

TUTORIAL REVIEWS

The attentional blink: A review of data and theory

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Under conditions of rapid serial visual presentation, subjects display a reduced ability to report the second of two targets (Target 2; T2) in a stream of distractors if it appears within 200–500 msec of Target 1 (T1). This effect, known as the attentional blink (AB), has been central in characterizing the limits of humans' ability to consciously perceive stimuli distributed across time. Here, we review theoretical accounts of the AB and examine how they explain key findings in the literature. We conclude that the AB arises from attentional demands of T1 for selection, working memory encoding, episodic registration, and response selection, which prevents this high-level central resource from being applied to T2 at short T1–T2 lags. T1 processing also transiently impairs the redeployment of these attentional resources to subsequent targets and the inhibition of distractors that appear in close temporal proximity to T2. Although these findings are consistent with a multifactorial account of the AB, they can also be largely explained by assuming that the activation of these multiple processes depends on a common capacity-limited attentional process for selecting behaviorally relevant events presented among temporally distributed distractors. Thus, at its core, the attentional blink may ultimately reveal the temporal limits of the deployment of selective attention.

Our visual environment constantly changes across the dimensions of both time and space. Within the first few hundred milliseconds of viewing a scene, the visual system is bombarded with much more sensory information than it is able to process up to awareness. To overcome this limitation, humans are equipped with filters at a number of different levels of information processing. For example, high-resolution vision is restricted to the fovea, with acuity drastically reduced at the periphery. Such front-end mechanisms reduce the initial input; however, they still leave the visual system with an overwhelming amount of information to analyze. To meet this challenge, the human attentional system prioritizes salient stimuli (targets) that are to undergo extended processing and discards stimuli that are less relevant for behavior after only limited analysis (Broadbent, 1958; Bundesen, 1990; Desimone & Duncan, 1995; Duncan, 1980; Kahneman, 1973; Neisser, 1967; Pashler, 1998; Shiffrin & Schneider, 1977; Treisman, 1969).

Given the vital role attention plays in visual cognition, it is not surprising that over the last 50 years, understanding the nature of the mechanisms involved in visual attention has been one of the major goals of both cognitive science and neuroscience (Miller, 2003). For the most part, this research has focused on understanding how humans process information distributed across space (see,

e.g., Pashler, 1998, for an extensive review). However, in the last 15 years, there has been intense interest among researchers in the mechanisms and processes involved in deploying attention across time (see Shapiro, Arnell, & Raymond, 1997, for an earlier review).

Here, we review research on temporal attention, specifically focusing on arguably the most widely studied effect in the field, the *attentional blink* (AB; Raymond, Shapiro, & Arnell, 1992). We begin by briefly discussing *rapid serial visual presentation* (RSVP; Potter & Levy, 1969), which is the paradigm primarily used to study the AB, and then describe the AB effect and theoretical accounts, both informal and formal, that have been put forward to explain the phenomenon. Following this, we examine how key findings in the literature fit with each model and conclude by highlighting the mechanisms that are most likely responsible for the AB. Research exclusively investigating the neural substrates of the AB will not be discussed here, since this has recently been summarized elsewhere (see Hommel et al., 2006; Marois & Ivanoff, 2005).

Early Investigations Into the Temporal Limits of Attention

In RSVP (Potter & Levy, 1969; see Figure 1A), stimuli appear sequentially at the same spatial location, for a frac-

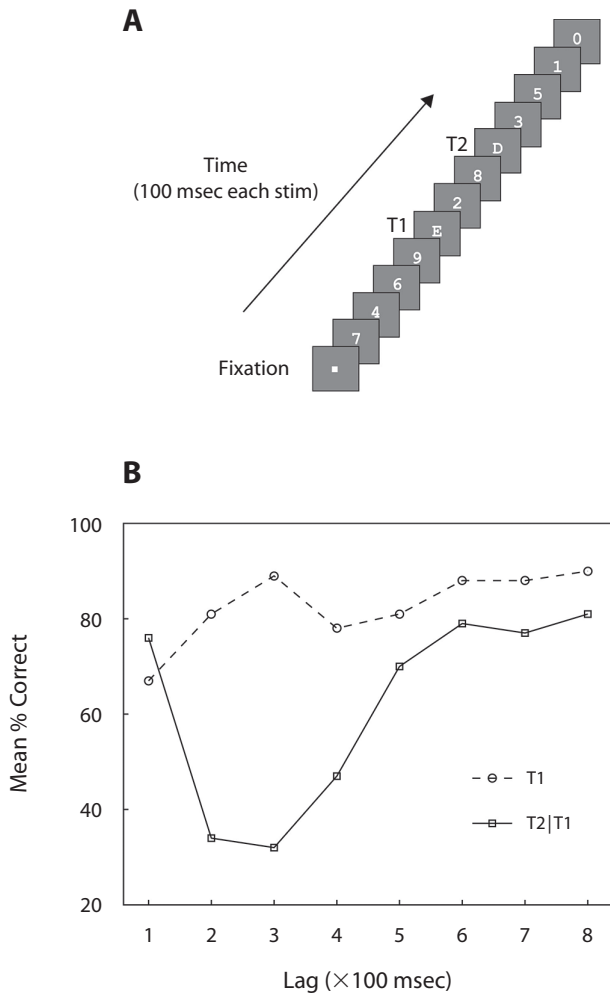


Figure 1. Standard attentional blink (AB) task and results. (A) Graphical depiction of a lag 3 trial (on which a large AB deficit is typically observed) in a standard AB task. In this task, subjects are required to search for two letter targets (Target 1, T1; Target 2, T2) among digit distractors and report them at the end of the stream. Typically, both T1 serial position and T1–T2 lag are varied across trials. Each gray frame represents the presentation of a stimulus for 100 msec. (B) Characteristic T1 and T2|T1 (T2 report given T1 correct) report accuracy as a function of T1–T2 lag. The AB corresponds to the impaired T2|T1 performance observed at lags 2–5, relative to lags 6–8 (or relative to T1 performance), whereas lag 1 sparing reflects the high T2|T1 accuracy at lag 1 (relative to lags 2–5). From “A Two-Stage Model for Multiple Target Detection in Rapid Serial Visual Presentation,” by M. M. Chun and M. C. Potter, 1995, *Journal of Experimental Psychology: Human Perception and Performance*, 21, p. 112, Table 1. Copyright 1995 by the American Psychological Association. Adapted with permission.

tion of a second each (e.g., 100 msec), and subjects are typically required either to report all the items presented (full report) or to report prespecified target item(s) and ignore the remaining distractor stimuli (partial report). The basic rationale behind RSVP paradigms is that, by stressing the temporal processing mechanisms to their limit, researchers are able to assess the rate at which information is analyzed and encoded (Chun & Wolfe, 2001; Coltheart, 1999).

A striking characteristic of temporal attention is that, even with RSVP rates of up to approximately 16 items/sec, the selection and encoding of a single target is quite easy. Lawrence (1971) found that, at this presentation rate, target report accuracy was approximately 70% with RSVP streams of words that contained a single target defined either featurally (uppercase letters) or categorically (animal word among nonanimal words). Similarly, in a seminal article, Potter (1975; see also Potter & Faulconer, 1975; Thorpe, Fize, & Marlot, 1996) demonstrated that, when subjects searched RSVP streams of scenes (8 items/sec), accuracy was comparable whether they looked for a particular stimulus that they had seen previously or one that had simply been described to them.

The results discussed above could be taken to suggest that target processing in RSVP is complete after only 100 msec. However, it can be shown that this is not the case when an additional target is added to an RSVP stream. In fact, at presentation rates of approximately 100 msec/item, subjects show a remarkable deficit in reporting the second (T2) of two different targets presented among distractors if it appears within approximately 200–500 msec of the first target (T1; Broadbent & Broadbent, 1987; Raymond et al., 1992). This effect is the AB (Figure 1B) and is an important discovery, since it helps characterize the limits of our ability to consciously perceive stimuli that are distributed across time (Sergent & Dehaene, 2004).

The Discovery of the Attentional Blink

Broadbent and Broadbent (1987) were the first to report an AB when they presented subjects with RSVP streams of words containing two targets defined by either category or letter case. On trials on which T1 was reported correctly, T2 performance was impaired if it appeared up to half a second after T1. Broadbent and Broadbent explained this result by proposing that although early perceptual features were extracted in parallel from RSVP streams, at short temporal intervals target identification processes interfered with one another, thereby resulting in the T2 deficit.

A similar posttarget processing deficit was found by Weichselgartner and Sperling (1987). In one of their experiments, subjects were presented with RSVP streams of digits at the rate of 100 msec/item, and their task was to name an outlined or bright digit (T1) and then the three stimuli that directly followed it. Subjects typically reported T1, the subsequent item, and then the stimulus that appeared 400 msec after the target. Weichselgartner and Sperling (see also Reeves & Sperling, 1986) took this pattern of results as evidence for the existence of two partially overlapping attentional processes: a fast-acting automatic process responsible for detecting (identifying) T1, as well as the item that directly followed it, and a slow effortful process that led to the recall of stimuli presented later in the stream.

Raymond et al. (1992), who first coined the term *attentional blink*, provided a crucial extension to the earlier work by demonstrating that the previously observed target-processing deficit was an attentional, rather than a

sensory, limitation. In their experiment, RSVP streams of black letter stimuli were presented at the rate of 100 msec/item, and the subjects were required to name a single white target letter (T1) and detect the presence/absence of the letter "X" as T2. Raymond et al. (1992) found that on trials on which T1 was reported correctly, T2 performance was impaired if it appeared within half a second of T1. Crucially, detection of T2 strongly improved when subjects ignored T1. This finding demonstrated that the effect was due to attentional, rather than sensory, limitations, since the same visual stimuli yielded different effects depending on task requirements. Two other important characteristics of the AB were also revealed in the Raymond et al. (1992) study. Whereas T2 accuracy was impaired if it appeared within 200–600 msec of T1 (and T1 required report), there was virtually no deficit when T2 was presented directly after T1, an effect now known as *lag 1 sparing* (see Figure 1; Potter, Chun, Banks, & Muckenhoupt, 1998; Visser, Bischof, & Di Lollo, 1999). In addition, T2 performance was strongly improved when T1 was followed by a blank gap in the RSVP stream, rather than by a distractor, suggesting that the stimulus following T1 plays a vital role in generating the AB. Before turning to a discussion of these and other key findings in the literature, we first will review theories of the AB and lag 1 sparing.

Theoretical Accounts of the Attentional Blink

Several theoretical accounts have been introduced to explain the AB. For the most part, theories of the phenomenon have been informal, with researchers simply describing the processes that underlie the effect. However, recently, a number of computational frameworks have explicitly modeled the T2 deficit and other relevant findings. In this section, we will begin by describing informal theories of the AB and then computational frameworks that formalize many of the ideas put forward in these purely descriptive accounts.

Informal Theories

Inhibition model. Raymond et al. (1992) proposed that the AB was the result of a suppressive mechanism that inhibited posttarget stimuli in order to reduce target and distractor featural confusion. This *gating theory* predicts that, when a dual-target RSVP stream is viewed, an attentional episode is triggered after the physical features (e.g., color, shape) of T1 are detected. The initiation of this attentional episode is likened to a gate opening to admit T1 for the purpose of identification. When an item follows T1, its features will also be processed along with those of T1, thus increasing the chance that the features of the two stimuli will be confused. For example, the color of T1 may be incorrectly bound to the identity of the subsequent stimulus. In order to reduce interference from posttarget stimuli and increase the probability that T1 will be correctly reported, the stimuli following T1 are suppressed at an early perceptual level. Raymond et al. (1992) likened this inhibitory process to the gate's closing. This model assumes that this attentional gate stays closed until identification (e.g., color and identity bound together) of T1 is complete, a process that Raymond et al.

(1992) hypothesized took approximately 500 msec. Thus, the AB arises because T2 is inhibited when it appears in close temporal proximity to T1. When T2 appears after T1 has been identified, the gate will no longer be closed, and as a result, T2 can be the subject of focused attention. Lag 1 sparing is said to occur when T1 is followed directly by T2, because both stimuli are admitted by the gate and undergo identification processes together. Furthermore, lag 1 sparing is deemed to be dependent on T1 and T2's not being separated by a distractor, rather than on the two target stimuli's appearing within 100 msec of one another.

Recently, Olivers and his colleagues (Olivers, van der Stigchel, & Hulleman, 2007; see also Olivers & Meeter, 2008) have revised and extended Raymond et al.'s (1992) inhibition hypothesis. They suggest that inhibition is not initiated to prevent color binding errors between T1 and T1 + 1 but, rather, to suppress distractors so that they do not interfere with target processing. This inhibition takes place at a relatively late stage of visual information processing, after the conceptual representations of the RSVP stimuli have been activated (but prior to working memory). In this revised inhibition framework, the T1 + 1 distractor is processed along with T1 because its temporal proximity to T1 confers on it the same attentional enhancement (boosting) that T1 receives. To prevent additional distractors from receiving this attentional boost and interfering with T1 processing, post-T1 + 1 stimuli are strongly suppressed, thereby impairing T2 performance at short T1–T2 lags. This model will be discussed in more detail in the Formal Theories section (the boost and bounce theory).

Interference theory. In Raymond et al.'s (1992) gating theory, it is the potential for featural confusion during T1 identification that leads to the AB. Subsequently, Shapiro, Raymond, and Arnell (1994) obtained results challenging the conclusion that the identification of T1 was necessary to elicit the T2 deficit, since they found that it also occurred when T1 only required detection. Consequently, Shapiro et al. (1994; see also Isaak, Shapiro, & Martin, 1999; Shapiro & Raymond, 1994) proposed the *interference theory* to account for their findings.

On the basis of Duncan and Humphreys's (1989) model of spatial visual search, Shapiro et al.'s (1994) interference theory assumes that, when an RSVP stream is viewed, initial perceptual representations are established for each stimulus. These representations are compared with selection templates (generated from the task instructions), and those stimuli that most closely match are selected and registered in visual working memory. Once in this store, each item is assigned a weighting based on the available space and its similarity to the templates. Typically, both targets, as well as the items that directly succeed each of them (T1 + 1 and T2 + 1), enter working memory, due to their temporal proximity to the targets. In working memory, items interfere with one another as retrieval processes are undertaken during report of the targets. In this model, an AB occurs when the targets are separated by a short interval, because T2 receives a diminished weighting due to the limited capacity of working memory, leaving it more open to interference from the other items in the store and,

therefore, reducing the likelihood of its being reported. Shapiro et al. (1994; see also Isaak et al., 1999; Shapiro & Raymond, 1994) suggested that an AB is not observed at long T1–T2 lags because visual working memory “may be flushed after sufficient time has passed with no demand made on it” (p. 371; although this aspect of the theory appears to be difficult to reconcile with the fact that both targets require report in the standard AB task at the end of the RSVP stream and, therefore, would both need to be maintained even at long lags). This framework suggests that lag 1 sparing occurs because stimulus interference is reduced when T2 appears directly after T1, since only three items enter visual short-term memory: T1, T2, and T2+1. Thus, according to this model, lag 1 sparing is determined by the characteristics of the T1+1 stimulus, rather than by the temporal gap between T1 and T2.

Bottleneck models. Chun and Potter (1995) presented a number of important findings for understanding the mechanisms responsible for the AB. First, they provided evidence that was inconsistent with Raymond et al.’s (1992) gating theory, since they observed a significant AB when the targets were defined categorically (searching for two black letter targets among black digit distractors) rather than perceptually (red target among black distractors), thereby demonstrating that the deficit can still arise even when there is no potential for a feature conjunction error between the color of T1 and the identity of the T1+1 stimulus. In addition, this result demonstrated that the AB was not the result of a task switch between the two targets, since both letters required identification and were drawn from the same stimulus set. Finally, by revealing how the AB was modulated by the extent to which the targets and distractors were both featurally and categorically similar, Chun and Potter’s (1995) study also highlighted the influence of target–distractor discriminability on this deficit (see also Dux & Coltheart, 2005; Maki, Bussard, Lopez, & Digby, 2003).

To account for their findings, Chun and Potter (1995) proposed a two-stage model of the AB. In Stage 1, a stimulus activates its stored conceptual representation. Recognition at this stage occurs rapidly, so the specific identities of most items in an RSVP stream are available (Potter, 1975, 1976, 1993), but this information is volatile and susceptible to both decay and overwriting by subsequent stimuli. Consistent with this notion, Giesbrecht and Di Lollo (1998; see also Dell’Acqua, Pascali, Jolicoeur, & Sessa, 2003; Giesbrecht, Bischof, & Kingstone, 2003) demonstrated that an AB occurs only if T2 is backward masked. To avoid being overwritten, stimuli must undergo the capacity-limited Stage 2 processing, during which they are encoded/consolidated into working memory. Stage 2 processing is initiated when relevant target features are identified in Stage 1, triggering a transient attentional response that leads to the target’s being encoded in working memory. The model explains the AB as being due to the severe capacity limitations of this second stage of processing. Consequently, when T2 is presented in close temporal proximity to T1, it must wait to be encoded into working memory until Stage 2 processing of T1 is completed and, therefore, is more susceptible to decay and interruption by distractors. Lag 1 sparing is said to occur when T2 appears

directly after T1 because, due to the slow temporal dynamics of the attentional system, T2 receives the same enhancement and access to Stage 2 processing as T1. Thus, this model predicts that lag 1 sparing is chiefly determined by the temporal distance between T1 and T2. There is now considerable neuroimaging and electrophysiological support for a two-stage framework, since several studies have demonstrated that whereas visual areas respond to both missed and reported T2s, parietal–frontal regions selectively respond to reported T2s (see Gross et al., 2004; Kranczioch, Debener, Schwarzbach, Goebel, & Engel, 2005; Marois, Yi, & Chun, 2004).

A similar bottleneck model was proposed by Ward, Duncan, and Shapiro (1996; see also Duncan, Ward, & Shapiro, 1994) to account for the T2 deficit they observed with a modified AB task. Ward et al. (1996) investigated the speed with which attention could be shifted to targets when these items were distributed across both time and space. In their experimental conditions, only the two targets, followed by their respective masks, were presented on distinct corners of an invisible diamond. By varying the *stimulus onset asynchrony* (SOA) between the two targets, Ward et al. (1996) demonstrated that report of T2 was impaired, relative to a control single-target condition, if it appeared within approximately 450 msec of T1. Ward et al. (1996) proposed the *attentional dwell time* hypothesis to account for their results and those of the standard AB. This hypothesis asserts that the two target objects compete for capacity-limited visual processing resources, with the winner of this competition undergoing extended processing at the expense of the loser. Because of its head start in the competition, T1 is typically the winner, thereby leaving T2 open to interference from the mask and, therefore, increasing the likelihood that it will go undetected. Since lag 1 sparing was not found by Ward et al. (1996), they did not incorporate an explanation of this effect into their theory. Indeed, it should be noted that in a detailed meta-analysis, Visser et al. (1999) concluded that lag 1 sparing is rarely observed under conditions in which the target stimuli are spatially displaced or there is a multi-dimensional attentional set switch between T1 and T2 (e.g., T1 is a categorically defined letter, whereas T2 is a color-defined digit; see also Potter et al., 1998).

Jolicoeur (1998, 1999; Jolicoeur & Dell’Acqua, 1998; see also Ruthruff & Pashler, 2001) extended the two-stage account of Chun and Potter (1995) to explain not only the AB, but also its relationship to the psychological refractory period (PRP; see Pashler, 1994). The PRP refers to subjects’ tendency to respond more slowly to the second of two sensory–motor tasks as the SOA between them is reduced. This Task 2 postponement is thought to result from a central capacity-limited stage that prevents two response selection processes from being performed concurrently (Pashler, 1994). Jolicoeur (1998) investigated the extent to which interference during response selection influenced the magnitude of the AB. In his experiments, a methodology similar to that employed by Raymond et al. (1992) was used, except that T1 required an immediate, rather than a delayed, response for some of the trials (T2 response was always offline). The inclusion of this speeded T1 task en-

sured that subjects performed response selection to Task 1 online, thereby creating at short lags a processing overlap between T1 response selection and T2 working memory encoding. Jolicœur's (1998) results revealed that a larger AB was elicited in speeded T1 trials, relative to unspeeded trials. Furthermore, the magnitude of the deficit increased as the T1 reaction times and number of T1 response alternatives increased. These findings provided clear evidence that response selection to Task 1 significantly exacerbates the AB. To account for these findings, Jolicœur (1998, 1999; Jolicœur & Dell'Acqua, 1998; see also Ruthruff & Pashler, 2001) proposed the *central interference theory*. This model is similar to Chun and Potter's (1995) theory, with the key difference being that in Jolicœur's (1998, 1999; Jolicœur & Dell'Acqua, 1998) framework, both response selection and working memory encoding require capacity-limited central processing.

Potter, Staub, and O'Connor (2002) proposed a further extension to the two-stage account (Chun & Potter, 1995), challenging the hypothesis that T1 gained privileged access to capacity-limited processing resources due to its temporal position. It had previously been found that at lag 1, performance for T2 was typically superior to that for T1 and that the report order of these two targets was often reversed (Chun & Potter, 1995; see also Hommel & Akyürek, 2005), hinting that T1 may not always be the first item to enter the bottleneck. To test this hypothesis, Potter et al. (2002; Bachmann & Hommuk, 2005) presented a word target in each of two concurrent, spatially displaced RSVP streams of symbol distractors (one stream above the other) to reduce the temporal lag between the targets without altering stimulus duration. The results demonstrated that when the targets were separated by 13–53 msec, report of T2 was superior to that of T1, which is a pattern of results opposite to that typically found in AB experiments. By contrast, at an SOA of 100 msec, performance was comparable for T1 and T2 (this is an example of lag 1 sparing occurring with spatially displaced targets), and at an SOA of 213 msec, the standard T2 deficit emerged. The superior report of T2 at very short SOAs indicated that T1 is not always consolidated before T2.

To account for their findings, Potter et al. (2002) proposed the two-stage competition model of visual attention. This theory postulates that targets compete in Stage 1 to gain access to the capacity-limited second stage of processing, with the first target that is initially identified entering the Stage 2 bottleneck first. The model explains the T1 deficit at very short lags (13–53 msec) as follows. When T1 is detected, an attentional window is opened, and the processes involved in initial identification begin. Crucially, when T2 appears very shortly after T1, it benefits from the attentional window's having already been opened and accrues resources faster than would have been the case if the attentional window had not been opened after T1 detection. As a result, T2 is identified more efficiently and enters the bottleneck before T1 (although the model is not specific about how resources would accrue faster for T2 than for T1 at these short SOAs). By contrast, at the presentation rates typically used in RSVP tasks—approximately 100 msec per item—T1 will have already

been identified and gained access to Stage 2 before T2 arrives, therefore resulting in a T2 deficit. According to this account, lag 1 sparing is dependent on T1 and T2's having an SOA of approximately 100 msec because, under these conditions, both the T1 and T2 Stage 1 representations are stable enough that attention (Stage 2 processing) can process both items without cost.

Although Chun and Potter's (1995) original hypothesis that the AB results primarily from a central bottleneck of information processing has been incorporated into several recent models of the AB, there is considerable debate regarding the number and location (along the information-processing pathways) of such bottlenecks. For one, Awh et al. (2004) challenged the hypothesis that the depletion of a capacity-limited *central* processing resource was responsible for the AB. These researchers suggested that rather than reflecting the competition for a single visual-processing channel (stage/resource), the AB arises from capacity-limited processing in a multitude of processing channels. Although they observed an AB when a face target temporally preceded a letter/digit target, no such T2 cost was observed when the order of target presentation was reversed (letter/digit first, then face). The data were explained by the hypothesis that face recognition engages both featural and configural information processing, thereby transiently preventing the featural channel from processing any subsequent letter/digit stimuli, whereas letter/digit identification relies only on the featural channel, thereby allowing the configural channel to process any subsequently presented face stimuli. However, Awh et al.'s findings of multiple bottlenecks have recently been questioned by Landau and Bentin (2008; see also Jackson & Raymond, 2006), with these researchers suggesting that Awh et al.'s results reflect the salience of face stimuli rather than the existence of multiple bottlenecks. Moreover, and as was mentioned earlier, the finding that drawing on the central processing stage of response selection affects the AB (Jolicœur, 1998, 1999; Jolicœur & Dell'Acqua, 1998; Ruthruff & Pashler, 2001) suggests that the deficit involves a central amodal bottleneck of information processing. This is further reinforced by studies in which an AB was observed even when the two target stimuli originated from different modalities (auditory vs. visual; e.g., Arnell & Duncan, 2002; Arnell & Jenkins, 2004; Arnell & Jolicœur, 1999; Arnell & Larson, 2002; but see Chun & Potter, 2001; Duncan, Martens, & Ward, 1997; Potter et al., 1998), although it has been difficult to rule out a task-switching account of this cross-modal attention deficit (Chun & Potter, 2001; Potter et al., 1998). Similarly, it is still unsettled as to whether this central amodal stage of information processing wholly encompasses the AB bottleneck or whether this deficit arises from processing limitations at both this amodal stage and at an earlier visual stage of information processing (Chun & Potter, 2001; Jolicœur, Dell'Acqua, & Crebolder, 2001; Ruthruff & Pashler, 2001).

A final extension to Chun and Potter's (1995; see also Potter et al., 2002) bottleneck theory is that offered up by Dux and Harris (2007a), who tested whether the encoding bottleneck also limited distractor inhibition. Dux and Harris (2007a; see also Drew & Shapiro, 2006) presented

subjects with RSVP streams similar to those employed by Chun and Potter (1995), with black letter targets appearing among black digit distractors. The crucial manipulation was that on half the trials, the items directly preceding ($T1 - 1$) and succeeding ($T1 + 1$) $T1$ were either identical to or different from one another. Dux and Harris (2007a) reasoned that if target selection involves distractor inhibition, repeating the distractors on either side of $T1$ would reduce the masking strength of the $T1 + 1$ distractor, since its representation would have been suppressed by the earlier presentation of the same character. This suppression of the $T1 + 1$ distractor would, in turn, improve $T1$ accuracy and, therefore, reduce the AB. This is indeed what Dux and Harris (2007a) observed, suggesting that distractor inhibition plays a key role in RSVP target selection (see also Dux, Coltheart, & Harris, 2006; Dux & Marois, 2008; Maki & Padmanabhan, 1994; Olivers & Watson, 2006). Importantly, in a subsequent experiment, in which they now repeated the distractors on either side of $T2$ rather than of $T1$, Dux and Harris (2007a) found that distractor repetition did not benefit $T2$ report when the $T2 - 1$ distractor was presented during the blink. These data suggest that distractor suppression is impaired by the AB bottleneck because it does not take place unless the distractor receives attention. Consistent with this notion, Dux and Harris (2007a) did observe distractor suppression when the $T2 - 1$ distractor was presented at lag 1, a position that favors attentional processing of that distractor along with $T1$ (see also Dux & Marois, 2008).

It should be noted that Drew and Shapiro (2006) also found that the AB was attenuated when $T1$ was temporally flanked by repeat distractors. However, these authors proposed an account different from that of Dux and his colleagues (Dux et al., 2006; Dux & Harris, 2007a), suggesting that this effect was caused by the same mechanism(s) as those responsible for repetition blindness (RB; see Kanwisher, 1987). RB refers to subjects' impaired ability to report both occurrences of a repeat stimulus in RSVP if they appear within 500 msec of one another and is typically thought to result from subjects' failure to register both repeat targets as episodically distinct objects, rather than from inhibition of $T2$ (see, e.g., Kanwisher, 1987). Dux et al. (2006) suggested that the distractor repetition effect and RB reflect different mechanisms, because the former does not occur between letter stimuli that differ only in case, whereas the latter is found under these conditions as well. In addition, Dux and Harris's (2007a) finding that the distractor repetition effect taps the same mechanism as the AB suggests that it differs from RB, since the processes underlying the AB and RB have been doubly dissociated (Chun, 1997b; see also Dux & Marois, 2007). Finally, Dux and Marois's (2008) distractor priming effects, described below, are convergent evidence for the inhibition of distractors in RSVP. Nevertheless, further research is required to isolate the mechanisms that give rise to the distractor repetition effect found in RSVP and to understand how it relates to RB.

Temporary loss of control hypothesis. All of the models discussed above, with the exception of the inhibition account (Olivers et al., 2007; Raymond et al., 1992),

are consistent with the notion that the AB results from the depletion of capacity-limited attentional resources by $T1$ processing, leaving too few of these resources available at short lags to be applied to $T2$. Employing an innovative paradigm, Di Lollo, Kawahara, Ghorashi, and Enns (2005; see also Olivers et al., 2007) provided data that posed a challenge for such $T1$ capacity-limited models. They presented subjects with RSVP streams that contained three successive targets (all of which required delayed report), with the third target ($T3$) appearing in a position where the blink is typically maximal—lag 2. When the targets were members of the same category (e.g., letters), there was no deficit in reporting $T3$ (Olivers et al., 2007, refer to this effect as “spreading of the [lag 1] sparing”), a result that is inconsistent with $T1$ resource depletion accounts of the AB. However, impaired $T3$ report was observed if $T2$ belonged to a category different from that of the other target stimuli (e.g., a computer symbol as opposed to letters).

Di Lollo et al. (2005) proposed the temporary loss of control (TLC) hypothesis to explain their results. This theory postulates that RSVP processing is governed by a filter configured to select targets and exclude distractors, which is endogenously controlled by a central processor that can execute only a single operation at a time (as was acknowledged by Di Lollo et al., 2005, this feature of the model adds a capacity-limited component to the framework). When a target is initially identified, the central processor switches from monitoring to consolidation processes, and the input filter is then under exogenous control (but see Nieuwenstein, 2006, for evidence that endogenous control is maintained during the AB). When $T2$ is drawn from the same category as $T1$, the input filter's configuration is unaltered, and as a result, this target is processed efficiently. If, however, $T2$ is drawn from a different category, it takes longer to process and is more susceptible to interruption from distractors. More important, the input filter's configuration is disrupted and needs to be reconfigured, resulting in all subsequent stimuli being processed less efficiently until this reconfiguration takes place. Di Lollo et al. suggested that both these sources of disruption contribute to the manifestation of the AB in this three-target paradigm. In addition, the same disruption-of-input-configuration account is invoked to explain the AB in a typical two-target RSVP paradigm, except that disruption here is triggered by the $T1 + 1$ distractor, instead of by a categorically distinct target. According to the TLC model, lag 1 sparing is observed with the sequential presentation of intracategory targets because such presentation does not disrupt the input filter. Thus, like the inhibition (Olivers et al., 2007; Raymond et al., 1992) and interference (Shapiro et al., 1994) theories, but contrary to bottleneck models (e.g., Chun & Potter, 1995), the TLC account predicts that lag 1 sparing is dependent on the nature of the $T1 + 1$ stimulus, rather than on $T1$ and $T2$'s appearing within 100 msec of one another.

Delayed attentional reengagement account. The delayed attentional reengagement account introduced by Nieuwenstein and his colleagues (Nieuwenstein, 2006; Nieuwenstein, Chun, van der Lubbe, & Hooge, 2005; Nieuwenstein & Potter, 2006; Nieuwenstein, Potter,

& Theeuwes, 2009) suggests that the AB reflects the dynamics of attentional selection. When a dual-target RSVP stream is viewed, the presentation of T1 elicits the deployment of top-down attentional resources to that stimulus. However, once target information ceases to be presented (due to the appearance of either a distractor or a blank gap in the RSVP stream), these resources are disengaged from the RSVP stream. In this model, the AB occurs because subjects are unable to rapidly reengage top-down attention to T2 shortly after it has been disengaged from the T1 stimulus. Lag 1 sparing results from attention's being sustained for T2 processing because T1 is followed by additional goal-relevant target information (T2) rather than by irrelevant information (distractor/blank gap). This model makes no specific prediction regarding the cause of lag 1 sparing—whether it is determined by the nature of the T1 + 1 stimulus or the temporal distance between T1 and T2—because, in this framework, both the duration of the T1 enhancement and the nature of the post-T1 information influence RSVP performance.

It follows from this theory that experimental conditions that help reengage or sustain attention after T1 processing should diminish the AB. Consistent with this prediction, Nieuwenstein et al. (2005; see also Maki, Frigen, & Paulson, 1997) demonstrated that if T2 is immediately preceded by a distractor that shares featural characteristics with that target—a manipulation that should help reengage attention prior to T2 presentation—the AB is reduced. Furthermore, the AB was virtually abolished when the task required report of all the RSVP stimuli, an experimental condition that is presumed to continuously engage attention throughout the RSVP stream (Nieuwenstein & Potter, 2006).

Hybrid models. A number of researchers have suggested that a combination of the mechanisms described above provides the most complete account of the AB. Vogel, Luck, and Shapiro (1998; see also Sergent, Baillet, & Dehaene, 2005; Vogel & Luck, 2002) demonstrated, using event-related potentials (ERPs), that T2 did not elicit a P300—a component believed to reflect the updating of working memory (Donchin, 1981; Donchin & Coles, 1988)—during the AB, suggesting that missed T2s do not enter the working memory store. To explain their results, Vogel et al. (1998; see also Maki, Couture, Frigen, & Lien, 1997; Maki, Frigen, & Paulson, 1997; Shapiro, Arnell, & Raymond, 1997) proposed a model that incorporated aspects from both the two-stage framework (Chun & Potter, 1995) and the interference theory (Shapiro & Raymond, 1994; Shapiro et al., 1994). This account suggests the existence of two processing stages, with stimuli first being conceptually processed before being selected to undergo capacity-limited encoding into visual working memory. Whether stimuli are selected for extended processing after their semantic representations are activated is determined by how closely they match target templates. Furthermore, due to interference between stimuli during preliminary conceptual processing, distractor items that appear in close temporal proximity to T2 are often incorrectly consolidated. Thus, both a bottleneck in working memory and

interference between the conceptual representations of stimuli are hypothesized to give rise to the AB.

More recently, Kawahara, Enns, and Di Lollo (2006) have also suggested that a combination of independent mechanisms contributes to the failure to identify T2 under RSVP conditions. Specifically, they proposed that a combination of Di Lollo et al.'s (2005) TLC hypothesis and bottleneck models provides the best account of the AB. They predicted that three factors determine target accuracy under RSVP conditions. Specifically, (1) switching from rejecting distractors to selecting targets when presented with T1 affects T1 performance, whereas both (2) the disruption of an input filter once T1 encoding has commenced and (3) an encoding bottleneck that delays T2 processing and leaves it susceptible to backward masking at short lags (due to online T1 processing) affect T2 performance.

Formal Theories

Gated auto-associator model. Chartier, Cousineau, and Charbonneau (2004) presented a framework to account for the AB observed when subjects searched a stream of green digit distractors for two red digit targets. This model predicts that from an input layer, stimuli are evaluated via two networks, with one performing number identification and feeding this information into an auto-associator (working memory), and another comparing the color of each stimulus with the target color specified by task instructions. Stimuli are admitted to and maintained in the auto-associator—as well as selected for report—on the basis of their weighting. An item's weighting is, in turn, determined by the extent to which an attentional gate is open when the stimulus is represented at the identification layer (i.e., perceptually presented). If the gate is open when the stimulus is identified, it will have a high weighting and be more likely to be admitted to the auto-associator and reported. This gating mechanism opens when the color comparison process recognizes that a stimulus is a target. However, this gating process is inhibited and, thus, becomes less efficient when another stimulus is being encoded into working memory. It is this inhibitory process, together with its slow recovery rate, that leads to T2's being weakly weighted at short T1–T2 lags, thereby giving rise to the AB. In this model, lag 1 sparing occurs because the time that the gate remains open exceeds the presentation duration of T1.

Corollary discharge of attention movement model. This model of the AB is an application of Taylor and Rogers's (2002) influential theory of attention control: the corollary discharge of attention movement (CODAM) model. Fragopanagos, Kockelkoren, and Taylor (2005) suggested that RSVP processing proceeds in the following manner: Initially, stimuli pass through input and object map modules before they reach a working memory module, where they become consciously available. Crucial to this model's account of the AB is the role played by an inverse model controller (IMC), which attentionally boosts items in the object map in order for them to be admitted into working memory. The AB occurs because this attentional boost is

withheld from T2 at short lags to prevent it from interfering with the encoding of T1. A monitor module inhibits the IMC until T1 processing is complete, which is determined by the monitor's continually comparing the target representation with a predictor of the current stimulus, which is computed via the current attentional control signal (corollary discharge) from the IMC. When T2 appears in close temporal proximity to T1, the IMC will be suppressed, because the target representation will be that for T1, whereas the corollary discharge will represent T2. As a result, T2 will not be attentionally enhanced and, therefore, will not enter working memory. At longer lags, both the target representation and corollary discharge will be those for T2, and hence, no T2 deficit will be observed. In this framework, lag 1 sparing results from the temporal dynamics of the IMC inhibition, which does not onset until after the T1+1 stimulus has been presented. Consequently, it too will be attentionally enhanced and enter working memory.

Locus coeruleus–norepinephrine model. Nieuwenhuis, Gilzenrat, Holmes, and Cohen (2005) suggested that the AB reflects the activation dynamics of the locus coeruleus (LC). The LC is a brain-stem nucleus that is thought to contain up to half of all noradrenergic neurons in the central nervous system (Berridge & Waterhouse, 2003). It projects widely to many areas of the cortex, and some have suggested that its innervation particularly influences regions involved in attentional processing (Nieuwenhuis, Aston-Jones, & Cohen, 2005). With respect to attentional tasks, it has been hypothesized that the presentation of a salient stimulus triggers LC neurons, causing the release of norepinephrine in brain areas innervated by the LC, thereby enhancing the responsivity of these areas to their input. This LC response is phasic, and the duration of norepinephrine modulation effects is also fleeting, lasting less than approximately 200 msec. After this initial firing, the LC enters into a refractory period where it does not respond to subsequent salient stimuli for approximately 500 msec. In Nieuwenhuis, Gilzenrat, et al.'s (2005) AB model, there are two major components: the LC and a behavior network that is made up of input, detection, and decision layers. On detection of a salient stimulus, the LC transiently adjusts the gain of the behavioral network by simulating the release of norepinephrine. Following this phasic response, the LC is suppressed and unavailable to enhance subsequent target processing, thus causing the AB. Importantly, the magnitude of this post-T1 suppression, and, thus, of the AB, is tied to the size of the LC phasic response to T1, giving this framework a capacity-limited component. As is the case with many of the models described above, lag 1 sparing results due to the temporal dynamics of the attentional enhancement, rather than to the nature of the T1+1 stimulus.

The global workspace model. The global workspace model is an influential theory of conscious perception and attentional control (Baars, 1989; Dehaene, Sergent, & Changeux, 2003). This framework has many similarities to the bottleneck theories described above and predicts that for individuals to become aware of a specific stimulus, the item must enter a global neuronal workspace. This

workspace activates neurons with long-distance axons capable of connecting distinct brain regions, such as those responsible for higher level processing and those involved in initial sensory analysis. Once a stimulus has activated a sufficient set of workspace neurons, activity becomes self-sustained, and this item can then be employed by a variety of neural areas via the long-distance connections. However, once activated by a stimulus, these workspace neurons inhibit neighboring workspace neurons, thus making them unavailable for subsequently presented stimuli. When two targets are presented in close temporal proximity, each of them goes through an initial sensory stage of information processing by distinct neuronal assemblies (*feed-forward sweep*) that do not inhibit one another. The AB results when these two neural assemblies compete for access to the global workspace (*top-down amplification*), with the winner's activity becoming self-sustained and triggering consciousness. Recovery from the AB occurs at later lags once the T1 brain-scale state has subsided, allowing the global workspace to become available for T2. Lag 1 sparing occurs due to the delayed onset of the workspace interneuronal inhibition, thereby allowing both T1 and T2 to enter consciousness. Thus, in this model, lag 1 sparing is governed by time rather than by the T1+1 item.

Boost and bounce theory. In the boost and bounce theory (Olivers & Meeter, 2008), capacity limits play no role in the generation of the AB. This model has two major stages: sensory processing and working memory. During sensory processing, both the perceptual features of a stimulus, such as its shape, color, and orientation, and its high-level representations, including semantic and categorical information, are activated. As stimuli are presented at the same spatial location in RSVP, each item's activation strength (during sensory processing) is influenced by those stimuli that appear around it, due to forward and backward masking. Working memory plays several roles in the model. First, it maintains task instructions, establishing an attentional set. Second, it stores encoded representations, where items to be reported have been linked to a response. Finally, and most important, working memory employs an input filter that enhances the processing of stimuli that match the target set and inhibits stimuli that do not (i.e., distractors). Specifically, the input filter inhibits the distractors presented before T1, thereby preventing them from gaining access to working memory, and attentionally enhances T1, which can therefore gain access to this store. Because of its temporal proximity to T1 and the dynamics of the enhancement, the T1+1 distractor also receives a strong attentional *boost* despite the fact that it is a distractor from a different stimulus set than the target. The attentional enhancement of a distractor stimulus—an item that does not require report—triggers strong but transient suppression (the *bounce*) of subsequently presented stimuli by the input filter to prevent the T1+1 distractor from entering working memory, thus causing the AB. According to this model, lag 1 sparing occurs due to the duration of the initial T1 boost, and this sparing extends to lags 2 and 3 of a continuous stream of targets (i.e., spreading of the sparing; Di Lollo et al.,

2005; Olivers et al., 2007) because no inhibitory signal is elicited when the T1+1 (and T1+2) stimulus originates from the same target set as does T1.

Episodic simultaneous type/serial token model. Bowman and Wyble (2007) presented a sophisticated theory of temporal attention and working memory known as the *simultaneous type/serial token* (STST) model and recently extended it to become the *episodic STST* framework (eSTST; Wyble, Bowman, & Nieuwenstein, 2009). This model borrows heavily from Chun and Potter's (1995; see also Chun, 1997b; Kanwisher, 1987) two-stage theory and suggests that the AB reflects processes involved in episodically distinguishing objects from one another. The eSTST account predicts that all the stimuli in the RSVP are identified at a conceptual stage (i.e., have their *type* representation activated). However, for these stimuli to be reported, they must have this identity information bound to a *token* in working memory that provides episodic information about the stimulus (e.g., the position of the item in the RSVP stream, relative to other stimuli). For a type representation to be bound to a token, it must be attentionally enhanced by a blaster that is transiently triggered when a target is detected in the RSVP stream, and it is the slow temporal dynamics of this blast that give rise to lag 1 sparing. However, because the binding process for a target is capacity limited at the stage of episodic registration (i.e., two stimuli cannot be encoded in the correct temporal position at the same time) and is susceptible to interference from other targets, the blaster is suppressed until T1 is linked to its specific token and consolidated into working memory. This T1-triggered blaster suppression prevents the T2 type from being bound to a token, thereby triggering an AB. Thus, the eSTST theory conceptualizes the AB as arising due to a kind of unconscious perceptual strategy that helps subjects overcome their capacity limits in episodic registration by suppressing the processing of future targets until T1 is episodically registered.

An important aspect of this theory, and a key difference between the STST and eSTST models, is that due to the interaction of excitatory and suppressive processes, the blaster is maintained in an enhanced mode when target items appear in an uninterrupted sequence. As a result, the identity encoding of several successive targets can be successfully performed, thereby giving rise to spreading of the sparing. However, the model further predicts that, under successive target conditions, there should be a high proportion of target report order reversals (it also explains other episodic errors, such as RB; Kanwisher, 1987), and T1 accuracy should be reduced because of increased competition between T1 and T2. Both of these predictions, which are not explicitly modeled by other AB frameworks, have been experimentally confirmed (Bowman & Wyble, 2007; see also Chun & Potter, 1995).

The attention cascade model. The formal models described above are all connectionist frameworks that involve many parameters and complex interactions between layers of artificial neurons to model the AB. Recently, Shih (2008) has put forward a mathematical model—the attention cascade theory—that has fewer parameters, yet provides a detailed account of the system that performs

RSVP processing (this model, however, does not make any predictions regarding the neural processes involved in the AB). Like the eSTST theory, this framework borrows heavily from the two-stage account of Chun and Potter (1995); however, it also incorporates characteristics of Shapiro et al.'s (1994; Shapiro & Raymond, 1994) interference theory. Specifically, the attention cascade model predicts that stimuli are initially processed along one of two channels: a mandatory pathway or a bottom-up salience pathway. Stimuli processed by the mandatory pathway activate conceptual long-term memory representations that are then passed into a *peripheral* sensory buffer. If a representation in this buffer matches the target template, it will trigger an attentional window and be enhanced. Following this enhancement, if there are enough encoding resources available, the target's representation will undergo encoding/consolidation where its strength will grow further, leading to its being passed into a decision processor (within working memory). Stimuli with strong bottom-up salience can also trigger the attentional window and enter directly into the decision processor (bottom-up salience pathway). Indeed, it is this component of the model that can account for the finding that salient distractors that share features with the target set can also trigger an AB for a subsequent target (Folk, Leber, & Egeth, 2002; Maki & Mebane, 2006). In Shih's framework, the AB occurs because the encoding/consolidation processor is capacity limited, causing a T2 appearing in close temporal proximity to T1 to wait for this encoding resource to become available, thereby leaving that T2 susceptible to interference and decay. Lag 1 sparing is said to result from the duration of the attentional enhancement, which extends beyond the presentation time of T1. Importantly, because the duration of this attentional enhancement window may vary according to task demands, it could encompass the successive presentations of several targets in an RSVP, allowing all of them to be reported successfully. Thus, Shih's model can also account for the spreading of the sparing results observed in three-target RSVP tasks. However, it should be noted that given that this model's account of the spreading of the sparing is dependent on task strategy, it is somewhat difficult to see how the framework explains this result under conditions in which uniform and varied trials are randomly intermixed, instead of being blocked (e.g., Olivers et al., 2007).

The threaded cognition model. In a recent computational model, Taatgen, Juvina, Schipper, Borst, and Martens (2009) have proposed that the AB reflects a protective production rule that prevents T2 from interfering with T1 consolidation. Borrowing from J. R. Anderson's (2007) adaptive control of thought—rational (ACT—R) architecture, the threaded cognition framework conceptualizes cognition as multiple processes that are threaded through a single processor (a single resource). With respect to dual-target RSVP search, this model predicts that target detection and consolidation can operate in parallel. However, due to default task allocation policies, target detection is held offline during the encoding of another target into working memory. Thus, at short T1—T2 lags, subjects have impaired T2 performance because they adopt an implicit

strategy to suppress detection of this T2 until consolidation is complete. The model accounts for lag 1 sparing and spreading of that sparing by assuming that the system recognizes that targets appearing directly after T1 require report and that this supersedes the control production rule that protects consolidation. As a result, detection is not suppressed for these target stimuli. It should be noted that although this model appears compatible with the item-based account of lag 1 sparing, it also has a temporal component, since the application of the production rule and consolidation rate are independent of the stimulus presentation rate. On the surface, this model bears strong similarity to the eSTST framework (both suggest that T2 detection is strategically suppressed during T1 processing). However, a key difference between the theories is that in the threaded cognition account, there are no capacity limitations; the AB merely reflects the application of an unnecessary protective rule. Indeed, Martens, Munneke, Smid, and Johnson (2006) have identified a group of subjects that apparently are immune to the AB deficit, and Taatgen et al. suggested that these individuals do not apply the production rule under RSVP conditions.

Comparison of Models of the Attentional Blink

Consideration of the theories described above indicates that virtually all of them provide adequate accounts of both the AB and lag 1 sparing. In addition, many of these models include mechanistically comparable stages of RSVP processing. Indeed, each theory predicts that at least one or more of the following processes lead to the AB.

1. Perceptual inhibition of post-T1 stimuli due to the potential for T1 and T1+1 featural confusion (gating theory).
2. Sustained postperceptual suppression triggered by the T1+1 distractor (boost and bounce theory).
3. Online T1 attentional depletion due to T1 working memory encoding/episodic registration/response selection (bottleneck theories, hybrid models, global workspace model, gated auto-associator model, CODAM, LCNE, attention cascade theory, eSTST¹).
4. Competition between target and distractor stimuli during offline retrieval from working memory (interference theory).
5. Disruption of an input filter by the T1+1 distractor (TLC, Kawahara, Enns, & Di Lollo's [2006] hybrid model).
6. Suppressed/delayed attentional enhancement of T2 (delayed attentional engagement, LCNE, CODAM, attention cascade theory, eSTST, threaded cognition model).

In this section, we will review key empirical results and how they fit with the mechanisms described above. Our summary and concluding remarks highlight the models that provide the most detailed account of the AB and associated findings.

Perceptual inhibition versus postperceptual selection in RSVP. As was previously discussed, there is little evidence to support Raymond et al.'s (1992) gating hypothesis that it is the potential for featural confusion between T1 and T1+1 that gives rise to the AB by triggering the perceptual inhibition of post-T1 stimuli. Shapiro

et al.'s (1994) finding that detection of T1 is sufficient to produce an AB and Chun and Potter's (1995) result that categorically defined targets also elicit the T2 deficit are inconsistent with this framework, since gating theory predicts that it is the potential for feature conjunction errors between the T1 and the T1+1 item that leads to the AB. It should also be noted that these results cannot be explained by the gated auto-associator model (Chartier et al., 2004), since this framework was designed only to explain ABs observed when both targets are defined by color. However, it should also be noted that the gated auto-associator model is not alone in terms of being limited to a particular task. Due to their specificity, many of the formal models described are able to account for the AB only under a particular set of conditions. For example, Wyble et al.'s (2009) eSTST framework models RSVP performance only when targets are defined categorically (e.g., two-digit targets among letter distractors).

Further evidence against gating theory and its prediction that the AB has a perceptual locus is that stimuli that are not reported from RSVP streams nevertheless undergo semantic/conceptual processing. Luck, Vogel, and Shapiro (1996) examined the extent to which missed T2s were processed in the AB, using ERPs, and observed an N400—a component associated with semantic processing—for a T2 in an AB task even when that target failed to be reported. Similarly, Shapiro, Driver, Ward, and Sorensen (1997) found in a three-target RSVP search that a missed T2 could conceptually prime report of a subsequent target. In addition, Marois, Yi, and Chun (2004) have demonstrated, using functional magnetic resonance imaging (fMRI), that distractor manipulations activate high-level visual areas in the brain (see also Kranczioch et al., 2005; Sergent et al., 2005). And it is not only missed T2s that undergo semantic analysis, but also distractors: Maki, Frigen, and Paulson (1997; see also Chua, Goh, & Hon, 2001) presented subjects with RSVP streams of words and found that a distractor that appeared in close temporal proximity to T2 could semantically prime that target's report. Thus, there is good evidence that in standard RSVP tasks, non-reported stimuli undergo considerable processing.

It should be noted, however, that under some conditions, initial processing of RSVP stimuli does display capacity limits. Like Luck et al. (1996), Giesbrecht, Sy, and Elliott (2007) examined the N400 for missed T2s in an RSVP and found that whereas an N400 was observed for T2 when T1 involved a low perceptual load (T1 spatially flanked by congruent distractors), it was completely suppressed for trials on which T1 involved a high perceptual load (T1 flanked by incongruent distractors). Similarly, in a series of studies, Dell'Acqua, Jolicœur, Robitaille, and Sessa (Dell'Acqua, Sessa, Jolicœur, & Robitaille, 2006; Jolicœur, Sessa, Dell'Acqua, & Robitaille, 2006a, 2006b) have found that the N2pc, a presemantic ERP component thought to be associated with visuospatial shifts of attention, is also influenced by the AB. In addition, Williams, Visser, Cunnington, and Mattingley (2008) have shown with fMRI that activity in the primary visual cortex is also sensitive to the AB. No current theory of the AB can fully account for these results. However, they can be ac-

commodated by assuming that in such nonstandard AB tasks—in which there are either spatially displaced target or distractor stimuli (as was the case in the studies described above)—the attentional resources devoted to target processing (see Lavie, 1995) can lead to missed/nonreported RSVP stimuli's being processed only up to early perceptual levels. These exceptions aside, the bulk of the evidence reviewed here does suggest that, at least for standard AB tasks, missed stimuli are processed postperceptually.

Postperceptual inhibition. The preceding section suggests that the perceptual inhibition of post-T1 stimuli due to the potential for a feature conjunction error between T1 and the T1 + 1 stimulus is not responsible for the AB. However, can gating theory be saved by assuming that such inhibition takes place at a postperceptual level of information processing and that it is elicited by the T1 + 1 distractor due to its interfering with T1 encoding (boost and bounce model; Olivers & Meeter, 2008)? A key prediction of the boost and bounce theory is that it is inhibition of post-T1 stimuli that gives rise to the AB. However, this hypothesis is inconsistent with the results of an individual-differences analysis of the AB (Dux & Marois, 2008; see also Martens et al., 2006) that suggested that subjects who inhibit post-T1 distractors actually exhibit enhanced T2 performance at short lags (attenuated ABs). These results not only are opposite to those predicted by post-T1 suppression accounts of the AB (Olivers & Meeter, 2008; Raymond et al., 1992); they also provide support for the hypothesis that a failure of distractor inhibition contributes to the AB (see Dux & Harris, 2007a).

Online T1 attentional depletion. The evidence that missed targets and distractors in RSVP undergo considerable processing is consistent with all of the AB models that conceptualize this deficit as a postperceptual phenomenon. Similarly, Dux and Marois's (2008; see also Dux & Harris, 2007a) individual-differences study is problematic only for models that predict that sustained suppression of all stimuli post T1 + 1, triggered by a post-T1 distractor, gives rise to the AB (e.g., Olivers & Meeter, 2008; Raymond et al., 1992). A point of greater conjecture among models of the AB, however, is whether the deficit is contingent on online T1 processing (i.e., T1 working memory encoding/episodic registration/response selection; see bottleneck theories, hybrid models, global workspace model, gated auto-associator model, CODAM, LCNE, attention cascade theory, eSTST, threaded cognition model) or whether it results from mechanisms that are independent of, and/or subsequent to, T1 encoding/episodic registration/response selection (see delayed attentional engagement, interference theory, TLC, the boost and bounce models). Given that several AB theories make very different predictions regarding the influence of T1 processing on T2 performance, numerous studies have examined the effect of T1 manipulations on the magnitude of the AB.²

Jolicœur's (1998, 1999) finding that an increase in the number of T1 response alternatives leads to larger AB magnitude supports the hypothesis that the extent to which subjects process T1 directly influences T2 performance and, thus, causes an AB. Further support for this hypoth-

esis has come from Ouimet and Jolicœur (2007; see also Akyürek, Hommel, & Jolicœur, 2007; Colzato, Spapé, Pannebakker, & Hommel, 2007), who have demonstrated that the AB is also larger when T1 working memory encoding load is increased. In addition, several studies have shown that increasing the masking strength of the T1 + 1 item (either perceptually or conceptually) and, thus, increasing the time required to process T1 lead to a larger T2 deficit (Dux & Coltheart, 2005; Grandison, Ghirardelli, & Egeth, 1997; McAuliffe & Knowlton, 2000; Raymond, Shapiro, & Arnell, 1995; Seiffert & Di Lollo, 1997; Shore, McLaughlin, & Klein, 2001; see also Marois, Chun, & Gore, 2004). Finally, an inverse correlation between T1 performance and AB magnitude has also been observed at the individual subject level (Dux & Marois, 2008; Martens et al., 2006; Seiffert & Di Lollo, 1997).

However, not all T1 manipulations have been shown to affect the magnitude of the AB. Ward, Duncan, and Shapiro (1997), for example, demonstrated that the AB was unaffected by whether subjects had to make "easy" or "difficult" size discriminations for T1. Similarly, Shapiro et al. (1994) reported no difference in AB magnitude between conditions in which subjects had simply to detect T1 and those in which they had to identify it. Furthermore, McLaughlin, Shore, and Klein (2001) found that manipulating the perceptual quality of the T1 stimulus by covarying the exposure duration of T1 and the T1 + 1 mask did not influence the size of the AB.

The mixed evidence that T1 manipulations influence the AB has led some researchers to postulate that T1 processing and the AB are not related. According to this view, previous results suggesting a link between T1 and AB magnitude can be accounted for by processes that are unrelated or subsequent to online T1 working memory encoding, episodic registration, or response selection, such as task switching between T1 and T2, post-T1 stimulus suppression, offline target retrieval, and attentional filter disruption (e.g., Potter et al., 1998; see also Enns, Visser, Kawahara, & Di Lollo, 2001; Kawahara, Zuvic, Enns, & Di Lollo, 2003; McLaughlin et al., 2001; Olivers & Meeter, 2008). In addition, it could be argued that some T1 manipulations, such as those employed in Jolicœur's speeded AB studies (Jolicœur, 1998, 1999), made the experimental task so different from standard AB paradigms that mechanisms related to the blink were no longer tapped. However, a number of studies have shown T1 effects on the AB without a task switch between T1 and T2, while holding T1 + 1 masking strength constant and requiring a delayed response to both targets. For example, in RSVP streams containing word targets, increasing the difficulty (and presumably, the duration) of T1 encoding by making this stimulus disyllabic instead of monosyllabic—a manipulation known to increase working memory difficulty for words (Baddeley, Thomson, & Buchanan, 1975; see also Coltheart & Langdon, 1998; Coltheart, Mondy, Dux, & Stephenson, 2004)—decreased T2 performance at short T1–T2 lags, but not at long T1–T2 lags (Olson, Chun, & Anderson, 2001). This suggests that increasing the phonological length of T1 delays T2's admittance to the encoding bottleneck. Similarly, Dux and Harris

(2007b) presented subjects with 2-D line drawings of objects and manipulated T1's in-plane orientation, which previous studies (e.g., Jolicœur, 1985) have demonstrated influences the time required to recognize objects. They observed that the magnitude of the AB was increased when T1 was presented at a 90° rotation (an orientation where objects take longer to name, relative to upright and upside-down objects; see Jolicœur, 1985), providing more evidence that online T1 processing influences the AB.

If T1 processing affects the AB, why did some of the studies fail to show an effect of this variable on the magnitude of this deficit? One potential reason for this discrepancy in the literature is that outlined by Olson et al. (2001; see also Visser, 2007, for a similar point of view), who claimed "that the blink is sensitive to only those variables that immediately affect attentional processing between initial sensory registration of a target and consolidation into working memory. The difficulty of tasks that can be performed on representations in working memory, after consolidation has been completed, do[es] not affect T2 performance" (p. 1117). In addition, Visser has suggested that strong masking manipulations of T1 may cut short T1 processing, thereby reducing its influence on the processing of subsequent targets. It is therefore possible that previous experiments that failed to show effects of T1 manipulations on the AB did so because they did not tap a stage of processing that occurred prior to or at the bottleneck. This point also raises an intrinsic limitation of the AB paradigm, which is that because it relies on accuracy, rather than reaction time, as a measure of performance, it is difficult to temporally pinpoint the different stages of processing that take place during that task and to identify which of these stages are the loci of interference in dual-target paradigms. Nevertheless, the discussed T1 effects are problematic for models that posit that it is the T1 + 1 stimulus that elicits the AB, rather than T1 processing (e.g., delayed attentional engagement, the boost and bounce model), since these hypotheses ascribe a limited role for a T1 bottleneck in the generation of the AB and, thus, would not predict that delaying T2's access to the bottleneck (through a T1 difficulty manipulation) would increase the magnitude of the T2 deficit (the exception is the threaded cognition model, which, despite its assumption of unlimited T1 processing capacity, predicts that T2 detection is suppressed by an unnecessary production rule until T1 processing is complete). Similarly, the T1 manipulation results are not easily reconcilable with Shapiro et al.'s (1994) interference theory, for although this account suggests that capacity-limited T1 processing underlies the AB, the limitation it proposes to be responsible for the deficit comes at the information-processing stage of memory retrieval, which takes place offline from the RSVP.

Lag 1 sparing: Time or T1 + 1 dependent? The discussion of T1 versus T1 + 1 processing also applies to lag 1 sparing, since there is a theoretical dispute as to whether the effect is determined by T1 and T2's having an SOA of 100 msec (e.g., Bowman & Wyble, 2007; Chun & Potter, 1995) or by the categorical identity of the T1 + 1 stimulus (e.g., Di Lollo et al., 2005). Bowman and Wyble (see also Nieuwenhuis, Gilzenrat, et al., 2005; Potter et al., 2002)

examined this question by reducing the stimulus exposure duration of the RSVP stimuli in a standard AB task (two letter targets among digits) to 54 msec. This manipulation created, at lag 2, a condition that pitted the lag at which AB is normally maximal against the time when lag 1 sparing is usually obtained (108 msec). Under these conditions, sparing was observed at lag 2, for, at this lag, T2 performance was comparable to that for both T1 (cited in Olivers & Meeter, 2008) and T2 performance at post-AB lags and was significantly greater than T2 performance at AB lags (post lag 2). Thus, Bowman and Wyble concluded that lag 1 sparing is time dependent.

Recently, however, Martin and Shapiro (2008) suggested that lag 1 sparing is determined by the nature of the T1 + 1 stimulus, rather than by the temporal distance between T1 and T2. In one of their experiments, subjects searched for two white letter targets among black letter distractors, with each stimulus appearing for 17 msec, followed by a temporal gap of 85 msec (102 msec per item RSVP rate). In two lag 1 (SOA of 100 msec) experimental conditions, a black digit was inserted 34 and 64 msec after T1 onset in the temporal gap between T1 and T2 (the T1–T2 SOA was maintained at 102 msec in these two conditions). These trials were compared with a control condition in which no distractor appeared in the temporal gap between the targets. Martin and Shapiro found that T2 performance at lag 1 was superior in the control condition, relative to the experimental conditions, and concluded that lag 1 sparing is determined by the nature of the stimulus that follows T1, rather than by the time that elapses between the two targets. However, it should be noted that even with the presence of the inserted distractor between the two targets at lag 1 (SOA, 100 msec), performance here did not significantly differ for T1 and T2, and T2 performance at lag 1 was always superior to that at lag 3 (the point where the AB was maximal). Thus, it is questionable whether lag 1 sparing really was absent in the distractor conditions, suggesting that lag 1 sparing is determined primarily by T1 and T2's having an SOA of approximately 100 msec.

Input filters and the T1 + 1 distractor. As was discussed above, the evidence that T1 processing plays an important role in generating the AB is problematic for theories that predict that the AB is independent of T1 encoding/episodic registration/response selection. However, if online capacity-limited T1 processing contributes to the AB, why did Di Lollo et al. (2005; see also Kawahara, Kumada, & Di Lollo, 2006; Olivers et al., 2007) find that three consecutive RSVP targets can be recalled equally well if they are members of the same stimulus category (uniform condition), but not if the middle target is a member of a different category (varied condition)? As the target number is increased for uniform trials, relative to the standard AB task, online T1 processing accounts predict that an AB will be observed under these conditions as well. Thus, spreading of the sparing appears problematic for AB models—namely, the bottleneck, hybrid, global workspace, gated auto-associator, CODAM, and LCNE models—that predict that online T1 capacity limitations underlie the AB.³

Dux, Asplund, and Marois (2008, 2009) have recently disputed the claim that the AB deficit is abolished under

uniform conditions. Specifically, they pointed out that although the studies of Di Lollo et al. (2005), Olivers et al. (2007), and Kawahara, Kumada, and Di Lollo (2006) showed that T3 and T1 performance did not differ in uniform trials, these authors also found that T3 accuracy increased and T1 performance decreased in uniform trials, relative to the varied trials, suggestive of a trade-off between T1 and T3 in the uniform condition. Such a T1–T3 trade-off is consistent with online T1 capacity limitations' contributing to the AB, but not with the hypothesis that it is the appearance of the T1+1 distractor (or the discontinuation of target information) that elicits the deficit, since these models ascribe a limited role for T1 processing in the blink (gating theory, boost and bounce, delayed attentional engagement, TLC, eSTST models, threaded cognition model).

To test their hypothesis that a T1–T3 trade-off underlies the disappearance of the AB under uniform conditions, Dux and his colleagues (Dux et al., 2008, 2009) manipulated, both exogenously and endogenously, the extent to which subjects devoted attention to T1 and examined its influence on the T1–T3 performance difference (AB). The logic they followed was that if the standard uniform effect was due to a trade-off between T1 and T3, making the T1 stimulus more salient would increase the attentional resources subjects devoted to its encoding and, thus, reduce the resources available to T3 encoding, leading to a T1–T3 performance difference.

To exogenously manipulate the attention devoted to T1, Dux et al. (2008) colored the targets and the post-T3 distractors red (pre-T1 distractors were white) so that T1 would exogenously capture attention due to its abrupt color onset (Maki & Mebane, 2006). Consistent with a trade-off between T1 and T3, this manipulation improved T1 performance but worsened T3 performance (but see Olivers, Spalek, Kawahara, & Di Lollo, 2009). As a further test of a T1–T3 performance trade-off under uniform conditions, Dux et al. (2009) endogenously manipulated the attentional resources subjects devoted to either T1 or T3 by varying their task relevance in separate blocks of trials. Specifically, in T1-relevant blocks of trials, subjects had to report T1 on all trials, whereas T3 (and T2) required report on only 50% of the trials (and vice versa for T3-relevant blocks of trials). Dux and colleagues predicted that more attention would be devoted to the target that was 100% task relevant, relative to the other two targets. Consistent with this prediction, T1 performance was superior to T3 performance in T1-relevant blocks (suggestive of an AB), and conversely, T3 performance was greater than T1 performance in T3-relevant blocks (a reversed AB!). Importantly, the exogenous and endogenous manipulations had a similar, although somewhat reduced, effect on T1 and T3 in the varied trials. This result suggests that the same processes are involved in sharing attentional resources between targets regardless of whether the targets are presented successively or separated by distractors, thus generalizing Dux et al.'s (2008, 2009) results to more typical AB paradigms. Nevertheless, further research is warranted in order to understand why subjects trade off T1 and T3 performance to a greater extent in the

uniform trials than in the varied trials. One particularly worthwhile hypothesis to explore is the possibility that attentional resources can be more easily manipulated across stimuli when these stimuli belong to the same attentional episode (as when targets are successively presented) than to different episodes (as when targets are separated by distractors).

Collectively, Dux et al.'s (2008, 2009) results are consistent with the hypothesis that the absence of an AB in standard uniform trials is the result of a trade-off between T1 and T3 performance, and they resonate with recent neuroimaging findings showing that targets share attentional resources during RSVP processing (e.g., Sergent et al., 2005; Shapiro, Schmitz, Martens, Hommel, & Schnitzler, 2006). They also fit very well with recent research from Dell'Acqua, Jolicœur, Luria, and Pluchino (2009), who also found an AB under uniform conditions when T3 report accuracy was conditionalized on correct report of T1 and T2. On the other hand, these findings are problematic for models that predict that it is exclusively the processing of the T1+1 distractor that gives rise to the blink (e.g., Di Lollo et al., 2005; Olivers & Meeter, 2008). Furthermore, Nieuwenstein et al.'s (2009) study suggests that the presence of the T1+1 distractor may not even be necessary for an AB to be observed, since they have shown that an AB can occur even when no stimuli follow T1 (no T1 masking), as long as T2 is presented briefly and heavily masked (T2 must be masked to obtain an AB unless there is a task switch between T1 and T2; Giesbrecht & Di Lollo, 1998; Kawahara et al., 2003). This is additional strong evidence against T1+1 distractor-based accounts of the AB (e.g., Di Lollo et al., 2005; Olivers & Meeter, 2008).

Impaired attentional enhancement of T2. Thus far, our examination of the literature suggests that the AB results, at least to some extent, from the devotion of capacity-limited attentional resources to T1, leaving too few of these resources available for processing T2 at short T1–T2 lags. Although the exact nature of this capacity-limited T1 processing has yet to be fully elucidated (is it identity encoding into working memory, episodic registration, response selection, a combination of these processes, or something else?), Nieuwenstein and his colleagues (Nieuwenstein, 2006; Nieuwenstein et al., 2005; for similar findings, see also Olivers & Meeter, 2008; Olivers et al., 2007; Wee & Chua, 2004) have demonstrated that an all-or-none, inflexible T1 bottleneck, where T2 waits for attentional resources until T1 has been fully processed, cannot be the mechanism responsible for the AB. Nieuwenstein et al. (2005) presented subjects with RSVP streams of black letter distractors and red digit targets and found that the AB was attenuated when a red distractor unexpectedly appeared one or two lags before T2. This effect was observed even when the cue and T2 were of different colors, as long as both colors were task relevant (e.g., search for a red T1 and a green T2 with the presentation of a red cue; Nieuwenstein, 2006). Importantly, these cuing manipulations had limited effects on T1 accuracy, suggesting that a trade-off between T1 and T2 was not responsible for the result. These findings suggest that the engagement of at-

tention to T2 is either suppressed or delayed by attentional processing of T1, since cuing attention just prior to the presentation of T2 reduces the AB (see also Chun, 1997a; Vul, Nieuwenstein, & Kanwisher, 2008). Thus, it appears that impaired/suppressed attentional enhancement of the T2 representation—a feature common to the delayed attentional engagement, LCNE, CODAM, eSTST, attention cascade, and threaded cognition models of the AB—plays a vital role in generating the T2 deficit.

The hypothesis that T1 processing suppresses the attentional enhancement of T2 also fits well with a range of other findings in the literature. Shapiro, Caldwell, and Sorensen (1997) demonstrated that the AB was substantially attenuated when T2 was a subject's own name, as compared with a different name. Similarly, A. K. Anderson and Phelps (2001) found that the AB was reduced when T2 was an emotionally arousing word. It may be the case that T2 performance was improved under these conditions because the strong bottom-up attentional saliency of these stimuli more than compensated for the attentional suppression brought about by T1 processing. In addition, a reduction in the extent to which attention is suppressed by T1 processing may explain cases in which distraction reduces the AB (Taatzgen et al., 2009; Wyble et al., 2009), findings that have been taken as evidence inconsistent with the predictions of online T1 processing accounts. For example, Olivers and Nieuwenhuis (2005; see also Olivers & Nieuwenhuis, 2006) found that the AB was reduced when subjects performed a concurrent auditory detection task. Similarly, Arend, Johnston, and Shapiro (2006) demonstrated that the AB was attenuated when the RSVP stream was presented on top of task-irrelevant moving dots (star-field motion). It may be the case that distraction relieves attentional suppression of T2 by preventing the over-commitment of attention toward T1 and the RSVP stream. In any event, the results discussed in this section suggest that although T1 processing may limit the subsequent encoding/episodic registration/response selection of T2 at short lags, a major factor giving rise to the AB deficit is the suppressed attentional enhancement of T2 when it appears in close temporal proximity to T1.

Theoretical Summary

Our examination of the theories and empirical studies discussed above suggests that the following processes may give rise to the AB. During a standard dual-target RSVP task, all the stimuli in the stream are processed both perceptually and conceptually, with semantic information about each of these stimuli available for further processing after this preliminary analysis (e.g., Chun & Potter, 1995; Luck et al., 1996; Maki, Frigen, & Paulson, 1997; Shapiro, Driver, et al., 1997). The strength of these initial representations is determined by their salience (A. K. Anderson & Phelps, 2001; Arnell, Killman, & Fijavz, 2007; Most, Chun, Widders, & Zald, 2005; Shapiro, Caldwell, & Sorensen, 1997; Smith, Most, Newsome, & Zald, 2006) and by the similarity (both perceptual and conceptual) between the target and distractor stimuli presented in the stream (Chun & Potter, 1995; Dux & Coltheart, 2005; Maki et al., 2003), with greater similarity between items

leading to greater masking and, therefore, weaker representations. On the basis of the attentional set established from the task instructions (e.g., Shapiro et al., 1994), distractor items are inhibited (e.g., Dux et al., 2006; Dux & Harris, 2007a; Olivers & Watson, 2006), and upon detection of a target (or a highly salient distractor; Arnell et al., 2007; Most et al., 2005; Smith et al., 2006), an attentional episode is triggered (e.g., Bowman & Wyble, 2007; Chun & Potter, 1995). This attentional episode leads to the enhancement of the representation of the target stimulus and the T1+1 target/distractor (if there is one), due to the temporal dynamics of the attentional deployment. Stimuli that are processed in the same attentional window compete to be admitted to higher stages of processing (e.g., Potter et al., 2005; Potter et al., 2002), with the winner of this competition undergoing episodic registration, working memory consolidation, and/or immediate response selection—all processes that require attention. Typically, under standard RSVP conditions and timing, it will be the T1 stimulus that wins this competition, due to its salience, its head start in preliminary identification relative to the T1+1 item, and, in the case in which T1+1 is a distractor, its task relevance. Because encoding, episodic registration, and response selection stages of processing are attentionally demanding, other stimuli that appear in close temporal proximity to T1 will not receive the same attentional enhancement (except when T2 appears at lag 1; see below) and access to working memory, leaving them vulnerable to decay and overwriting (e.g., Chun & Potter, 1995; Giesbrecht & Di Lollo, 1998). Put differently, as attention is devoted to encoding/registering/responding to the first target, this limits/suppresses the attention that is available to enhance subsequently presented targets (e.g., Bowman & Wyble, 2007; Chun & Potter, 1995) and the ability of the system to inhibit distractors (e.g., Dux & Harris, 2007a; Dux & Marois, 2008). All these factors contribute to the generation of the AB. Under conditions in which T2 appears at lag 1, this will typically lead to sparing of T2, because both T1 and T2 will be attentionally enhanced and undergo high-level processing simultaneously. Nevertheless, at lag 1, there will still be competition between the target stimuli, which may result in the superior report of T2, as compared with T1 (e.g., Bowman & Wyble, 2007; Chun & Potter, 1995; Potter et al., 2002), and significant numbers of temporal order swaps between the two targets (e.g., Bowman & Wyble, 2007; Chun & Potter, 1995; Hommel & Akyürek, 2005).

Conclusion

The AB is a robust phenomenon that has been demonstrated across a wide range of experimental conditions. Our review of the literature suggests that the AB reflects the competition between targets for attentional resources, not only for working memory encoding, episodic registration, and response selection (and perhaps additional processes that have yet to be identified), but also for the enhancement of target representations and the inhibition of distractors. T1 processing renders these attentional resources temporarily unavailable for subsequent stimuli, thereby impairing the report of T2 at short T1–T2 lags. Of

the current models, those proposed by Wyble et al. (2009, eSTST; Bowman & Wyble, 2007; see also Chun & Potter, 1995) and Shih (2008, attention cascade model) accommodate the largest number of empirical findings, since they incorporate capacity-limited T1 processing (episodic registration in the eSTST and encoding in the attentional cascade model), which leads to the impaired attentional enhancement for subsequent targets at short T1–T2 lags. Although Dehaene et al. (2003), Chartier et al. (2004), Fragopanagos et al. (2005), Nieuwenhuis, Gilzenrat, et al. (2005), Nieuwenstein et al. (2009), and Taatgen et al. (2009) present somewhat related hypotheses, these models fail to incorporate mechanisms that account for several important findings (e.g., those related to T1+1 and T2+1 masking in the RSVP stream). Having said this, no current theory can fully account for all the findings related to this complex phenomenon.

Irrespective of which model best fits the AB literature, our theoretical summary seems to argue for a multifactorial origin of this processing deficit: Attentional selection, working memory encoding, episodic registration, response selection, attentional enhancement and engagement, and distractor inhibition have all been implicated in limiting multitarget performance in RSVP. However, it is also possible that these multiple processes rely on a common capacity-limited attentional resource and that it is this resource that underlies the AB deficit. According to this view, the process that is responsible for the trade-off between T1 and T3 performance in the serial target experiments of Dux et al. (2008, 2009) is the same as that which underlies the AB impairment in the distractor-less design of Nieuwenstein et al. (2009) or the attenuating effect of distraction in the experiments of Olivers and Nieuwenhuis (2005, 2006)—namely, the deployment of selective attention. The more attention deployed for T1, because it is more salient or more task relevant or requires more encoding into working memory, the less available for processing subsequent targets. Similarly drawing attention away from T1, either by cuing a distractor prior to T2 (e.g., Nieuwenstein, 2006) or by including distracting tasks (see above), may alleviate the T2 deficit. The neuroimaging evidence that AB manipulations recruit the frontal–parietal attentional networks of the brain (Hommel et al., 2006; Marois & Ivanoff, 2005) adds further weight to the view that, first and foremost, the AB represents a deficit of selective attention. Attention, after all, is generally regarded as the mechanism by which behaviorally relevant items, such as targets, are selectively processed over other items, such as distractors (Pashler, 1998). Thus, any stages of information processing that are involved in achieving that behavioral goal may be the recipients of attention and, hence, contribute to the AB deficit.

Although appealing in its simplicity, this selective attention account of the AB—like all the models elaborated above—does not encapsulate all of the characteristics of this deficit. But as was mentioned previously, it is unlikely that a single mechanism can explain the myriad of AB findings. Moreover, even if the AB can be attributed to more than one process, the extent to which each of these processes contributes to the deficit remains to be determined.

In particular, there have been few attempts to distinguish, in both the theoretical and experimental literatures, between the factors that cause the AB (i.e., are essential for its occurrence) and those that merely modulate its magnitude. Evidently, more research is needed to further understand the cognitive mechanisms that give rise to the AB and, more important, the implications of this fundamental temporal processing limitation for visual awareness.

AUTHOR NOTE

This work was supported by an ARC grant (DP0986387) to P.E.D. and NIMH (R01 MH70776) and NSF (0094992) grants to R.M. We thank Howard Bowman, Roberto Dell'Acqua, Mark Nieuwenstein, Adriane Seiffert, and Kimron Shapiro for helpful comments. Correspondence concerning this article should be addressed to P. E. Dux, School of Psychology, University of Queensland, 463 McElwain Building, St. Lucia, QLD 4072, Australia (e-mail: paul.e.dux@gmail.com).

REFERENCES

- AKYÜREK, E. G., HOMMEL, B., & JOLICŒUR, P. (2007). Direct evidence for a role of working memory in the attentional blink. *Memory & Cognition*, *35*, 621-627.
- ANDERSON, A. K., & PHELPS, E. A. (2001). Lesions of the human amygdala impair enhanced perception of emotionally salient events. *Nature*, *411*, 305-309.
- ANDERSON, J. R. (2007). *How can the human mind occur in the physical universe?* New York: Oxford University Press.
- AREND, I., JOHNSTON, S., & SHAPIRO, K. (2006). Task-irrelevant visual motion and flicker attenuate the attentional blink. *Psychonomic Bulletin & Review*, *13*, 600-607.
- ARNELL, K. M., & DUNCAN, J. (2002). Separate and shared sources of dual-task cost in stimulus identification and response selection. *Cognitive Psychology*, *44*, 105-147.
- ARNELL, K. M., & JENKINS, R. (2004). Revisiting within-modality and cross-modality attentional blinks: Effects of target–distractor similarity. *Perception & Psychophysics*, *66*, 1147-1161.
- 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., KILLMAN, K. V., & FIJAVZ, D. (2007). Blinded by emotion: Target misses follow attention capture by arousing distractors in RSVP. *Emotion*, *7*, 465-477.
- ARNELL, K. M., & LARSON, J. M. (2002). Cross-modality attentional blinks without preparatory task-set switching. *Psychonomic Bulletin & Review*, *9*, 497-506.
- AWH, E., SERENCES, J., LAUREY, P., DHALIWAL, H., VAN DER JAGT, T., & DASSONVILLE, P. (2004). Evidence against a central bottleneck during the attentional blink: Multiple channels for configural and featural processing. *Cognitive Psychology*, *48*, 95-126.
- BAARS, B. (1989). *A cognitive theory of consciousness*. New York: Cambridge University Press.
- BACHMANN, T., & HOMMUK, K. (2005). How backward masking becomes attentional blink: Perception of successive in-stream targets. *Psychological Science*, *16*, 740-742.
- BADDELEY, A. D., THOMSON, N., & BUCHANAN, M. (1975). Word length and the structure of short-term memory. *Journal of Verbal Learning & Verbal Behavior*, *14*, 575-589.
- BERRIDGE, C. W., & WATERHOUSE, B. D. (2003). The locus coeruleus–noradrenergic system: Modulation of behavioral state and state dependent cognitive processes. *Brain Research Reviews*, *42*, 33-84.
- BOWMAN, H., & WYBLE, B. P. (2007). The simultaneous type, serial token model of temporal attention and working memory. *Psychological Review*, *114*, 38-70.
- BROADBENT, D. E. (1958). *Perception and communication*. London: Pergamon.
- 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.

- BUNDESEN, C. (1990). A theory of visual attention. *Psychological Review*, **97**, 523-547.
- CHARTIER, S., COUSINEAU, D., & CHARBONNEAU, D. (2004). A connectionist model of the attentional blink effect during a rapid serial visual task. In *Proceedings of the 6th International Conference on Cognitive Modelling* (pp. 64-69). Mahwah, NJ: Erlbaum.
- CHUA, F. K., GOH, J., & HON, N. (2001). Nature of codes extracted during the attentional blink. *Journal of Experimental Psychology: Human Perception & Performance*, **27**, 1229-1242.
- CHUN, M. M. (1997a). Temporal binding errors are redistributed by the attentional blink. *Perception & Psychophysics*, **59**, 1191-1199.
- CHUN, M. M. (1997b). Types and tokens in visual processing: A double dissociation between the attentional blink and repetition blindness. *Journal of Experimental Psychology: Human Perception & Performance*, **23**, 738-755.
- 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.
- CHUN, M. M., & POTTER, M. C. (2001). The attentional blink and task-switching. In K. Shapiro (Ed.), *The limits of attention: Temporal constraints in human information processing* (pp. 20-35). Oxford: Oxford University Press.
- CHUN, M. M., & WOLFE, J. M. (2001). Visual attention. In B. Goldstein (Ed.), *Blackwell handbook of perception* (pp. 272-310). Oxford: Blackwell.
- COLTHEART, V. (ED.) (1999). *Fleeting memories: Cognition of brief visual stimuli*. Cambridge, MA: MIT Press.
- COLTHEART, V., & LANGDON, R. (1998). Recall of short word lists presented visually at fast rates: Effects of phonological similarity and word length. *Memory & Cognition*, **26**, 330-342.
- COLTHEART, V., MONDY, S., DUX, P. E., & STEPHENSON, L. (2004). Effects of orthographic and phonological word length on memory for lists shown at RSVP and STM rates. *Journal of Experimental Psychology: Learning, Memory, & Cognition*, **30**, 815-826.
- COLZATO, L. S., SPAPÉ, M., PANNEBAKKER, M. M., & HOMMEL, B. (2007). Working memory and the attentional blink: Blink size is predicted by individual differences in operation span. *Psychonomic Bulletin & Review*, **14**, 1051-1057.
- DEHAENE, S., SERGENT, C., & CHANGEUX, J. P. (2003). A neuronal network model linking subjective reports and objective physiological data during conscious perception. *Proceedings of the National Academy of Sciences*, **100**, 8520-8525.
- DELL'ACQUA, R., JOLICÉUR, P., LURIA, R., & PLUCHINO, P. (2009). Re-evaluating encoding-capacity limitations as a cause of the attentional blink. *Journal of Experimental Psychology: Human Perception & Performance*, **35**, 338-351.
- DELL'ACQUA, R., PASCALI, A., JOLICÉUR, P., & SESSA, P. (2003). Four-dot masking produces the attentional blink. *Vision Research*, **43**, 1907-1913.
- DELL'ACQUA, R., SESSA, P., JOLICÉUR, P., & ROBITAILLE, N. (2006). Spatial attention freezes during the attentional blink. *Psychophysiology*, **43**, 394-400.
- DESIMONE, R., & DUNCAN, J. (1995). Neural mechanisms of selective visual attention. *Annual Review of Neuroscience*, **18**, 193-222.
- DI LOLLO, V., KAWAHARA, J., GHORASHI, S. M. S., & ENNS, J. T. (2005). The attentional blink: Resource depletion or temporary loss of control. *Psychological Research*, **69**, 191-200.
- DONCHIN, E. (1981). Surprise! ... Surprise? *Psychophysiology*, **18**, 493-513.
- DONCHIN, E., & COLES, M. G. H. (1988). Is the P300 component a manifestation of context updating? *Behavioural & Brain Sciences*, **11**, 357-374.
- DREW, T., & SHAPIRO, K. (2006). Representational masking and the attentional blink. *Visual Cognition*, **13**, 513-528.
- DUNCAN, J. (1980). The locus of interference in the perception of simultaneous stimuli. *Psychological Review*, **87**, 272-300.
- DUNCAN, J., & HUMPHREYS, G. (1989). Visual search and stimulus similarity. *Psychological Review*, **96**, 433-458.
- DUNCAN, J., MARTENS, S., & WARD, R. (1997). Restricted attentional capacity within but not between sensory modalities. *Nature*, **387**, 808-810.
- DUNCAN, J., WARD, R., & SHAPIRO, K. (1994). Direct measurement of attentional dwell time in human vision. *Nature*, **369**, 313-315.
- DUX, P. E., ASPLUND, C. L., & MAROIS, R. (2008). An attentional blink for sequentially presented targets: Evidence in favor of resource depletion accounts. *Psychonomic Bulletin & Review*, **15**, 809-813.
- DUX, P. E., ASPLUND, C. L., & MAROIS, R. (2009). Both exogenous and endogenous target salience manipulations support resource depletion accounts of the attentional blink: A reply to Olivers, Spalek, Kawahara, and Di Lollo (2009). *Psychonomic Bulletin & Review*, **16**, 219-224.
- DUX, P. E., & COLTHEART, V. (2005). The meaning of the mask matters: Evidence of conceptual interference in the attentional blink. *Psychological Science*, **16**, 775-779.
- DUX, P. E., COLTHEART, V., & HARRIS, I. M. (2006). On the fate of distractor stimuli in rapid serial visual presentation. *Cognition*, **99**, 355-382.
- DUX, P. E., & HARRIS, I. M. (2007a). On the failure of distractor inhibition in the attentional blink. *Psychonomic Bulletin & Review*, **14**, 723-728.
- DUX, P. E., & HARRIS, I. M. (2007b). Viewpoint costs occur during consolidation: Evidence from the attentional blink. *Cognition*, **101**, 47-58.
- DUX, P. E., & MAROIS, R. (2007). Repetition blindness is immune to the central bottleneck. *Psychonomic Bulletin & Review*, **14**, 729-734.
- DUX, P. E., & MAROIS, R. (2008). Distractor inhibition predicts individual differences in the attentional blink. *PLoS ONE*, **3**, e3330.
- ENNS, J. T., VISSER, T. A. W., KAWAHARA, J.-I., & DI LOLLO, V. (2001). Visual masking and task switching in the attentional blink. In K. Shapiro (Ed.), *The limits of attention: Temporal constraints in human information processing* (pp. 65-81). Oxford: Oxford University Press.
- FOLK, C. L., LEBER, A. B., & EGETH, H. E. (2002). Made you blink! Contingent attentional capture produces a spatial blink. *Perception & Psychophysics*, **64**, 741-753.
- FRAGOPANAGOS, N., KOCKELKOREN, S., & TAYLOR, J. G. (2005). A neurodynamic model of the attentional blink. *Cognitive Brain Research*, **24**, 568-586.
- GIESBRECHT, B., BISCHOF, W. F., & KINGSTONE, A. (2003). Visual masking during the attentional blink: Tests of the object substitution hypothesis. *Journal of Experimental Psychology: Human Perception & Performance*, **29**, 238-255.
- GIESBRECHT, B., & DI LOLLO, V. (1998). Beyond the attentional blink: Visual masking by object substitution. *Journal of Experimental Psychology: Human Perception & Performance*, **24**, 1454-1466.
- GIESBRECHT, B., SY, J. L., & ELLIOTT, J. C. (2007). Electrophysiological evidence for both perceptual and post-perceptual selection during the attentional blink. *Journal of Cognitive Neuroscience*, **19**, 2005-2018.
- 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.
- GROSS, J., SCHMITZ, F., SCHNITZLER, I., KESSLER, K., SHAPIRO, K., HOMMEL, B., & SCHNITZLER, A. (2004). Long-range neural synchrony predicts temporal limitations of visual attention in humans. *Proceedings of the National Academy of Sciences*, **101**, 13050-13055.
- HOMMEL, B., & AKYÜREK, E. (2005). Lag-1 sparing in the attentional blink: Benefits and costs of integrating two events into a single episode. *Quarterly Journal of Experimental Psychology*, **58A**, 1415-1433.
- HOMMEL, B., KESSLER, K., SCHMITZ, F., GROSS, J., AKYÜREK, E., SHAPIRO, K., & SCHNITZLER, A. (2006). How the brain blinks: Towards a neurocognitive model of the attentional blink. *Psychological Research*, **70**, 425-435.
- ISAAK, M. I., SHAPIRO, K. L., & MARTIN, J. (1999). The attentional blink reflects retrieval competition among multiple rapid serial visual presentation items: Tests of an interference model. *Journal of Experimental Psychology: Human Perception & Performance*, **25**, 1774-1792.
- JACKSON, M. C., & RAYMOND, J. E. (2006). The role of attention and familiarity in face identification. *Perception & Psychophysics*, **68**, 543-557.
- JOLICÉUR, P. (1985). The time to name disoriented natural objects. *Memory & Cognition*, **13**, 289-303.
- JOLICÉUR, P. (1998). Modulation of the attentional blink by on-line response selection: Evidence from speeded and unspeeded Task₁ decisions. *Memory & Cognition*, **26**, 1014-1032.
- JOLICÉUR, P. (1999). Concurrent response-selection demands modulate the attentional blink. *Journal of Experimental Psychology: Human Perception & Performance*, **25**, 1097-1113.
- JOLICÉUR, P., & DELL'ACQUA, R. (1998). The demonstration of short-term consolidation. *Cognitive Psychology*, **32**, 138-202.

- JOLICÉUR, P., DELL'ACQUA, R., & CREBOLDER, J. M. (2001). The attentional blink bottleneck. In K. Shapiro (Ed.), *The limits of attention: Temporal constraints in human information processing* (pp. 82-99). Oxford: Oxford University Press.
- JOLICÉUR, P., SESSA, P., DELL'ACQUA, R., & ROBITAILLE, N. (2006a). Attentional control and capture in the attentional blink paradigm: Evidence from human electrophysiology. *European Journal of Cognitive Psychology*, **18**, 560-578.
- JOLICÉUR, P., SESSA, P., DELL'ACQUA, R., & ROBITAILLE, N. (2006b). On the control of visual spatial attention: Evidence from human electrophysiology. *Psychological Research*, **70**, 414-424.
- KAHNEMAN, D. (1973). *Attention and effort*. Englewood Cliffs, NJ: Prentice Hall.
- KANWISHER, N. (1987). Repetition blindness: Type recognition without token individuation. *Cognition*, **27**, 117-143.
- KAWAHARA, J.-I., ENNS, J. T., & DI LOLLO, V. (2006). The attentional blink is not a unitary phenomenon. *Psychological Research*, **70**, 405-413.
- KAWAHARA, J.-I., KUMADA, T., & DI LOLLO, V. (2006). The attentional blink is governed by a temporary loss of control. *Psychonomic Bulletin & Review*, **13**, 886-890.
- KAWAHARA, J.-I., ZUVIC, S. M., ENNS, J. T., & DI LOLLO, V. (2003). Task switching mediates the attentional blink even without backward masking. *Perception & Psychophysics*, **65**, 339-351.
- KRANCZIOCH, C., DEBENER, S., SCHWARZBACH, J., GOEBEL, R., & ENGEL, A. K. (2005). Neural correlates of conscious perception in the attentional blink. *NeuroImage*, **24**, 704-714.
- LANDAU, A. N., & BENTIN, S. (2008). Attentional and perceptual factors affecting the attentional blink for faces and objects. *Journal of Experimental Psychology: Human Perception & Performance*, **34**, 818-830.
- LAVIE, N. (1995). Perceptual load as a necessary condition for selective attention. *Journal of Experimental Psychology: Human Perception & Performance*, **21**, 451-468.
- LAWRENCE, D. H. (1971). Two studies of visual search for word targets with controlled rates of presentation. *Perception & Psychophysics*, **10**, 85-89.
- LUCK, S. J., VOGEL, E. K., & SHAPIRO, K. L. (1996). Word meanings can be accessed but not reported during the attentional blink. *Nature*, **383**, 616-618.
- MAKI, W. S., BUSSARD, G., LOPEZ, K., & DIGBY, B. (2003). Sources of interference in the attentional blink: Target-distractor similarity revisited. *Perception & Psychophysics*, **65**, 188-201.
- MAKI, W. S., COUTURE, T., FRIGEN, K., & LIEN, D. (1997). Sources of the attentional blink during rapid serial visual presentation: Perceptual interference and retrieval competition. *Journal of Experimental Psychology: Human Perception & Performance*, **23**, 1393-1411.
- MAKI, W. S., FRIGEN, K., & PAULSON, K. (1997). Associative priming by targets and distractors during rapid serial visual presentation: Does word meaning survive the attentional blink? *Journal of Experimental Psychology: Human Perception & Performance*, **23**, 1014-1034.
- MAKI, W. S., & MEBANE, M. W. (2006). Attentional capture triggers an attentional blink. *Psychonomic Bulletin & Review*, **13**, 125-131.
- MAKI, W. S., & PADMANABHAN, G. (1994). Transient suppression of processing during rapid serial visual presentation: Acquired distinctiveness of probes modulates the attentional blink. *Psychonomic Bulletin & Review*, **1**, 499-504.
- MAROIS, R., CHUN, M. M., & GORE, J. C. (2004). A common parieto-frontal network is recruited under both low visibility and high perceptual interference conditions. *Journal of Neurophysiology*, **92**, 2985-2992.
- MAROIS, R., & IVANOFF, J. (2005). Capacity limits of information processing in the brain. *Trends in Cognitive Sciences*, **9**, 296-305.
- MAROIS, R., YI, D. J., & CHUN, M. M. (2004). The neural fate of consciously perceived and missed events in the attentional blink. *Neuron*, **41**, 465-472.
- MARTENS, S., MUNNEKE, J., SMID, H., & JOHNSON, A. (2006). Quick minds don't blink: Electrophysiological correlates of individual differences in attentional selection. *Journal of Cognitive Neuroscience*, **18**, 1423-1438.
- MARTIN, E. W., & SHAPIRO, K. L. (2008). Does failure to mask T1 cause lag-1 sparing in the attentional blink? *Perception & Psychophysics*, **70**, 562-570.
- MCAULIFFE, S. P., & KNOWLTON, B. J. (2000). Dissociating the effects of featural and conceptual interference on multiple target processing in rapid serial visual presentation. *Perception & Psychophysics*, **62**, 187-195.
- 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.
- MILLER, G. A. (2003). The cognitive revolution: A historical perspective. *Trends in Cognitive Sciences*, **7**, 141-144.
- MOST, S. B., CHUN, M. M., WIDDERS, D. M., & ZALD, D. H. (2005). Attentional rubbernecking: Cognitive control and personality in emotion-induced blindness. *Psychonomic Bulletin & Review*, **12**, 654-661.
- NEISSER, U. (1967). *Cognitive psychology*. New York: Appellton-Century-Crofts.
- NIEUWENHUIS, S., ASTON-JONES, G., & COHEN, J. D. (2005). Decision making, the P3, and the locus coeruleus-norepinephrine system. *Psychological Bulletin*, **131**, 510-532.
- NIEUWENHUIS, S., GILZENRAT, M. S., HOLMES, B. D., & COHEN, J. D. (2005). The role of the locus coeruleus in mediating the attentional blink: A neurocomputational theory. *Journal of Experimental Psychology: General*, **134**, 291-307.
- NIEUWENSTEIN, M. R. (2006). Top-down controlled, delayed selection in the attentional blink. *Journal of Experimental Psychology: Human Perception & Performance*, **32**, 973-985.
- NIEUWENSTEIN, M. R., CHUN, M. M., VAN DER LUBBE, R. H. J., & HOOGE, I. T. C. (2005). Delayed attentional engagement in the attentional blink. *Journal of Experimental Psychology: Human Perception & Performance*, **31**, 1463-1475.
- NIEUWENSTEIN, M. R., & POTTER, M. C. (2006). Temporal limits of selection and memory encoding: A comparison of whole versus partial report in rapid serial visual presentation. *Psychological Science*, **17**, 471-475.
- NIEUWENSTEIN, M. R., POTTER, M. C., & THEEUWES, J. (2009). Unmasking the attentional blink. *Journal of Experimental Psychology: Human Perception & Performance*, **35**, 159-169.
- OLIVERS, C. N. L., & MEETER, M. (2008). A boost and bounce theory of temporal attention. *Psychological Review*, **115**, 836-863.
- OLIVERS, C. N. L., & NIEUWENHUIS, S. (2005). The beneficial effect of concurrent task-irrelevant mental activity on temporal attention. *Psychological Science*, **16**, 265-269.
- OLIVERS, C. N. L., & NIEUWENHUIS, S. (2006). The beneficial effects of additional task load, positive affect, and instruction on the attentional blink. *Journal of Experimental Psychology: Human Perception & Performance*, **32**, 364-379.
- OLIVERS, C. N. L., SPALEK, T. M., KAWAHARA, J.-I., & DI LOLLO, V. (2009). The attentional blink: Increasing target salience provides no evidence for resource depletion. A commentary on Dux, Asplund, and Marois (2008). *Psychonomic Bulletin & Review*, **16**, 214-218.
- OLIVERS, C. N. L., VAN DER STIGCHEL, S., & HULLEMAN, J. (2007). Spreading the sparing: Against a limited-capacity account of the attentional blink. *Psychological Research*, **71**, 126-139.
- OLIVERS, C. N. L., & WATSON, D. G. (2006). Input control processes in rapid serial visual presentations: Target selection and distractor inhibition. *Journal of Experimental Psychology: Human Perception & Performance*, **32**, 1083-1092.
- OLSON, I. R., CHUN, M. M., & ANDERSON, A. K. (2001). Effects of phonological length on the attentional blink for words. *Journal of Experimental Psychology: Human Perception & Performance*, **27**, 1116-1123.
- OUMET, C., & JOLICÉUR, P. (2007). Beyond Task 1 difficulty: The duration of T1 encoding modulates the attentional blink. *Visual Cognition*, **15**, 290-304.
- PASHLER, H. (1994). Dual-task interference in simple tasks: Data and theory. *Psychological Bulletin*, **116**, 220-244.
- PASHLER, H. (1998). *The psychology of attention*. Cambridge, MA: MIT Press.
- POTTER, M. C. (1975). Meaning in visual search. *Science*, **187**, 965-966.
- POTTER, M. C. (1976). Short-term conceptual memory for pictures. *Journal of Experimental Psychology: Human Learning & Memory*, **2**, 509-522.
- POTTER, M. C. (1993). Very short-term conceptual memory. *Memory & Cognition*, **21**, 156-161.
- POTTER, M. C., CHUN, M. M., BANKS, B. S., & MUCKENHOUT, M.

- (1998). Two attentional deficits in serial target search: The visual attentional blink and an amodal task-switch deficit. *Journal of Experimental Psychology: Learning, Memory, & Cognition*, **24**, 979-992.
- POTTER, M. C., DELL'ACQUA, R., PESCIARELLI, F., JOB, R., PERESOTTI, F., & O'CONNOR, D. H. (2005). Bidirectional semantic priming in the attentional blink. *Psychonomic Bulletin & Review*, **12**, 460-465.
- POTTER, M. C., & FAULCONER, B. A. (1975). Time to understand pictures and words. *Nature*, **253**, 437-438.
- POTTER, M. C., & LEVY, E. I. (1969). Recognition memory for a rapid sequence of pictures. *Journal of Experimental Psychology*, **81**, 10-15.
- POTTER, M. C., STAUB, A., & O'CONNOR, D. H. (2002). The time course of competition for attention: Attention is initially labile. *Journal of Experimental Psychology: Human Perception & Performance*, **28**, 1149-1162.
- 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.
- RAYMOND, J. E., SHAPIRO, K. L., & ARNELL, K. M. (1995). Similarity determines the attentional blink. *Journal of Experimental Psychology: Human Perception & Performance*, **21**, 653-662.
- REEVES, A., & SPERLING, G. (1986). Attention gating in short-term visual memory. *Psychological Review*, **93**, 180-206.
- RUTHRUFF, E., & PASHLER, H. E. (2001). Perceptual and central interference in dual-task performance. In K. Shapiro (Ed.), *The limits of attention: Temporal constraints in human information processing* (pp. 100-123). Oxford: Oxford University Press.
- SEIFFERT, A., & DI LOLLO, V. (1997). Low-level masking in the attentional blink. *Journal of Experimental Psychology: Human Perception & Performance*, **23**, 1061-1073.
- SERGENT, C., BAILLET, S., & DEHAENE, S. (2005). Timing of the brain events underlying access to consciousness during the attentional blink. *Nature Neuroscience*, **8**, 1391-1400.
- SERGENT, C., & DEHAENE, S. (2004). Is consciousness a gradual phenomenon? Evidence for an all-or-none bifurcation during the attentional blink. *Psychological Science*, **15**, 720-728.
- 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. (1994). Temporal allocation of visual attention: Inhibition or interference? In D. Dagenbach & T. H. Carr (Eds.), *Inhibitory mechanisms in attention, memory and language* (pp. 151-188). Boston: Academic Press.
- 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.
- SHAPIRO, K. [L.], SCHMITZ, F., MARTENS, S., HOMMEL, B., & SCHNITZLER, A. (2006). Resource sharing in the attentional blink. *NeuroReport*, **17**, 163-166.
- SHIFFRIN, R. M., & SCHNEIDER, W. (1977). Controlled and automatic human information processing: II. Perceptual learning, automatic attending, and a general theory. *Psychological Review*, **84**, 127-190.
- SHIH, S. I. (2008). The attention cascade model and the attentional blink. *Cognitive Psychology*, **56**, 210-236.
- SHORE, D. I., McLAUGHLIN, E. N., & KLEIN, R. (2001). Modulation of the attentional blink by differential resource allocation. *Canadian Journal of Experimental Psychology*, **55**, 318-324.
- SMITH, S. D., MOST, S. B., NEWSOME, L. A., & ZALD, D. H. (2006). An "emotional blink" of attention elicited by aversively conditioned stimuli. *Emotion*, **6**, 523-527.
- TAATGEN, N. A., JUVINA, I., SCHIPPER, M., BORST, J. P., & MARTENS, S. (2009). Too much control can hurt: A threaded cognition model of the attentional blink. *Cognitive Psychology*, **59**, 1-29. doi:10.1016/j.cogpsych.2008.12.002
- TAYLOR, J. G., & ROGERS, M. (2002). A control model of the movement of attention. *Neural Networks*, **15**, 309-326.
- THORPE, S., FIZE, D., & MARLOT, C. (1996). Speed of processing in the human visual system. *Nature*, **381**, 520-522.
- TREISMAN, A. M. (1969). Strategies and models of selective attention. *Psychological Review*, **76**, 282-299.
- VISSER, T. A. W. (2007). Masking T1 difficulty: Processing time and the attentional blink. *Journal of Experimental Psychology: Human Perception & Performance*, **33**, 285-297.
- VISSER, T. A. W., BISCHOF, W. F., & DI LOLLO, V. (1999). Attentional switching in spatial and nonspatial domains: Evidence from the attentional blink. *Psychological Bulletin*, **125**, 458-469.
- VOGEL, E. K., & LUCK, S. J. (2002). Delayed working memory consolidation during the attentional blink. *Psychonomic Bulletin & Review*, **9**, 739-743.
- 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.
- VUL, E., NIEUWENSTEIN, M., & KANWISHER, N. (2008). Temporal selection is suppressed, delayed, and diffused during the attentional blink. *Psychological Science*, **19**, 55-61.
- WARD, R., DUNCAN, J., & SHAPIRO, K. (1996). The slow time-course of visual attention. *Cognitive Psychology*, **30**, 79-109.
- WARD, R., DUNCAN, J., & SHAPIRO, K. (1997). Effects of similarity, difficulty, and nontarget presentation on the time course of visual attention. *Perception & Psychophysics*, **59**, 593-600.
- WEE, S., & CHUA, F. K. (2004). Capturing attention when attention blinks. *Journal of Experimental Psychology: Human Perception & Performance*, **30**, 598-612.
- WEICHELGARTNER, E., & SPERLING, G. (1987). Dynamics of automatic and controlled visual attention. *Science*, **238**, 778-780.
- WILLIAMS, M. A., VISSER, T. A., CUNNINGTON, R., & MATTINGLEY, J. B. (2008). Attenuation of neural responses in primary visual cortex during the attentional blink. *Journal of Neuroscience*, **8**, 9890-9894.
- WYBLE, B., BOWMAN, H., & NIEUWENSTEIN, M. (2009). The attentional blink provides episodic distinctiveness: Sparing at a cost. *Journal of Experimental Psychology: Human Perception & Performance*, **35**, 787-807.

NOTES

1. Here, we list the eSTST model as a framework consistent with online T1 resource depletion accounts of the AB, because this theory suggests that episodic registration of T1 is capacity limited and that this registration causes the attentional blaster to be suppressed when T2 is presented at short lags. It should be noted, however, that Wyble et al. (2009) viewed their account as one that implicates a perceptual strategy at the origin of the deficit.

2. Note that, here, we discuss the influence of T1 processing on the AB, rather than the influence of T1 difficulty on the AB. This is an important distinction because a significant problem with accuracy data is that they do not allow one to fully elucidate whether incorrect responses reflect short- or long-duration processing (the same can be said of correct responses). Put differently, just because subjects show reduced performance on a T1 task, relative to another task, does not mean that they devoted the same amount of attention/time to the two conditions; it may be the case that they devoted less to the former because it was too difficult (i.e., they quit processing the stimulus early and moved on to a subsequent task—T2). Thus, T1 capacity-limited models of the AB do not make directional predictions regarding the influence of T1 difficulty on the T2 deficit (difficult T1s could lead to bigger or smaller ABs); rather, their directional predictions apply to the influence on the AB of the attention subjects devote to T1.

3. It should be noted that some frameworks that do incorporate some type of capacity-limited T1 processing, such as the eSTST model (limited capacity in the number of stimuli that can be episodically registered as distinct items at a time) and the attention cascade model (limited capacity in the number of items that can be encoded at a time), are not inconsistent with spreading of the sparing, since the attentional enhancement of targets in these models is sustained under uniform conditions.