Prediction and final temporal errors are used for trial-to-trial motor corrections

Joan López-Moliner[1], Cécile Vullings[2], Laurent Madelain[2], Robert J. van Beers[3]

[1]Vision and Control of Action (VISCA) Group, Department of Cognition, Development and Psychology of Education, Institut de Neurociències, Universitat de Barcelona, Passeig de la Vall d'Hebron 171 08035 Barcelona, Catalonia, Spain

[2]CNRS, CHU Lille, UMR 9193, SCALab Sciences Cognitives et Sciences Affectives, Université de Lille, Lille, France

[3]Department of Human Movement Sciences, Vrije Universiteit Amsterdam, Amsterdam, The Netherlands and Radboud University, Donders Institute for Brain, Cognition and Behaviour, Nijmegen, The Netherlands

Abstract

Correction on the basis of previous errors is paramount to sensorimotor learning. While corrections of spatial errors have been studied extensively, little is known about corrections of previous temporal errors. We tackled this problem in different conditions involving arm movements (AM), saccadic eye movements (SM) or button presses (BP). The task was to intercept a moving target at a designated zone (i. e. no spatial error) either with the hand sliding a pen on a graphics tablet (AM), a saccade (SM) or a button press (BP) that released a cursor moving ballistically for a fixed time of 330 ms. The dependency of the final temporal error on action onset varied from "low" in AM (due to possible online corrections) to "very high" in the other conditions (i.e. open loop). The lag-1 cross-correlation between action onset and the previous temporal error were close to zero in all conditions suggesting that people minimized temporal variability of the final errors across trials. Interestingly, in conditions SM and BP, action onset did not depend on the previous temporal error. However, this dependency was clearly modulated by the movement time in

the AM condition: faster movements depended less on the previous actual temporal error.

An analysis using a Kalman filter confirmed that people in SM, BP and AM involving fast

movements used the prediction error (i.e. intended action onset minus actual action onset)

for next trial correction rather than the final target error. A closer look at the AM condition

revealed that both error signals were used and that the contribution of each signal follows

different patterns with movement time: as movement progresses the reliance on the

prediction error decreases non-linearly and that on the final error increases linearly.

Author summary

Many daily life situations (e.g. dodging an approaching object or hitting a moving target)

require people to correct planning of future movements on the basis of previous temporal

errors. This is paramount to learning motor skills. However the actual temporal error can

be difficult to measure or perceive: imagine, for example, a baseball batter that swings and

misses a fastball. Here we show that in these kinds of situations people can use an internal

error signal to make corrections in the next trial. This signal is based on the discrepancy

between the actual action onset and the expected one. The relevance of this error decreases

with the movement time of the action in a particular way while the final actual temporal

error gains relevance for the next trial with longer motor durations.

Introduction

Timing errors of actions are ubiquitous in daily-life and learning from these errors to

improve planning of future movements is of great importance. Suppose you are batting in a

baseball game and you just missed a fast ball by 50 ms. Assuming you validly expect

another fast ball, how and how much you should you correct for this error in the next

movement may depend on different factors. You could use an estimate of this temporal

error (between the bat and the ball) and try to react earlier if you were late. However your

measurement of this error can be noisy. Since the movement time of your hitting

movement can be quite constant you could alternatively rely on correcting the start of the

swing relative to some relevant moment (e.g. ball motion onset). In this study, we address

on what basis one corrects for temporal errors under different situations of uncertainty

about the final temporal error and the possibility of correction during the movement.

Correcting on the basis of previous errors is one of the hallmarks of motor learning (1,2)

and many studies have addressed how people correct for spatial errors when there is some

external perturbation (e.g. with force-fields or distorted visual feedback) (3-7) or in

situations without perturbations (8).

It is known that larger uncertainty on the observed spatial error leads to smaller

corrections (5,9,10). This is either because one would weight the final sensed error less

relative to some internal prediction of the error, as predicted by Bayesian frameworks (11)

(see Fig1A), or because the noise added by the movement execution is relatively large

compared to the noise in planning that movement (8). The possibility of control while

unfolding the action could also affect the relevance of the final temporal error. For instance,

in open loop actions or when the movement time is very stable (e.g. the baseball example

or in saccadic eye movements) the time of action onset becomes relevant to the final

temporal error (i.e. they are highly correlated) and one could weight the final error less and

base the corrections on some prediction error between the intended and actual action

onset (Fig1A). This can be so especially in fast movements in which predictive components

are important. Alternatively, both prediction and final errors can be used in combination to

specify the next trial correction. We consider these possibilities in this study.

We know that predictions based on forward models (12) are important for correction

mechanisms in general. That is, discrepancies between the prediction and some feedback,

be it internal or sensed (13), are the key for mainstream computational models of motor

learning (14,2) to explain the corrections of saccadic movements (15) or fast arm

movements which are too brief to benefit from the final sensory feedback. In particular, in

conditions where humans are aware of perturbations, errors based on internal predictions

can even override final target spatial errors (16) leading to the distinction between

different kinds of errors: aiming errors (i.e. discrepancy between the planned and final

positions) and target errors (i.e. target vs final position discrepancy), which are important

in motor learning models (17).

Here, we resort to a similar distinction: errors based on the discrepancy between internally

predicted and sensed action onset (prediction error) and temporal errors based on the

experienced sensory feedback at the end of the movement. We expect a different

contribution of each error type in the next trial correction depending on how fast the

movements are (i.e. prediction error being more relevant in faster movements). We test

this hypothesis by using temporal corrections in an interception task.

We will consider the situation in which errors arise when inappropriate motor commands

are issued (execution errors) as opposed to errors caused by external changes (18,5). In

order to see the extent of the corrections, we exploit the properties of the time series of

action onset in arm movements, saccadic eye movements and button-presses to study how

people correct when the initial prediction error at action onset (see Fig1A) contributes

differently to the final sensory temporal error with respect to a moving target in the

different conditions. In the button press condition, a keypress released a fixed movement

cursor to intercept the target. In this condition and in the eye movements condition the

prediction error is highly correlated with the final temporal error. However, the former

error can be perceived with high perceptual uncertainty in the eye movements condition

due to the variability of saccadic reaction time and the temporal and spatial distortions at

the time of saccades starting about 50 ms before saccade onset and up to 50ms after

saccade offset, a phenomenon often termed saccadic suppression (19,20). Finally, arm

movements with different movement times will enable us to determine whether the

relative contribution of either type of error depends on the movement time. A model based

on a Kalman filter will be used to obtain an estimate of the predicted action onset and

therefore, the prediction error. We show that both prediction error relative to action onset

and final temporal error relative to the target can be used in combination for trial-to-trial

corrections. The contribution of each error signal follows a specific time course since action

onset.

Methods

Arm movement experiment

Participants

15 subjects (age range 22-33, 11 males) participated in the experiment. Twelve of them were right-handed and three were left-handed as by self-report. All of them had normal or corrected-to-normal vision, and none had evident motor abnormalities. All subjects gave written informed consent. The study was approved by the local research ethics committee.

Apparatus

Participants sat in front of a graphics tablet (Calcomp DrawingTablet III 24240) that recorded movements of a hand-held stylus. Stimuli were projected from above by a Mitsubishi SD220U ceiling projector onto a horizontal back-projection screen positioned 40 cm above the tablet. Images were projected at a frame rate of 72 Hz and a resolution of 1024 by 768 pixels (60 x 34 cm). A half-silvered mirror midway between the back-projection screen and the tablet reflected the images shown on the visual display giving participants the illusion that the display was in the same plane as the tablet. Lights between the mirror and the tablet allowed subjects to see the stylus in their hand. Virtual moving targets were white dots on a black background (shown red on white in Fig 1A). A custom program written in C and based on OpenGL controlled the presentation of the stimuli and registered the position of the stylus at 125 Hz. The software ran on a Macintosh Pro 2.6 GHz Quad-Core computer. The set-up was calibrated by aligning the position of the stylus

129   with dots appearing on the screen, enabling us to present visual stimuli at any desired

130   position of the tablet.

Procedure

To start each trial, subjects had to move the stylus to the home position (grey dot in Fig

1B). After a random period between 0.8 and 1.2 seconds, a moving target that consisted of

a white dot of 1.2 cm diameter appeared moving rightwards (or leftwards for left-handed

subjects). Targets could move at one of three possible constant speeds (20, 25 or 30 cm/s),

interleaved across the session. The target moved towards two vertical lines of 2 cm height

and separated by 1.2 cm. The space between the lines was aligned with the home position

(Fig 1B). Subjects had to hit the target (i.e. passing through it) at the moment the target was

between the two vertical lines. Because we instructed participants to hit the target in the

interception zone, we only had temporal errors associated to responses, except for the

trials in which subjects missed the zone (less than 2%). The starting position of the target

was determined by the initial time to contact (i.e. time for the target to reach the

interception zone) value, which was 0.8 s for all target speeds. Auditory feedback was

provided (100ms beep at 1000Hz) whenever the absolute temporal error between the

hand and the target was shorter than 20 ms when the hand crossed the target's path

between the two lines. Each subject completed 360 trials.

Data analysis

The individual position data time series were digitally low-pass filtered with a Butterworth

filter (order 4, cut-off frequency of 8 Hz) for further analysis. Hand tangential velocity was

computed from the filtered positional data by three-point central difference calculation.

For each trial, we then computed the time of arm movement onset, the peak velocity, the

movement time (elapsed time from the hand movement onset until the hand crossed the

target's path), and the temporal error with respect to the target. Movement onset was

computed offline by using the A algorithm reported in (21) on the tangential velocity of the

hand.

Fig 1. (A) Action onset and its reliability to predict the relevant task variable: temporal error

with respect to the moving target. The uncertainty in determining the planning of the action

onset (hidden variable) is illustrated by the orange Gaussians, while the execution (or

measurement) noise is denoted by the blue Gaussians centered at the actual action onset.

Different variability in the planning of action onset or its measurement is denoted by the type

of line (dashed: lower noise; solid: higher noise). The prediction error is the difference

between the planned (or predicted) and actual action onset. The top row illustrates a slow

movement after action onset (longer duration until crossing the target) and the bottom row a

fast movement. One would expect larger corrections when the measurement noise of the

actual action onset is lower (blue dashed curves) relative to the planned noise (solid orange

curves). The decay of the relevance of the prediction error after action onset is denoted by the

green line, while the increasing relevance of the final temporal error for next trial is denoted

by the red line. These particular trends are based on the assumption that the quadratic sum of

both lines would sum up to one. (B) Illustration of the experimental tasks: arm movements

(top) and eye movements (bottom).

Button press experiment

Participants

Eight participants (age range 23-32, 5 males) participated in this experiment. All of them

had normal or corrected-to-normal vision, and none had evident motor abnormalities. All

subjects were right handed and gave written informed consent. The study was approved by

the local research ethics committee.

Apparatus

Stimuli were shown on a Philips CRT-22 inch (Brilliance 202P4) monitor at a frame rate of

120 Hz and a resolution of 1024 by 768 pixels. The viewing distance was about 60cm and

the head was free to move. A custom program written in C and based on OpenGL controlled

the presentation of the stimuli and registered the time of the button-presses by sampling

an ancillary device at 125 Hz. The software was run on a Macintosh Pro 2.6 GHz Quad-Core

computer.

Procedure

The stimuli were the same as in the Arm Movement experiment except for the fact that the motion was presented on the fronto-parallel plane. In this experiment subjects had to press a button that initiated the release of a moving cursor from the home position. Subjects had to press the button timely so that the cursor would hit the target when passing between the two vertical lines (interception zone). The movement time of the cursor from the home position to the interception zone was 312 ms and its velocity profile was extracted from an actual arm movement. In this experiment the time of the button-press determined completely the final temporal error. Subjects took the same number of trials and sessions as in the Arm movement experiment.

Eye Movement experiments

Participants

Fifteen participants (age range 18–47, 7 males, including two authors) participated in the experiments. Among them, ten (age range 18–46, 4 males) participated in th first experiment (termed knowledge of results, KR) and twelve (age range 23-47, 5 males) participated in the second one (knowledge of performance, KP). They had normal or corrected-to-normal vision. All participants gave written informed consent. The study was approved by the local research ethics committee.

Apparatus

Stimuli were generated using the Psychophysics Toolbox extensions for Matlab® (22,23) and displayed on a video monitor (Iiyama HM204DT, 100 Hz, 22''). Participants were seated on an adjustable stool in a darkened, quiet room, facing the center of the computer screen at a viewing distance of 60 cm. To minimize measurement errors, the participant's head movements were restrained using a chin and forehead rest, so that the eyes in primary gaze position were directed toward the center of the screen. Viewing was binocular, but only the right eye position was recorded in both the vertical and horizontal axes. Eye movements were measured continuously with an infra-red video-based eye tracking system (Eyelink®, SR Research Ltd., 2000 Hz) and data were transferred, stored, and analyzed via programs written in Matlab®. The fixation point that was used as a home position for the gaze was a 0.4 deg×0.4 deg square presented always on the bottom left quadrant of the screen. The target was a 0.4 deg of diameter disk, and the interception area was a goal box of 0.6 deg of diameter. The interception area was located 12 deg to the right of the home position (see Fig 1B). All stimuli were light grey (16 cd/m^2 luminance) displayed against a dark grey background (1.78 cd/m^2 luminance). Before each experimental session, the eye tracker was calibrated by having the participant fixate a set of thirteen fixed locations distributed across the screen. During the experiment the subject had to look at the center of the screen for a one-point drift check every fifty trials. If there was any gaze drift, the eye tracker was calibrated again.

Procedure

A session consisted of 390 discrete trials lasting between 2 and 2.45 secs. Each trial started with the subject looking at the fixation point for a period randomly varying between 700 and 1100ms. Participants were instructed to make a saccade to intercept the target, that was moving downward towards the interception area, at the time it was within the interception area. Targets moved with a constant velocity of either 20, 25 or 30 deg/s. Target velocities were interleaved across trials in both the KR and KP-interleaved experiments. In the KP-blocked condition, the targets' velocities were presented in three consecutive 130-trial blocks (in a pseudo-random order counterbalanced across participants). The same participants experienced both KP conditions; the order was counterbalanced across subjects. The time to contact the interception area was 600 ms since target onset, and the target starting point was therefore depended on the actual target velocity. The occurrence of a saccade was crudely detected when the online eye velocity successively exceeded a fixed threshold of 74 deg/s. If the offset of an ongoing saccade was detected before the target reached the interception zone the target was extinguished at the next frame, i.e. within the next 10ms (offline measurements revealed that the target disappeared on average 2 ms after the time of the actual saccade offset). If the target center was aligned with the goal box before a saccade was detected we extinguished the target. Therefore, participants never saw the target after it had reached the interception zone. We delivered an auditory feedback (100 ms beep at 1000 Hz) if the eye landed within 3 deg of the interception area with an absolute temporal error smaller than 20 ms. To this end, the actual saccade onset- and offset-time and position were

computed immediately after the saccade using the real-time Eyelink algorithm with a 30

deg/s velocity and 8000 deg/s^2 acceleration thresholds (on average we retrieved these

values 12 ms after the end of the saccade). In the first experiment (KR), participants did not

receive explicit feedback on their performance other than the auditory one. In the second

experiment (KP), the actual temporal error was displayed numerically in milliseconds at

the end of each trial (KP). For offline analyses, a human observer validated each saccade

manually. Saccades with an amplitude gain smaller than 0.5 or a duration longer than 100

ms were discarded.

Analysis

Testing for the optimality of corrections: autocorrelation analysis

It is known that the serial dependence of consecutive movement errors depends on the

amount of trial-by-trial correction (24). If participants are trying to make temporal

corrections based on the prediction error we should be able to see a serial dependence of

the action onset (T^s) in both simulated and behavioral data that will depend on β, the

fraction of correction. Suppose that no corrections are made whatsoever. In this case, we

expect that consecutive initiation times will be similar to the previous one. The absence of

correction would be revealed by a significant positive lag-1 autocorrelation function

(acf(1)) of the action onset under the assumption that planning noise accumulates from

trial to trial. On the contrary, if one aims at correcting for the full observed error (β=1) then

consecutive movements will tend to be on opposite sides of the average response because

one corrects not only for the error in planning but also for the random effects of execution

noise. In both scenarios (β=0 and β=1) there is an unnecessarily large temporal variability

due to different causes. If one does not correct, not only will previously committed errors

persist but also previous planning errors will accumulate across trials increasing the

variability much like when one repeatedly reaches out for static targets. If one does fully

correct, the variability due to changes in the planned time will be larger than if smaller

corrections were made. In either case the process is not optimal in the sense that the

temporal error is more variable than necessary. When corrections are large enough to

compensate for random variability but not too large to make the behavior unstable, then

the temporal error variance is minimal and the correction fraction is optimal. For such

fractions of corrections, acf(1) of the temporal errors will be zero (8). In our case

participants can correct by changing the action onset, so we are interested in the cross-

correlation function (ccf(1)) between action onset at trial *i* and the relevant target error at

trial *i-1*. Note that for the button press condition action onset is perfectly correlated with

the final error and for eye movements the correlation is very high, therefore the ccf(1)

would be undistinguishable from the acf(1) of either the actual error or action onset.

Similarly, a zero cross-correlation ccf(1) would denote an optimal change of the time of

action onset to correct for the previous error.

Dependency on the previous actual temporal error

We analyzed the dependency of the time of action onset in the current trial on the temporal

error with respect to the target in the previous trial in the different conditions by fitting

linear mixed-effect models (LMMs), which enable us to easily analyze the effects of the

previous trial on the current response. In the model, the action onset time was the

dependent variable and the previous target temporal error, the independent variable. Both

intercept and slope were allowed to vary as random effects across subjects Both intercept

and slope were allowed to vary as random effects across subjects. We used the lmer

function (v.1.0–6) (25) from R software

Simulations and process modelling

In order to estimate the prediction error relative to action onset we used a Kalman filter to

estimate the predicted action onset time before the actual observation. For the Kalman

filter to work, one needs knowledge of the sources of variability (process and measurement

noise). To get further insight into the variance of the generative process of the action onset,

we implemented the temporal corrections at the action onset across simulated trials in

which we manipulated different sources of variability: process variability and

measurement (i.e. motor) variability. The process variance in the time of action onset is

captured by the following expression and mainly accounts for variability of sensory origin:

$$V_t = \left(\frac{\sigma_x}{v}\right)^2 + \sigma_t^2 \qquad (1)$$

The first term is velocity dependent and the second one corresponds to a timing variability

(26). σ_x is the spatial variability about the target position at action onset and v is the target

speed. Uncertainty caused by measuring target speed may likely contribute to the timing or

velocity dependent variability. However, in practice both sources of variability are difficult

to tease apart because an error in misjudging the target position would be

indistinguishable from a timing error. In each simulated trial i the generation of an

intended action onset τ is a stochastic process where τ_i, the planned action onset at trial i,

is updated according to:

$$\tau_{i+1} = \tau_i - \beta e_i + q, \quad q \sim N(0, V_t) \qquad (2)$$

where β is a learning rate or, in our case, the simulated fraction of error (e) correction and

q is the process noise related to eq. 1. The actual action onset T^s is simulated by adding

measurement noise (produced by motor noise) to the intended action onset:

$$T_i^s = \tau_i + r, \quad r \sim N(0, \sigma_m^2) \qquad (3)$$

where r is the execution noise (added noise from when the motor command is issued until

movement onset). The final temporal error e at trial i is given by:

$$e_i = T^T - (T_i^s + T_i^m) \qquad (4)$$

where T^T is the time at which the target is centred within the interception zone and T_i^m is

the movement time. Without loss of generality, we set T_i^m and T^T to zero.

Modeling the corrections. Using the equations introduced above, we modeled a trial-to-trial

correction of the time of initiation, assuming that all the final temporal error is fully caused

by the time of action initiation T^s. This was certainly the case in the eye movement

conditions and button press conditions – because in our case the time to reach the target

was fixed once the button was triggered - while for arm movements there is some room for

online corrections by adjusting the movement time. We modeled 16 different correction

fractions from 0.06 to 1 by increments of 0.06 (range: 0.06-0.96) and four values of r ($\sigma_m =$

0.022, 0.05 0.1 and 0.2 s). We set σ_x to 1 cm and σ_t to 0.05 s. These values were used with

three target velocities: 20, 25 and 30 m/s resulting in a mean process noise variance of

0.0042 s^2. These choices were guided by values reported in previous studies (26,27). If the

simulated time at trial i was shorter (i.e responding too early) than a target value (e.g. 0

ms) by some magnitude e_i, the value of the intended time onset (τ_{i+1}) was increased by βe

on the next trial, or decreased if the observed time was too long. We ran 1000 simulations

for each combination of β and r. Each simulation consisted of a series of 360 responses or

trials in which speed was interleaved (but note that the time the target took to reach the

interception zone was the same for all speeds, so target speed changes between

consecutive trials are not a problem for making trial-by-trial corrections).

Estimation of process and measurement variances

The fraction of correction β can be estimated from the behavioural data through the

Kalman gain (K) (9). The Kalman filter estimates the planned action onset as the hidden

state from the actual (noisy) action onsets. In order to know K one possibility is to estimate

the process (V_t) and measurement (σ_m^2) variances (28). In steady state (which in our

experiments was approached after a few trials), K can be approximated by the following

expression (5):

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$$K = \frac{V_t}{V_t + \sigma_m^2} \qquad (5)$$

Since V_t and σ_m^2 are known in the simulations, this expression approximates the

corresponding optimal correction fractions for the different values of simulated motor

(measurement) noise: K= 0.09, 0.29, 0.61 and 0.86 starting with largest value of σ_m^2.

This is not as straightforward when analyzing the behavioral data since both parameters

are unknown. In order to estimate the process noise variance V_t in the different

experimental conditions, we proceeded as follows: first we fitted a linear model to the

process noise variance in the simulated data based on terms that could be obtained from

the observed data (both simulated and behavioral). Second, we used the fitted model to

predict the process variance in the experiments.

The linear model contained three terms plus their interactions. Two of the terms come

from the decomposition of the actual temporal variance into estimates of $(\sigma_x/v)^2$ and σ_t^2

which may contain measurement noise because they were estimated from the observed

simulated data. The third term was the ccf(1) of the action onsets. When we fitted the

model to the process noise variance in the simulated data the model accounted for the 94%

of the variance.

In order to obtain the two first terms of the linear model in both simulated and behavioral

data, we fitted the following model (29) to the total spatial variability:

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$$SD_x = \sqrt{\sigma_x^2 + (v\sigma_t)^2} \qquad (6)$$

We estimated σ_x and σ_t^2 for each series of 360 trials in the simulations and for each

participant and condition. Fig S1A shows the simulated process variance against the

predicted process variance from the model. Fig S1B shows the estimated process variance

in the human data based on the linear model used to fit the process variance in the

simulated data. The process variance is plotted against the whole observed temporal

variance. Fig S1C shows how the whole temporal variance is decomposed according to

equation eq. 1.

Once we had estimated the process variance V_t, the measurement noise was the only free

parameter when fitting the Kalman filter to the behavioral data.

The Kalman filter model

We applied a Kalman filter model to determine the degree of correction based on the

prediction error. As shown in eq. 3 the actual action onset T^s is a noisy realization of the

predicted action onset τ. We can rewrite eq. 2 as:

$$\tau_i = \tau_{i-1} + c_i + q \qquad (7)$$

where c_i is a correction factor that has to be determined by the Kalman filter. But, how does

the Kalman filter work out the magnitude of the correction? The Kalman estimates c_i

recursively by combining a predicted action onset (i.e. a priori) and the observation of

action onset that has been corrupted by noise T^s. After movement onset at trial i, the

Kalman filter estimates a posterior time of action onset (denoted by the hat operator):

$$\hat{\tau}_i = \tau_i + K_i(T_i^s - \tau_i) \qquad (8)$$

The posterior will be used as a predicted action onset time in trial $i+1$, becoming τ_i in

(eq. 7). K_i is called the Kalman gain and reflects the fraction of correction of the prior time

of action onset. If $K = 0$ no change is made in the planning for the next trial; alternatively, if

$K = 1$ the whole difference between the prediction and the observed action onset is

accounted for in the posterior. We will refer to the difference between T^s and τ as

prediction error.

In order to compute K, the Kalman filter takes into account the uncertainty of the

prediction and the one of the observation.

$$K_i = P_i(P_i + \sigma_m^2)^{-1} \qquad (9)$$

where P_i is the uncertainty in the prediction of the planned onset time before the

observation of action onset takes place. Note the equivalence with eq. 5. This *a priori*

uncertainty is also obtained from the posterior estimate of the uncertainty, $\hat{P}$, in trial *i-1*:

$$P_i = \hat{P}_{i-1} + V_t \qquad (10)$$

The Kalman filter will correct the internal estimate (i.e. predicted action onset) by a

fraction K of the prediction error $T^s - \tau$. However, although the prediction error is highly

correlated with the final temporal target error in some conditions, the prediction error is

not the task-relevant error shown in eq. 4. We analysed the correction with respect to

action onset because we are interested in how people correct in the planning phase.

The planning of the action onset should aim at minimizing the expected final temporal

error $(e(\tau) = 0)$ which can be stated as:

$$e = T^T - \tau + T^m \qquad (11)$$

In order to be accurate across all observed responses we need that:

$$T^T = T^s - T^m \qquad (12)$$

Substituting eq. 12 in eq. 11:

$$e = (T^s - T^m) - (\tau + T^m) = T^s - \tau \qquad (13)$$

which is the prediction error with respect to action onset that the Kalman filter is

correcting. This equation shows that, given some constraints in the distribution of

movement time T^m (i.e. shifted mean with respect to T^s), correcting for the prediction

error is equivalent to correcting for the final temporal error. This is true on average, since

for individual trials the prediction error does not necessarily correspond to the final error.

Parameter estimation. In order to estimate the predicted action onset time (τ) σ_t^2, the

measurement noise was the only free parameter. σ_t^2 was determined by minimizing the

negative log-likelihood of the actual action onset given the estimated (planned) action

onset computed by the Kalman filter in each participant and condition.

Fig 2. (A) Example of action onset times for the different conditions. Different conditions are color-coded (see legend in Fig2B). Each response series corresponds to a single participant. The two examples of the arm movement condition correspond to a fast (top-left) and slow (top-right) participant. The action onset time is centered at zero (by substracting the mean) to optimize panel space. (B) Mean lag-1 cross-correlation functions, ccf(1), between the time of action onset at trial t and actual temporal error at trial t-1 for the different conditions. Error bars denote the 95%-CI of the correlation coefficients. (C) (Simulated data) The temporal error variance as a function of the simulated fraction of correction β for the four different levels of simulated execution noise. The arrows point to the value of β that corresponds to the minimum variance. As can be noted, this fraction of correction is similar to

428   *the simulated gain (which in turn depends on the level of execution noise, see legend in panel*

429   *D). The largest gain (i.e. K=0.86) requires larger corrections in order to minimize the*

430   *variance. (D) (Simulated data) The acf(1) values of action onset in the simulated data against*

431   *the amount of correction. As can be seen, the acf(1) should be near zero to be optimal for each*

432   *gain.*

Results

Are temporal corrections optimal?

Assuming that open-loop control schemes are used to execute the movements, we expect a

modulation of the initiation times by prior temporal errors but also that the time of action

initiation relative to the interception time is not statistically different across different

target velocities. That is, relevant decision variables regarding the action onset would

mainly rely on temporal estimates of the remaining time to contact from the action

initiation. An ANOVA on the linear mixed model in which action onset was the dependent

variable, target speed (fixed effect as continuous variable), conditions (fixed effect as

factor) and subjects treated as random effects failed to report a significant effect of target

speed on action onset ($F<1$, $p=0.96$) and only condition was significant ($F=53$, $p<0.001$).

The interaction was not significant ($F<1$, $p=0.473$).

Fig 2A shows examples of series of observed action onset times from the different

experimental conditions. From the different series we first computed the lag-1 cross-

correlation function (ccf(1)) between the action onset in trial t and the temporal error in

trial t-1. To qualify as "optimal correction", ccf(1) between previous target error and action

onset must be zero (or very close to zero). Fig 2B shows the mean lag-1 cross-correlation

function ccf(1) between the time of action onset and previous error for the different

conditions. These values are consistent with participants changing their action onset

optimally or near optimally. ccf(1) values for arm movements and eye movements (KP-

interleaved) were very low but significantly different from zero.

We conducted the same analysis on the simulated data. First, Fig 2C shows how the fraction

of correction modulates the overall temporal variance. The correction fraction for which

the temporal variance is minimal is the optimal correction fraction. This fraction is

different for the different levels of simulated execution or motor noise (measurement

noise) that correspond to the different Kalman gains. Importantly the values of optimal

correction correspond to values of ccf(1) (or acf(1) in the simulations) very close to zero

(Fig 2D). From the different data patterns shown in Fig 2 we can be quite confident that

participants corrected by changing the time of action onset in an optimal way or close to an

optimal way.

463   Dependency on the previous temporal error

Autocorrelation indicates how consecutive points tend to be around the mean (e.g. if one

overcorrects then consecutive points will likely be on opposite sides), but does not indicate

which fraction of the previous actual error is being accounted for in the change of action

onset in the present response. In order to get an estimate of this magnitude we ran the

Linear Mixed Model (described in the methods sections). The time of action initiation at

each trial was fitted as a function of the previous final temporal error. The slope denotes

how much the previous error is considered. Fig 3A (red dots for the Arm movement

condition and boxplot) shows the values of the slopes. The larger slopes were found in the

arm movement condition and the average slope was significantly different from zero

(slope=0.12 fraction/trial, t=5.39, p<0.0001). For the remaining conditions, only in the Eye

movements (KP-interleaved) the slope was significantly different from zero (slope=0.07

fraction/trial, t=3.78, p=0.004). The distribution of individual slopes in the Arm

movements condition reveals an interesting and clear positive linear relation between the

movement time and the dependency on the previous temporal error (Fig 2A main panel).

Participants with slower arm movements modified more the action onset in the present

trial more as a function of the previous interception temporal error. Movement time in

saccades did not have enough variability across subjects to observe a similar distribution

and cursor movement time was fixed in the button press condition. The corrections in the

Button press and Eye movements conditions (KR and KP-blocked) did not rely on the

previous temporal error with respect to the target (slopes not different from zero).

Fig 3. (A) Dependency (slope in the linear model) of the action onset on the previous actual

temporal error as a function of movement time in the Arm movement condition. Each dot

corresponds to an individual participant. (inset) The slopes (boxplot) corresponding to the

The question then is how do people correct in these conditions? One possibility (depicted

in Fig 1A) is that people corrected the aimed action onset, not based on the final temporal

error but on the difference between the planned action onset and the actual action onset

(i.e. the prediction error). Since we could not measure this prediction error in the

experiment, we had to model correcting based on this error to infer how large these

corrections were. We used a Kalman filter model to estimate the Kalman gain, that is the

fraction of the prediction error that is used to update the aimed action onset for the next

trial.

Corrections based on the action onset prediction error

Unlike in the simulations, both the process (V_t) and measurement (σ_m^2) variances are

unknown in the experiments. The Kalman filter requires knowledge of these variances in

order to estimate the optimal Kalman gain (K). We inferred the process noise variance (V_t)

in the experimental data by predicting this variability from the linear model that was used

to fit the process noise variance in the simulations.

The standard deviation of the process noise for the eye movements (KR) was larger than in

the other conditions (58 ms versus 37 ms in Button press, 38 ms in Arm movements, 44 ms

in Eye movements KP-Blocked and KP-interleaved). Only the difference between KR and

Arm movements reached significance (corrected p=0.009). For the hand movements

condition, we found a clear difference between slower and faster (<400 ms of movement

time) participants with the latter being more variable (49 ms versus 17 ms, p<0.0001).

The Kalman filter was fitted to the time series based on the action onset with the

measurement noise variance (σ_m^2) as the only free parameter.

Fig 4. (A) Relation between the contribution to the correction of the action onset based on the

previous temporal error (Target error contribution) and the contribution to the correction

based on the prediction error (i.e. based on the Kalman gain) for the Arm movement

condition. Each dot corresponds to a different participant and the shape corresponds to the

classification of the movement time duration (fast: less than 0.5 s; slow larger than 0.5 s). The

target error contribution between 0 and 1 corresponds to the actual fraction of correction of

the final error obtained from the slopes shown in Fig 3A by using a linear the model. The

linear model was obtained with the simulated data and allowed to estimate the

corresponding slope for each (simulated) fractions of corrections. The Kalman gain

contribution is obtained by scaling the Kalman gain between 0 and 1. The grey line

corresponds to the predicted relation assuming that both correction fractions are combined

into a quadratic sum (equation 14) (B). Evolution in time of the correction fractions (i.e.

relevance given to prediction and final error). The shift of the two lines (prediction and

feedback-based correction) depends only on the time at which the final temporal error will

start to be considered for correction in the next trial (about 171 ms in the figure). See text and

equation 15 for the computation of the two lines in panel B.

The individual as well as the mean Kalman gains for participants and conditions are shown

in Fig 3B together with the value of the ccf(1). This plot shows that different values of

correction with respect to the action onset prediction error (i.e. Kalman gain) can

correspond to optimal or near optimal corrections. The estimated magnitude of the

measurement noise was very similar among conditions and its standard deviation ranged

from 33 ms (Arm movements) to 35 ms (Eye movements KP-Blocked). No difference was

significant. There was no difference between slow (31 ms) and fast (34 ms) participants in

the arm movement condition. Thus, the differences in process to measurement variance

ratio that determines the Kalman gain are due to differences in the process noise. Fig 3B

also shows the difference between slow and fast participants in the Arm movement

conditions with the parameters corresponding to the fast group being very similar to those

of the Eye movements and Button press conditions. This is consistent with people

correcting less based on the prediction error (difference between planned action onset and

actual action onset) when they moved more slowly.

This high Kalman gain (i.e. use of prediction error) would be expected in the Eye movement

conditions because the sensory feedback of the final temporal error can be noisy. However

this is not the case for the Button press condition in which participants could perfectly

perceive the error. Since the final temporal error is fully explained by the time of action

onset in this condition, it seems that the correction based on the prediction error rather

than the final error seems to be based on the reliability (or correlation) of the prediction

error with respect to the final temporal error.

Relation between prediction error and final temporal error

Based on the auto-correlations, people make corrections that minimize the temporal

variability across trials. However, in some conditions there is no dependency on the

previous temporal error. One possibility that would explain this apparent contradiction is

that people correct based on some combination of the prediction error and the final

temporal error, with this combination being modulated by the movement time. This would

explain the difference in Kalman gain between slow and fast movement times in the Arm

movement condition. The prediction error (actual onset minus planned onset times) would

be weighted more heavily in the next trial for short movement times with a progressively

decay in favor of the final temporal error (relative to the target) as movement time

increased (Fig. 1A). The estimated Kalman gains support this hypothesis, but to further

explore this possible use in combination of both error signals we plotted the relation

between the (normalized) dependency on the actual previous temporal error and the

Kalman gain contribution (1 meaning that all the estimated Kalman gain is used for

correction) for the different subjects who participated in the Arm movements condition

(Fig 4A). Interestingly, the relation between the corrections fractions based on both types

of errors resembles a specific type of combination described by the grey line in Fig4A. This

line denotes a quadratic sum of the two error signals contributions (x and y axes). For

example, if we take any point (x,y) along the grey line (e.g. x=0.7, y=0.72), the quadratic

sum $\sqrt{(x^2 + y^2)}$ adds to one.

Faster subjects (circles in Fig 4A) show larger Kalman gains (as in Fig 3B) and the

dependency on the previous target error is small. This trade-off changes for slow

participants (triangles in Fig 4A). This transition is well described by the grey line that

corresponds to a quadratic sum of the two fractions of correction:

$$\omega = (\beta^2 + K^2)^{1/2} \qquad (14)$$

where β and K correspond to the contributions of the actual temporal error and prediction

error respectively and ω denotes the trade-off between the two error signals.

Some models of cue combination (30) have used the expression represented by eq. 14 to

define the combined reliability (i.e. reciprocal of variance) from the individual reliabilities.

In this sense equation eq. 14 captures the maximum likelihood estimation of the combined

reliability. Such a combined use of error signals (not contemplated in our model) results in

an increased precision with respect to using either signal alone. However, one important

assumption is that the individual reliabilities are independent (i.e. uncorrelated). This is

not the case in our two temporal error signals. The prediction error and the final target

error are highly correlated in our conditions (see Fig S2). Due to the magnitude of the

correlation, there is very little or no benefit in correcting based on integrating or combining

both error signals (31) (see Fig S3). The expression in eq. 14 then should not be interpreted

as a weighted combination but as denoting the trade-off between the two error signals.

Fig 4A does not contain any temporal information concerning how the relevance of each

signal β and K evolves over time. The modulation of the relative relevance in next trial by

the movement time (Fig. 4B) sheds some light on how both types of errors (the prediction

error based on K and the final temporal error based on β) evolve over time. Fig 4B simply

plots the same points shown in Fig 4A as a function of the corresponding movement time

for each point. As can be seen, the relevance of the prediction error decreases in a non-

linear manner, while the contribution of the target error increases linearly. Moreover, Fig

4B shows that the increase of β takes place after some critical movement time or

sensorimotor delay δ_{sm}. We can formulate this trend (red lines in Fig 4B) according to the

following piecewise function:

604
$$\beta_t = \begin{cases} 0, & \text{if } T^m \leq \delta_{sm} \\ aT^m, & \text{otherwise} \end{cases} \qquad (15)$$

605   where T^m is the movement time. On the other hand, we can obtain how the corresponding

606   fraction of correction given to prediction (K) as a function of the change in β across time

607   from eq. 14. The value of K is expected to decrease with time according to this expression

608   (green line in Fig 4B):

609
$$K_t = (1 - \beta_t^2)^{(1/2)} \qquad (16)$$

610   We only adjusted δ_{sm} in Fig 4B so that the dcrease of K_t and simultaneous increase of β_t

611   minimized squared errors across the red and green dots in Fig 4B. The parameter a in

612   eq. 15 will then depend on δ_{sm}. In our case a=3.58. The obtained value of δ_{sm} was 170 ms.

613   This value imposes a lower temporal bound on the movement time from which the final

614   temporal error with respect to the target will be considered to be corrected for in next trial.

Discussion

We show that people minimize the temporal variance across trials when correcting for temporal errors. This is concluded from the structure of temporal correlations (32): we report near zero lag-1 ccf between action onset and the previous temporal error in an interception task. However, which error signal is predominantly used to correct differs depending on the condition and duration of the movement of the action: slower movements showed larger dependency on the previous final temporal error with respect to the target in the Arm movement condition. In most conditions (Eye movements, Button press and fast Arm movements), people strongly rely on the prediction error at action onset rather than the actual temporal error with respect to the target to change the planned initiation time in the next trial. This is based on the high values of the Kalman gain, which denotes that the prior (planned) action onset will be shifted in the next trial by a large fraction (about 0.8) of the prediction error (i.e. difference between the prior and actual action onset). In the Arm movements condition, fast movements were initiated later, therefore their planning could have been more robust than slow movements (initiated earlier) increasing the reliability of the prediction error. The reliance on the prediction error has obvious advantages when the final sensory feedback is noisy. This is the case in the Eye movements conditions where perception of the sensory temporal error signal can be noisy. Correcting from the prediction error at action onset is possible under some restrictions (e.g. open loop), as there is a clear correlation between the prediction error and the final temporal error (Fig S2). Due to the possibility of making corrections during

the movement, the correlation is lower in the Arm movement condition, and this is the

condition in which we find less contribution of the prediction error (slower movements).

Models of motor learning would predict less correction (e.g. lower learning rate) when the

sensory feedback is more uncertain (5,10,11,9) or if error signals are perceived less

relevant (7,33). The Bayesian explanation is that the sensory error feedback is weighted

less in favor of internal state estimates (34). This is usually the case when studies focus on

the reliability of the final task-relevant error. Our findings, however, show that the picture

can be more complex. We found the same amount of correction in the Eye movements and

Button press conditions while the final temporal error signal is likely perceived with very

different uncertainty, as the Button press is more reliable. Our results show that

corrections in these two conditions are executed in a very similar way (similar Kalman

gains and dependency on the previous temporal error). The way temporal errors are

corrected (mainly in the Button press condition) challenges some of the assumption of

current models, and merely considering the final sensory error might not suffice, at least

when temporal errors are relevant. The control at the time of the button press seems to be

an important factor as to which error signal will be used to correct in the next trial.

Prediction errors have been mostly regarded as relevant for online corrections when the

final sensory feedback would arrive too late to make useful corrections. Here we show that

prediction errors can be useful for trial-by-trial corrections after the actual error is known.

This process is apparently not affected by low uncertainty of the final error (e.g. Button

press condition) because it does not override the use of the prediction error at action onset.

Interestingly, it seems that the contribution of the final error for next trial correction

depends on the movement time. We found that the final target error will start to be

weighted for correction in the next trial by movement times close to 200 ms. This is

consistent with the value that has recently been reported for online spatial corrections

when there is a target movement (35). The model (Fig 4B) also predicts that, as movement

progresses, the reliability of the prediction error at action onset decreases reaching a

minimum after 400 ms. This time course of the contribution of the prediction error for the

next trial parallels the shift from prediction to sensory signals in online correction of

spatial errors of arm movements (12).

The evolution of the contirbution given to prediction and final errors suggests that the

system has some access to or knowledge about the noise that is added from the time of

action onset. This would be in agreement with previous work showing that the motor

system is able to model the temporal uncertainty of the movement time when

programming reaching movements under temporal constraints (36).

The relevance of the prediction error in trial-to-trial temporal corrections is mostly

noticeable in the Eye movements condition. The behavioral plasticity of the saccadic

system has been well established in the temporal domain: saccade latencies may be

strongly affected by a number of factors such as temporal stimulus arrangement (37),

stimulus properties (38,39), urgency (40), expectations (41) or reinforcement

contingencies (42). Moreover, studying saccades directed toward a moving target revealed

that the saccadic system takes into account both the saccade latency and duration, and is

able to adjust to experimentally induced perturbations (43). Our current results shed a new

light on the underlying adaptive process revealing that the temporal error is integrated on

a trial-to-trial basis to adjust the saccade-triggering. It is noteworthy that these conclusions

nicely echoe ones from saccade adaptation studies in which the adjustment of saccade

amplitude has been well accounted for by postulating a Bayesian integration in which the

weight associated with each piece of information is adjusted depending on the sensory

evidence available (2).

Concerning the eye movements, we found a small but significant dependency of the final

temporal error on whether the target speed was interleaved and knowledge of

performance based on the final error was provided (KP-interleaved). Although the

correlation between the final temporal error and prediction error is slightly weaker in the

interleaved condition (slope of 0.75 Fig S2) than in the other two Eye movements

conditions (slope of 0.80 and 0.81), the differences are not significant. The Kalman gain was

not significantly smaller in this condition (KP-interleaved) compared to when the speed

was blocked (KP-Blocked), which denotes also a strong weight of the prediction error.

However, the condition of a non-stationary environment (variable speed across trials)

could have encouraged a larger contribution of the final error. Note that knowledge of the

magnitude of the error was not provided in the KR condition in which the speed was also

interleaved. Conditions of stationary environment can be an important factor that also

contributes to how the final error is weighted. In addition to stationary stimuli conditions,

the temporal restrictions on which feedback is provided can also change. One limitation of

our study is that we have used a relatively constant temporal window for participants to hit

the moving target and the feedback was given with respect to a fixed temporal window.

The learning rates or correction fractions might be also tuned to this temporal requirement

and different learning rates could have been observed by varying the temporal window on

which feedback was provided. For example, lax temporal constraints would lead to smaller

learning rates. Recent studies have shown that different sensitivities to execution errors

arise in motor learning depending on the stationary conditions of the environment (44).

From our study we do not know whether the specific weighting pattern of the two signals

can be generalized to other conditions, such as non-stationary environments in which the

temporal constraints are not constant. Future studies will have to address whether flexible

learning rates also apply to the temporal domain.

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References

1. Pearce JM, Hall G. A model for pavlovian learning: Variations in the effectiveness of
conditioned but not of unconditioned stimuli. Psychological review. 1980;87(6):532.

2. Shadmehr R, Smith MA, Krakauer JW. Error correction, sensory prediction, and
adaptation in motor control. Annu Rev Neurosci. 2010;33:89–108.

3. Thoroughman KA, Shadmehr R. Learning of action through adaptive combination of
motor primitives. Nature. 2000;407(6805):742–7.

4. Donchin O, Francis JT, Shadmehr R. Quantifying generalization from trial-by-trial
behavior of adaptive systems that learn with basis functions: Theory and experiments in
human motor control. J Neurosci. 2003 Oct;23(27):9032–45.

5. Burge J, Ernst M, Banks MS. The statistical determinants of adaptation rate in human
reaching. J Vis. 2008;8(4):20.1–19.

6. de la Malla C, López-Moliner J, Brenner E. Dealing with delays does not transfer across
sensorimotor tasks. J Vis. 2014;14(12).

7. Herzfeld DJ, Vaswani PA, Marko MK, Shadmehr R. A memory of errors in sensorimotor
learning. Science. 2014 Sep;345(6202):1349–53.

8. van Beers RJ. Motor learning is optimally tuned to the properties of motor noise. Neuron.
2009;63(3):406–17.

9. van Beers RJ. How does our motor system determine its learning rate? PloS one.
2012;7(11):e49373.

10. Kording KP, Tenenbaum JB, Shadmehr R. The dynamics of memory as a consequence of
optimal adaptation to a changing body. Nature neuroscience. 2007;10(6):779.

11. Wei K, Körding K. Uncertainty of feedback and state estimation determines the speed of
motor adaptation. Frontiers in computational neuroscience. 2010;4.

12. Wolpert DM, Ghahramani Z, Jordan MI. An internal model for sensorimotor integration.
Science. 1995;269(5232):1880–2.

13. Desmurget, Grafton. Forward modeling allows feedback control for fast reaching
movements. Trends Cogn Sci. 2000 Nov;4(11):423–31.

14. Shadmehr R, Krakauer JW. A computational neuroanatomy for motor control. Exp Brain
Res. 2008 Mar;185(3):359–81.

15. Xu-Wilson M, Chen-Harris H, Zee DS, Shadmehr R. Cerebellar contributions to adaptive
control of saccades in humans. Journal of Neuroscience. 2009;29(41):12930–9.

16. Mazzoni P, Krakauer JW. An implicit plan overrides an explicit strategy during
visuomotor adaptation. J Neurosci. 2006 Apr;26(14):3642–5.

17. Taylor JA, Ivry RB. Flexible cognitive strategies during motor learning. PLoS
computational biology. 2011 Mar;7(3):e1001096.

18. Diedrichsen J, Hashambhoy Y, Rane T, Shadmehr R. Neural correlates of reach errors. J
Neurosci. 2005 Oct;25(43):9919–31.

19. Binda P, Cicchini GM, Burr DC, Morrone MC. Spatiotemporal distortions of visual
perception at the time of saccades. Journal of Neuroscience. 2009;29(42):13147–57.

20. Bremmer F, Kubischik M, Hoffmann K-P, Krekelberg B. Neural dynamics of saccadic
suppression. Journal of Neuroscience. 2009;29(40):12374–83.

21. Teasdale N, Bard C, Fleury M, Young DE, Proteau L. Determining movement onsets from
temporal series. Journal of motor behavior. 1993;25(2):97–106.

22. Brainard DH. The psychophysics toolbox. Spatial vision. 1997;10:433–6.

23. Pelli DG. The videotoolbox software for visual psychophysics: Transforming numbers
into movies. Spatial vision. 1997;10(4):437–42.

24. van Beers RJ, Brenner E, Smeets JBJ. Random walk of motor planning in task-irrelevant
dimensions. J Neurophysiol. 2013;109(4):969–77.

25. Bates D, Maechler M, Bolker B, Walker S. Fitting linear mixed-effects models using lme4.
Journal of Statistical Software. 2015;67:1:48.

26. Brenner E, Smeets JBJ. Sources of variability in interceptive movements. Exp Brain Res.
2009;195(1):117–33.

27. López-Moliner J, Supèr H, Keil M. The time course of estimating time-to-contact:
Switching between sources of information. Vision Res. 2013;92:53–8.

28. Mehra R. On the identification of variances and adaptive kalman filtering. IEEE
Transactions on automatic control. 1970;15(2):175–84.

29. Linares D, Holcombe AO, White AL. Where is the moving object now? Judgments of
instantaneous position show poor temporal precision (sd = 70 ms). J Vis. 2009;9(13):9.1–
14.

30. Ernst MO, Banks MS. Humans integrate visual and haptic information in a statistically
optimal fashion. Nature. 2002;415:429–33.

31. Oruç I, Maloney LT, Landy MS. Weighted linear cue combination with possibly
correlated error. Vision research. 2003;43(23):2451–68.

32. van Beers RJ, Meer Y van der, Veerman RM. What autocorrelation tells us about motor
variability: Insights from dart throwing. PLoS One. 2013;8(5):e64332.

33. Wei K, Körding K. Relevance of error: What drives motor adaptation? Journal of
neurophysiology. 2009;101(2):655–64.

34. Kording K. Decision theory: What "should" the nervous system do? Science. 2007
Oct;318(5850):606–10.

35. Franklin DW, Reichenbach A, Franklin S, Diedrichsen J. Temporal evolution of spatial
computations for visuomotor control. Journal of Neuroscience. 2016;36(8):2329–41.

36. Hudson TE, Maloney LT, Landy MS. Optimal compensation for temporal uncertainty in
movement planning. PLoS Comput Biol. 2008 Jul;4(7):e1000130.

37. Kalesnykas R, Hallett P. The differentiation of visually guided and anticipatory saccades
in gap and overlap paradigms. Experimental Brain Research. 1987;68(1):115–21.

38. Madelain L, Krauzlis RJ, Wallman J. Spatial deployment of attention influences both
saccadic and pursuit tracking. Vision research. 2005;45(20):2685–703.

39. Harwood MR, Madelain L, Krauzlis RJ, Wallman J. The spatial scale of attention strongly
modulates saccade latencies. Journal of neurophysiology. 2008;99(4):1743–57.

40. Montagnini A, Chelazzi L. The urgency to look: Prompt saccades to the benefit of
perception. Vision research. 2005;45(27):3391–401.

41. Anderson AJ, Carpenter RHS. Changes in expectation consequent on experience,
modeled by a simple, forgetful neural circuit. Journal of Vision. 2006;6:822–35.

42. Vullings C, Madelain L. Control of saccadic latency in a dynamic environment: Allocation
of saccades in time follows the matching law. Journal of neurophysiology. 2017;jn–00634.

43. Fleuriet J, Goffart L. Saccadic interception of a moving visual target after a
spatiotemporal perturbation. J Neurosci. 2012 Jan;32(2):452–61.

44. Herzfeld DJ, Shadmehr R. Motor variability is not noise, but grist for the learning mill.
Nat Neurosci. 2014 Feb;17(2):149–50.

Fig S1. (A) The simulated process variance is plotted as a function of the prediction made from

the linear model based on σ_t^2, $(\sigma_x/v)^2$ and the ccf(1). (B) The predicted process variance

obtained from the linear model based on the simulated data as a function of the whole

temporal variance in the different experimental conditions. (C) Decomposition of the whole

temporal variance into the two factors of eq. 1. One can see the part of the variability that

comes from spatial (σ_t^2) and temporal ($(\sigma_x/v)^2$) origin.

Fig S2. Density between the final temporal error (x-axis) and prediction error (y-axis) that is computed from the Kalman filter for each condition. The density plot includes all participants. The bar plot shows the slope of the fitted (grey) line for each condition. The slope for the Arm movement condition is shown without separating fast and slow movement time. However the slope was significantly smaller for slower movements (0.49 versus 0.72, p=0.01.)

Fig S3. Expected reliability of a weighted linear combination of two cues c1 and c2 with

simulated reliabilities r1=1 and r2=2. See (31) for details about the computation of the

combined reliability for correlated cues. The x-axis denotes the contribution of cue 1. Five

different correlations between c1 and c2 are shown. For the range of correlations observed

between the prediction error and the target error (Fig S2) the expected benefit from

integration is very little or null.