PDF hosted at the Radboud Repository of the Radboud University Nijmegen

The following full text is a publisher's version.

For additional information about this publication click this link.
http://hdl.handle.net/2066/125880

Please be advised that this information was generated on 2019-02-09 and may be subject to change.
Behavioral/Cognitive

Saccade Target Selection Relies on Feedback Competitive Signal Integration

Joke P. Kalisvaart, André J. Noest, Albert V. van den Berg, and Jeroen Goossens

Radboud University Nijmegen Medical Centre, Donders Institute for Brain, Cognition and Behaviour, Department of Cognitive Neuroscience, Section Biophysics, 6500 HB Nijmegen, The Netherlands, and Developmental Biology Department, Utrecht University, 3584 CH Utrecht, The Netherlands

It is often assumed that decision making involves neural competition, accumulation of evidence “scores” over time, and commitment to a particular alternative once its scores reach a critical decision threshold first. So far, however, neither the first-to-threshold rule nor the nature of competition (feedforward or feedback inhibition) has been revealed by experiments. Here, we presented two simultaneously flashed targets that reversed their intensity difference during each presentation and instructed human subjects to make a saccade toward the brightest target. All subjects preferentially chose the target that was brightest during the first stimulus phase. Unless this first phase lasted only 40 ms, this primacy effect persisted even if the second, reversed-intensity phase lasted longer. This effect did not result from premature commitment to the initially dominant target, because a strong target imbalance in the opposite direction later drove nearly all responses toward that location. Moreover, there was a nonmonotonic relation between target imbalance and primacy: increasing the target imbalance beyond 40 cd/m² caused an attenuation of primacy. These are the hallmarks of hysteresis, predicted by models in which target representations compete through strong feedback. Reaction times were independent of the choice probability. This dissociation suggests that target selection and movement initiation are distinct phenomena.

Introduction

To explain how saccadic responses compete for selection and execution, current decision-making theories assume that sensory evidence is noisy and that it is accumulated over time to reach a decision bound (Ratcliff and McKoon, 2008). Different types of models make different assumptions about how evidence is combined (Smith and Ratcliff, 2004): “evidence scores” for two or more alternatives can accumulate independently (Cousineau, 2004), compete with feedforward inhibitory interaction (Laming, 1966; Link and Heath, 1975; Nosofsky and Palmeri, 1997; Palmeri, 1997), or compete via feedback inhibition (Usher and McClelland, 2001; Wang, 2002; Wong and Wang, 2006). So far, it has been impossible to single out one among the many types of visual decision-making models: despite different architectures, they predict similarly optimal choice behavior in static two-alternative forced choice tasks (model mimicry; van Zandt and Ratcliff, 1995; Bogacz et al., 2006).

To distinguish between feedforward and feedback cross-inhibition, we introduced a new manipulation: we studied saccadic choices between pairs of briefly flashed targets the intensities of which were swapped during presentation (Fig. 1A,B). Under such conditions, circuits with feedback competition that exceed a certain critical strength are sensitive to the initial stimulus bias but relatively resistant to the later bias reversal. This characteristic behavior (hysteresis) occurs because the initial “winner” maintains its advantage by recurrent inhibition of its competitor (Noest et al., 2007; Furman and Wang, 2008). These models thus predict that subjects typically choose the target with the highest initial strength provided that the initial differences outlast the time constant of the feedback creating the hysteresis and the stimuli are not strong enough to overcome this hysteresis (Fig. 1C). In contrast, feedforward integration models predict that for balanced durations of the two stimulus epochs, early and late biases are equally effective (because the total evidence for both targets is the same), whereas leaky-integrator models even predict that later biases dominate the final evidence scores. This implies that subjects should either show no preference at all or show a preference for the target that is strongest at the end, unless they make an irreversible choice for the initially strongest target before the stimulus ends (premature choice commitment). Indeed, all accumulator models can produce primacy (i.e., a choice preference for the initially strongest target) if they include some thresholding mechanism that induce choice commitment. We probed the contribution of such thresholding mechanisms by appending strong intensity biases toward the initially weakest target.

We found robust preferences for the initially strongest target, which reversed only if the first stimulus epoch was sufficiently short and the second epoch outlasted the first one by a sufficient amount. The basic primacy effect was attenuated at larger stimulus biases and could even be inverted completely if the stimuli were immediately followed by a strong intensity bias toward the
and the overall stimulus durations were systematically varied. In a modified version of this paradigm, the intensities of the left and right target were reversed (reversal trials) or remained unaltered (control trials, data not shown). In either case, both targets remained present for another 40–120 ms until the screen turned black. For each reversal condition, control stimuli comprised of different intensities (ΔI) appeared. After a delay of 40–120 ms (D2), the intensities of the left and right target was the same only when the durations of the first and second epoch were the same. High-to-low and low-to-high intensity reversals could be unambiguous or ambiguous. In reversal trials, the mean intensity of the left and right target remained unaltered. In reversal trials, however, the intensities of the left and right target were reversed compared with the first target epoch. In both cases, targets were then displayed for another 40, 80, or 120 ms (D3) until the screen turned black. For each reversal condition, control stimuli were presented with the same total duration (i.e., D1 + D3).

Subjects were always instructed to first look at the fixation ring and then make a saccade to the most intense target as quickly and accurately as possible. They received no feedback about their performance. Target intensities and durations of the first and second target presentation epoch were manipulated systematically across trials and experimental sessions.

Target intensities in Experiments 1–3 ranged from 4.5 to 36.6 cd/m². Background luminance was 0.273 cd/m². The intensity difference (ΔI) between the left and the right target on any given trial could be large (4.5 vs 36.6 cd/m²), medium (7.8 vs 29.1 cd/m²), small (12.3 vs 22.4 cd/m²), or zero (both targets 16.7 cd/m²). Thus, in control trials, target luminance could be unambiguous or ambiguous. In reversal trials, the mean intensity of the left and right target was the same only when the durations of the first and second epoch were the same. High-to-low and low-to-high intensity changes were achieved with a time constant of ~10 ms, limiting the shortest practical presentation time to 40 ms.

Experiments 1–3 established the conditions under which primacy arises, but could not discriminate between two very different mechanisms that can generate primacy: hysteresis due to strong feedback competition or premature commitment due to an absorbing bound (see Introduction). Testing whether primacy can still be reverted does discriminate between two very different model types. Primacy due to absorbing bounds is by definition irreversible and should increase monotonically with increasing ΔI. Conversely, primacy due to hysteric integration dynamics should be reversible by a sufficiently strong stimulus. In Experiment 4, we therefore presented reversal stimuli across an extended range of target contrasts (i.e., from 22.5 vs 33.6 cd/m² up to 1.5 vs 77.5 cd/m²). The durations D1 and D2 were kept fixed at 50 ms. Targets in the 100 ms

Materials and Methods

Subjects

Ten adult human subjects (5 male, 5 female) participated in the experiments. All had normal or corrected-to-normal visual acuity. Subjects were informed about the experimental procedures and gave informed consent before the start of the experiments. Procedures were approved by the Radboud University Medical Centre. Subjects J.R., V.G., J.G., and J.K. were experimenters; all other subjects were kept unaware of the aim of the study.

Setup

Subjects were seated in a darkened room at 80 cm of a projection screen on which stimuli were back projected. In the first three experiments, an LCD projector (model DLA-S10E; JVC.) with a refresh rate of 75 Hz and a maximum luminance level of ~45 cd/m² was used. To present targets at higher contrast and luminance levels in Experiment 4, we used a digital light processing projector with a maximum luminance level of ~300 cd/m² and a refresh rate of 60 Hz (P1265 model DXN0702; Acer). Luminance levels were measured with a luminance meter (model LS-100; Minolta).

A chinrest was used to minimize head movements. Eye movements were measured with the scleral search coil technique (Remmel Laboratories). Coils were inserted after one drop of topical anesthetic (oxybuprocaine hydrochloride 0.4%; Thea Pharma). Once the coil was in place, a drop of artificial tear (methylcellulose 0.5%; Thea Pharma) and a bandage lens (a large contact lens with a strength of zero dioptries) were applied to minimize ocular discomfort. Use of the bandage lens doubled the measuring time with the coil to ~1 h per session (Sprunger et al., 2008). Eye position signals were low-pass filtered, amplified, and sampled at 500 Hz per channel. The spatial resolution of the horizontal and vertical eye position signals was better than 0.1° (root mean square measure).

Paradigms

In the first three experiments, each trial consisted of four epochs (Fig. 1). At the beginning of each trial, a fixation ring with diameter of 0.5° was presented at the center of the screen. Then, after a random period of 400–1200 ms, the fixation ring disappeared and two filled, circular targets with a diameter of 0.5° were presented simultaneously at 10° to the left and right of the center. After a variable delay (D1) of 40, 80, or 120 ms, this first set of targets was replaced by a second set of targets at the same locations. In control trials, the intensity of both the left and right target remained unaltered. In reversal trials, however, the intensities of the left and right target were reversed compared with the first target epoch. In both cases, targets were then displayed for another 40, 80, or 120 ms (D3) until the screen turned black. For each reversal condition, control stimuli were presented with the same total duration (i.e., D1 + D3).

Subjects were always instructed to first look at the fixation ring and then make a saccade to the most intense target as quickly and accurately as possible. They received no feedback about their performance. Target intensities and durations of the first and second target presentation epoch were manipulated systematically across trials and experimental sessions.

Target intensities in Experiments 1–3 ranged from 4.5 to 36.6 cd/m². Background luminance was 0.273 cd/m². The intensity difference (ΔI) between the left and the right target on any given trial could be large (4.5 vs 36.6 cd/m²), medium (7.8 vs 29.1 cd/m²), small (12.3 vs 22.4 cd/m²), or zero (both targets 16.7 cd/m²). Thus, in control trials, target luminance could be unambiguous or ambiguous. In reversal trials, the mean intensity of the left and right target was the same only when the durations of the first and second epoch were the same. High-to-low and low-to-high intensity changes were achieved with a time constant of ~10 ms, limiting the shortest practical presentation time to 40 ms.

Experiments 1–3 established the conditions under which primacy arises, but could not discriminate between two very different mechanisms that can generate primacy: hysteresis due to strong feedback competition or premature commitment due to an absorbing bound (see Introduction). Testing whether primacy can still be reverted does discriminate between two very different model types. Primacy due to absorbing bounds is by definition irreversible and should increase monotonically with increasing ΔI. Conversely, primacy due to hysteric integration dynamics should be reversible by a sufficiently strong stimulus. In Experiment 4, we therefore presented reversal stimuli across an extended range of target contrasts (i.e., from 22.5 vs 33.6 cd/m² up to 1.5 vs 77.5 cd/m²). The durations D1 and D2 were kept fixed at 50 ms. Targets in the 100 ms

Materials and Methods

Subjects

Ten adult human subjects (5 male, 5 female) participated in the experiments. All had normal or corrected-to-normal visual acuity. Subjects were informed about the experimental procedures and gave informed consent before the start of the experiments. Procedures were approved by

initially weakest target. This near-complete transition from primacy to recency shows that the observed primacy cannot be attributed to premature commitment to the initially strongest target. Simulations showed that all these results were best described by a feedback cross-inhibition model. Reaction times were independent of the changes in choice probability and instead depended on the initial intensity of the selected target.
ambiguous condition had an intensity of 27.8 cd/m². Control stimuli lasted 50 ms and target contrasts ranged from 7.2 versus 49.5 cd/m² to 26.0 versus 29.7 cd/m². Background luminance was 0.18 cd/m².

To further probe the extent to which primacy can be undone by later stimulus evidence, Experiment 4 also included two forced reversal conditions in which the stimulus presentation ended with a bright pulse at the location of the initially weakest target. More specifically, in the 50-50–67 ms forced reversal condition, a 50–50 ms reversal stimulus was immediately followed by a third, 67 ms epoch in which there was a strong bias toward the initially weakest target. In the 83–67 ms forced reversal condition, 83 ms of initial bias was immediately followed by 67 ms of reversed extreme bias. In both cases, the initial bias was 49.5 versus 7.2 cd/m² and the final, reversed bias was 7.2 versus 283 cd/m². For subjects J.K. and J.G., we also tested forced reversal conditions with an initial bias of 13.7 versus 40.0 cd/m² and 3.6 versus 62.7 cd/m², respectively, being nullified in Experiment 4 challenge mechanisms that attribute our primacy effects to premature commitment caused by absorbing bounds, we consider the class of accumulator models that have absorbing bounds but no hysteresis in their evidence accumulation. This whole class of models can then be tested by comparing the forced reversal conditions with their unforced counterparts. Key in this analysis is the following notion: at the end of $D_o$ (which is either the end of the whole stimulus or the start of forcing $D_f$ phase) a fraction of the accumulated-evidence trajectories already reached the boundary of the initially strongest target (unknown fraction $A_o$) or the boundary for the other target (unknown fraction $A_f$), leaving only a limited fraction (1 − $A_o$ − $A_f$) of formally “undecided” trajectories that can still be influenced by a later forcing stimulus (Fig. 5A). If we knew for sure that the bright $D_f$ forcing pulse were strong enough to drive all “surviving” trajectories to (or closest to) the boundary of the initially weakest target, the probability of still choosing the initially strongest target in the forced reversal condition would provide a direct measure of $A_o$. Unfortunately, we cannot be sure that the forcing stimulus is indeed sufficiently strong to achieve this. We can, however, show that the paired probabilities from the two experimental conditions must satisfy an inequality that—if violated by our data—rejects this whole class of bounded accumulator models:

Let $F_o$ denote the probability of still choosing the initially strongest target in the forced reversal condition. This observed probability then sets the upper limit for the unknown fraction of trials, $A_o$, in which the accumulated-evidence trajectories reached the absorbing bound of the initially strongest target before the end of $D_o$. Note that we can only conclude that $F_o \geq A_o$ because, as we noted above, we cannot be sure that the $D_f$ forcing stimulus is strong enough. However, from the time- and polarity-antisymmetric nature of the reversal stimulus and from the observed behavior in the (unforced) reversal condition, we can now derive further constraints on $A_o$.

For simplicity, we first assume absence of response biases. This assumption is eventually lifted, because it can be shown to only make our test more conservative. Now let $R_o$ denote the observed probability of choosing the initially dominant target in the (unforced) reversal condition. Next, keep in mind that $R_o$ is the sum of $A_o$ and the probability that an undecided trajectory ends up being closer to the boundary of the initially strongest target (Fig. 5A). Next, we know that $A_o \approx A_f$. This holds because at the end of $D_o$, the trajectories tend to be closer to the boundary of the initially brightest target (as verified by the control condition). Therefore, given that $D_f = D_o$ and given the same (but reversed) $\Delta I$, a drift to the other boundary during $D_f$ takes more time. This, in combination with the assumed lack of hysteresis and bias in this model class, implies that at most one half of the undecided trajectories can eventually contribute to choosing the initially strongest target. That gives us the starting point for some algebra which relates $R_o$ to $F_o$:

$$R_o \leq A_o + \frac{1}{2} (1 − A_o − A_f) = \frac{1}{2} (1 + A_o − A_f) > \frac{1}{2} (1 + F_o)$$

(3)

or:

$$F_o \geq 2 R_o − 1$$

(4)

Therefore, we can test whether the measured $R_o$ and $F_o$ combinations violate this inequality.

The only remaining step required for the inequality test of Equation 4 is to account for the effect of choice biases that were clearly present in our data. We can do this by averaging the “raw” $R_o$ and $F_o$ values plotted in Figure 4 for the two mirror-symmetric stimulus conditions per subject. This results in more bias-resistant measures of $R_o$ and $F_o$:

$$\bar{R}_o = \frac{1}{2} (R_o^+ + 1 − R_o^-)$$

and

$$\bar{F}_o = \frac{1}{2} (F_o^+ + 1 − F_o^-)$$

(5)

Where the $^+$ and $^−$ superscripts refer to initial stimulus biases of $+\Delta I$ and $−\Delta I$, respectively. Using the modified $\bar{R}_o$ and $\bar{F}_o$ variables in this test.
not only reduces the effect of subject-dependent biases (which are difficult to estimate precisely), but the remaining “second-order” bias effects only makes our test more conservative (at least in a wide regime of practical interest, \( \tilde{F}_i < 0.5 \), because of convexity properties of the psychometric functions).

SEs for \( \tilde{R}_i \) and \( \tilde{F}_i \) were obtained from error-propagation rules:

\[
SE(\tilde{R}_i) = \sqrt{\frac{\tilde{R}_i (1 - \tilde{R}_i)}{4N_R} + \frac{R_i (1 - R_i)}{4N_F}}
\]

\[
SE(\tilde{F}_i) = \sqrt{\frac{\tilde{F}_i (1 - \tilde{F}_i)}{4N_R} + \frac{F_i (1 - F_i)}{4N_F}}
\]

Where the \( N \)s denote the total numbers of trials per test condition, with the subscripts \( R \) and \( F \) referring to the reversal conditions and forced reversal conditions, respectively, and with the + and − superscripts referring to the two different intensity biases as before. Using normal approximation, we then tested the null hypothesis from Equation 4 against the alternative hypothesis (i.e., \( F_i < 2 \tilde{R}_i - 1 \)) by applying the following one-sided \( z \) test:

\[
z = \frac{2\tilde{R}_i - 1 - \tilde{F}_i}{SE}
\]

where \( SE = \sqrt{A \cdot SE(\tilde{R}_i)^2 + SE(\tilde{F}_i)^2} \) (7)

Chronometric response functions. Saccade latencies were also analyzed as a function of \( \Delta I \). For each subject, we first subtracted the average reaction time measured in the ambiguous control condition pooled across rightward and leftward saccades. This was done because the average latency relative to target onset differed greatly among subjects (range: 250–600 ms). The resulting chronometric response functions were quantified by fitting linear regression lines to the data:

\[
\Delta RT = \alpha \cdot \Delta I + \beta
\]

One-way ANCOVA, with \( \Delta I \) as the predictor and \( \Delta RT \) (in ms) as the response and condition as the grouping variable, was used to test for differences between the chronometric curves. Correction for multiple testing was achieved by using a Tukey–Kramer test for multiple comparisons.

Results
Balanced durations
In the first experiment, two targets were presented simultaneously for 80 or 160 ms, and the initial intensity bias reversed halfway during stimulus presentation (i.e., \( D_1 = D_2 = 40 \) ms or \( D_1 = D_2 = 80 \) ms). In control trials, the intensities remained the same across \( D_1 \) and \( D_2 \). Figure 2 shows the probability of rightward saccades as a function of the initial rightward intensity bias (\( \Delta I \), positive if the right target was brightest at stimulus onset). In both control conditions (Fig. 2A, C), subjects discriminated target intensities easily; the psychometric curves had steep, positive slopes and saccades were almost always directed toward the brightest target at the largest intensity differences (\( \Delta I = \pm 36 \) cd/m\(^2\)). For purely ambiguous control stimuli (\( \Delta I = 0 \) cd/m\(^2\)), choice probabilities scattered around 50%. The psychometric curves obtained in the 80–80 ms reversal condition (Fig. 2B) also had steep, positive slopes, indicating a significant preference for the initially strongest target in all subjects (\( t \) test, \( p < 0.0001 \)). Subjects even maintained a clear preference for the initially brightest target in the 40–40 ms reversal condition (Fig. 2D; slopes > 0; \( t \) test, \( p < 0.001 \) for all subjects). Compared with the control conditions, the slopes were somewhat reduced (80–80 ms condition: \( t \) test, \( p < 0.05 \) for all subjects), or reduced considerably (40–40 ms reversal condition: \( t \) test, \( p < 0.01 \) for all subjects except V.G.), indicating that target selection was not exclusively based on the initial intensity differences; subsequent stimulus information also had a significant influence on the eventual decision, even though the changes in target luminance were not consciously perceived.

Unbalanced durations
To probe further the contributions of initial and later stimulus epochs, two additional experiments were performed in which durations \( D_1 \) and \( D_2 \) were manipulated. In Experiment 2, we first kept the total stimulus duration fixed at \( D_1 + D_2 = 160 \) ms and swapped the targets’ intensities at \( D_1 = 40, 80, \) or 120 ms after stimulus onset. Choices in the control condition (Fig. 3A) and 80–80 ms reversal condition (Fig. 3C) replicated the primacy effects of Experiment 1 with more subjects. In the new 120–40 ms reversal condition (Fig. 3B), that is, if \( D_1 \) lasted three times longer than \( D_2 \), choice behavior became almost indistinguishable from the control condition in all subjects. In contrast, if \( D_1 \) was three times shorter than \( D_2 \) in the 40–120 ms reversal condition, the subjects’ choice behaviors diverged considerably (Fig. 3D). Three subjects (I.B., A.M., and J.K.) still showed a significant, albeit weaker primacy effect (slopes > 0; \( t \) test, \( p < 0.05 \)), but the other four subjects (J.R., V.G., D.B., and D.A.) now showed a significant preference for the target that was most intense at the end (slopes < 0; \( t \) test, \( p < 0.01 \)). However, even in these four subjects, the choice probabilities were not completely inverted compared with the 120–40 ms condition. To summarize these changes, Figure 3E shows the averaged slopes of the psychometric curves as a function of \( \Delta D = D_1 - D_2 \). Note the systematic increase in slopes with increasing \( \Delta D \). The asymmetric effect of intensity difference
in $D_1$ versus $D_2$ is reflected in the fact that the trend line does not pass through the origin.

Because Experiment 2 kept the total duration constant, one cannot conclude that $\Delta D$ is the only or even the mechanistically relevant variable controlling the choice process. Indeed, strong hysteresis is only expected if $D_1$ exceeds the (probably subject-dependent) feedback time constant. To test this, Experiment 3 used the smallest practical $D_1$ (40 ms) and varied $D_2$ among 40, 80, and 120 ms to quantify the potential breakdown of the primacy that we found at larger $D_1$. Figure 3F, G illustrates the behavior of two different subjects in this experiment. Note that their choices were consistently influenced by $D_2$, with a clear breakdown of primacy at $D_2 \geq 80$ ms in one of them (Fig. 3G).

This behavior is quantified in Figure 3H for all four subjects. Note the systematic decrease in slope of the psychometric curves with increasing $D_2$: the longer the $D_2$, the weaker the preference for the initially most intense target. For two subjects, the sign of the slopes even flipped from positive to negative if $D_2$ exceeded $D_1$, indicating a preference for the target with the highest final intensity.

Thus, a brief (40-80 ms) initial bias consistently dominated the responses if $D_1$ lasted an equal amount of time or longer than $D_2$. However, stimulus information in $D_2$ was not simply ignored. In Experiment 2, the longer that $D_2$ exceeded $D_1$, the more often subjects responded to the target with the highest final intensity and, for the short $D_1$ durations in Experiment 3, some but not all subjects showed a transition from primacy to recency.
**Strong target imbalance**

As noted in the Introduction, observing primacy does not necessarily imply that the dynamics of evidence accumulation produces hysteresis. In fact, any decision process that can commit to a particular choice before the stimulus ends can produce primacy independently of its dynamic properties (Fig. 1C). The capability for such premature commitment is common to a wide range of decision models, which share the crucial assumption of becoming committed to a particular choice as soon as the evidence (accumulated via some model-dependent process) for that particular alternative hits a fixed bound or threshold. However, the crucial distinction is that dynamic hysteresis due to feedback cross-inhibition creates primacy without irreversible commitment: sufficiently strong input should be able to “override” hysteresis, even when such input arrives well after the internal dynamics has settled into one of its hysteretic states. We therefore decided to present equal-duration reversal stimuli across an extended range of target imbalances. This yields two clearly different predictions for models with or without strong feedback competition. In the absence of feedback competition or if the feedback is not strong enough to create hysteresis, primacy due to absorbing bounds should increase systematically with target imbalance, so the psychometric curves should increase monotonically from saturation at \( p = 0 \) to saturation at \( p = 1 \). Primacy due to strong feedback competition, however, should produce non-monotonic curves because primacy will eventually decrease if the target imbalance becomes sufficiently large. This prediction holds because the input during \( D_2 \) will start to exceed the strength of the hysteresis induced during \( D_1 \), and the absence of commitment (absorbing boundaries) will allow the decision process to evolve for at least the full duration of the stimulus. Some of the psychometric curves from Experiments 1–3 may already hint toward such nonmonotonic choice behavior, but testing this prediction over a wider range of target imbalances is clearly called for. For this purpose, we switched to a different projector capable of producing higher contrasts and luminances (see Materials and Methods).

Figure 4 shows the results from all six subjects who participated in this fourth experiment. Dashed curves are the psychometric curve fitted to the control data (Equation 1, circles). For these brief, 50 ms stimuli, all subjects correctly chose the most intense target in the condition with the highest intensity and correct responses decreased to \(~50\%\) for very small contrast. In the 50-50 ms reversal condition (Equation 1, squares), subjects showed a consistent primacy effect that increased with target imbalance up to \( \sim 40 \text{ cd/m}^2 \). This regime corresponds to the capability of rightward saccades (\( \pm \text{SEM} \)) as a function of the (initial) intensity difference between the left and the right target in Experiment 4. Each panel shows the results from one subject. Circles and dotted lines indicate the 50 ms control condition with a fitted psychometric curve (Equation 1). Squares and solid lines indicate the 50-50 ms reversal trials (\( n = 100 \) for each \( \Delta I \) with fitted curves following Equation 2. Note the nonmonotonic nature of these psychometric response functions. Arrows indicate a significant decrease in the primacy effect (Fischer’s exact test, \( p < 0.05 \)). Forced reversal stimuli induced a near-complete inversion of the primacy effect. Diamonds indicate the 50-50-67 ms forced reversal condition. Triangles indicate the 83-67 forced reversal condition.

** Forced reversal**

Although the basic primacy effect could indeed be reduced by stronger target contrasts (approximately for \( |\Delta I| > 40 \text{ cd/m}^2 \)), these findings could not provide a strict upper limit on the possible contribution of premature choice commitment or a strict lower limit on the contribution of dynamic hysteresis. To obtain those limits, we investigated to what extent the primacy effect breaks down when later-arriving stimulus evidence in favor of the opposite target location is very strong. Experiment 4 therefore included “forced reversal” trials in which \( D_2 \) was immediately followed by a \( D_3 \) epoch in which the reversed target balance was made even stronger than the reversed imbalance during \( D_2 \). As explained in the Materials and Methods and illustrated in Figure 5A, any residual probability of choosing the initially...
brightest target in this forced reversal condition then sets an upper limit on how much of the primacy seen without the forcing $D_3$ phase is caused by premature choice commitment. Conversely, the amount of primacy in the unforced condition that can be attributed to hysteresis is at least as large as the reduction in primacy that occurs when appending the forcing $D_3$ phase to the stimulus.

For efficiency, and to ensure that forced reversal trials were relatively rare occurrences in any given block of trials (17%), we only used a target contrast that produced a strong primacy effect, as measured in pilot experiments. The duration of the forcing $D_3$ phase was set at 67 ms. Note that if $D_3$ were too short to reach maximal choice forcing, it would merely reduce the test’s power to reject the hypothesis of absorbing bounds being responsible for primacy effects. As shown in Figure 4 (isolated diamonds near $p = 0$ and $p = 1$), adding the $D_3$ forcing phase to the stimulus completely abolished primacy and drove all subjects to choose almost exclusively the target with final dominance. The same result was obtained with the 83–67 ms forcing condition (Fig. 4, triangles; see Materials and Methods). These findings strongly suggest that the contribution of premature choice commitment was extremely small.

As outlined in the Materials and Methods, the simplest approach to quantifying how strong these data challenge mechanisms that attribute primacy effects to premature commitment is to focus on a slightly more specific class of models with absorbing bounds, namely those without hysteresis in their evidence accumulation. This restriction still includes all “standard” decision models, such as “bounded drift-diffusion” (Gold and Shadlen, 2007). All models that we are aware of that use bounded integration, leaky or not, and independently of their mutual interactions as long as they are too weak to cause hysteresis. This whole class of models can then be tested quantitatively by comparing the probability $F_i$ of still choosing the initially strongest target in forced reversal conditions with the probability $R_i$ of choosing the initially strongest target in the corresponding unforced reversal conditions. More specifically, we could test whether the measured $R_i$ and $F_i$ combinations violate the inequality (see Materials and Methods for details): $F_i \geq 2 R_i - 1$. If so, our data would provide strong evidence against the very wide class of models with absorbing bounds and no hysteresis. One can in fact derive less conservative inequalities for the $R_i$, $F_i$ combinations when focusing on specific models within this highly diverse class, but this is beyond the scope of the present paper and proved unnecessary for our present purposes.

To visualize the inequality test, we plotted the averaged $R_i$ and $F_i$ data pairs from each subject (see Materials and Methods) in the scatter plot in Figure 5 (error bars indicate ±1 SEM). Note that the data points would have to fall within the gray area delineated by the line $F_i = 2 R_i - 1$ to satisfy the predictions of absorbing bound models without hysteresis. It is quite clear, however, that nearly all data from the 50–50–67 ms forcing condition fell well outside of that region (Fig. 5B). A one-sided z test on each data pair (see Materials and Methods) indeed showed a highly significant violation of inequality 4 ($p < 0.00015$) for all but one of our subjects. When we performed the test with the $F_i$’s derived from the 83–67 ms forcing condition (Fig. 5C), we also found highly significant violations of inequality 4 ($p < 0.00005$), except for subject J.E. As can be seen in Figure 4, the lack of statistical significance for J.E. in these tests stems from a single $R_i$ outlier data point. If we would have based the test for this subject on his full dataset (i.e., sampling from the fit-function based on all data points), this test too would have strongly violated inequality 4.

**Model simulations**

The choice data from Experiment 4 provide strong evidence against the broad class of accumulator models with absorbing...
bounds. Here, we simulated feedforward and feedback competition models without absorbing bounds and tested their performance against our observations. Figure 6 shows a schematic drawing of the feedforward (Fig. 6A) and feedback (Fig. 6B) architectures. The inputs, representing the intensities of the two targets, pass through a leaky integrator and a nonlinear Naka-Rushton compression stage \( s = x^2/(x^2 + c) \) for \( x > 0 \), otherwise \( s = 0 \); \( c = 1 \). The difference between the two models is in the type of cross-inhibition only. Choices were assigned by comparing the units’ response magnitude at stimulus offset. An additional integration step (shown transparently) in the feedforward model can—for this reversal condition—“repair” its failure to produce primacy without absorbing bounds. C. Output of the left (green) and right (red) decision units from the feedback model as a function of time for leftward and rightward choices (columns) in the three different \( (D_1, D_2) \) timing conditions of Experiment 3 (rows). Model parameters for the simulations in A–C, Integration constant \( T_i = 80 \text{ ms} \); strength of the cross-inhibition, \( \gamma = 3.33 \) for feedback cross-inhibition, \( \lambda = 1.0 \) for feedforward cross-inhibition; Variance of the Gaussian white noise on the inputs was 20% of the mean.

Figure 6 shows the choice behavior of the two models (same format as Figs. 2, 3). Five different conditions were simulated. The first three panels in each row show simulations of the reversal conditions from Experiment 2 (from left to right, 40–120 ms, 80–80 ms and 120–40 ms, respectively). The fourth panel shows the 50–50 ms reversal condition that was tested over a larger contrast range in Experiment 4. The two curves in each graph show the results obtained with two different parameter sets, representing two subjects that showed distinctly different choice behavior in the 40–120 ms reversal condition of Experiment 2 (gray: D.A.-like subject; black: J.K.-like subject).

The feedback model (Fig. 7A) generally prefers the initially most intense target. However, in the 40–120 ms reversal condition (Fig. 7A, left), the presence of hysteresis depends critically on the integration constant \( T_i \) and the cross-inhibition strength (\( \gamma \)). With a longer \( T_i \) and a weaker \( \gamma \), the hysteresis disappears (note that the absolute values of \( T_i \) should not be taken literally because they scale with the shape parameter \( c \) of the sigmoid function). For high target contrasts (Fig. 7A, right), the model
shows the observed decline of the primacy effect as such strong stimuli are able to “override” the hysteresis. Response biases as observed in the experiments could be induced, for example, by asymmetries in input gains for the left and right decision units (data not shown).

For the feedforward model, a direct readout of the competition stage at stimulus offset (Fig. 7B) always predicts a preference for the target that is most intense at the end. Note, however, that this failure to replicate the basic primacy effect can be “repaired” by adding an accumulator stage that integrates the output of the competition stage without any further interactions or absorbing bounds (Fig. 6A). Indeed, when the choices are based on the total time integral of its competition stage output (Fig. 7C), primacy occurs when the two stimulus epochs are equal or the first epoch is longer. Nevertheless, with this augmented version of the feedforward model, it remains impossible to produce primacy in the 40-120 ms reversal condition (as seen in some of our subjects in Experiments 2 and 3). This is due to the absence of absorbing bounds and absence of hysteresis in the evidence accumulation. Both versions of the feedforward model also fail to account for the observed decline in primacy at large target contrasts (Experiment 4) regardless of its parameter settings. Thus, in short, neither the competition stage nor the added accumulator stage of the feedforward model were able to fully capture all key features of the saccadic choice behavior that were revealed across our different reversal conditions.

Figure 7. Model simulations of the different reversal conditions. A, Choice behavior of the feedback model. B, C, Choice behavior of the feedforward model in the competition stage (B) and the accumulator stage (C) presented in the same format. Each panel shows the probability of choosing the right target as a function of rightward bias at stimulus onset ($\Delta$, here in arbitrary units) for different timing conditions, as indicated in the top right corner. Note the larger range of $\Delta$ values on the abscissa of the rightmost panels, corresponding to the larger contrast range that was tested in Experiment 4. Black and gray symbols represent simulation results obtained with two different sets of parameters. Values in each set were chosen to either mimic the behavior of subject D.A. (gray) or subject J.K. (black), because these two subjects clearly showed opposite choice behavior in the 40-120 ms reversal condition of Experiments 2 and 3. Model parameters were obtained by manual adjustment: $T_1 = 50$ ms and $T_2 = 25$ ms (gray and black curves, respectively); $\gamma = 3.25$ and $\gamma = 7.00$ for feedback cross-inhibition (gray and black curves, respectively), $\lambda = 1$ for feedforward cross-inhibition; Variance of the Gaussian white noise on the inputs was 60% of the mean. Each data point consists of 500 simulated trials.

Reaction time

The choice patterns in Experiments 2 and 3 showed remarkably different dependencies on the same stimulus set between our subjects. This offers a unique opportunity to investigate whether reaction times for the alternatives follow similarly different patterns in the same subjects or if they are dissociated from choice probabilities altogether. We wondered, for example, if the relation between reaction time and choice probability followed some simple constraints such as: (1) the mean latencies of saccades toward the most frequently chosen target are systematically shorter than the mean latencies of saccades toward the competing target, and (2) increases in choice preference are associated with increases in this latency difference.

The data from Experiments 1–3 showed, however, that there was no significant correlation between choice preference and
mean latency difference of leftward versus rightward saccades in the control conditions (Fig. 8A; \( r = 0.11, t \) test, \( p = 0.28 \) over all subjects and conditions). For most data from the reversal conditions (Fig. 8B), we did find a significant correlation between choice preference and latency difference (\( r = 0.65, t \) test, \( p < 0.0001 \)), but only for timing conditions that produced a preference for the initially brightest target (Fig. 8B, small symbols, thin regression line). For timing conditions that produced a preference for the target with the highest final intensity (i.e., negative slopes of psychometric curves, \~{}20% of the data), we actually found an opposite relationship between latency and choice. Under these latter conditions (Fig. 8B, large symbols, thick regression line), the mean latencies of rightward saccades increased significantly compared with the mean latencies of leftward saccades as the preference for the right target increased and vice versa if preference for the left target increased (\( r = -0.68, t \) test, \( p < 0.0001 \)).

This remarkable dissociation between choice probability and saccade latency is further illustrated in Figure 8C–E for the four subjects that showed nearly inverted choice preferences in the 80–80 ms and 40–120 ms (gray) reversal conditions in Experiment 2 (Fig. 3B, D). Note that for each saccade direction, the changes in reaction time as a function of initial target contrast were the same under both timing conditions (Fig. 8D, E; ANCOVA, \( F = 0.24, p > 0.8 \) and \( F = 1.88, p > 0.1 \), respectively), whereas the changes in choice probability were opposite (Fig. 8C, \( t \) test, \( p < 0.0001 \)). This shows that the reaction times depended strongly on the initial target contrast and on the direction of the ensuing saccade, but not on the probability of choosing either target. Saccade latency was short compared with the ambiguous control condition if the initially stronger target was chosen (i.e., \( \Delta RT < 0 \)) but long if the initially weaker target was chosen (i.e., \( \Delta RT > 0 \)), and these latency differences increased as a function of the initial intensity difference regardless of the choice probability.

This relation between reaction time and the initial intensity of the selected target (or its initial intensity difference with the other target) was found across all experiments and all conditions (Fig. 9). In Experiments 1–3, the chronometric response functions obtained in reversal conditions (Fig. 9B) were actually quite similar to the ones obtained in control conditions (Fig. 9A). This is quantified in Figure 9C for all test conditions applied in Experiment 1 using the slopes of the chronometric response functions (see Materials and Methods, above; Eq. 8). Because the chronometric functions for rightward and leftward saccades were practically mirror images, we pooled these responses by inverting the slope of the rightward responses. Neither the 80–80 ms nor the 40–40 ms reversal condition produced significantly different latency effects compared with the control conditions (ANCOVA, \( F = 1.18, p > 0.2 \) and \( F = 0.5, p > 0.4 \), respectively). Similar results were obtained in Experiments 2 and 3 (Fig. 9D, E, respectively). In both experiments, changes in \( D_1 \) and \( D_2 \) had very little influence on the saccade latencies. This contrasts markedly with the robust influences on choice behavior (Fig. 3). In fact, the chronometric functions remained invariant to both the presence and timing of the reversals (ANCOVA: no significant differences in Experiment 2, \( F = 0.28, p > 0.8 \) and Experiment 3, \( F = 1.25, p > 0.2 \)). This similarity between control and reversal data in Experiments 1–3 shows that a revision of the initial decision by later evidence did not lead to an extra delay in saccade reaction time. In fact, in Experiment 4, when we forced saccades in the opposite direction of the initially strongest target, thus breaking down the hysteresis built up during the initial part of the stimulus, the latencies were somewhat shorter than the ones in the corresponding unforced condition (mean ± SEM latency difference for movements to the same location: 26 ± 4 ms, paired \( t \) test, \( p < 0.0001 \); and for movements to the opposite location: 17 ± 5 ms, \( p < 0.01 \); data not shown). This latency reduction by late stimulus evidence also refutes the notion that initial information alone might have been the determining factor in reaction times. The latter is further corroborated by the significant differences in slopes of the chronometric function for the control versus reversal conditions obtained in Experiment 4 (Fig. 9F; ANCOVA: \( F = 47.53, p < 0.0001 \)), for which the mean target intensity levels were larger than in Experiments 1–3 (i.e., 27.8 cd/m² vs 16.7 cd/m², respectively; see Materials and Methods).
initially strongest target and concluded that this could be due to feedback competition.

One might argue that feedforward models do predict primacy if subjects adopted a low decision threshold reached shortly after stimulus onset (Ludwig, 2009) or if the integration of evidence were bounded (Kiani et al., 2008). However, our finding that the primacy effect decreases at higher target contrasts (Fig. 4) cannot be explained by any feedforward model with irreversible decision thresholds. Moreover, the very low probabilities of still choosing the initially strongest target in the forced reversal condition (Fig. 5) show that even for feedback models that do allow dynamic hysteresis, absorbing bounds play no significant role in controlling the well defined saccadic choice behavior that occurred in Experiment 4. These robust findings are corroborated by some of the results obtained by Tsetsos et al. (2012), who found that motion-direction discrimination in one of their subjects showed a primacy-to-recency transition for short- versus long-duration stimuli. Given the evidence against choice commitment, dominance of the initial stimulus also contradicts many other model types: high-leakage models (Kiani et al., 2008) and most “urgency” models (Ditterich, 2006; Cisek et al., 2009) give more weight to the final stimulus (unless the urgency signal rose faster than our 40 ms minimal duration; Standage et al., 2011). Likewise, recent gating models (Purcell et al., 2010; Schall et al., 2011) prevent accumulation of the initial, low-amplitude part of the sensory input.

In theory, it is possible that a different mechanism triggered a saccade to the bright forcing target even though the original mechanism had already committed to the other target. In particular, one might worry that, due to subsiding fixation activity, the late forcing pulses were able to elicit so-called express saccades (Dorris et al., 1997; Bell et al., 2006). There was, however, no evidence for this; the latencies of individual saccades were always >150 ms relative to the onset of the forcing pulse, which rules out any specific involvement of the express pathways. We also examined the actual eye movement traces. Prior commitment to the initially strongest target in a parallel pathway would predict that saccades toward the forcing target are substantially influenced by (preparatory) activity related to the other impending saccade. However, as shown in Figure 10, the metrics and kinematics of saccades in the forced condition were actually very similar to those in the control and unforced condition. This supports our assumption that they resulted from the same saccadic mechanism.

Interestingly, we found that reaction times depended on the initial stimulus contrast and on the direction of the ensuing saccade (Fig. 9), but not on the probability of choosing either target (Fig. 8). In fact, by manipulating the timing of the intensity swaps in Experiments 2 and 3, we could nearly invert the relation between reaction time and choice probability from faster to slower.

**Discussion**

We studied saccadic decision making using two simultaneously (≤160 ms) presented targets with intensity reversals occurring at different moments in time. Using this novel approach, we found a robust primacy effect: subjects always preferred the target that was brightest during the early part of each stimulus even if longer-lasting, opposite differences were present in the second part of the stimulus. This primacy effect collapsed when: (1) the duration of the early stimulus phase was reduced to ≤40 ms, (2) the target intensity differences were large, or (3) the reversal stimulus was followed by a strong stimulus bias toward the initially weakest target. The latter two findings show that the basic primacy effect did not result from premature commitment to the initially dominant target. A decision model that assumes feedback cross-inhibition, however, fully described the observed choice behavior.

There is some prior evidence that decision making is biased toward early visual information. In a motion discrimination task, Ludwig et al. (2005) found that subjects rely mostly on information provided in the beginning, a finding that they later explained with a time-varying decision threshold (Ludwig, 2009). Disturbing motion pulses also have more effect when they occur early in a trial (Huk and Shadlen, 2005; Kiani et al., 2008; Tsetsos et al., 2012). Furthermore, using a search task in which target intensities jittered, Tsetsos et al. (2011) found a preference for the

![Figure 9](image-url)
reactions for the more likely outcome in some of our subjects. Such a remarkable dissociation between reaction time and choice behavior has, to our knowledge, never been reported. Instead, it is typically found that changes in choice probability go hand in hand with opposite changes in reaction time (Palmer et al., 2005; Chittka et al., 2009, but see Niwa and Ditterich, 2008, who found another example of a dissociation: reaction times that varied across conditions without changes in choice probability and vice versa). Current decision-making theories explain these findings from the first-to-threshold principle, but this principle cannot account for the choice behaviors observed in Experiment 4 (Figs. 4, 5), the decoupling of choices and reaction times (Fig. 8), and the remarkable invariance of the chronometric functions to both the presence and timing of the reversals observed in Experiments 1–3 (Fig. 9A–E). We thus conclude that reaction time and choice are determined by separate mechanism rather than by a single-stage competition process. This fits in previous conceptual schemes (Findlay and Walker, 1999), which propose that “when” and “where” are determined by parallel but hierarchically organized pathways.

Physiological studies clearly support the notion that target selection and movement initiation are distinct phenomena. In monkeys performing visual search, visually responsive cells in frontal eye fields, lateral intraparietal sulcus, and superior colliculus discriminate between target and distractors (frontal eye field: Schall and Hanes, 1993; lateral intraparietal sulcus: Schall and Hanes, 1993; Ipata et al., 2006; Thomas and Paré, 2007; superior colliculus: Basso and Wurtz, 1997; McPeek and Keller, 2002), but the latency with which these cells discriminate the target from distractors is unrelated to the timing of the ensuing movement. In fact, the selection takes place independently of movement execution (Schall et al., 1995; Thompson et al., 1996; Thompson et al., 1997; Murthy et al., 2001; Sato and Schall, 2003; Juan et al., 2004; Murthy et al., 2009).

The chronometric functions from all of our experiments indicated that reaction times decreased systematically with increasing initial intensity of the selected target (Figs. 8, 9). One might speculate, therefore, that the reaction times might have been determined by the target contrasts at stimulus onset. Such a theory would be consistent, for example, with the contrast-dependent spike timing recently observed in primary visual cortex (Lee et al., 2010) and is further supported by Experiments 1–3, which showed that neither the presence nor the timing of the intensity reversals had a significant influence on the chronometric response curves (Fig. 9C–E). However, in Experiment 4, we found that the chronometric functions for the control and reversal conditions were clearly different (Fig. 9F) and that the later forcing stimulus actually shortened the saccade latencies, indicating that later stimulus information did affect the reaction times. A possible interpretation of these effects is that increases in stimulus intensity decrease the delays in visual processing (Bell et al., 2006).

The present literature leaves some uncertainty about how the target-selection stage is read out. In our simulations, we tested two different approaches. Either the integrated output of the decision units was compared, choosing the channel with the highest value as the current winner, or the output levels of the decision units at stimulus offset were compared. The latter could rely, for example, on the activity levels of cells that keep the latest choice in working memory (Kojima and Goldman-Rakic, 1982; Chafee and Goldman-Rakic, 1998). For the feedforward model, the type of readout heavily influenced its choice behavior, but due to the absence of hysteresis in the evidence accumulation, neither type could be reconciled with the experimental data (Fig. 7B, C). For the feedback model, only a readout at stimulus offset produced predictions that were qualitatively consistent with the observed behavior (Fig. 7A). Results from the integrated output (data not shown) failed to account for the nonmonotonic nature of the psychometric curves in the 50–50 ms reversal conditions of Experiment 4.

Bollimunta and Ditterich (2012) obtained physiological evidence from monkey lateral intraparietal sulcus for the presence of a feedback inhibition component in the random-dot motion-direction discrimination task. At the same time, the presence of feedback inhibition could not be ruled out. Here, we did not simulate hybrid models. It is possible, and perhaps even likely, that feedback inhibition also contributes to the visual target-selection process that we have studied here. However, having ruled out a significant contribution of premature choice commitment in Experiment 4, the essence of our findings is that the observed primary effects can only be accounted for by a model that includes sufficiently strong feedback cross-inhibition.

We conclude that saccadic responses to competing visual targets are best described by a model featuring a competitive choice mechanism based on feedback cross-inhibition that exerts executive control (possibly mediated by the substantia nigra; Hikosaka and Wurtz, 1983) over the initiation of upcoming saccades.
References


