It's not the spoon that bends: Internal states of the observer determine serial dependence

Ayberk Ozkirli, David Pascucci

Laboratory of Psychophysics, Brain Mind Institute, École Polytechnique Fédérale de Lausanne (EPFL), Lausanne, Switzerland

Keywords: serial dependence, history biases, perceptual uncertainty, decision-making, optimality, Bayesian perception

Corresponding author: ayberk.ozkirli@gmail.com

Abstract

Traditional views suggest that human perception handles uncertainty using optimal strategies. For

instance, when prior stimuli are more reliable than current ones, perceptual decisions rely more on

the past, leading to stronger serial dependence. Here, we report findings that challenge this view. We

asked human observers to reproduce the average orientation of an ensemble of stimuli under varying

stimulus uncertainty. Contrary to optimal strategies, we found that serial dependence is stronger when

prior stimuli are more uncertain. We hypothesize that fluctuations in stimulus uncertainty may

influence internal states of observers, such as participants expectations about uncertainty and beliefs

about their own performance. A striking finding is that manipulating these internal states through

rigged feedback can yield drastic effects on serial dependence, even when external input (i.e., stimulus

uncertainty) remained constant. Our findings suggest that phenomena like serial dependence can be

better understood by considering internal states of the observer, beyond fixed computations and

optimal strategies.

Introduction

No one likes uncertainty. Yet, uncertainty is not only a persistent aspect of our environment, but also an extremely volatile one, evolving swiftly and sometimes with dramatic consequences. Imagine a pilot navigating through rapidly changing weather conditions, facing the risk of making the wrong decisions by relying on suddenly unclear visual cues.

According to traditional models of perception and decision-making, the brain deals with uncertainty in a near-optimal and Bayesian way[1–9]. For example, when a stimulus is unclear or ambiguous, the brain leverages more reliable sources of information, such as prior experiences, to guide its interpretation[9–12]. A central tenet of these models is that uncertainty is computed from each stimulus, and then multiple stimuli is integrated, with a greater weight to the more reliable ones[2,4,10,13,14]. Models of this kind, often described within the framework of the 'ideal observer'[15], can explain performance in a variety of perceptual tasks, suggesting that, in most cases, humans can effectively combine information from multiple stimuli to reduce uncertainty in an optimal way.

However, not all human behavior strictly adheres to Bayesian and optimality principles[16–19]. Recent research on serial dependence, the phenomenon whereby perceptual decisions are biased toward previous stimuli[20,21], provides a clear example. Ideal observer models predict that serial dependence is stronger when the previous stimulus is more reliable than the current one, as the optimal strategy is to rely more on more reliable stimuli[10]. Nonetheless, existing results are rather mixed. Some studies show that only the present uncertainty, not the past, influences serial dependence[22–24], while others report increased serial dependence when the uncertainty on the previous trial is high, rather than low[25,26]. In sum, human perceptual decisions do not seem to weigh prior stimuli in the same way as ideal observers would.

The reasons for such discrepancies can be manifold, as multiple factors contribute to serial dependence. These include the interplay between attractive and repulsive forms of serial dependence[27–30], modulations due to higher-level cognitive functions, such as confidence, attention, working-memory and expectations[20,27–29,29–34], among others. These factors may all affect the way uncertainty is handled in perceptual decision-making. For instance, a change in stimulus uncertainty might be internally perceived as a change in the state of the world[32] —e.g., an uncertain stimulus is likely to be followed by another uncertain one. Consequently, individuals may adjust how they integrate information over time, particularly when experiencing moments of high uncertainty. These changes in integration strategies may ultimately impact the strength of serial dependence, beyond the predictions offered by ideal observer models alone. The contribution of these factors, such as participants expectations about uncertainty and beliefs about their own performance —termed here

as 'internal states'— cannot be adequately captured by models predicting serial dependence by

relying exclusively on stimulus uncertainty.

In this study, we focused on the role of internal states in serial dependence and how they may,

at least partially, explain patterns in behavior that deviate systematically from ideal observer models.

In a first step, we tested how transitions between different levels of uncertainty affect serial

dependence in an orientation averaging task, where participants reproduced the average orientation

of an ensemble of stimuli (Figure 1A). In two experiments, we found that serial dependence increased

when the previous stimulus was uncertain (e.g., an ensemble with a wider distribution of orientations)

and was the largest when both the current and previous stimuli were uncertain—a pattern nearly

opposite from that predicted by the ideal observer. To explain this behavior, we hypothesized that the

persistence of high uncertainty across consecutive moments induces an internal state that promotes

serial dependence—a hypothesis formalized in an accompanying model. Building on this hypothesis,

we predicted that, even under constant stimulus uncertainty, changes in internal states could still

influence serial dependence. To test this prediction, we conducted a final experiment where we

manipulated performance feedback. Strikingly, serial dependence was observed only when

participants believed their performance was poor due to manipulated feedback, despite the level of

stimulus uncertainty, performance, and the feedback received on the most recent trial remaining

constant.

In summary, our findings suggest that phenomena like serial dependence are largely flexible,

contingent upon internal states triggered by the need to mitigate uncertainty, and cannot be reduced

to ideal observer models that rely solely on stimulus parameters.

Results

The effect of transitions in stimulus uncertainty on serial dependence

In serial dependence, perceptual decisions are biased towards the stimulus shown on the preceding

trial[21,35]. According to an ideal observer, the bias should be larger when the current stimulus is more

uncertain than the previous one, and reduced when the current stimulus is more reliable[10] (see Figure

S3A in Supplementary Material).

To test these predictions, we asked sixteen observers to reproduce the perceived average

orientation of an ensemble of 36 Gabor stimuli (Figure 1A). On randomly interleaved trials, the

ensemble orientations were either drawn from a normal distribution with a standard deviation (σ) of

8° (*high* σ) or all identical (*low* $\sigma = 0$°). These two conditions carried different degrees of uncertainty,

as reflected in performance, with a larger standard deviation of adjustment errors in the *high* σ (12.56

± 2.69°) compared to the *low* σ condition (6.77 ± 1.18°; *high* vs. *low*: t(15) = 12.31, *p* < .001, Cohen's
d' = 3.08, paired t-test).

Figure 1. Main paradigm and results of Experiment 1a-b. A) Participants reproduced the average orientation of an ensemble of 36 Gabors shown in the periphery of the visual field (see Methods). B) We manipulated uncertainty by intermixing ensembles with high and low orientation variability (e.g., standard deviation [σ] of the orientation distribution, σ = 8° vs. 0° in Experiment 1a, σ = 15° vs. 5° in Experiment 1b). C) The first panel shows serial dependence in the high-to-low (in orange) and low-to-low conditions (in grey). The second panel shows serial dependence in the high-to-high (in blue) and low-to-high conditions (in green). A circular running average (sliding window size of 35°) is computed on the folded adjustment errors of all participants as a function of the absolute difference in average orientation between the previous and current ensemble (Δ; |previous minus current|, see Methods). Shaded areas are 1 standard deviation of the running average. The curves represent the average first Derivative of Gaussian (δoG) fit across participants. The bar plots in the third panel show the average half-amplitude parameter (α) of δoG fits across participants, for each combination of prior and current uncertainty. Error bars represent standard errors of the mean, corrected for within-subject repeated-measures designs[36]. D) Results from Experiment 1b where the two σ used were 15° and 5°. The color coding is the same as in C.

We estimated the strength of serial dependence using the half-amplitude parameter from a δoG fit (α, see Methods), considering all possible combinations of current and past uncertainty, in a full-factorial design. A repeated-measures ANOVA with factors 'Uncertainty on the preceding trial'

(*low* vs. *high*) and 'Uncertainty on the current trial' (*low* vs. *high*) revealed no effect of the current

trial uncertainty (F(1, 15) = 2.22, *p* = .16, η_p^2 = 0.13), but a significant main effect of the uncertainty

on the preceding trial (F(1, 15) = 5.21, *p* = .037, η_p^2 = 0.26) as well as a significant interaction (F(1,

15) = 21.19, *p* < .001, η_p^2 = 0.59). The interaction was driven by the serial dependence occurring when

both the previous and current stimuli were of high uncertainty (the *high-to-high* condition), which

was significantly larger when compared to all the other conditions (*high-to-high* vs. average of the

other conditions: t(15) = 3.837, *p* = .0016, Cohen's *d'* = 0.959; see Figure 1C).

This finding was inconsistent and even nearly opposite to what would be expected by the ideal

observer with classic optimal integration (see Supplementary Material). In other words, observers did

not exhibit stronger biases when the prior stimulus was more reliable; instead, they exhibited the

strongest bias toward past stimuli when uncertainty was high in both the current and previous trials.

One aspect worth noting is that the interaction term was significant: when the current

uncertainty was low, we did not find any effect of the previous trial uncertainty (Figure 1C, first

panel). This may be attributed to ceiling performance and weak serial dependence in the low

uncertainty condition, where all stimuli shared the same orientation. Another possibility is that *low* σ

stimuli also carried stronger repulsive effects that contrasted attractive serial dependence[27–29,37,38],

since repulsive biases are often observed after well-visible stimuli that can be represented with high

precision[39–41].

To address these concerns, we performed a follow-up experiment with 16 new participants,

increasing the standard deviations (σ) in both the *high* (σ = 15°) and *low* (σ = 5°) uncertainty

conditions. As in the first experiment, observers were less precise in the *high* σ (standard deviation of

adjustment errors: 16.54 ± 4.90°) compared to the *low* σ condition (11.05 ± 2.78°; *high* vs. *low*: t(15)

= 7.37, p < .001, *d'* = 1.84, paired t-test).

Upon increasing σ in both conditions, a repeated-measures ANOVA revealed a significant

effect of the preceding trial (F(1, 15) = 11.698, *p* = 0.0038, η_p^2 = 0.44), but no significant effect of the

current trial uncertainty (F(1, 15) = 0.796, *p* = 0.386, η_p^2 = 0.05) or the interaction (F(1, 15) = 0.217,

p = .65, η_p^2 = 0.014). Therefore, when uncertainty was increased in both conditions compared to

Experiment 1a, serial dependence was once again overall stronger when the preceding stimulus was

more uncertain, regardless of the current trial uncertainty. Further analyses indicated that these effects

strengthen with an increasing number of consecutive high-uncertainty trials in the past (see Figure

S2), thereby excluding the possibility that the observed differences stemmed from other confounding

factors associated with the representation of average orientations on the immediately preceding trial

(e.g., heightened repulsive adaptation-like aftereffects following more precise orientation

representations).

The patterns observed in Experiment 1a-b, despite starkly contradicting the predictions of

ideal observer models of serial dependence (see Figure S3A), can be more accurately captured

through a straightforward yet effective conceptual extension of these models. Indeed, a model that

incorporates the impact of sequences of uncertain stimuli on the observer internal states can

effectively adjust the trial-by-trial tendency to integrate information over time, outperforming the fit

of a simple ideal observer in both Experiment 1a and 1b (see *State-dependent model of serial*

dependence in Supplementary Material).

The key concept here is that serial dependence is not solely determined by a fixed computation

of uncertainty based on the stimulus; rather, it depends on internal states of the observer, that evolve

as a function of the history of stimulus uncertainty. This led us to formulate a key prediction: even

when stimulus uncertainty remains constant, internal states of an observer can entirely dictate the

strength of serial dependence. To directly test this prediction, we conducted a final experiment, which

also served to definitively rule out any other explanation of the results based on differences between

the physical parameters of the stimuli across conditions, such as adaptation-like effects and

orientation bias.

Internal states modulate serial dependence

We tested 24 new participants in a paradigm where we fixed the ensemble σ to 15° and introduced

trial-by-trial feedback. The goal of the feedback was to manipulate internal states of the observer by

changing their beliefs about their own performance, and thus the experienced uncertainty in different

blocks of trials although the external uncertainty provided by the stimulus always remained constant

(see Methods). To this end, we used two types of blocks: *low feedback* blocks, where observers

received more frequent feedback indicating poor performance ("below average"), and *high feedback*

blocks, where they received more frequent feedback indicating good performance ("above average").

In both types of blocks, low and high feedback trials were intermixed with an equal number of trials

in which participants received neutral ("average") feedback. Importantly, within each block, the

feedback indicating poor or good performance still reflected the actual performance levels (see

Methods). We focused on 'block' effects on serial dependence —i.e., modulations due to the overall

frequency of *low* or *high feedback* in a block—and thus, to control for any effect of the specific

feedback received on the most recent trial, we restricted our analysis to trials following neutral

feedback, which was received with equal frequency in both block types.

Participants performed the task with comparable levels of performance between the two block

types (standard deviation of adjustment errors: *low*: 13.73 ± 3.59°, *high*: 14.20 ± 3.44°, *low* minus

high: t(23) = -1.65, *p* > .05, *d'* = -0.34). Strikingly, however, and in line with our predictions, serial

dependence was significantly larger in the *low feedback* compared to the *high feedback* blocks (*low*

minus *high*: t(23) = 2.44, *p* = .023, *d' =* 0.50). Thus, even though the stimulus uncertainty was fixed, and performance was comparable between conditions, manipulating internal states of the observer led to clear differences in serial dependence: under the internal belief of overall worse performance in a sequence of trials, the tendency to integrate current and prior stimuli increased.

Figure 2. Results of Experiment 2. Serial dependence in the two types of blocks (red: *low feedback*, blue: *high feedback*). *Low feedback* blocks contained a higher number of poor performance feedback trials. *High feedback* blocks contained more good performance feedback trials. In both blocks, the two types of feedback were intermixed with equal frequency of neutral feedback trials (see Methods). Serial dependence was assessed on trials following neutral feedback to mitigate any potential impact of feedback on the immediately preceding trial. The graph on the left depicts moving averages of folded errors and the best fitting δoG, similar to Figure 1. Bar plots on the right represent the average of the δoG half-amplitude parameter (α) across participants in the two conditions. Error bars represent standard errors of the mean. Note that performance (i.e., the standard deviation of adjustment errors) remained constant across conditions, despite evidently different levels of serial dependence (see main text).

Discussion

In this study, we investigated the impact of uncertainty on serial dependence in perceptual decisions. Existing frameworks, based on ideal observer models, predict stronger serial dependence when prior stimuli are more reliable than current ones[10]. Contrary to this prediction, we observed an increase in serial dependence when prior stimuli were more uncertain. We hypothesized that this phenomenon could be due to internal states of the observer that modulate serial dependence beyond the mere effects of stimulus uncertainty. We directly tested this hypothesis in a final experiment, where we manipulated internal states, namely participants' beliefs about their own performance in a block of trials, using feedback, and found drastic effects on serial dependence, even when the uncertainty in the stimulus remained constant.

Prior research has modeled serial dependence following the classic assumptions of optimal and Bayesian models[5,42,43]. These models posit that serial dependence is influenced by internal priors

about external world stability and by the relative reliability between these priors and current sensory

evidence[38,43,44]. However, empirical validation of these models has lacked a systematic assessment

across all possible conditions of relative uncertainty. Through a full factorial design, we demonstrate

that these models provide only a loose approximation, as they fail to account for a crucial aspect

observed in two experiments (Experiments 1a-b): the increase in bias when the uncertainty of prior

stimuli is high (see Figure 1).

We argue that our findings can be explained in the context of temporal expectations and

adaptable internal beliefs[25,32]. Even in the absence of regularities and temporal structures, the brain

constantly uses recent experiences to infer the current state of the environment[37,45–47], a core notion

of predictive theories of the brain[37,45,47,48]. These internal beliefs act as downstream, top-down

predictions that are fed back and shape processing as early as in primary visual cortex[49]. Crucially,

internal beliefs encompass not only specific stimulus features, but also the level of uncertainty

associated with encountering a stimulus[32]. Thus, the occurrence of an uncertain stimulus may prompt

a change in the internal states of the observer, leading to a greater tendency towards serial dependence

as a means to mitigate uncertainty.

This interpretation is consistent with the notion that, while classic models assume a fixed

structure of prior beliefs[38,43,44], actual beliefs are dynamic and contingent on internal states that adapt

and reset when changes in the environment are detected[25,50]. In contrast to static priors, indeed, we

show that a simple extension of an ideal observer model, which incorporates internal states, provides

a more accurate approximation of the data (see *State-dependent model of serial dependence* in

Supplementary Material). This idea is reminiscent of Kalman filters that adapt and self-tune to

changes in input data[42,51,52], which may provide a promising modeling framework for future studies.

One may question the extent to which the findings of Experiment 1a-b could be accounted for

by alternative explanations and more complex models. For instance, previous studies have

highlighted the existence of both attractive and repulsive forms of serial dependence[27,28,37,38,53]. The

interaction and relative prevalence of these opposing biases may be influenced by various factors,

including the stimulus uncertainty, with lower uncertainty favoring repulsive effects. Although we

cannot dismiss the potential role of a repulsive component, which might have countered attractive

serial dependence in conditions of low uncertainty, our supplementary analyses revealed patterns that

cannot be explained by this. Specifically, we observed that attractive serial dependence following an

uncertain stimulus increased in strength as a function of the number of consecutive uncertain stimuli

occurring before (see Supplementary Material for details). Moreover, Experiment 2, where stimulus

uncertainty remained constant, provided clear and unequivocal evidence that internal states,

independent of stimulus parameters, can significantly influence the strength and even the

manifestation of serial dependence. This finding aligns with existing research that explores factors

beyond the stimulus in understanding serial dependence, including confidence[54], attention[20], memory

load[55], time-on-task[56], task demand[28,29,31,37,57], stimulus predictability[58], as well as internal (e.g.,

neural) sources of uncertainty[43].

As noted earlier, other studies have also reported patterns that diverge from those predicted

by simple ideal observer models. For example, Ceylan et al.[22] found similar trends to our study, with

the strongest serial dependence occurring in high-to-high uncertainty conditions. However, they did

not observe significant differences between this and all other conditions, as we did. The reasons for

this inconsistency are unclear and may stem from various factors, including differences in stimuli

(single stimuli vs. ensembles) and the manipulation of uncertainty (e.g., changes in low-level features

such as spatial frequency or noise vs. variability in ensembles), as well as differences in sensitivity

(e.g., in[22], the four combinations were tested only in one out of three blocks of trials, with a reduced

number of trials per condition compared to the current study). Nevertheless, the presence of similar

trends across multiple studies[22–26] suggests that serial dependence cannot be solely explained by ideal

observer and Bayesian models in their standard form. Building on this notion, recent research has

used the term 'partial Bayesian inference' to describe the computations underlying serial

dependence[24]. The term 'partial' implies the involvement of other contributing factors, and as

demonstrated here, one such factor relates to the internal states of the observer, whose influence can

nearly reverse the patterns predicted by ideal models.

Systematic deviations from ideal observer models, like the one reported here, do not

necessarily imply that such models are incorrect, but highlight the need to move beyond simple and

classic models that rely solely on basic aspects of the stimulus[32,33]. In the present context, differences

in serial dependence can be found even without manipulating the stimulus parameters: what matters

is the internal state of the observer. Similarly, in other domains like spatial vision, classic models that

rely solely on local stimulus properties and feedforward processing may fail to explain phenomena

like crowding and ensemble perception[59–62].

In summary, we demonstrate that serial dependence is modulated by internal states of the

observer, providing, at least a partial explanation of why the observed patterns typically deviate from

the predictions of existing models. These results corroborate the view of serial dependence as a two-

stage process, where effects are shaped not only by the stimulus but also by higher-level cognitive

processes, in a recurrent hierarchical system where past experience plays a role at multiple interacting

stages[28].We argue that taking into account these aspects may not only lead to more accurate and

generalizable predictions of serial dependence, but also to a better understanding of its functional

roles, as flexible states of serial dependence may be more useful than a hard-wired and fixed

mechanism. Integrating prior and current stimuli can be advantageous, particularly when anticipating

moments of uncertainty in perception. However, integration may not be compulsory, especially when

we can rely on our ability to discriminate between changes[58].

Methods

Ethics statement

348   The study was approved by Commission cantonale d'éthique de la recherche sur l'être humain

349   (protocol number 2021-02270) under the Declaration of Helsinki[63], except for preregistration.

Apparatus

351   Stimuli were presented on a gamma-corrected VG248QE monitor (resolution: 1920 x 1080 pixels,

352   refresh rate: 120 Hz) and were generated with custom-made scripts written in MATLAB (R2013a)

353   and the Psychophysics Toolbox[64], on a Windows-based machine. Participants sat at approximately 57

354   cm from the computer screen, with the head held stable on a chin rest. The experiments were

355   performed in a dark room.

Participants

357   62 healthy participants (20 in Experiment 1a, 17 in Experiment 1b; age range of 18-37 years; 5

358   females in Experiment 1a, 9 females in Experiment 1b, and 15 females in Experiment 2), from the

359   EPFL and the University of Lausanne, took part in the study for a monetary reward (25 CHF/hour).

360   Participants were naïve to the purpose of the experiments. Written informed consent was collected

361   from all participants beforehand.

Stimuli and procedures

The stimuli and procedure are depicted in Figure 1. Each trial started with a dark red fixation spot

appearing for 300-1000 ms (randomly sampled in steps of 100 ms). Participants were then presented

with a display of 36 oriented Gabor stimuli for 500 ms (peak contrast: 25% Michelson, spatial

frequency: 2 cycles/°, Gaussian envelope: 0.225°, phase = 0.25). Gabor stimuli were presented on a

gray background (62.66 cd/m^2) and positioned inside an annulus centered at the fovea (internal radius

of 4.825°; external radius of 9.175°). No stimuli were presented in the central part of the display to

avoid systematic biases toward the center during mean orientation estimates[62,65]. Inside the annulus,

Gabor stimuli were arranged at equidistant locations (approximately 2.2° with a 1° random jittering)

inside a 12x12° grid. To prevent after-effects, a noise mask was presented within the same annulus

for 500 ms immediately after the Gabor ensemble. Following a blank interval of 1000 ms, participants

reproduced the perceived average orientation by adjusting a response tool. The response tool
consisted of two symmetric dots, each with a diameter of 0.25°, positioned approximately 1.5° apart,
marking the endpoints of an imaginary line. The orientation of the response tool was initialized
randomly, and participants were instructed to rotate the imaginary line using the mouse and confirm
their response with a left click.

On each trial, the orientation of the Gabor stimuli was drawn from a normal distribution with
the mean randomly selected over the full 0-179° orientation range, in steps of 1°. Experiment 1a
involved two randomly intermixed conditions in which the standard deviation (σ) of the ensemble's
orientations was either 0° (*low* σ) or 8° (*high* σ). In Experiment 1b, σ was increased to 5° (*low* σ) and
15° (*high* σ). In Experiment 2, it was fixed at 15°.

In Experiment 2, each trial was followed by written feedback, shown at the center after 1 s
from the response: "below average", "average", or "above average". To establish the criteria for
assigning feedback, we first determined an error threshold based on the mean absolute error (11°)
estimated from the group of participants in Experiment 1b, in the condition with σ = 15°. The
frequency was manipulated across different blocks. In the *low feedback* blocks, we assigned "below
average" feedback on 75% of the trials where the error exceeded the threshold, and "average"
feedback on the remaining 25% of those trials. Conversely, for trials in which the error was below
the threshold, we assigned "above average" feedback on only 25% and "average" feedback to 75% of
them. The opposite assignment pattern was applied in the *high feedback* blocks. In other words,
"below average" feedback was given on only 25% of trials where the error was above the threshold,
while "average" feedback was assigned to the remaining 75% of those trials. For trials with errors
below the threshold, "above average" feedback was assigned to 75% and "average" feedback to 25%
of them. As a result of this manipulation, participants were more likely to receive "below average"
(indicating poorer performance) in the *low feedback* blocks and "above average" (indicating better
performance) in the *high feedback* blocks, despite their actual performance potentially remaining
consistent.

An important aspect of this manipulation was that the frequency of trials with "average"
feedback was the same in both blocks. This allowed us to examine serial dependence based on the
overall block condition (*low* vs. *high feedback*) focusing exclusively on trials following neutral
feedback, thus controlling for the specific feedback received on the preceding trial.

To effectively manipulate internal states, it was also important that the feedback was perceived
as reliable and reflective of the actual performance. According to our debriefing with participants at
the end of Experiment 2, 23 out of 25 of them reported having had poorer performance in some

blocks, and 22 of those reported that this was due to difficulty difference across blocks (even though
the stimulus σ was constant).

All participants received verbal and written instructions at the beginning of each experiment
and completed a series of practice trials under the experimenter's guidance. The data from the practice
trials were not included in the analysis. In total, participants completed 400 trials in all three
experiments. The trials were divided into 10 blocks for Experiment 1a and 1b, and 4 for Experiment
2. The *high* and *low feedback* conditions alternated across the blocks, with the order of the conditions
counterbalanced among participants. Participants were instructed to keep their gaze fixed at the center
of the screen throughout each block. All experiments lasted about 1 hour.

Preprocessing of adjustment errors
Adjustment errors were computed as the acute angle between the response and the stimulus on each
trial. We first excluded trials where errors were larger than 1.5 interquartile ranges above the upper
quartile or below the lower quartile (separately for each experimental condition) and reaction times
larger than 10 seconds. The first trial of each block, as well as trials following a removed trial, were
also excluded. Overall, less than 10% of trials were removed in all experiments. In both Experiment
1a and 2, one participant was excluded due to a large overall negative bias (< -5°). In addition,
participants were excluded if their biases in a specific condition of interest were 1.5 interquartile
ranges larger than the upper quartile or smaller than the lower quartile within that condition (3 in
Experiment 1a and 1 in Experiment 1b). The final analyses included 16 participants in Experiment
1a, an independent group of 16 participants in Experiment 1b, and 24 participants in Experiment 2.

In addition to the data cleaning procedure mentioned above, we also evaluated the presence
of orientation biases, another typical step in preprocessing orientation adjustment errors for serial
dependence analysis[28]. Orientation biases—i.e., the non-uniform distribution of errors over the
orientation space— can introduce noise and artifacts into the estimation of serial dependence. Given
that orientation biases may vary depending on uncertainty[66], we examined their presence and the
necessity for their removal in each condition of Experiments 1a-b, where uncertainty varied (see
Supplementary Material). We found a clear orientation bias only in the condition with σ = 0°
(Experiment 1a, see Figure S1). This finding aligns with previous research indicating that while
orientation bias is typically apparent with a single stimulus (or all identical stimuli), its magnitude
can vary significantly and be substantially reduced in the context of more heterogeneous ensembles[66].
The bias was removed by taking the residuals of a *malowess* function in MATLAB. Note that even
without this additional step, the results remained consistent (see Supplementary Material).

Analysis of serial dependence

In Experiment 1a and 1b, we separated the conditions based on the preceding and current stimulus uncertainty, which resulted in four categories: *low-to-low*, *high-to-low*, *low-to-high*, and *high-to-high*. We fitted the first derivative of a Gaussian function (δoG) to the errors as a function of the difference in orientation between the previous and current average orientation of the ensemble (Δ). The δoG has the following form:

$$error = \Delta\alpha wce^{-(w\Delta)^2}$$

[1]

where c is a constant $c = \frac{\sqrt{2}}{e^{-0.5}}$ and w is the inverse of the curve width. The half-amplitude parameter α quantifies the deviation of the errors, in degrees, from the actual orientation, as a function of Δ: positive values of α indicate a systematic deviation of errors towards the orientation of the preceding stimulus, and negative values indicate a deviation away —i.e., repulsion. The parameter w was estimated using an aggregate participant for each condition[31,67]. Then, α was estimated for each participant and condition separately, using the respective aggregate w for that condition. For statistical purposes, repeated-measures ANOVA and t-tests were performed on the estimated α, depending on the experimental design and comparison of interest. For ANOVA, the effect sizes were reported in terms of partial-eta squares (0.01 to 0.059 indicating a small effect size, 0.06 to 0.139 indicating a medium effect size, and 0.14 or higher indicating a large effect size). For t-tests, we reported Cohen's d (0.2 to 0.49 indicating a small effect size, 0.50 to 0.79 indicating a medium effect size, and 0.8 or higher indicating a large effect size).

Acknowledgments

This research was supported by funding from the Swiss National Science Foundation (grant no. PZ00P1_179988 and PZ00P1_179988/2 to DP). The funders had no role in study design, data collection, and analysis. The authors would like to thank Bianca Ribeiro Riccioppo and Zehra Merchant for helping with the collection of the data.

Data availability

The data will be made available in online repositories upon publication.

Author contributions

The authors contributed equally to this work.

References

1. Diamond, M. E. Perceptual uncertainty. *PLOS Biol.* **17**, e3000430 (2019).

2. Ernst, M. O. & Banks, M. S. Humans integrate visual and haptic information in a statistically optimal fashion. *Nature* **415**, 429–433 (2002).

3. Gardner, J. L. Optimality and heuristics in perceptual neuroscience. *Nat. Neurosci.* **22**, 514–523 (2019).

4. Kersten, D., Mamassian, P. & Yuille, A. Object perception as Bayesian inference. *Annu Rev Psychol* **55**, 271–304 (2004).

5. Knill, D. C. & Pouget, A. The Bayesian brain: the role of uncertainty in neural coding and computation. *Trends Neurosci.* **27**, 712–719 (2004).

6. Körding, K. P. & Wolpert, D. M. Bayesian integration in sensorimotor learning. *Nature* **427**, 244–247 (2004).

7. Maloney, L. T. & Zhang, H. Decision-theoretic models of visual perception and action. *Vision Res.* **50**, 2362–2374 (2010).

8. Mamassian, P. Uncertainty and Confidence in Visual Perception. *-Percept.* **2**, 183–183 (2011).

9. Stocker, A. A. & Simoncelli, E. P. Noise characteristics and prior expectations in human visual speed perception. *Nat. Neurosci.* **9**, 578–585 (2006).

10. Cicchini, G. M., Mikellidou, K. & Burr, D. The functional role of serial dependence. *Proc. R. Soc. B* **285**, 20181722 (2018).

11. Feldman, J. Tuning your priors to the world. *Top. Cogn. Sci.* **5**, 13–34 (2013).

12. Stone, J. v., Kerrigan, I. s. & Porrill, J. Where is the light? Bayesian perceptual priors for lighting direction. *Proc. R. Soc. B Biol. Sci.* **276**, 1797–1804 (2009).

13. Alais, D. & Burr, D. The Ventriloquist Effect Results from Near-Optimal Bimodal Integration. *Curr. Biol.* **14**, 257–262 (2004).

14. Cicchini, G. M., Arrighi, R., Cecchetti, L., Giusti, M. & Burr, D. C. Optimal encoding of interval timing in expert percussionists. *J. Neurosci. Off. J. Soc. Neurosci.* **32**, 1056–1060 (2012).

15. Geisler, W. S. Contributions of ideal observer theory to vision research. *Vision Res.* **51**, 771–781
(2011).

16. Adler, W. T. & Ma, W. J. Comparing Bayesian and non-Bayesian accounts of human confidence
reports. *PLOS Comput. Biol.* **14**, e1006572 (2018).

17. Morales, J. *et al.* Low attention impairs optimal incorporation of prior knowledge in perceptual
decisions. *Atten. Percept. Psychophys.* **77**, 2021–2036 (2015).

18. Rahnev, D. & Denison, R. N. Suboptimality in Perceptual Decision Making. *Behav. Brain Sci.*
1–107 (2018).

19. Stengård, E. & Berg, R. van den. Imperfect Bayesian inference in visual perception. *PLOS*
Comput. Biol. **15**, e1006465 (2019).

20. Fischer, J. & Whitney, D. Serial dependence in visual perception. *Nat. Neurosci.* **17**, 738–743
(2014).

21. Pascucci, D. *et al.* Serial dependence in visual perception: A review. *J. Vis.* **23**, 9 (2023).

22. Ceylan, G., Herzog, M. H. & Pascucci, D. Serial dependence does not originate from low-level
visual processing. *Cognition* **212**, 104709 (2021).

23. Gallagher, G. K. & Benton, C. P. Stimulus uncertainty predicts serial dependence in orientation
judgements. *J. Vis.* **22**, 6–6 (2022).

24. You, F.-H., Gong, X.-M. & Sun, Q. Serial dependencies between form orientation and motion
direction are asymmetric. *Front. Psychol.* **14**, (2023).

25. Akaishi, R., Umeda, K., Nagase, A. & Sakai, K. Autonomous mechanism of internal choice
estimate underlies decision inertia. *Neuron* **81**, 195–206 (2014).

26. Fritsche, M. To Smooth or not to Smooth: Investigating the Role of Serial Dependence in
Stabilising Visual Perception [dissertation]. *Radboud Univ.* (2016).

27. Fritsche, M., Mostert, P. & de Lange, F. P. Opposite effects of recent history on perception and
decision. *Curr. Biol.* **27**, 590–595 (2017).

28. Pascucci, D. *et al.* Laws of concatenated perception: Vision goes for novelty, decisions for perseverance. *PLOS Biol.* **17**, e3000144 (2019).

29. Ceylan, G. & Pascucci, D. Attractive and repulsive serial dependence: The role of task relevance, the passage of time, and the number of stimuli. *J. Vis.* **23**, 8 (2023).

30. Bae, G.-Y. & Luck, S. J. Serial dependence in vision: Merely encoding the previous-trial target is not enough. *Psychon. Bull. Rev.* **27**, 293–300 (2020).

31. Fischer, C. *et al.* Context information supports serial dependence of multiple visual objects across memory episodes. *Nat. Commun.* **11**, 1–11 (2020).

32. Beierholm, U., Rohe, T., Ferrari, A., Stegle, O. & Noppeney, U. Using the past to estimate sensory uncertainty. *eLife* **9**, e54172 (2020).

33. Denison, R. N., Adler, W. T., Carrasco, M. & Ma, W. J. Humans incorporate attention-dependent uncertainty into perceptual decisions and confidence. *Proc. Natl. Acad. Sci.* **115**, 11090–11095 (2018).

34. Ceylan, G. & Pascucci, D. Expectations and cognitive control modulate history biases in perceptual decisions. *J. Vis.* **23**, 5392 (2023).

35. Manassi, M., Murai, Y. & Whitney, D. Serial dependence in visual perception: a meta-analysis and review. *Journal of Vision* (in press).

36. Cousineau, D. Confidence intervals in within-subject designs: A simpler solution to Loftus and Masson's method. *Tutor. Quant. Methods Psychol.* **1**, 42–45 (2005).

37. Pascucci, D. & Plomp, G. Serial dependence and representational momentum in single-trial perceptual decisions. *Sci. Rep.* **11**, 9910 (2021).

38. Fritsche, M., Spaak, E. & de Lange, F. P. A Bayesian and efficient observer model explains concurrent attractive and repulsive history biases in visual perception. *eLife* **9**, e55389 (2020).

39. Bae, G.-Y. & Luck, S. J. Interactions between visual working memory representations. *Atten. Percept. Psychophys.* **79**, 2376–2395 (2017).

40. Bliss, D. P., Sun, J. J. & D'Esposito, M. Serial dependence is absent at the time of perception but

increases in visual working memory. *Sci. Rep.* **7**, 14739 (2017).

41. Manassi, M., Liberman, A., Kosovicheva, A., Zhang, K. & Whitney, D. Serial dependence in

position occurs at the time of perception. *Psychon. Bull. Rev.* 1–9 (2018).

42. Burr, D. & Cicchini, G. M. Vision: Efficient Adaptive Coding. *Curr. Biol.* **24**, R1096–R1098

(2014).

43. van Bergen, R. S. & Jehee, J. F. Probabilistic representation in human visual cortex reflects

uncertainty in serial decisions. *J. Neurosci.* **39**, 8164–8176 (2019).

44. Kalm, K. & Norris, D. Visual recency bias is explained by a mixture model of internal

representations. *J. Vis.* **18**, 1 (2018).

45. Kok, P. & de Lange, F. P. Predictive coding in sensory cortex. in *An introduction to model-based*

cognitive neuroscience 221–244 (Springer, 2015).

46. Rosen, R. Anticipatory Systems. in *Anticipatory Systems: Philosophical, Mathematical, and*

Methodological Foundations (ed. Rosen, R.) 313–370 (Springer New York, New York, NY,

2012). doi:10.1007/978-1-4614-1269-4_6.

47. Trapp, S., Pascucci, D. & Chelazzi, L. Predictive brain: Addressing the level of representation by

reviewing perceptual hysteresis. *Cortex* (2021).

48. Friston, K. The free-energy principle: a unified brain theory? *Nat. Rev. Neurosci.* **11**, 127–138

(2010).

49. Yon, D. *et al.* Stubborn predictions in primary visual cortex. *J. Cogn. Neurosci.* **35**, 1133–1143

(2023).

50. Nassar, M. R., Wilson, R. C., Heasly, B. & Gold, J. I. An Approximately Bayesian Delta-Rule

Model Explains the Dynamics of Belief Updating in a Changing Environment. *J. Neurosci.* **30**,

12366–12378 (2010).

51. Pascucci, D., Rubega, M. & Plomp, G. Modeling time-varying brain networks with a self-tuning

optimized Kalman filter. *PLOS Comput. Biol.* **16**, e1007566 (2020).

52. Gekas, N., McDermott, K. C. & Mamassian, P. Disambiguating serial effects of multiple
timescales. *J. Vis.* **19**, 24–24 (2019).

53. Sheehan, T. C. & Serences, J. T. Attractive serial dependence overcomes repulsive neuronal
adaptation. *PLoS Biol.* **20**, e3001711 (2022).

54. Samaha, J., Switzky, M. & Postle, B. R. Confidence boosts serial dependence in orientation
estimation. *J. Vis.* **19**, 25–25 (2019).

55. Markov, Y., Tiurina, N. & Pascucci, D. A matter of load: prior history effects scale with visual
working memory load. Preprint at https://doi.org/10.31234/osf.io/6h2cy (2023).

56. Barbosa, J. & Compte, A. Build-up of serial dependence in color working memory. *Sci. Rep.* **10**,
10959 (2020).

57. Houborg, C., Kristjánsson, Á., Tanrıkulu, Ö. D. & Pascucci, D. The role of secondary features in
serial dependence. *J. Vis.* **23**, 21 (2023).

58. Blondé, P., Kristjánsson, Á. & Pascucci, D. Tuning perception and decisions to temporal context.
iScience 108008 (2023) doi:10.1016/j.isci.2023.108008.

59. Doerig, A. *et al.* Beyond Bouma's window: How to explain global aspects of crowding? *PLoS*
Comput. Biol. **15**, e1006580 (2019).

60. Doerig, A., Schmittwilken, L., Sayim, B., Manassi, M. & Herzog, M. H. Capsule networks as
recurrent models of grouping and segmentation. *PLoS Comput. Biol.* **16**, e1008017 (2020).

61. Tiurina, N., Markov, Y., Choung, O.-H., Herzog, M. H. & Pascucci, D. Unlocking crowding by
ensemble statistics. *Curr. Biol.* **32**, 4975-4981.e3 (2022).

62. Tiurina, N., Markov, Y., Whitney, D. & Pascucci, D. The functional role of spatial anisotropies
in ensemble perception. Preprint at https://doi.org/10.31234/osf.io/yd3n8 (2023).

63. World Medical Association. World Medical Association Declaration of Helsinki: Ethical
Principles for Medical Research Involving Human Subjects. *JAMA* **310**, 2191–2194 (2013).

64. Brainard, D. H. The psychophysics toolbox. *Spat. Vis.* **10**, 433–436 (1997).

65. Pascucci, D., Ruethemann, N. & Plomp, G. The anisotropic field of ensemble coding. *Sci. Rep.*

11, 1–10 (2021).

66. Tomassini, A., Morgan, M. J. & Solomon, J. A. Orientation uncertainty reduces perceived

obliquity. *Vision Res.* **50**, 541–547 (2010).

67. Cicchini, G. M., Benedetto, A. & Burr, D. C. Perceptual history propagates down to early levels

of sensory analysis. *Curr. Biol.* (2020).

601