

Evidence for strategic suppression of irrelevant activation in the Simon task



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ABSTRACT

It is well known that the Simon effect usually decreases with an increasing response time. According to a prominent account this is due to a gradually increasing suppression of irrelevant location-induced activation. What is open, however, and what was investigated in the present study, is to what extent this selective suppression can be adjusted strategically. We hypothesized that strategic suppression should depend on the availability of information about the inhibitory demands. Therefore, in two experiments the demand was modulated by varying the delay between a spatial cue and the target. In the first experiment, where target delay was randomized, there was a negative Simon effect for the longer delays. In a second experiment, where delay was blocked, the Simon effect remained positive. However, the overall Simon effect was larger than in Experiment 1. Together, our results show that the strength of selective suppression can be adjusted strategically, but that this does not necessarily lead to a smaller Simon effect.

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1. Introduction

Dealing with opposing response tendencies is an important aspect of goal-directed behavior, and several experimental conflict paradigms, such as the Stroop task (Steinhauser & Hübner, 2009; Stroop, 1935), or the Flanker task (Eriksen & Eriksen, 1974; Hübner, Steinhauser, & Lehle, 2010), have been developed for investigating the involved processes. In these paradigms internal stimulus features are commonly used to induce a response conflict. However, a conflict can also be induced by more external features such as location. This is obvious in the *Simon task* (Proctor, 2011; Simon, 1969), where a spatial response (e.g. pressing a 'left' or 'right' button) to a target stimulus is required, depending on the target's value on a non-spatial dimension (e.g., its color or form). If the stimulus is then presented either to the left or right of fixation, responses are usually faster and more reliable when the target position is ipsilateral to the required response, compared to when it is contralateral. This *Simon effect*, which is in the focus of the present study, demonstrates that task irrelevant spatial information can have a substantial influence on response selection (for an overview see Hommel, 2011).

A widely acknowledged account of the Simon effect is the dual-route idea (De Jong, Liang, & Lauber, 1994; Kornblum, Hasbroucq, & Osman, 1990), which assumes an indirect or controlled route, where task-relevant information is intentionally translated into the required

response, and a direct route, along which the location of a stimulus automatically activates the spatially corresponding response code. The basic Simon effect is then explained by assuming that the automatic activation of the response code via the direct route has a facilitating or interfering influence on response selection, depending on the respective congruency of the trial.

A specific characteristic of the typical Simon effect is that it decreases with an increasing response time (RT), and that this holds for both latencies and accuracy. That the congruency effect in accuracy decreases with RT is common to all conflict paradigms. It is large for short RTs, because of a high proportion of impulsive responses to the irrelevant information. The proportion then decreases with an increasing RT. In the latencies, the Simon effect usually also decreases with RT. This, however, is opposite to other conflict paradigms, such as the Flanker task, where the corresponding congruency effect in the latencies usually increases with RT (e.g. Hübner et al., 2010). How the congruency effect changes with RT can be examined by considering so-called *delta functions* (e.g. De Jong et al., 1994; Ridderinkhof, 2002a), which directly specify the effect size as function of RT. The decreasing Simon effect in the latencies with RT is then reflected by a negative slope of the corresponding delta function.

The relation between effect size and RT, as reflected by the delta functions, is very helpful for inferring the involved mechanisms (for an overview see van den Wildenberg et al., 2010). To explain the characteristic time course of the Simon effect in the latencies, the *temporal overlap* hypothesis has been proposed. The idea is that the automatic activation induced by stimulus location decays with time (Hommel, 1993, 1994). Accordingly, on trials on which task-relevant information

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is processed slowly, response translation takes place, at least partly, after the irrelevant location-induced activation has decayed, so that the period of interference is relatively short. In contrast, on trials where the relevant information is processed rapidly, the translation process overlaps temporally to a large extent with the irrelevant activation, which produces a substantial interference.

Early support for the temporal-overlap hypothesis was provided by Simon, Acosta, Mewaldt, and Speidel (1976). In their study they presented relevant as well as irrelevant information at the beginning of a trial, but the response had to be withheld until a go-signal (tone) appeared at a variable delay. As expected, the Simon effect decreased with an increasing delay, and was absent at a delay of 250 ms. Vallesi and Umiltà (2009) replicated this result, but also found that the Simon effect did not decrease completely when the relevant information occurred after the irrelevant location information.

An important question with respect to the temporal-overlap idea is whether the location-induced activation decays passively, or whether some strategic suppression is necessary. Hommel (1994) investigated this question by manipulating not only the temporal overlap between irrelevant spatial activation and response selection, but also the proportion of congruent trials, which is known to affect the strategy of processing. Because no effect of proportion on the time-course of the Simon effect was found, Hommel (1994) concluded that spatial activation decays passively and not due any strategic suppression.

However, in contrast to the passive-decay hypothesis, Ridderinkhof (2002a, 2002b) proposed that irrelevant location-induced activation is suppressed actively. In his *activation-suppression account* he assumes that, to prevent unwanted responses to location, one needs to selectively inhibit the response code that is automatically activated by the irrelevant location. Because it is assumed that suppression builds up gradually, though, it is more effective for relatively slow than for fast responses. Moreover, the strength of suppression depends on inhibitory demands. For instance, if the proportion of incongruent trials increases (e.g. Stürmer, Leuthold, Soetens, Schröter, & Sommer, 2002), then selective suppression must be stronger. That suppression is indeed stronger under such conditions is reflected by a larger negative slope of the corresponding delta functions in the latencies (Ridderinkhof, 2002b).

Burle, van den Wildenberg, and Ridderinkhof (2005) tested the activation-suppression account by laterally presenting an irrelevant cue before, simultaneously with, or after a bilateral target. They hypothesized that activation followed by selective suppression should produce a negative effect when the location cue precedes the relevant information. Indeed, the Simon effect was positive for cues presented simultaneously with or after the target, but negative for cues appearing before the target. The fact that the Simon effect was negative if the spatial information was presented before the target strongly supports the idea that irrelevant activation is suppressed. At the same time, however, it questions the assumption that the strength of suppression depends on the inhibitory demands, at least that the strength is adjusted appropriately. Obviously, suppression was too strong in Burle et al.'s (2005) experiment, otherwise there should have been no negative Simon effect.

The aim of the present study was to further investigate the details of activation suppression, especially, to what extent its strength can be adjusted strategically. The fact that the Simon effect was negative in Burle et al.'s (2005) study indicates that activation suppression was stronger than needed. However, this does not imply that the strength of activation suppression cannot be adjusted strategically. It should be noted that the passive-decay hypothesis and the activation-suppression hypothesis are not mutually exclusive. Therefore, it is possible that both processes are involved in the Simon task, and that suppression is used to accelerate the decay of activation. Thus, if one assumes that location-induced activation always decays passively, then the inhibitory demand decreases with an increasing delay between the spatial cue and target. Accordingly, less suppression is needed for a long delay than for a

short one. However, for an appropriate adjustment of suppression the inhibitory demand must be known in advance. This was not the case in Burle et al. (2005), because the delays were randomized. In contrast, in a similar experiment with blocked delays, Vallesi and Umiltà (2009) observed only positive Simon effects. This suggests that the participants in Burle et al.'s study applied more suppression than needed to prevent the Simon effect, at least for some delays.

In the present study these ideas were tested directly by conducting two experiments in which the delay between a spatial cue and the target was varied. Our procedure was similar to that of Experiment 2 in Vallesi and Umiltà's (2009) study, where first a peripheral spatial cue was presented that was then replaced by the target after a certain delay. However, instead of producing a further onset signal with the appearance of the target, we presented the target by deleting one cue component. As Vallesi & Umiltà remarked, an onset-target might produce an additional and delay-independent activation of the spatial code, which reduces the delay effect.

More specifically, in our experiments we used left facing and right facing arrows (<, and >) as stimuli. Spatial cues were constructed by superimposing both arrow types and by presenting them at one side of the display (see Fig. 1). After a certain delay one of the two arrows was removed, and the participants had to indicate the pointing direction of the remaining one. By this procedure task-relevant information could be delayed without producing a second onset signal that might produce an unwanted refresh of the location-induced activation. Because with our procedure a delay of zero would have led to a qualitatively different location cue, we used only non-zero delays. However, our shortest delay was 33 ms, which might be considered as practically equivalent to a delay of zero.

In our first experiment, where the delays were randomized across trials, we tried to replicate the results of Burle et al. (2005) with our modified procedure. Then, for comparison, the delays were blocked in Experiment 2. If activation suppression is relatively inflexible, then the Simon effect should be similar in both experiments. However, if participants can use the delay information in the blocked condition to strategically adapt their suppression strength accordingly, then performance should differ. In both experiment performance was examined by analyzing not only mean data, but also delta functions for latencies and accuracy.

2. Experiment 1

In our first experiment several delays between a spatial cue and the target were randomized. First of all, we expected the Simon effect to decrease with an increasing delay. Moreover, if an appropriate adjustment of selective activation suppression should not be possible because of the randomized delays, then the Simon effect should be negative for the longer delays. The dynamics of activation suppression for the individual delays should be examined by also inspecting the corresponding delta functions.

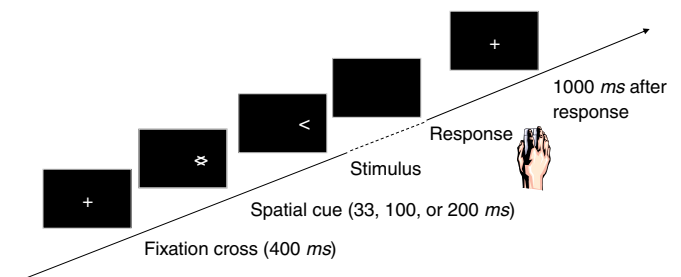


Fig. 1. Procedure of Experiment 1.

2.1. Method

2.1.1. Participants

17 students (mean age of 24 years; 5 male) from the Universität Konstanz, Germany, participated in the experiment. All had normal or corrected-to-normal vision and were paid 8 € for their participation.

2.1.2. Stimuli, apparatus, and procedure

Stimuli were right or left facing arrows (<, >). They were presented in white against a black background on an 18" color-monitor with a resolution of 1280 × 1024 pixels, and a refresh rate of 60 Hz. Participants were seated in front of the screen at a viewing distance of approximately 60 cm. Each trial started with the presentation of a fixation cross at the center of the screen for 400 ms, followed by a spatial cue consisting of two superimposed arrows, one facing to the left, the other facing to the right (see Fig. 1). The arrows appeared randomly at a distance of 2.2 cm left or right of the fixation cross, and were presented for 33, 100, or 200 ms, after which the task-irrelevant arrow was removed. These delays were chosen, because we expected the largest effects in this range (Simon et al., 1976). Delay was randomized across trials. The target arrow remained on screen until response. 1000 ms after the response the next trial started.

The task was to indicate the pointing direction of the target arrow by pressing one of the two mouse buttons (left button for left facing arrow; right button for right facing arrow) with the index or middle finger of the right hand, respectively. The participants were instructed to respond as fast as possible without making many errors. On every incorrect trial, an auditory feedback was provided. At the end of each block the mean error rate and RT in that block was displayed. If the mean error rate exceeded 10% the participants were asked to be more accurate.

There was 1 practice block (30 trials), in which no spatial pre-cue was presented, and 10 experimental blocks (96 trials/block). Every block contained an equal number of congruent and incongruent trials. On congruent trials, the cue (target) position was ipsilateral to the required response, while on incongruent trials it was contralateral.

2.2. Results

Responses faster than 100 ms or slower than 2000 ms were excluded from analysis (<0.2% of all data).

The latencies of correct responses were analyzed by a two-factor ANOVA for repeated measurements on the factors *congruency* (congruent, or incongruent), and *delay* (33, 100, or 200 ms). The analysis revealed a significant main effect of *delay*, $F(2, 32) = 18.9$, $p < .001$, $\eta_p^2 = .541$. RTs decreased with an increasing delay (412, 398, 391 ms). However, there was also a significant interaction between *delay* and *congruency*, $F(2, 32) = 12.2$, $p < 0.001$, $\eta_p^2 = .433$. The congruency effect was positive ($\Delta 14$ ms) for the 33 ms delay, but negative for the 100 and 200 ms delays ($\Delta -14$ ms, $\Delta -9$ ms). Congruency was significantly negative for the two longer delays, $F(1, 16) = 5.29$, $p < 0.05$, $\eta_p^2 = .252$.

Mean error rate was 6.02%. Subjecting the error rates to an ANOVA of the same type as for the latencies also revealed a significant main effect of *delay*, $F(2, 32) = 42.9$, $p < .001$, $\eta_p^2 = .728$. The error rate increased with an increasing delay (1.67%, 5.51%, and 10.9%). *Congruency* did not produce a reliable effect nor a significant interaction with *delay*. Numerically, however, there was a positive Simon effect for the shortest delay ($\Delta 0.69$), and a negative one for the longer delays ($\Delta -2.10\%$; $\Delta -1.52\%$).

2.2.1. Delta functions

To compute the delta functions for the latencies (Fig. 2), we first computed the cumulative distribution functions for the RTs of correct responses for each congruency condition (congruent, incongruent), and delay (33, 100, 200 ms) by quantile-averaging (.1, .3, .5, .7, .9)

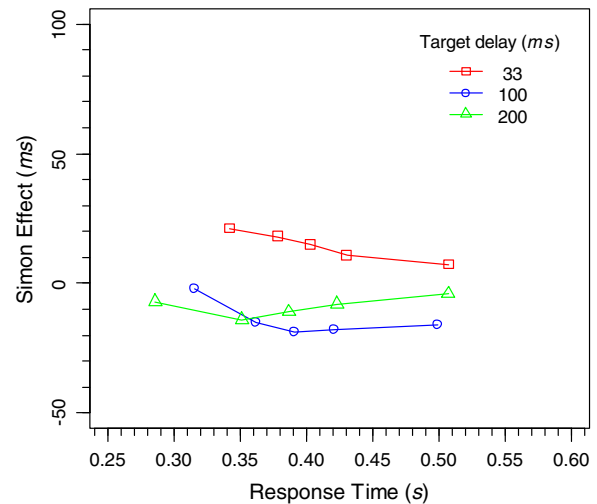


Fig. 2. Delta functions in the latencies of correct responses in the different target-delay conditions in Experiment 1.

the data (Ratcliff, 1979). Delta functions were then obtained by calculating the differences of the corresponding quantiles between the congruent and incongruent conditions and relating them to their respective average RTs (see Fig. 2). For a statistical analysis, the quantiles for the individual delay conditions and delays were entered into three-factor ANOVAs for repeated measurements on the factors *congruency* (congruent, or incongruent), *delay* (33, 100, or 200 ms), and *quantile* (1 to 5). We only report relevant results involving the factor *quantile*.

The analysis revealed a significant three-way interaction between all factors, $F(8, 128) = 2.69$, $p < 0.01$, $\eta_p^2 = .144$, indicating that the variation of the Simon effect with RT differed between the delay conditions. Further analyses showed that the Simon effect in the latencies changed significantly with RT for delay 33 ms, $F(4, 64) = 2.82$, $p < 0.05$, $\eta_p^2 = .150$, and for delay 100 ms, $F(4, 64) = 4.07$, $p < 0.01$, $\eta_p^2 = .203$, but not for delay 200 ms (see Fig. 2). Moreover, the variation did not differ reliably between the two shortest delays.

To obtain delta functions for accuracy, we first calculated conditional accuracy functions for each participant and condition. For this objective the corresponding data were sorted into five 20% bins. Then the proportion of correct responses and the mean RT for each bin were computed and the resulting values were averaged across participants. Delta functions were then constructed by calculating

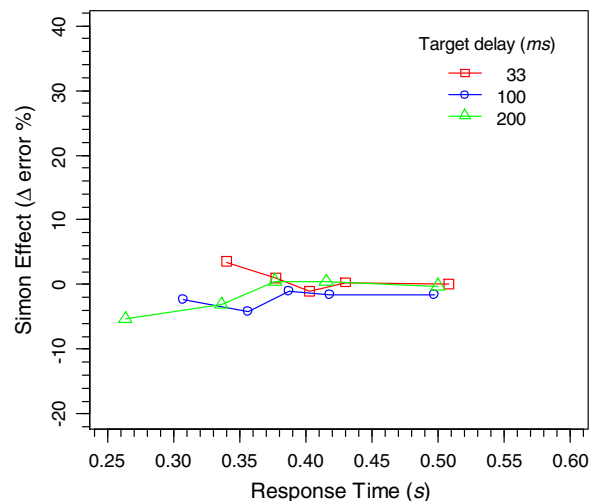


Fig. 3. Delta functions for accuracy in the different target-delay conditions in Experiment 1.

the differences in accuracy of the corresponding bins between congruent and incongruent conditions, and relating them to their averaged RT (see Fig. 3). Statistical analyses were performed analogously to those for the RT delta functions.

As can be seen in Fig. 3, the delta functions in accuracy were rather flat. Accordingly, there was no significant variation of the Simon effect with RT.

2.3. Discussion

The results of this experiment show that the delays had a substantial impact on the Simon effect. It was positive for the shortest delay, but negative for the two longer delays. Thus, our results are similar to those of Burle et al. (2005). The fact that the Simon effect was negative for the longer delays indicates that there was selective activation suppression. At the same time, however, the negative effects suggest that the strength of suppression was not optimal for each of the individual delays. Rather it seems that the participants chose a suppression strength that was presumably appropriate for the shortest delay, but too strong for the longer ones.

That selective activation suppression was rather strong is also reflected by the delta functions. If we consider the functions for the latencies (Fig. 2), then we see that for the shortest delay the Simon effect decreased monotonically with RT and almost reached zero for the slowest responses. For the 100 ms delay, however, overall suppression strength was so strong that the Simon effect was already absent for the fastest responses, and then became increasingly negative until it leveled off.

In the mean error rates no reliable Simon effect occurred, although numerically the pattern was similar to that in the mean RTs. If we consider the delta functions for accuracy (Fig. 3), then we see that the Simon effect was absent not only for the slow responses, but also for the faster ones. The fact that the Simon effect was practically absent even for the fastest responses indicates that impulsive responses were largely prevented right from the beginning of processing. Because selective suppression is assumed to affect mainly slow responses, this prevention must have been due to some other process. It is conceivable, for instance, that the susceptibility for spatial information was generally reduced (van den Wildenberg et al., 2010).

Taken together, the present results indicate that there was selective suppression, but that its strength was not adapted to the inhibitory demands of the individual delay conditions. However, this does not imply that suppression can principally not be adapted strategically. Rather, it could be that adaptation was not possible in the present case, because the delay for a given trial was not known in advance. Consequently, participants chose a strength that met the average demands across delays. Whether the strength of suppression can be controlled strategically, if the inhibitory demand is known in advance on each trial, was tested in the next experiment.

3. Experiment 2

This experiment was similar to our first one, except that delay was blocked. If the results in Experiment 1 were indeed due to a general inability to adapt the strength of selective suppression to the trial-based inhibitory demands, then the same results as in Experiment 1 should be observed. However, if the strength can be adjusted strategically, given the demands are known in advance on each trial, then blocking the delays should allow an appropriate adjustment. Accordingly, it can be expected that an individual and optimal suppression strength will be applied for each delay. If this is the case, and if we further assume that location-induced activation cannot fully be suppressed for the shortest delay, then the Simon effect should also decrease with an increasing delay. However, it should approach zero, and not be negative, even for the longest delay.

3.1. Method

17 students (mean age of 24 years; 3 male) from the Universität Konstanz, Germany, participated in the experiment. All had normal or corrected-to-normal vision and were paid 8 € for their participation. Apparatus and stimuli were same as in Experiment 1. The procedure was also similar, except that the delays were blocked. Additional to the practice block there were 5 experimental blocks (64 trials/block) for each of the three delays. The order of delays was balanced across participants.

3.2. Results

Responses faster than 100 ms or slower than 2000 ms were excluded from data analysis (<0.3% of all data).

The latencies of correct responses were subjected to a two-factor ANOVA for repeated measurements on the factors *congruency* (congruent, or incongruent), and *delay* (33, 100, or 200 ms). The analysis revealed significant main effects of both *congruency*, $F(1, 16) = 13.7$, $p < .01$, $\eta_p^2 = .462$, and *delay*, $F(2, 32) = 6.19$, $p < .01$, $\eta_p^2 = .279$. Responses were faster for congruent compared to incongruent stimuli (407 ms versus 423 ms), and speed increased with an increasing delay (422, 417, 407 ms). A further test revealed that the mean RT was also significantly different between the two longest delays, $F(1, 16) = 8.44$, $p < .05$, $\eta_p^2 = .345$. However, there was a significant interaction between *congruency* and *delay*, $F(2, 32) = 18.9$, $p < .001$, $\eta_p^2 = .541$, indicating that the congruency effect varied with delay ($\Delta 34$, $\Delta 3$, $\Delta 11$ ms). A further analysis revealed that the Simon effect was no longer significant when the shortest delay was excluded, $F(1, 16) = 2.00$, $p = 0.177$, $\eta_p^2 = .088$.

Mean error rate was 6.75%. Subjecting the error rates to an ANOVA of the same type as for the response times also revealed a significant main effect of *congruency*, $F(1, 16) = 22.0$, $p < 0.001$, $\eta_p^2 = .579$, and of *delay*, $F(2, 32) = 3.60$, $p < .05$, $\eta_p^2 = .184$. The error rate was smaller for congruent than for incongruent stimuli (4.97% versus 8.53%), and increased with delay (5.85%, 6.71%, 7.69%). However, *congruency* interacted significantly with *delay*, $F(2, 32) = 3.38$, $p < .05$, $\eta_p^2 = .175$, indicating that the congruency effect was larger for the 33 ms delay than for the other delays ($\Delta 5.82\%$, $\Delta 2.18\%$, $\Delta 2.70\%$). However, a further analysis revealed that the Simon effect remains significant after excluding the shortest delay, $F(1, 16) = 4.58$, $p < 0.05$, $\eta_p^2 = .223$.

3.2.1. Delta functions

Delta functions were computed and analyzed in the same way as in Experiment 1. The analysis revealed a significant three-way interaction between all factors, $F(8, 128) = 3.73$, $p < 0.001$, $\eta_p^2 = .189$, which indicates that the variation of the Simon effect with RT differed between the delay conditions. Further analyses showed that the variation was significant for delay 100 ms, $F(4, 64) = 4.40$, $p < 0.01$, $\eta_p^2 = .216$, and marginally significant for delay 33 ms, $F(4, 64) = 2.27$, $p = 0.071$, $\eta_p^2 = .124$. The Simon effect did not vary reliably for delay 200 ms.

Fig. 5 shows the delta functions for accuracy. As can be seen, the Simon effect was relatively large for the fastest responses, especially for the shortest delay. It then decreased quickly with RT and approached zero. The significant three-way interaction between all factors, $F(8, 128) = 3.68$, $p < 0.001$, $\eta_p^2 = .187$, indicates that the decrease of the Simon effect in accuracy with RT differed between the delay conditions. Further analyses revealed that it was significant for delay 33 ms, $F(4, 64) = 23.4$, $p < 0.001$, $\eta_p^2 = .594$, and for delay 100 ms, $F(4, 64) = 5.96$, $p < 0.01$, $\eta_p^2 = .271$, but not for delay 200 ms.

3.2.2. Comparison with Experiment 1

For comparison, the mean data of the present experiment were combined with those of Experiment 1 and subjected to three-factor ANOVAs for the within-participant factors *congruency* (congruent, or

incongruent), and *delay* (33, 100, or 200 ms), and the between-participant factor *experiment* (randomized delay, or blocked delay). We report only results involving the factor *experiment*.

For the latencies the analysis revealed a significant interaction between *experiment* and *congruency*, $F(1, 32) = 11.9$, $p < .01$, $\eta_p^2 = .270$. The Simon effect was smaller for randomized delays than for blocked ones. This is even the case when we compare the absolute effect sizes, i.e. ignore their sign, $F(1, 32) = 5.75$, $p < .05$, $\eta_p^2 = .152$. Moreover, the interaction between all three factors was far from significance, $F(2, 64) = .107$, $p = .899$, $\eta_p^2 = .003$. This implies that the Simon effect for the shortest delay was also smaller under randomized delays, compared to blocked ones, which we nevertheless confirmed separately, $F(1, 32) = 16.5$, $p < .001$, $\eta_p^2 = .340$.

For the error rates the analysis also revealed a significant interaction between *experiment* and *congruency*, $F(1, 32) = 15.7$, $p < .001$, $\eta_p^2 = .330$. The overall Simon effect was small and negative for randomized delays, but large and positive for blocked ones. Moreover, there was an interaction between *experiment* and *delay*, $F(2, 64) = 18.8$, $p < .001$, $\eta_p^2 = .371$. This was due to the fact that the error rate increased only slightly with delay in the blocked delay condition, but substantially in the randomized condition.

We also examined the differences of the delta functions between the two experiments by comparing the corresponding functions for each delay. In the latencies, merely the functions for delay 100 ms differed significantly, $F(4, 128) = 3.23$, $p < .05$, $\eta_p^2 = .092$. In accuracy, the functions for delay 33 ms differed reliably, $F(4, 128) = 16.0$, $p < .001$, $\eta_p^2 = .333$, as did the functions for delay 100 ms, $F(4, 128) = 3.18$, $p < .05$, $\eta_p^2 = .091$.

3.3. Discussion

Our results show that the Simon effect also varied with blocked delays. However, different from Experiment 1, the effect remained positive. There was a substantial Simon effect for the shortest delay, whereas for the longer delays it was reliable only in the error rates. Most importantly, it remained positive. This is in contrast to Burle et al.'s (2005) results and our Experiment 1, where strong negative Simon effects were found. Because delay was blocked in the present experiment, the results support our idea that the strength of suppression can be adjusted according to the inhibitory demands for the individual delay conditions, but only if the delay is known in advance.

Different from our expectation, though, the Simon effect did not monotonically decrease with an increasing delay. Rather, at least numerically, the effect was stronger for the 200 ms delay than for the 100 ms delay. However, because mean RT was reliably shorter for the 200 ms delay, compared to the 100 ms delay, this tendency could have been due to some kind of tradeoff.

The variation of the Simon effects with RT for the individual delays can be seen by inspecting the corresponding delta functions (Figs. 4 and 5). The functions for the latencies show that the Simon effect for the shortest delay first increased with RT and then decreased. However, it remained relatively strong across the whole RT range. These data are similar to those of Baroni, Pellicano, Lugli, Nicoletti, and Proctor (2012), who also presented lateralized arrows as targets, but without a prior spatial cue. The similarity between these data sets confirms our assumption that our shortest delay is practically equivalent to a delay of zero. For the longer delays the Simon effect was smaller, compared to the short delay, and the corresponding delta functions were concave rather than convex. That is, the effect first decreased to a minimum and then increased. However, this variation was reliably only for delay 100 ms.

If we consider the delta functions for accuracy (Fig. 5), then we see that they are rather different from those in the previous experiment. The Simon effect was rather large for the fastest responses, especially for the shortest delay. However, the effect quickly vanished with RT for all conditions. The large Simon effect in accuracy for fast responses

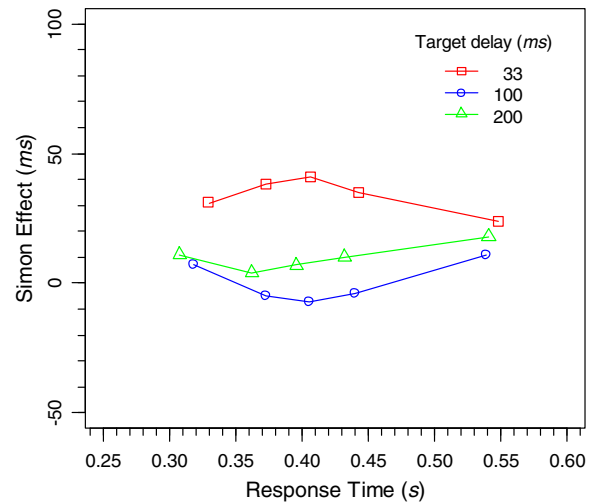


Fig. 4. Delta functions for latencies of correct responses in the different target-delay conditions in Experiment 2.

indicates that the readiness for responding impulsively to irrelevant spatial information was relatively large, compared to our first experiment with randomized delays.

4. General discussion

The aim of the present study was to further investigate selective activation suppression (Ridderinkhof, 2002a, 2002b) in the Simon task, especially to what extent its strength can strategically be adjusted to inhibitory demands. We assumed that location-induced activation automatically decays with time and that, therefore, the inhibitory demand decreases with an increasing temporal separation of irrelevant activation and response selection (Hommel, 1993, 1994). Thus, one way to modulate the inhibitory demand is to vary the delay between an irrelevant spatial cue and the target. By applying such a procedure, Burle et al. (2005) found a strong negative Simon effect when the cue preceded the target, which speaks for selective suppression, but questions that it can strategically be adjusted optimally. However, in that study the delays were randomized so that no information about the inhibitory demand on a given trial was available in advance. In a similar experiment but with blocked delays Vallesi and Umiltà (2009) found that the Simon effect remained positive. Thus,

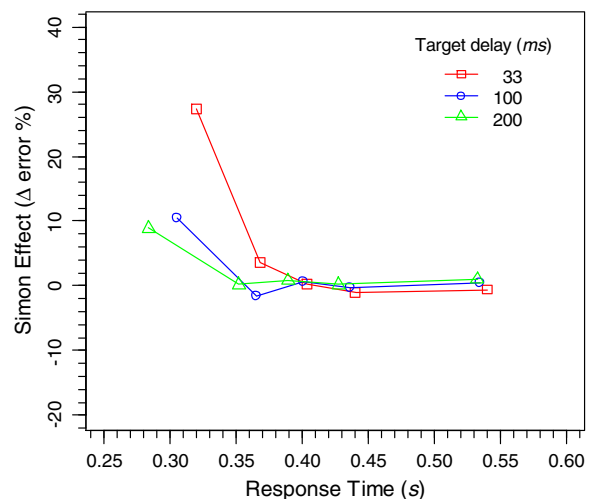


Fig. 5. Delta functions for accuracy in the different target-delay conditions in Experiment 2.

our idea was to examine whether strategic suppression is possible if such information is provided.

In our first experiment, three delays were randomized in a block of trials to see whether Burle et al.'s (2005) results can be replicated with a slightly modified procedure (e.g., offset stimuli and different delays). As a result, the Simon effect was positive only for the shortest delay, but negative for the two longer delays, confirming Burle et al.'s outcome. Thus, our data also indicated that location-induced activation was not suppressed in dependence of the inhibitory demands of the individual delays. Rather, it seemed that the same strength of suppression was applied for all delays. The strength was largely appropriate for the shortest delay, but for the longer delays, which required less inhibition, it was too high.

If location-induced activation can be suppressed strategically, but only if the inhibitory demands are known in advance, then the Simon effect should vary differently with delay when the delays are blocked. Whether this is the case was tested in Experiment 2. The results show that the Simon effect was again large and positive for the shortest delay. However, and most importantly, although the effect also decreased for the longer delays, it remained positive. This supports the idea of strategic activation suppression. The result can be interpreted in the sense that selective suppression was relatively strong for the shortest delay. Nevertheless, because suppression takes some time to come into effect, and is probably also limited in strength, it could not prevent that irrelevant activation strongly affected response selection. For the longer delays, the demands were lower, because location-based activation had partly decayed and suppression had more time to act. Since the participants knew the demands in advance, they could adjust their suppression strength accordingly, so that the Simon effect was absent in RT and relatively weak in the error rates.

How selective suppression proceeded in the different experiments and conditions can, at least partly, be inferred from the delta functions. If we consider Fig. 4, then, taken together, the functions for the different delays suggest that irrelevant spatial activation increased after cue onset and was then suppressed for some time. Interestingly, with randomized delays the dynamics of suppression was rather similar (see Fig. 2). It merely seems that suppression started somewhat earlier and continued for a longer period, which suggests that participants had some control of the timing of suppression. The most striking difference of the delta functions for the latencies, however, was that they were shift downwards for randomized delays (Experiment 1). This could indicate that suppression was generally stronger in this case. However, given that selective suppression takes some time to take effect, as assumed by the activation–suppression model, one would have expected a larger difference in shape of the delta functions between the experiments, especially for slow responses. Yet, only one (delay 100 ms) of the three delta functions differed significantly between the experiments.

Thus, it seems that some other mechanism was also involved in producing the differences in performance between the experiments. It has been suggested that the readiness for making fast impulsive responses to irrelevant information is also under voluntary control, at least to some extent, and that this readiness is primarily reflected by the shape of the delta functions for accuracy (e.g. van den Wildenberg et al., 2010). The more impulsive responses to location are allowed, the larger the congruency effect in the error rates for fast responses. That the readiness for impulsive responses indeed differed between our experiments can be seen by comparing the corresponding delta functions for accuracy. For randomized delays there were only few impulsive responses to location (see Fig. 3), i.e. the Simon effect was practically absent even for the fastest responses, whereas for blocked delays many such fast responses occurred, which produced a correspondingly large Simon effect (Fig. 5), especially for the shortest delay.

Alternatively, it is conceivable that under randomized delays the flow of information along the direct route was generally blocked.

Such a mechanism would not only account for the different patterns of the delta functions for accuracy between experiments, but also for the vertical shift of the delta functions for the latencies, i.e. for the fact that the Simon effect was generally smaller under randomized delays. In any case, the analyses of the delta functions strongly suggest that two mechanisms were applied by the participants to adapt to the different demands in our experiments.

Thus, taken together, our results show that suppression strength can be adjusted according to the demands, given they are known in advance. Moreover, they suggest that also the readiness to respond impulsively to location is under strategic control, and that this readiness is reduced if there is uncertainty about target arrival. However, our results also indicate that the different adjustment do not necessarily produce the smallest possible Simon effects. The comparison of the Simon effect between the two experiments revealed that it was reliably larger under blocked delays than under randomized ones. This was even the case when we consider the absolute effect sizes (i.e. ignore the sign). Thus, the possibility to adjust suppression to the individual delays did not lead to a smaller overall Simon effect, as one might have expected, but to a larger one.

How can this unexpected result be explained? Implicitly, one might have assumed that the participants' goal was to prevent the Simon effect. This, however, was obviously not the case. Rather, our data suggest that the participants' goal was to meet our requirements concerning accuracy. This makes sense, because, first of all, they presumably did not notice the size of the Simon effect. Furthermore, what they certainly noticed, also due to our feedback procedure, was their error rate. Accordingly, the overall error rate was rather similar between the two experiments (6% versus 7%). For the blocked delays (Experiment 2), a similar error rate also occurred for the individual delays (6%, 7%, 8%). For the randomized delays (Experiment 1), however, the individual error rates differed substantially, i.e. they increased considerably with delay (2%, 6%, 11%).

These results indicate that location-induced activation had to be utilized to a lesser extent and suppressed more strongly in the random condition to perform as reliably, on average, as in the blocked condition, which also resulted in a smaller Simon effect. This interpretation also explains why the Simon effect for the shortest delay was larger in the blocked condition than in the randomized one. Although it was obviously possible in the blocked condition to suppress location-induced activation more strongly, it was not necessary, because the required accuracy was already achieved with less suppression. Moreover, it should be noticed that activation suppression also produces costs on congruent trials. Thus, it should not always be maximized. These considerations suggest that selective activation suppression was not used in our experiments to minimize the Simon effect, but to control the level of accuracy. Nevertheless, our results support the idea that activation suppression can be adjusted strategically.

To conclude, the present study demonstrates that the effect of irrelevant location information in the Simon task can strategically be controlled, depending on the situational demands. Our results suggest that two mechanisms are applied. One mechanism, selective activation suppression, develops in time and is used to reduce the irrelevant activation. This mechanism can effectively be adapted to the specific demands of individual delays, given they are known in advance. The other mechanism controls the utilization of location information. The use of location information for response selection is reduced if the adaptation of the first mechanism to individual conditions is largely restricted, which is the case with randomized delays. In any case, the primary goal of these strategic modulations is not to minimize the Simon effect, but to meet the situational requirements with respect to accuracy.

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