur's would predict that short manual RTs would be associated with reduced ABs, and this was not observed.

Independence of AB Magnitude

Intriguingly, there was no relationship between AB magnitude and overall performance on RSVP and non-RSVP tasks. If higher overall RSVP target accuracy suggests more efficient RSVP target processing and bottleneck models of the AB are correct that faster consolidation of T1 should reduce the wait at the bottleneck, one would expect a smaller AB for those with high single-target, T1, and T2 accuracy, as well as shorter naming and manual RTs. These relations were not observed.

The only variable that predicted AB magnitude was T1 accuracy, where the participants with higher T1 accuracy rates produced larger ABs. If high T1 accuracy suggests more efficient T1 processing, bottleneck models would suggest that individuals with higher T1 accuracy should show a smaller AB, not a larger one. Indeed, manipulations of T1 difficulty have been shown to modulate the magnitude of the AB collapsed across participants, where more difficult T1 tasks produced larger ABs (e.g., Seiffert & Di Lollo, 1997). However, McLaughlin et al. (2001) and Shore et al. (2001) suggested that individual differences in T1 performance are due primarily to the difference in resources allocated to T1 by different individuals, where more resources to T1 lead to better T1 performance and, thus, worse T2 accuracy in the AB window. Therefore, this model predicts that greater T1 accuracy will be associated with greater AB magnitude, as was observed here. However, in the present study, T1 accuracy and T2 accuracy were significantly and positively associated (see Table 3), suggesting that individual ability, not a T1/T2resource trade-off, was responsible for the relationship between T1 accuracy and the AB magnitude. Also, if the positive association between T1 accuracy and AB magnitude results from some participants using more resources to process T1 at the expense of T2, as has been suggested by McLaughlin et al. (2001), the relationship between T1 accuracy and AB magnitude should become even stronger when the AB is calculated using only T1 correct trials, given that one can be sure that attentional resources were

deployed to T1 on these trials. However, the relationship between AB magnitude and T1 accuracy was eliminated when T2 accuracy was conditionalized upon correct T1 report (i.e., there was no relationship between T1 accuracy and AB magnitude when T2 accuracy was calculated using T1 correct trials only). Instead, the results suggest that T1 accuracy and AB magnitude were positively correlated simply because an AB could be observed only on trials in which T1 was properly attended (i.e., no AB should be observed when T1 was missed). Thus, participants who fail to attend to T1 more often will show lower T1 accuracy and a smaller AB, unless all T1 misses are removed from the AB calculation.

Conclusions

This study is the first to look for relationships involving individual differences on tasks that are potential correlates of AB magnitude and RSVP target accuracy. Limitations in the processing efficiency of stimulus consolidation into working memory and/or response selection operations have been put forward as explanations of the AB (e.g., Chun & Potter, 1995; Jolicœur, 1998, 1999; Jolicœur & Dell'Acqua, 1998, 1999). Interestingly, individual differences on tasks requiring these mental processes were *not* found to predict individual differences in AB magnitude. However, individual differences in these tasks *did* predict single-target RSVP accuracy, T1 accuracy, and T2 accuracy averaged across lags. This leaves an intriguing situation in which it is not clear why some participants produce a large AB and others a small AB on the same task.

The tasks tested as correlates of the AB in the present study followed directly from AB models that stress the importance of stimulus identification, consolidation, and response selection limitations (e.g., Chun & Potter, 1995; Jolicœur, 1999). It is, of course, possible, if not likely, that AB magnitude is related to other cognitive performance measures that were not tested here. For example, when fMRI scans were taken while participants performed an AB task, Marois, Chun, and Gore (2000) and Marcantoni, Lepage, Beaudoin, Bourgouin, and Richer (2003) observed involvement of the lateral frontal cortex (thought to be associated with working memory functioning; see, e.g., Owen, Evans, & Petrides, 1996) and the anterior cingulate (linked to performance monitoring; see, e.g., Carter et al., 1998). Thus, individual differences in executive control and/or working memory measures may have some predictive utility and merit further investigation.

Interestingly, Olivers and Nieuwenhuis (2005) have shown recently that the magnitude of the AB was larger when participants were told to "try hard" during the AB task than when they were instructed to distract themselves with another mental activity. Although task instructions were manipulated between participants in Olivers and Nieuwenhuis's study, so that it is not possible to know how a given individual's AB magnitude changed across instruction conditions, the results raise the interesting possibility that the AB individuals produce on any given occasion have more to do with their temporary mental set and context (a state condition) than with their informationprocessing or attentional abilities (a trait condition). In the present study, we observed fairly substantial shared variance in AB magnitude across the four stimulus types, suggesting a relatively consistent AB magnitude. However, the four AB tasks were run back to back in a single session, so transient state factors (aside from stimulus type) are not likely to have varied greatly. Interestingly, however, McLaughlin et al. (2001) observed a correlation of +.66 when correlating AB magnitude found using the typical RSVP procedures with the same participants' AB magnitude found when removing all elements from the RSVP stream except the two targets and the items that immediately trailed them. The use of the different methodology and the fact that the two AB tasks were performed approximately 4 weeks apart suggest that the AB may be a stable within-subjects individual-differences characteristic when task instructions are kept constant. The present results suggest that AB magnitude was roughly as stable across stimulus types as T1, T2, and single-target RSVP accuracy, which showed significant correlations with several of the cognitive performance measures used here. Therefore, we suggest that the stability of the AB observed here is sufficient to expect significant correlations with the other cognitive performance measures but that this expectation was not realized because AB magnitude is actually unrelated to these performance measures.

Existing models of the AB (e.g., Arnell et al., 2004; Chun & Potter, 1995; Jolicœur, 1999) suggest that the AB reflects a fundamental information-processing bottleneck on conscious stimulus identification and consolidation. Such models then implicitly assume that the AB observed for a given individual depends on the information-processing abilities of the individual, which, aside from practice effects, should be more or less consistent for a given task and set of stimuli. However, the present results show that although these abilities are good predictors of an individual's RSVP target performance, they are not good predictors of an individual's AB magnitude—a pattern of results that is not consistent with an information-processing bottleneck. Bottleneck models appear to have much explanatory power when examined with the typical mean differences approach. Using a correlational, individualdifferences approach allows one to directly test models of the AB in a way that is different from the typical methodology of group means testing. This study is a first step in that direction. Its results suggest the need for future studies examining the stability of the AB within an individual and useful individual-difference predictors of AB magnitude.

REFERENCES

- ANDERSON, A. K. (2005). Affective influences on the attentional dynamics supporting awareness. *Journal of Experimental Psychology: General*, 134, 258-281.
- ARNELL, K. M., & DUNCAN, J. (2002). Shared and separate sources of dual-task cost in stimulus identification and response selection. <u>Cog-</u> nitive Psychology, 44, 105-147.
- ARNELL, K. M., HELION, A. M., HURDELBRINK, J. A., & PASIEKA, B. (2004). Dissociating sources of dual-task interference using human electrophysiology. *Psychonomic Bulletin & Review*, **11**, 77-83.

- ARNELL, K. M., JOANISSE, M. F., KLEIN, R. M., BUSSERI, M. A., & TANNOCK, R. (2006). Decomposing RAN: The contribution of object recognition and attention to rapid naming and reading. Manuscript submitted for publication.
- ARNELL, K. M., & JOLICŒUR, P. (1999). The attentional blink across stimulus modalities: Evidence for central processing limitations. *Journal of Experimental Psychology: Human Perception & Performance*, **25**, 630-648.
- ARNELL, K. M., & LARSON, J. M. (2002). Cross-modality attentional blinks without preparatory task-set switching. *Psychonomic Bulletin* & *Review*, 9, 497-506.
- BROADBENT, D. E., & BROADBENT, M. H. P. (1987). From detection to identification: Response to multiple targets in rapid serial visual presentation. *Perception & Psychophysics*, 42, 105-113.
- CARTER, C. S., BRAVER, T. S., BARCH, D. M., BOTVINICK, M. M., NOLL, D., & COHEN, J. D. (1998). Anterior cingulate cortex, error detection, and the online monitoring of performance. *Science*, 280, 747-749.
- CHEUNG, V., CHEN, E. Y. H., CHEN, R. Y. L., WOO, M. F., & YEE, B. K. (2002). A comparison between schizophrenia patients and healthy controls on the expression of attentional blink in a rapid serial visual presentation (RSVP) paradigm. *Schizophrenia Bulletin*, 28, 443-458.
- CHUN, M. M., & POTTER, M. C. (1995). A two-stage model for multiple target detection in rapid serial visual presentation. *Journal of Experimental Psychology: Human Perception & Performance*, 21, 109-127.
- DENCKLA, M. B., & RUDEL, R. [G.] (1974). Rapid "automatized" naming of pictured objects, colors, letters and numbers by normal children. *Cortex*, 10, 186-202.
- DUNCAN, J., MARTENS, S., & WARD, R. (1997). Restricted attentional capacity within but not between sensory modalities. *Nature*, 387, 808-810.
- GIESBRECHT, B. L., & DI LOLLO, V. (1998). Beyond the attentional blink: Visual masking by item substitution. *Journal of Experimental Psychology: Human Perception & Performance*, **24**, 1454-1466.
- GRANDISON, T. D., GHIRARDELLI, T. G., & EGETH, H. E. (1997). Beyond similarity: Masking of the target is sufficient to cause the attentional blink. *Perception & Psychophysics*, **59**, 266-274.
- GREEN, C. S., & BAVELIER, D. (2003). Action video game modifies visual selective attention. *Nature*, 423, 534-537.
- HILLSTROM, A. P., SHAPIRO, K. L., & SPENCE, C. (2002). Attentional limitations in processing sequentially presented vibrotactile targets. *Perception & Psychophysics*, **64**, 1068-1082.
- HUSAIN, M., SHAPIRO, K., MARTIN, J., & KENNARD, C. (1997). Abnormal temporal dynamics of visual attention in spatial neglect patients. *Nature*, 385, 154-156.
- JOLICŒUR, P. (1998). Modulation of the attentional blink by on-line response selection: Evidence from speeded and unspeeded Task₁ decisions. *Memory & Cognition*, 26, 1014-1032.
- JOLICŒUR, P. (1999). Concurrent response selection demands modulate the attentional blink. *Journal of Experimental Psychology: Human Perception & Performance*, **25**, 1097-1113.
- JOLICEUR, P., & DELL'ACQUA, R. (1998). The demonstration of shortterm consolidation. *Cognitive Psychology*, **36**, 138-202.
- JOLICEUR, P., & DELL'ACQUA, R. (1999). Attentional and structural constraints on memory encoding. *Psychological Research*, 62, 154-164.
- KAVCIC, V., & DUFFY, C. J. (2003). Attentional dynamics and visual perception: Mechanisms of spatial disorientation in Alzheimer's disease. *Brain*, **126**, 1173-1181.
- KEIL, A., & IHSSEN, N. (2004). Identification facilitation for emotionally arousing verbs during the attentional blink. *Emotion*, 4, 23-35.
- LAHAR, C. J., ISAAK, M. I., & MCARTHUR, A. D. (2001). Age differences in the magnitude of the attentional blink. <u>Aging, Neuropsychology</u>, & Cognition, 8, 149-159.
- LI, C. R., LIN, W., CHANG, H., & HUNG, Y. (2004). A psychophysical measure of attention deficit in children with attention-deficit/ hyperactivity disorder. *Journal of Abnormal Psychology*, **113**, 228-236.
- LI, C. R., LIN, W., YANG, Y., HUANG, C., CHEN, T., & CHEN, Y. (2002). Impairment of temporal attention in patients with schizophrenia. *NeuroReport*, 13, 1427-1430.
- LUCK, S. J., VOGEL, E. K., & SHAPIRO, K. L. (1996). Word meanings can be accessed but not reported during the attentional blink. <u>*Nature*</u>, 383, 616-618.

- MACIOKAS, J. B., & CROGNALE, M. A. (2003). Cognitive and attentional changes with age: Evidence from attentional blink deficits. *Experimental Aging Research*, 29, 137-153.
- MARCANTONI, W. S., LEPAGE, M., BEAUDOIN, G., BOURGOUIN, P., & RICHER, F. (2003). Neural correlates of dual task interference in rapid visual streams: An fMRI study. *Brain & Cognition*, 53, 318-321.
- MAROIS, R., CHUN, M. M., & GORE, J. (2000). Neural correlates of the attentional blink. *Neuron*, 28, 299-308.
- MCLAUGHLIN, E. N., SHORE, D. I., & KLEIN, R. M. (2001). The attentional blink is immune to masking induced data limits. *Quarterly Journal of Experimental Psychology*, 54A, 169-196.
- MONDOR, T. A. (1998). A transient processing deficit following selection of an auditory target. *Psychonomic Bulletin & Review*, **5**, 305-311.
- OLIVERS, C. N. L., & NIEUWENHUIS, S. (2005). The beneficial effect of concurrent task-irrelevant mental activity on temporal attention. *Psychological Science*, **16**, 265-269.
- OWEN, A. M., EVANS, A. C., & PETRIDES, M. (1996). Evidence for a two-stage model of spatial working memory processing within the lateral frontal cortex: A positron emission tomography study. *Cerebral Cortex*, 6, 31-38.
- POTTER, M. C., CHUN, M. M., BANKS, B. S., & MUCKENHOUPT, M. (1998). Two attentional deficits in serial target search: The visual attentional blink and an amodal task-switch operation. *Journal of Experimental Psychology: Learning, Memory, & Cognition*, 24, 979-992.
- RAYMOND, J. E., SHAPIRO, K. L., & ARNELL, K. M. (1992). Temporary suppression of visual processing in an RSVP task: An attentional blink? *Journal of Experimental Psychology: Human Perception & Performance*, 18, 849-860.
- ROKKE, P. D., ARNELL, K. M., KOCH, M. D., & ANDREWS, J. T. (2002). Dual-task attention deficits in dysphoric mood. *Journal of Abnormal Psychology*, **111**, 370-379.
- SCHNEIDER, W., ESCHMAN, A., & ZUCCOLOTTO, A. (2002). *E-Prime* user's guide. Pittsburgh: Psychology Software Tools.
- SEIFFERT, A. E., & DI LOLLO, V. (1997). Low-level masking and the attentional blink. *Journal of Experimental Psychology: Human Perception & Performance*, 23, 1061-1073.
- SHAPIRO, K. L., ARNELL, K. M., & RAYMOND, J. E. (1997). The attentional blink. *Trends in Cognitive Sciences*, 1, 291-296.
- SHAPIRO, K. L., CALDWELL, J., & SORENSEN, R. E. (1997). Personal names and the attentional blink: A visual "cocktail party" effect. *Journal of Experimental Psychology: Human Perception & Performance*, 23, 504-514.
- SHAPIRO, K. L., DRIVER, J., WARD, R., & SORENSEN, R. E. (1997). Priming from the attentional blink: A failure to extract visual tokens but not visual types. *Psychological Science*, 8, 95-100.
- SHAPIRO, K. L., RAYMOND, J. E., & ARNELL, K. M. (1994). Attention to visual pattern information produces the attentional blink in rapid serial visual presentation. *Journal of Experimental Psychology: Human Perception & Performance*, 20, 357-371.
- SHORE, D. I., MCLAUGHLIN, E. N., & KLEIN, R. M. (2001). Modulation of the attentional blink by differential resource allocation. *Canadian Journal of Experimental Psychology*, 55, 318-324.
- SOTO-FARACO, S., & SPENCE, C. (2002). Modality-specific auditory and visual temporal processing deficits. *Quarterly Journal of Experimen*tal Psychology, 55A, 23-40.
- SOTO-FARACO, S., SPENCE, C., FAIRBANK, K., KINGSTONE, A., HILL-STROM, A. P., & SHAPIRO, K. (2002). A crossmodal attentional blink between vision and touch. *Psychonomic Bulletin & Review*, 9, 731-738.
- VAN SELST, M., & JOLICEUR, P. (1994). A solution to the effect of sample size on outlier elimination. *Quarterly Journal of Experimental Psychology*, 47A, 631-650.
- VOGEL, E. K., LUCK, S. J., & SHAPIRO, K. L. (1998). Electrophysiological evidence for a postperceptual locus of suppression during the attentional blink. *Journal of Experimental Psychology: Human Perception & Performance*, 24, 1656-1674.
- WARD, R., DUNCAN, J., & SHAPIRO, K. [L.] (1997). Effects of similarity, difficulty, and nontarget presentation on the time course of visual attention. *Perception & Psychophysics*, **59**, 593-600.

NOTES

1. It is possible that target accuracy depends not only on how easily the participant can process the target, but also on the resources that the participant invests in processing that target (see McLaughlin et al., 2001). To the extent that higher target accuracy is the result of greater resource investment, one may, instead, expect a positive relationship between T1 accuracy and AB magnitude and a positive relationship between single-target accuracy and AB magnitude.

2. Pilot testing was used for both single- and dual-task RSVP programs to achieve presentation durations that resulted in approximately 70% detection accuracy for target present/absent judgments.

3. Although luminance and contrast were equivalent for all letters, digits, and objects in the present experiment (all were presented in black), luminance and contrast differed element to element in the color RSVP streams. The additional luminance and contrast information may have been used more or less effectively by at least some participants, thus differentiating color RSVP scores from the letter, digit, and object scores.

4. When conditionalized AB magnitude scores were used as the criterion in all of the regression analyses, instead of unconditionalized AB magnitude scores, the pattern of results remained the same, except that T1 accuracy was no longer a significant predictor. When all non-RSVP tasks were used as predictors of conditionalized AB magnitude, they explained just over 8% of the variability in AB magnitude (p > .63), with no significant unique predictors. When T1, T2, and single-target accuracy were added, just over 11% of the variability in AB magnitude was explained, again with no significant unique predictors. The overall AB magnitude was 30.4% (68.3% - 37.9%) when conditionalizing on T1 accuracy.

> (Manuscript received September 27, 2004; revision accepted for publication July 19, 2005.)