



City Research Online

City, University of London Institutional Repository

Citation: Whiteley, L., Yarrow, K., Haggard, P. & Rothwell, J. C. (2006). Biases in the perceived timing of perisaccadic perceptual and motor events. *Attention, Perception and Psychophysics*, 68(7), pp. 1217-1226. doi: 10.3758/bf03193722

This is the unspecified version of the paper.

This version of the publication may differ from the final published version.

Permanent repository link: <https://openaccess.city.ac.uk/id/eprint/331/>

Link to published version: <https://doi.org/10.3758/bf03193722>

Copyright: City Research Online aims to make research outputs of City, University of London available to a wider audience. Copyright and Moral Rights remain with the author(s) and/or copyright holders. URLs from City Research Online may be freely distributed and linked to.

Reuse: Copies of full items can be used for personal research or study, educational, or not-for-profit purposes without prior permission or charge. Provided that the authors, title and full bibliographic details are credited, a hyperlink and/or URL is given for the original metadata page and the content is not changed in any way.

Biases in the perceived timing of perisaccadic perceptual and motor events

Kielan Yarrow^{1,2,*}, Louise Whiteley^{2,3}, Patrick Haggard^{2,3} & John C Rothwell¹

1. Sobell Department of Motor Neuroscience and movement disorders,
Institute of Neurology,
U.C.L.

2. Institute of Cognitive Neuroscience,
U.C.L.

3. Department of Psychology,
U.C.L.

Running head: Time biases for perisaccadic events

Production Number: P360

* Author for correspondence:

Kielan Yarrow,
Sobell Department of Motor Neuroscience &
Movement Disorders,
Institute of Neurology,
8-11 Queen Square,
London WC1N 3BG.

Tel: +44 (0)20 7679 1154
Fax: +44 (0)20 7813 2835
Email: k.yarrow@ion.ucl.ac.uk

Abstract

Subjects typically experience the temporal interval immediately following a saccade as longer than a comparable control interval. One explanation of this effect is that the brain antedates the perceptual onset of a saccade target to around the time of saccade initiation. This could explain the apparent continuity of visual perception across eye movements. This “antedating” account was tested in three experiments in which subjects made saccades of differing extents then judged either the duration or the temporal order of key events. Post-saccadic stimuli underwent subjective temporal lengthening and had early perceived onsets. A temporally advanced awareness of saccade completion was also found, independently of antedating effects. These results provide convergent evidence supporting antedating, and differentiating it from other temporal biases.

Introduction

In everyday life, people regularly make rapid, saccadic movements of the eyes to bring objects of interest onto the high acuity fovea. Saccades raise serious computational problems that the visual system must overcome (Bridgeman, Van der Heijden, & Velichkovsky, 1994). Perisaccadic biases in spatial vision have provided insights into the way the brain solves these problems (Schlag & Schlag-Rey, 2002). Recent data suggest that temporal judgements are also biased when we make saccades to fixate new targets. Subjects consistently overestimate the duration of a post-saccadic stimulus compared to the same stimulus seen at fixation, an illusion termed “saccadic chronostasis” (Yarrow, Haggard, & Rothwell, 2004a; Yarrow, Haggard, Heal, Brown, & Rothwell, 2001; Yarrow, Johnson, Haggard, & Rothwell, 2004b). In a typical experiment, subjects saccade to a target that changes form or colour during the saccade. Subjects judge whether the new target stimulus was presented for a longer or shorter time than subsequently presented reference stimuli, and these judgements are used to determine a point of subjective equality (PSE; the point at which target and reference stimuli are perceived to have identical durations). The same task performed at fixation forms a control. Reduced PSEs in saccade compared to control conditions imply temporal overestimation of the post-saccadic stimulus.

One finding from previous saccadic chronostasis experiments is particularly noteworthy. When saccades of different extents are made, the size of the illusion changes; the duration of the illusion reflects the duration of the preceding saccade (Yarrow et al., 2001). This suggests the following explanation. Both retinal blur and active suppression degrade visual input during saccades (Ross, Morrone, Goldberg, & Burr, 2001) leaving a gap in perception corresponding to the saccade duration. The

brain may simply assume that the information in the post-saccadic image has remained constant across the saccade. Hence the perceived onset of the saccade target is effectively antedated to a moment just prior to saccade initiation, helping to provide the visual continuity we experience. The illusion arises following many different kinds of saccade with partially separable neural substrates, and has an effect size that typically exceeds the duration of the saccade by around 50 ms (Yarrow et al., 2004b). These observations suggest that an efference copy signal originating in a subcortical region such as the superior colliculus could act as a trigger or time marker for the antedating process. The same signal may initiate receptive field shifts occurring in cells in parietal cortex and other areas, which might in turn influence conscious visual perception (Duhamel, Colby, & Goldberg, 1992; Goldberg, Bisley, Powell, Gottlieb, & Kusunoki, 2002; Sommer & Wurtz, 2002; Umeno & Goldberg, 1997; Walker, Fitzgibbon, & Goldberg, 1995).

Two key objections have been made to this *antedating* account. These two problems are described below in some detail because the experiments and analyses presented later were intended to provide new positive evidence for antedating that is not subject to these concerns.

Perceived duration is an indirect measure of perceived event timing.

Previous studies of saccadic chronostasis have used interval judgements like those employed to investigate how humans assess the duration of perceptual epochs (Allan, 1979; Allan, 1998). However, an interval is bounded by the events marking its onset and offset (see Figure 1). Few studies have looked for consistent temporal effects using measures of both interval and event timing. The extent to which these

different measures are biased together might provide valuable insights about the fine-grain structure and coherence of conscious temporal experience. In the current context, *interval* judgement tasks can only provide indirect evidence that a post-saccadic *event* such as the visual onset of a saccade target stimulus is antedated. This leaves open the possibility that the changes in perceived interval duration do not reflect antedating of postsaccadic events, but rather alternative processes such as changes in internal clock speed (Hodinott-Hill, Thilo, Cowey, & Walsh, 2002, but see Yarrow et al., 2004a). However, another type of task that investigates event timing could provide direct evidence for the antedating of events: the temporal order judgement (TOJ).

FIGURE 1 ABOUT HERE

In TOJ tasks, subjects judge the relative timing of two stimuli or events presented in close temporal proximity. These judgements can be used to determine a point of subjective simultaneity (PSS) where the events appear to have occurred at the same moment. The resulting PSS may be biased, with factors such as sensory modality, stimulus intensity, attentional allocation and even recent sensory experiences influencing when each stimulus is perceived to occur (Fujisaki, Shimojo, Kashino, & Nishida, 2004; Hirsh & Sherrick, 1961; Jaskowski, 1999; Shore, Spence, & Klein, 2001; Spence, Shore, & Klein, 2001; Stone et al., 2002).

A few studies have used TOJ tasks to study temporal perception in the context of saccadic eye movements. Park, Schlag-Rey, & Schlag (2003) had observers judge the order of two 3 ms spots flashed near the target of a saccade and found a temporal advantage for the spot perceived as closest to the saccade end point. Deubel, Irwin, &

Schneider (1999) had subjects perform a rather different task, judging their position of gaze at the time a ring stimulus was flashed. Subjects showed a marked bias that varied with stimulus position, interpreted by the authors as a tendency to feel that the eyes had moved to the target early, at the moment the shift of spatial attention that preceded the saccade occurred. Diamond (2003) found a bias in the same direction as Deubel et al. (1999). His observers used a seven-point scale to judge the timing of a briefly flashed green bar (equiluminant against a red background) relative to a saccade. Two out of three perceived the bar as delayed, although the bias tended to disappear for stimuli presented during the saccade. Finally, Volkman & Moore (1978) had observers report whether a grating was displayed before, during or after a saccade. No simple direction of temporal bias could be determined from the complex pattern of results that emerged.

It is challenging to relate these studies to the hypothesis that the target of a saccade is antedated. A particular difficulty arises because the temporal order judgement that is being made is typically between a brief visual reference and the saccade itself, rather than the post-saccadic target stimulus. Temporal biases arise when motor acts are judged relative to brief sensory stimuli (Dassonville, 1995; McCloskey, Colebatch, Potter, & Burke, 1983). In chronostasis experiments, however, the illusion biases judgements about a visual stimulus (the saccade target) not a motor act. Consequently, the question of whether the perceived time of post-saccadic events is antedated remains open. By asking specific questions about the timing of either the post-saccadic stimulus or the motoric act of saccading, we can both seek direct evidence for antedating and begin to dissect the various biases that may have been conflated in previous investigations.

Evidence favouring antedating relies on incorrect assumptions about saccadic suppression

In typical chronostasis experiments, the points of subjective equality reported in saccade conditions are not simply calculated using the duration for which the target stimulus appeared on the screen in each trial. These PSEs incorporate an additional correction based on the time between the (presumably unseen) target change and the target foveation at the end of the primary saccade. The rationale for this correction is as follows. The target stimulus changes into its target state during the subject's primary saccade, at a time when perception is degraded (Ross et al., 2001). This suggests that it is not perceived to a degree compatible with the initiation of a mental timing operation until it is actually foveated (or at least para-foveated, in the case of trials where a corrective saccade follows the primary saccade). Hence the time for which the stimulus was on screen during the primary saccade (the period from stimulus change to saccade termination) is subtracted from presentation times before PSEs are calculated. The effects that are reported (the difference between control and saccade PSEs) can therefore be broken down into two components: 1) an increase in perceived duration relative to the on-screen presentation time, and 2) the correction.

Although saccadic suppression is a robust phenomenon, only magnocellular input is strongly suppressed, and even this suppression is not complete (Ross et al., 2001). There are cases where visual information presented during a saccade is processed quite effectively (e.g. Eggert, Ditterich, & Straube, 1999; Hunt, Chapman, & Kingstone, submitted for publication). If the mid-saccadic target change occurring in chronostasis experiments is clearly perceived, then the correction procedure may not be justified, with two implications. Firstly, the magnitude of the saccadic

chronostasis effect will be overestimated (although not eliminated). This problem can be addressed by analysing the data without applying the correction to demonstrate that a reliable effect still exists. A second implication, however, is more critical for the antedating hypothesis. The saccade size effect is an important result because it demonstrates that perceived time is being adjusted in direct response to a key motor parameter of the eye movement itself. In the original experiment reporting this effect (Yarrow et al., 2001, Experiment 1) the change to the target stimulus was triggered a set proportion of the distance into the saccade. This means that the size of the correction varied in the short and long saccade conditions, being larger in the latter case. Hence, if the correction is unwarranted, the saccade size difference may have been artificially enhanced¹. In this case, the link between the visual illusion and saccadic motor control would be undermined. In the experiments that follow, we adjusted trigger times such that our correction was constant across conditions, eliminating this potential artefact.

The current experiments

In light of these concerns, we designed three convergent, parallel experiments to investigate antedating in the saccadic chronostasis illusion and its relation to the perceived timing of saccades. By using both interval and temporal order judgement tasks, we aimed to cross validate our previous findings on saccadic chronostasis and counter explanations and criticisms that are tied to a specific experimental task.

The first experiment aimed to replicate the finding of a chronostasis effect that scales with saccade size, and used a standard interval duration judgement (*chronostasis experiment*; cf. Yarrow et al., 2001). An adjustment was made to the

timing of the mid-saccadic stimulus change to equate the correction procedure across conditions. The second experiment used a procedure very similar to the first. Again, a visual change in the target stimulus was triggered during a saccade. Now, however, subjects judged whether they first perceived the post-saccadic visual stimulus before or after a brief auditory tone (*audio-visual TOJ experiment*). The antedating hypothesis predicts a bias to perceive the visual onset of the saccade target to be shifted back to an earlier time. Moreover, this bias should again scale with saccade duration. The third experiment also used a temporal order judgement. In this experiment, however, the visual stimulus remained constant and subjects made their temporal order judgements relative to the time point at which they considered their eyes had arrived at the saccade target (*audio-saccade TOJ experiment*). This replicates the situation that most previous studies of the relation between saccades and time perception have used (Deubel et al., 1999; Diamond, 2003; Volkman & Moore, 1978). In contrast to these studies, the audio-saccade TOJ experiment used an auditory reference rather than a visual one. This change is not trivial, given recent evidence that transient visual events are themselves subject to perisaccadic temporal biases, whereas transient auditory events are not (Morrone, Ross, & Burr, 2005). The auditory-saccade TOJ experiment also assessed biases with saccades of different extents, further differentiating it from previous work. Judgements about the timing of actions relative to sensory events are often biased (see above) and we wished to establish whether such a bias existed alongside the chronostasis effect and could be distinguished from it.

Materials and methods

Participants. The same 18 subjects (13 male, mean age 29.2, SD 6.6) completed all three experiments. A further two subjects were tested and subsequently rejected because of high trial to trial variability in one or more experiments (logistic regression $p > 0.05$ for combined short/long saccades in either control or saccade conditions).

Apparatus. Subjects sat before a 22" CRT colour monitor refreshing at 120 Hz. Eye to screen distance was maintained at 41 cm using an adjustable chin and forehead rest. Horizontal eye movements were recorded from the left eye using an infra-red eye tracker (Applied Science Laboratories Eye-trac model 310) and sampled at 200 Hz (12 bit A/D card; National Instruments DAQ 1200). Visual stimuli were black or grey crosses and filled squares on a white background (average luminance 91 cd/m²) subtending 0.6° of visual angle. Auditory stimuli in TOJ experiments were pure tones of 600 Hz pitch and 25 ms duration, delivered from a piezoelectric speaker behind the subject's head.

Design. Each experiment employed a two factor (2 x 2) repeated-measures design. The first factor *eye status* compared trials involving saccades to constant fixation (control) trials. The second factor *saccade size* varied the position of visual stimuli to produce saccades of either 10° or 50° extent, with constant fixation trials completed at matched eccentricity ($\pm 5^\circ$, $\pm 25^\circ$). Subjects completed 75 randomly interleaved trials from each condition in a single block of 300 trials. Rejected trials (see procedure, below) were repeated at the end of the block. Subjects completed the three experiments in a counterbalanced order.

FIGURE 2 ABOUT HERE

Procedure. A schematic of the experimental procedures is shown in Figure 2.

Chronostasis experiment. In saccade trials, subjects initially fixated a peripheral black cross. They began the trial with a mouse key press, at which point a second black cross appeared on the opposite side of the screen. Subjects maintained fixation at the initial black cross until it became grey (500 ms after the key press) then made a speeded saccade to the second black cross. The eye movement triggered this black cross to be replaced with a black square when the saccade had travelled a set proportion of the distance to target (30% in the 10° saccade condition, 80% in the 50° condition; see our earlier methodological comments). The square remained on screen for a variable duration (125-875 ms) then disappeared, to be replaced by an identical square (the reference stimulus, 500 ms duration) after 500 ms. Subjects indicated whether they saw the first square for more or less time than the second square. The first square's duration was selected randomly on each trial from a condition-specific distribution containing values between 125 and 875 ms in 25 ms increments. The distribution was initially uniform, in the region 300-700 ms, but was updated after each accepted trial according to the generalized P'olya urn model (Rosenberger & Grill, 1997; type IV, $k = 8$). This procedure produces many values close to the point of subjective equality.

In the saccade conditions, each trial's actual target stimulus duration value could be corrected post hoc to match the duration for which the stimulus was seen after the primary saccade landed at or near the target (the same correction employed in our previous studies). This was accomplished by subtracting the time the eye was in motion following the triggered change to a square (not including any subsequent corrective saccades). We produced a corrected data set in this manner, but also

retained an uncorrected data set for comparison. PSEs were then obtained from both data sets using logistic regression. Saccade start/end points were calculated automatically. Velocity was calculated based on the difference between samples n and $n+3$, and saccades were typically taken to start (end) at sample $n+1$ when this velocity rose above (fell below) 120 (75) degrees per second.² Estimates were superimposed on saccade traces and displayed trial by trial along with other key saccade statistics. Trials were rejected automatically for a number of reasons, most critically: 1) when the first saccade did not exceed 90% of the total distance recorded (summed across all detected saccades), and 2) where the stimulus change at the saccade target was not triggered during the first saccade. The programme maintained a running average of reaction time (RT) and saccade duration for the last ten trials of a given condition (initially set at 200/50 ms and 200/130 ms for the RT/duration of the 10° and 50° conditions respectively). Trials were also rejected when eye movements were initiated either anticipatorily or too slowly ($>$ running average RT + 300 ms from cross greying). Subsequent to the experiment, each accepted saccade trace was examined and classified as containing either a single saccade or a primary saccade plus one or more corrective saccades (defined as additional saccades occurring within 300 ms of primary saccade termination).

In control (constant fixation) trials, subjects initially fixated a cross at equivalent eccentricity. It turned grey 500 ms after the subject's key press for a duration determined by the running average RT from the relevant saccade condition, then disappeared for a duration determined by the relevant running average saccade duration. The variable-duration black square then appeared (still at fixation) with subsequent stimulus presentation and subject responses as per saccade trials. Hence foveal stimulation in control trials was very similar to that experienced during saccade

trials. Control trials were rejected if a saccade was detected. Direction of saccade (saccade conditions: left to right and vice versa) or position of the fixation cross (control conditions: left or right) alternated every trial.

Auditory-Visual TOJ experiment. The procedure differed from that employed in the chronostasis experiment in the following respects. Only the first black square was displayed, its offset (after 500 ms) marking the end of the trial. An auditory beep sounded once on each trial. Subjects judged whether the beep occurred before or after they first saw the black square. The time of delivery for the beep was randomly selected. In control trials, it came from an adaptive distribution (see above) with a possible range from 375 ms before black square onset to 375 ms after black square onset, but initially containing values only from -200 to +200 ms. Delivery times and responses were entered into logistic regressions to determine points of subjective simultaneity. In saccade conditions, the randomly selected delivery time was *targeted* based on the relevant running average values for RT and saccade duration. Recorded delivery times were then adjusted such that the beep was correctly localised relative to the moment the black square was first seen with static eyes (the end of the primary saccade). An uncorrected data set was also maintained, in which the beep was localised relative to the moment the black square appeared on screen. In both control and saccade conditions, no beeps were delivered prior to the point at which the fixation cross greyed, meaning that few delivery times as extreme as -375 ms actually occurred.

Auditory-Saccade TOJ experiment. The procedure followed was identical to that of the auditory-visual TOJ experiment with the following exceptions. In saccade conditions, the black target cross did not change into a black square during the saccade. Instead, it remained a black cross. Subjects judged whether the beep occurred

before or after their eyes first arrived at the target (i.e. the end of their eye movement). In control conditions, the grey cross and subsequent brief blank were followed by the reappearance of the black cross. Subjects judged whether the beep occurred before or after this black cross reappeared.

Results

Data verification. Table 1 provides a summary of the extent and duration of primary saccades and the timing features of saccade-contingent display changes made in each experiment. Data from the main analysis are shown on the left hand side of the table. As expected, the manipulation of saccade extent produced a large difference in saccade durations between short and long saccades (mean change 77 ms). However, the timing of the mid-saccadic display change (occurring in the first two experiments) remained constant for short and long eye movements when assessed relative to the end of the primary saccade. More specifically, all trigger times preceded saccade termination by approximately 30 ms.

Temporal judgements. Results based on the judgements made in all three experiments are displayed in Figure 3. In the chronostasis experiment (top panel) PSEs in control conditions were reduced relative to the reference stimulus duration of 500 ms. That is, subjects showed a general trend to overestimate the first stimulus duration. PSEs were further reduced in both saccade conditions relative to their respective controls. This difference is particularly clear when the standard correction for saccadic suppression is employed, but is also evident in the uncorrected data. Crucially, this saccade-related reduction was around 45 ms greater for long saccades

than for short saccades. Two statistical analyses were conducted, one for the corrected data and one for the uncorrected data. For the corrected data, A 2 x 2 repeated measures ANOVA showed significant main effects of eye status (control versus saccade; $f = 8.25$, $df = 1, 17$, $p = 0.011$) and saccade size ($f = 4.53$, $df = 1, 17$, $p = 0.048$) and a significant interaction ($f = 8.26$, $df = 1, 17$, $p = 0.011$). Pairwise follow-ups indicated a significant difference between the long and short saccade conditions ($t = 2.73$, $df = 17$, $p = 0.014$) but no difference between control conditions. For the uncorrected data, the ANOVA did not yield a significant main effect of eye status (control versus saccade; $f = 3.6$, $df = 1, 17$, $p = 0.075$) but did yield a significant effect of saccade size ($f = 4.5$, $df = 1, 17$, $p = 0.05$) and a significant interaction ($f = 8.8$, $df = 1, 17$, $p = 0.009$). Pairwise follow-ups indicated a significant difference between long and short saccade conditions ($t = 2.7$, $df = 17$, $p = 0.014$). Although not significant as a main effect, the difference between control and saccade conditions was significant between the long saccade condition and its matched control ($t = 2.6$, $df = 17$, $p = 0.019$).

FIGURE 3 ABOUT HERE

The middle panel of Figure 3 shows PSSs for the auditory-visual TOJ experiment. In the control conditions, the positive PSSs indicate that the beep had to be presented after the onset of the black square to be judged simultaneous with it. In the saccade conditions employing the standard correction, the negative PSSs indicate that the beep had to be presented well before target foveation to subjectively coincide with the perceptual onset of the post-saccadic black square. The uncorrected saccade data show that subjective simultaneity occurred when the beep was presented before

the target had even appeared on screen. PSSs for short and long saccade conditions differed by over 50 ms, with control conditions differing by 22 ms. ANOVAs were conducted on both corrected and uncorrected data sets and yielded identical patterns of significance. There were significant main effects of eye status (corrected: $f = 12.15$, $df = 1, 17$, $p = 0.003$; uncorrected: $f = 6.2$, $df = 1, 17$, $p = 0.023$) and saccade size (corrected: $f = 22.57$, $df = 1, 17$, $p < 0.001$; uncorrected: $f = 19.8$, $df = 1, 17$, $p < 0.001$) and also significant interactions (corrected: $f = 5.36$, $df = 1, 17$, $p = 0.033$; uncorrected: $f = 4.5$, $df = 1, 17$, $p = 0.048$). Follow-ups indicated significant differences between the two saccade conditions (corrected: $t = 4.11$, $df = 17$, $p = 0.001$; uncorrected: $t = 3.8$, $df = 17$, $p = 0.001$) but also between the two control conditions ($t = 3.29$, $df = 17$, $p = 0.004$).

In the auditory-saccade TOJ experiment (lower panel of Figure 3) PSSs were close to zero in control conditions, indicating near veridical temporal order judgements. The substantial negative PSSs in saccade conditions indicate that the beep had to be presented well before the eyes arrived at their target to be judged simultaneous with this event. Short and long saccade conditions yielded almost identical biases. An ANOVA revealed significant main effects of eye status ($f = 12.90$, $df = 1, 17$, $p = 0.002$) and saccade size ($f = 6.98$, $df = 1, 17$, $p = 0.017$) which also interacted significantly ($f = 4.83$, $df = 1, 17$, $p = 0.042$). Follow-ups showed a significant difference between the two control conditions ($t = 4.78$, $df = 17$, $p < 0.001$) but not between saccade conditions.

Corrective saccade artefact. In all three experiments, short and long saccade conditions resulted in significantly different percentages of trials containing one or more corrective saccades (mean across subjects: chronostasis experiment, short =

22.7%, long = 62.8%, $t = 8.64$, $df = 17$, $p < 0.001$; auditory-visual experiment, short = 22.0%, long = 62.3%, $t = 10.99$, $df = 17$, $p < 0.001$; auditory saccade experiment, short = 23.4%, long = 59.8%, $t = 7.84$, $df = 17$, $p < 0.001$).

In order to assess the importance of this potential artefact, PSEs/PSSs were recalculated for each subject using only those trials that did not contain any corrective saccades. Mean PSEs/PSSs were compared in each experiment based only on the subset of participants for whom these single-saccade PSEs/PSSs could be reliably determined (logistic regression $p < 0.05$) in both long and short saccade conditions. If differences between short and long saccade conditions still emerged, they could not have resulted from an increase in corrective saccades. These PSEs employed the standard correction for saccadic suppression and are plotted in Figure 4. The related saccade extent/timing statistics are shown in the right hand side of Table 1.

FIGURE 4 ABOUT HERE

For the subset of nine participants with reliable single-saccade PSEs in both conditions, the critical decrease in PSEs from short to long saccade conditions was numerically comparable to that in the full analysis (41 ms) but did not reach statistical significance due to the reduced power ($t = 1.478$, $df = 8$, $p > 0.05$; power = 0.36 to detect a 45 ms difference). For the auditory-visual experiment, a subset of 12 participants with reliable single-saccade PSSs showed a significant decrease from short to long saccade conditions similar to that obtained in the full analyses (60 ms decrease, $t = 4.08$, $df = 11$, $p = 0.002$). Finally, in the auditory-saccade experiment, for the subset of 13 participants with reliable single-saccade PSSs, scores now decreased

significantly from the short to the long saccade conditions (25 ms decrease, $t = 2.886$, $df = 12$, $p = 0.014$).

Discussion

The chronostasis experiment demonstrated a subjective temporal lengthening of post-saccadic stimuli compared to identical stimuli viewed at fixation. The effect was greater following large saccades than following smaller saccades, replicating our previous work (Yarrow et al., 2001). Although the difference in PSEs between the two saccade conditions was not numerically as large as the difference in saccade durations, it was reliable. Moreover, this result cannot have been an artefact of our correction technique, as target changes were triggered at a similar time relative to the end of the saccade in both saccade conditions.

While we cannot rule it out entirely, it seems unlikely that an increase in the frequency or size of corrective saccades from the short to the long saccade conditions was responsible for the reduction in PSEs. Although lacking in power, our comparison based on single saccade trials showed a trend towards lower PSEs in the long saccade condition even when no corrective saccades were made. This trend was of a magnitude very similar to the effect found in our main analysis. In our previous work (Yarrow et al., 2001) we obtained a large difference in PSEs when comparing saccades of 22 and 55 degrees extent. The difference found in that experiment was actually larger than the one obtained here, but the difference in proportions of trials containing corrective saccades from the short to the long saccade condition was much less striking (increase from 69% to 75%; Yarrow, 2003).

A previous study (Yarrow et al., 2001) found that an increase in saccade duration produced an almost numerically matched increase in the size of the chronostasis effect. In contrast, the present study found that the effect of saccade amplitude on chronostasis was numerically smaller (45 ms) than the difference in saccade durations (79 ms). We offer the following speculation relating to the trigger times we employed. Saccadic suppression is greater at the beginning of a saccade than towards its termination (Diamond, Ross, & Morrone, 2000). Although we know of no experiments investigating saccadic suppression for saccades as large as the ones employed here, or comparing it across saccades of differing extents, it seems possible that suppression was less complete in our long saccade condition, where the change of stimulus occurred very late. Subjects may have occasionally clearly perceived the genuine onset of the post-saccadic square stimulus and failed to antedate it in the typical manner, diluting the overall effect. To summarise, we introduced a new correction procedure to eliminate potential artefacts in estimating the numerical magnitude of the chronostasis effect. However, the new procedure would tend to reduce any effect of saccade amplitude on time perception. We nevertheless found that saccade amplitude significantly affected chronostasis.

Turning briefly to the control conditions of the chronostasis experiment, we found a bias in both conditions, implying an extended perception of the first stimulus relative to the second. This is an example of the time order error (Hellstroem, 1985). Calculating chronostasis effects as the difference between control and saccade conditions takes account of this bias, which should be constant across conditions.

In the auditory-visual TOJ experiment, PSSs were substantially reduced in saccade conditions compared to control conditions. The TOJ method measures the difference between the perceived times of two events. If a difference is found,

however, the TOJ method cannot determine which of the two events is perceived as shifted in time, and which is stable. Thus, when a saccade was made, either the perception of the beep was delayed by around 100 ms, or the onset of the post-saccadic (square) stimulus was perceived to have occurred correspondingly early.

Two considerations might suggest the former interpretation. Firstly, stimuli presented in unattended sensory modalities (and/or at unattended spatial locations) are typically judged to arrive later than simultaneously presented stimuli in attended modalities/locations (prior entry effect; e.g. Spence et al., 2001). Secondly, there is good behavioural and neurophysiological evidence for a mandatory link between saccadic eye movements and shifts of spatial attention to the saccade target (Deubel & Schneider, 1996; Moore & Armstrong, 2003). Hence attention might have been more focussed on the saccade target stimulus (and consequently less focussed on the beep) in saccade conditions. Note however that to our knowledge there is no evidence directly linking saccades with changes in the *cross-modal* distribution of attention; the fact that attention is shifting spatially does not actually imply that it is being withdrawn from the auditory modality, although this is a possibility.

Despite this possibility, we favour the second interpretation, that the post-saccadic stimulus was antedated, for the following reasons. Prior entry biases are typically much smaller than the effect observed here (Johnson & Haggard, 2003; Spence et al., 2001). Furthermore, a prior entry account cannot straightforwardly explain why the perceived time of the beep should vary with saccade size. Our analysis of single saccade trials gave no reason to believe that the presence of corrective saccades was responsible for this result, which was significant even when all corrective saccade trials were removed. In particular, the possibility that large saccades require a greater allocation of attention than small saccades is undermined

by the failure to obtain a saccade size effect of the same magnitude in the auditory-saccade experiment discussed next. Hence we favour a saccade target antedating account of both experiments, although we recognise that an additional independent prior entry effect might perhaps sum with the chronostasis effect in TOJ tasks.

The auditory-saccade TOJ experiment showed a slightly different pattern of results from the other two experiments. Large negative PSSs in both saccade conditions imply that either the beep was perceptually delayed relative to the sensation of saccade termination, or the sensation of saccade termination was perceptually advanced relative to the beep. The relevance of the control conditions is less clear here than in the other two experiments, because the control comparison was between an auditory stimulus and a visual stimulus whereas the experimental comparison was between an auditory stimulus and a movement. Judgements about the moment of saccade termination may have depended partially upon visual cues (fixation of the saccade target) but might also depend on efferent or afferent information related to production of the saccade.

Results from our reanalysis based on the presence or absence of corrective saccades further complicate interpretation of this experiment. For trials containing only single saccades a significant but numerically small difference between short and long saccades now emerged, in contrast to analysis of the data set as a whole. This finding suggest that subjects' judgements about the moment their eyes stopped moving might sometimes have been biased forwards by the presence of subsequent corrective saccades (i.e. making a corrective saccade makes you feel that your eyes have finished moving later). Given the preponderance of such saccades in the large saccade condition, this would explain why a saccade size effect only materialised when corrective saccade trials were excluded.

We offer the following interpretation of the auditory-saccade TOJ results as a whole. In saccade conditions, subjects displayed an advanced awareness of their eye position, consistent with previous reports for arm movements and pursuit (Dassonville, 1995; Van Beers, Wolpert, & Haggard, 2001). This bias may have been supplemented by the prior entry effect discussed in relation to the auditory-visual TOJ experiment. Both biases predict similar effects in short and long saccade conditions. Because judgements in the saccade condition of this experiment could also be based in part on visual factors (subjects might infer the timing of the end of their eye movement based on their foveation of the saccade target) some influence of antedating is possible. Such an influence might have yielded the small but reliable saccade size difference in our single saccade analysis, although it is also possible that the advanced awareness of eye position was greater for large saccades.

The auditory-saccade TOJ experiment shared some features with previous work investigating the timing of visual events relative to saccades, so a brief comparison seems appropriate. Deubel et al. (1999) have conducted the most directly comparable study. They had subjects judge the position of their gaze (whether they were looking at the pre-saccadic or post-saccadic target) at the time a ring stimulus was flashed. Subjects showed a bias in the same direction as that reported here using an auditory stimulus: the ring had to be flashed well before the saccade in order to seem coincident with pre-saccadic fixation. This bias was most striking when the ring appeared at the post-saccadic fixation position, was reduced when it appeared at an opposite position, and was absent when it appeared at the central pre-saccadic position. Like the auditory-saccade TOJ experiment reported here, interpretational difficulties emerge because the subjects may have been relying on either efferent/afferent information relating to their eye movement, or visual information, or

both, in order to determine their position of gaze. The authors suggested that subjects mistakenly believed they had moved their eyes when they had in fact simply shifted their visual attention prior to an eye movement. Presumably, this mistake was particularly compelling when attention had already shifted to the position of the flashed ring stimulus, accounting for the differences observed when the flashed stimulus appeared at different locations. If we apply such an account to the auditory-saccade TOJ experiment reported here, we would predict an effect that grows in line with saccade duration (because the judgement we used related to the moment at which the saccade ended, and the shift of visual attention would be expected to precede the beginning of the eye movement by a constant amount). This account received only limited support from our data.

One further unpredicted and interesting result emerged from the two TOJ experiments. In both experiments, control conditions employed an auditory-visual TOJ task. Within each experiment, control PSSs were significantly reduced in the long saccade condition compared to the short saccade condition. No actual saccade was being made in these conditions; the main difference was in the orbital eccentricity of the fixated sequence of visual stimuli. Hence this result could imply that stimuli that are peripherally located in egocentric space have faster perceptual latencies than stimuli at the body midline, even when both are foveated. The finding recalls results obtained in tactile TOJ experiments with crossed or uncrossed hands, which also suggest that the egocentric spatial localisation of a stimulus influences (and therefore precedes) the determination of stimulus timing (Yamamoto & Kitazawa, 2001). Spatial localisation also appears to precede temporal localisation when perception is biased at the time of saccades (Park et al., 2003). However, the current result needs further investigation because visual stimulation was not identical just before the

appearance of the critical visual stimulus. The blank period before stimulus onset had a duration based on the running average saccade duration from the equivalent saccade condition, and was hence longer in the long saccade control condition than in the short saccade control condition. The implication here is that a fixation blank period of approximately 130 ms decreases the perceptual latency of a subsequent visual stimulus compared to a fixation blank period of approximately 50 ms. If this interpretation is correct, it is interesting to note that no compatible effect was found in the control condition of the chronostasis experiment. This would suggest that interval judgements and temporal order judgements may dissociate with respect to how prior visual stimulation affects the onset of a target (see Jaskowski, 1999, for a related discussion focussing on dissociations between TOJ and reaction time data).

To summarise our main findings, our first two experiments provided converging evidence for the antedating of stimuli perceived following a saccadic eye movement. Different tasks assessing both interval and event-based timing yielded consistent results. That is, measures of saccadic chronostasis based on judgements of *intervals* result from changes in the perceived time of the *events* that bound those intervals. Given this convergence of two psychophysical methods, and the common effect of saccade duration on perceptual timing, we suggest that the target of a saccade is antedated towards the time of saccade initiation. It may complement other processes that give rise to perceptual continuity across saccades, such as saccadic suppression and visual masking (Campbell & Wurtz, 1978; Ross et al., 2001). Antedating is not, however, the only temporal bias that needs to be considered in the context of saccades. In addition to possible prior entry effects, we discerned a strong bias during movement comparisons to judge the end of a saccade as having occurred earlier than was actually the case. This bias may relate to the classic anticipatory awareness of

actions (Dassonville, 1995; Haggard, Newman, & Magno, 1999; McCloskey et al., 1983). The motoric bias in our final experiment had a similar direction and magnitude to the perceptual (antedating) bias in our first experiment, but was less dependent upon saccade extent. Dissociations between action awareness and the visual experiences that accompany movements have been reported before (Johnson & Haggard, 2005). Further investigations will need to take account of these various biases, and perhaps others, in order to provide a full explanation of temporal perceptions across eye movements.

Acknowledgements

This work was supported by a Wellcome Trust project grant to JCR and PH. Additional funding came from a Royal Society equipment grant to PH.

Footnotes

1) We originally addressed this concern in a control experiment in which the trigger time for the display change was varied by 85 ms within a large saccade. If subjects perceived the mid saccadic stimulus change and this percept informed their duration judgements, their duration judgements should have been affected by an equivalent amount. This manipulation yielded only a small and non significant (11 ms) effect on duration judgements, but the negative nature of the result makes experimental power (0.71 in this case) an issue.

2) On some trials these thresholds were automatically raised in response to high signal noise in the eyetracker.

References

- Allan, L. G. (1979). The perception of time. *Perception and Psychophysics*, 26, 340-354.
- Allan, L. G. (1998). The influence of the scalar timing model on human timing research. *Behavioural Processes*, 44, 101-117.
- Bridgeman, B., Van der Heijden, A.-H. C., & Velichkovsky, B. M. (1994). A theory of visual stability across saccadic eye movements. *Behavioral and Brain Sciences*, 17, 247-292.
- Campbell, F. W. & Wurtz, R. H. (1978). Saccadic omission: why we do not see a grey-out during a saccadic eye movement. *Vision Research*, 18, 1297-1303.
- Dassonville, P. (1995). Haptic localization and the internal representation of the hand in space. *Experimental Brain Research*, 106, 434-448.
- Deubel, H. & Schneider, W. X. (1996). Saccade target selection and object recognition: evidence for a common attentional mechanism. *Vision Research*, 36, 1827-1837.
- Deubel, H., Irwin, D. E., & Schneider, W. X. (1999). The subjective direction of gaze shifts long before the saccade. In W. Becker, H. Deubel, & T. Mergner (Eds.), *Current Oculomotor Research: Physiological and Psychological Aspects* (pp. 65-70). New York: Plenum.

Diamond, M. R., Ross, J., & Morrone, M. C. (2000). Extraretinal control of saccadic suppression. *Journal of Neuroscience*, 20, 3449-3455.

Diamond, M. R. (2003). *The effect of saccades on visual sensitivity and time perception*. Unpublished doctoral thesis, The University of Western Australia, Nedlands, Western Australia.

Duhamel, J. R., Colby, C. L., & Goldberg, M. E. (1992). The updating of the representation of visual space in parietal cortex by intended eye movements. *Science*, 255, 90-92.

Eggert, T., Ditterich, J., & Straube, A. (1999). Intrасaccadic target steps during the deceleration of primary saccades affect the latency of corrective saccades. *Experimental Brain Research*, 129, 161-166.

Fujisaki, W., Shimojo, S., Kashino, M., & Nishida, S. (2004). Recalibration of audiovisual simultaneity. *Nature Neuroscience*, 7, 773-778.

Goldberg, M. E., Bisley, J., Powell, K. D., Gottlieb, J., & Kusunoki, M. (2002). The role of the lateral intraparietal area of the monkey in the generation of saccades and visuospatial attention. *Annals of the New York Academy of Sciences*, 956, 205-215.

Haggard, P., Newman, C., & Magno, E. (1999). On the perceived time of voluntary actions. *British Journal of Psychology*, 90, 291-303.

Hellstroem, A. (1985). The time-order error and its relatives: Mirrors of cognitive processes in comparing. *Psychological Bulletin*, 97, 35-61.

Hirsh, I. J. & Sherrick, C. E. J. (1961). Perceived order in different sense modalities. *Journal of Experimental Psychology*, 62, 423-432.

Hodinott-Hill, I., Thilo, K. V., Cowey, A., & Walsh, V. (2002). Auditory chronostasis: Hanging on the telephone. *Current Biology*, 12, 1779-1781.

Hunt, A. R., Chapman, C. S., & Kingstone, A. Chronostasis: Taking a long look at action and time perception. *Submitted for publication*.

Jaskowski, P. (1999). Reaction time and temporal-order judgment as measures of perceptual latency: The problem of dissociations. In G. Aschersleben, T. Bachmann, & J. Mueseler (Eds.), *Cognitive contributions to the perception of spatial and temporal events* (pp. 265-282). Amsterdam: Elsevier.

Johnson, H. & Haggard, P. (2003). The effect of attentional cueing on conscious awareness of stimulus and response. *Experimental Brain Research*, 150, 490-496.

Johnson, H. & Haggard, P. (2005). Motor awareness without perceptual awareness. *Neuropsychologia*, 43, 227-37.

McCloskey, D. I., Colebatch, J. G., Potter, E. K., & Burke, D. (1983). Judgements about onset of rapid voluntary movements in man. *Journal of Neurophysiology*, 49, 851-863.

Moore, T. & Armstrong, K. M. (2003). Selective gating of visual signals by microstimulation of frontal cortex. *Nature*, 421, 370-373.

Morrone, M. C., Ross, J., & Burr, D. (2005). Saccadic eye movements cause compression of time as well as space. *Nature Neuroscience*, 8, 950-954.

- Park, J., Schlag-Rey, M., & Schlag, J. (2003). Spatial localization precedes temporal determination in visual perception. *Vision Research*, 43, 1667-1674.
- Rosenberger, W. F. & Grill, S. E. (1997). A sequential design for psychophysical experiments: an application to estimating timing of sensory events. *Statistics In Medicine*, 16, 2245-2260.
- Ross, J., Morrone, M. C., Goldberg, M. E., & Burr, D. C. (2001). Changes in visual perception at the time of saccades. *Trends in Neurosciences*, 24, 113-121.
- Schlag, J. & Schlag-Rey, M. (2002). Through the eye, slowly: delays and localization errors in the visual system. *Nature Reviews Neuroscience*, 3, 191-215.
- Shore, D. I., Spence, C., & Klein, R. M. (2001). Visual prior entry. *Psychological Science*, 12, 205-212.
- Sommer, M. A. & Wurtz, R. H. (2002). A pathway in primate brain for internal monitoring of movements. *Science*, 296, 1480-1482.
- Spence, C., Shore, D. I., & Klein, R. M. (2001). Multisensory prior entry. *Journal of Experimental Psychology: General*, 130, 799-832.
- Stone, J. V., Hunkin, N. M., Porrill, J., Wood, R., Keeler, V., Beanland, M. et al. (2002). When is Now? Perception of simultaneity. *Proceedings of the Royal Society of London Series B: Biological Sciences*, 268, 31-38.
- Umeno, M. M. & Goldberg, M. E. (1997). Spatial processing in the monkey frontal eye field. I. Predictive visual responses. *Journal of Neurophysiology*, 78, 1373-1383.

Van Beers, R. J., Wolpert, D. M., & Haggard, P. (2001). Sensorimotor integration compensates for visual localization errors during smooth pursuit eye movements. *Journal of Neurophysiology*, 85, 1914-1922.

Volkman, F. C. & Moore, R. K. (1978). Saccadic Eye Movements and the Perception of a Clear and Continuous Visual World. In J.C.Armington, J. Krauskopf, & B. R. Wooten (Eds.), *Visual Psychophysics and Physiology* (pp. 353-362). New York: Academic Press.

Walker, M. F., Fitzgibbon, E. J., & Goldberg, M. E. (1995). Neurons in the monkey superior colliculus predict the visual result of impending saccadic eye movements. *Journal of Neurophysiology*, 73, 1988-2003.

Yamamoto, S. & Kitazawa, S. (2001). Reversal of subjective temporal order due to arm crossing. *Nature Neuroscience*, 4, 759-765.

Yarrow, K., Haggard, P., & Rothwell, J. C. (2004a). Action, arousal, and subjective time. *Consciousness and Cognition*, 13, 373-390.

Yarrow, K. (2003). *Temporal illusions subsequent to movement*. Unpublished doctoral thesis, University College London, London, U.K.

Yarrow, K., Haggard, P., Heal, R., Brown, P., & Rothwell, J. C. E. (2001). Illusory perceptions of space and time preserve cross-saccadic perceptual continuity. *Nature*, 414, 302-305.

Yarrow, K., Johnson, H., Haggard, P., & Rothwell, J. C. E. (2004b). Consistent chronostasis effects across saccade categories imply a subcortical efferent trigger. *Journal of Cognitive Neuroscience*, 16, 839-847.

Figure legends

Figure 1. Schematic showing the physical relationship between temporal intervals and the events that bound them. For both instantaneous and extended stimuli, it is possible to make judgements about both intervals (bounded durations) and events (moments of occurrence/onset/offset).

Figure 2. Schematic of experimental procedure. (a). Sequence of stimuli presented in saccade and control conditions of the three experiments. (b). Timing of the saccade in both long and short saccade conditions and timing of the auditory stimulus for the two temporal order judgement experiments relative to the sequence shown in part a.

Figure 3. Temporal judgement results. Error bars denote standard errors of the mean. Top panel: Mean points of subjective equality between variable and fixed-length stimuli in the chronostasis experiment. Middle/bottom panels: Mean points of subjective simultaneity in the auditory visual TOJ experiment (middle) and the auditory-saccade TOJ experiment (bottom). Positive values on the Y axis indicate presentation of the auditory stimulus after the appropriate referent.

Figure 4. Temporal judgement results for the subset of participants who provided reliable estimates based only on single saccade trials (trials without a corrective saccade). Top panel: Mean points of subjective equality between variable and fixed-length stimuli in the chronostasis experiment. Middle/bottom panels: Mean points of subjective simultaneity in the auditory visual TOJ experiment (middle) and the

auditory-saccade TOJ experiment (bottom). Positive values on the Y axis indicate presentation of the auditory stimulus after the appropriate referent.

TABLE 1.

Mean saccade extent (in degrees visual angle), mean saccade duration (in milliseconds) and mean time of saccade-contingent display change (“trigger time” in milliseconds from saccade onset, with the corresponding size of the correction typically applied in chronostasis experiments shown in brackets) as a function of saccade size for all three experiments. Data from the subsets of participants who could be assessed using only single saccade trials are also shown.

Experiment	Complete Data Set						Single Saccade Trials Only					
	10° Saccade			50° Saccade			10° Saccade			50° Saccade		
	Extent	Duration	Trigger Time (Correction)	Extent	Duration	Trigger Time (Correction)	Extent	Duration	Trigger Time (Correction)	Extent	Duration	Trigger Time (Correction)
Chronostasis	8.0	48	19 (29)	46.0	127	100 (27)	8.3	46	17 (29)	46.9	118	90 (28)
Auditory-Visual TOJ	8.7	52	18 (34)	48.6	128	94 (34)	8.6	53	17 (36)	48.0	129	94 (35)
Auditory-Saccade TOJ	8.7	51	n/a	47.3	126	n/a	8.5	52	n/a	46.9	131	n/a

Figure 1 (Yarrow et al)

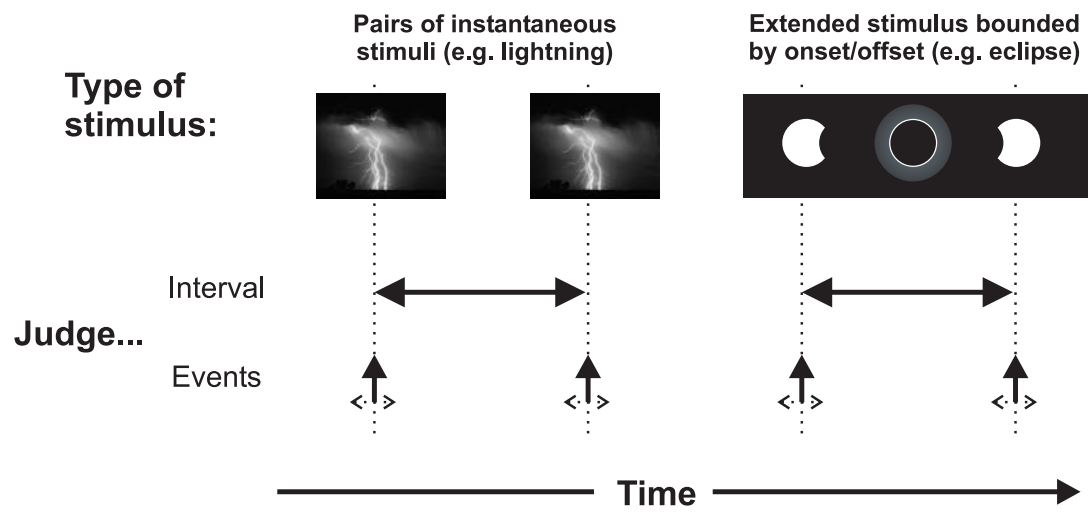


Figure 2 (Yarrow et al)

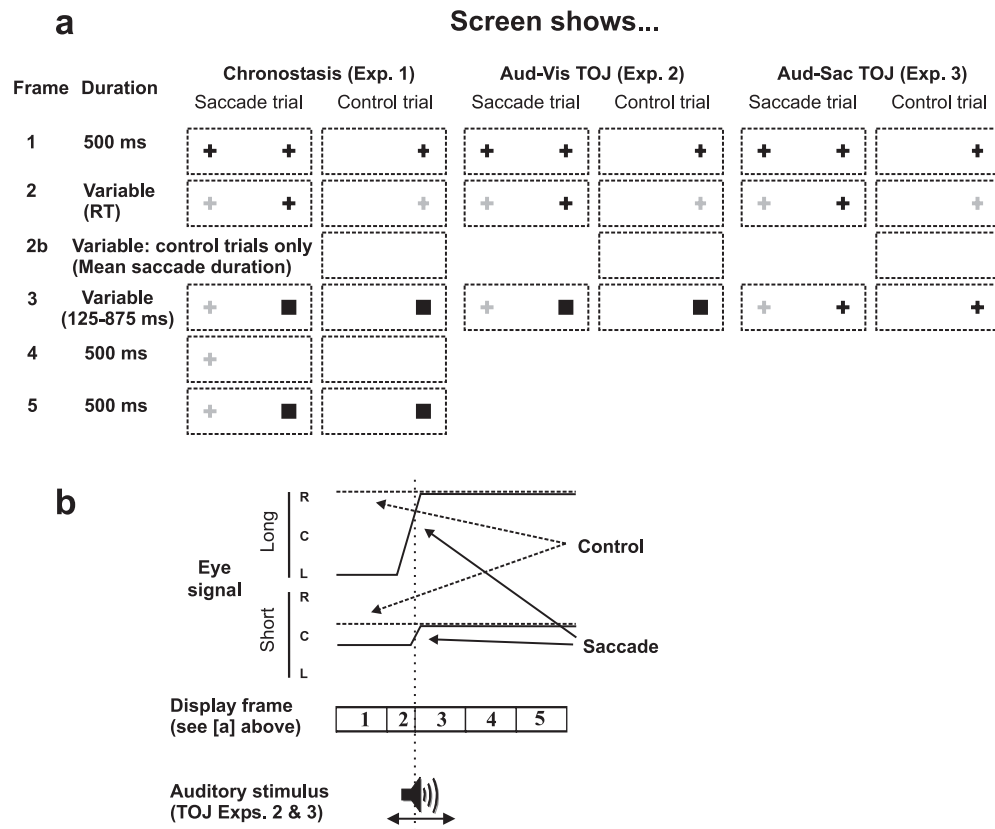


Figure 3 (Yarrow et al)

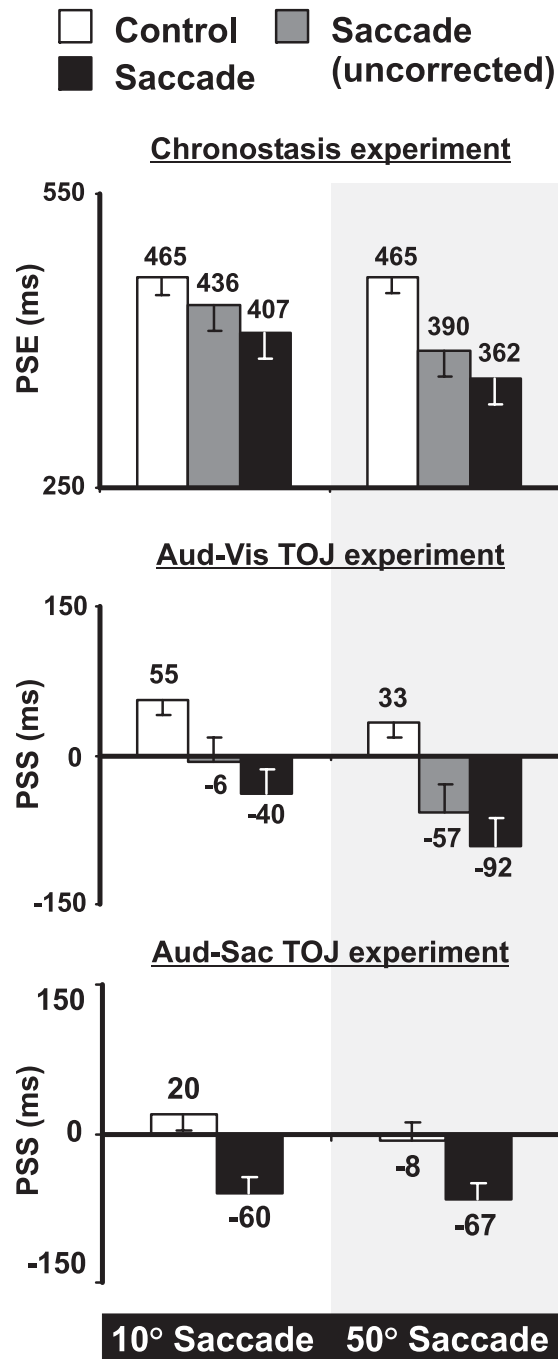


Figure 4 (Yarrow et al)

