Deliberation gated by opportunity cost adapts to context with urgency

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Abstract

Finding the right amount of deliberation, between insufficient and excessive, is a hard decision making problem that depends on the value we place on our time. Average-reward, putatively encoded by tonic dopamine, serves in existing reinforcement learning theory as the stationary opportunity cost of time, and of deliberation in particular. However, this cost often varies with environmental context that can change over time. Here, we introduce an opportunity cost of deliberation estimated adaptively on multiple timescales to account for non-stationary contextual factors. We use it in a simple decision-making heuristic based on average-reward reinforcement learning (AR-RL) that we call *Performance-Gated Deliberation* (PGD). We propose PGD as a strategy used by animals wherein deliberation cost is implemented directly as urgency, a previously characterized neural signal effectively controlling the speed of the decision-making process. We show PGD outperforms AR-RL solutions in explaining behaviour and urgency of non-human primates in a context-varying random walk prediction task and is consistent with relative performance and urgency in a context-varying random dot motion task. We make readily testable predictions for both neural activity and behaviour and call for an integrated research program in cognitive and systems neuroscience around the value of time.

9 Keywords: primate decision-making, reinforcement learning, urgency, opportunity cost

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symbol	quantity	
t	within-trial time	
k	trial index	
S_t	within-trial state at time t	
$oldsymbol{S}_t$	state sequence up to time t	
R_k	reward of k th trial	
T_k	duration of k th trial	
$t_k^{ m dec}$	decision time of k th trial	
$\mathcal{C}_t^{\mathrm{del}}$	within-trial opportunity cost of deliberation	
$r_{\rm max}$	maximum reward acheiveable in a trial	
b_t	belief of correct report given \boldsymbol{S}_t	
\bar{r}_t	expected reward for reporting at time t	
$\mathcal{C}_t^{\mathrm{com}}$	within-trial opportunity cost of commitment	
ho	stationary reward rate	
$ ho^*$	optimal stationary reward rate	
α	context parameter	
$ ho_{lpha}$	context-conditioned stationary reward rate	
T_{α}	context-conditioned stationary average trial duration	
$\hat{ ho}_k^{ au}$	reward history filtered through a timescale, τ	
$ au_{ m long}$	a long timescale over which to estimate ρ	
$\tau_{\rm context}$	a context-specific timescale over which to estimate ρ_{α}	
ν	tracking cost sensitivity	
K	subjective reward scale factor	
$T_{\rm block}$	characteristic duration of a trial block	
c	auxiliary deliberation cost rate	
N_t	tokens difference	
p	jump probability of random walk, $p \ge 1/2$	

Table I. Symbol glossary. Highlighted in gray are parameters of the PGD model presented in this paper.

10

INTRODUCTION

Humans and other animals make a wide range of decisions throughout their daily lives. Any particular action usually arises out of a hierarchy of decisions involving a careful balance between resources, including one that is always limited: time. The cost of *spending* time depends on its value, a construct that relies on comparing against the alternative things an agent could potentially do with it. Estimating time's value is not straightforward for a number of reasons. There are alternative choices at multiple decision levels, e.g. moving on from a job and moving on from a career, and each level requires its own evaluation. Moreover, the value of alternatives needs to be tracked as they may change over time depending on the context in which a decision is made. For example, animals will learn to value a given food resource differently depending on whether it is encountered during times of plenty versus scarcity. The agent's knowledge of and ability to track context thus influences the value it assigns to possible alternatives.

²³ These are significant, practical complications of making decisions contingent on *opportu*-

3

²⁴ nity costs [1], the formal economic concept capturing the value of the alternatives lost by ²⁵ committing a limited resource to a given use. The opportunity cost of time is nevertheless ²⁶ well-studied in decision-making theory. It plays the role of a reference reward in defini-²⁷ tions of relative value, most notably as the average reward in average-reward reinforcement ²⁸ learning (AR-RL) [2].

In neuroscience, AR-RL was first proposed to extend the reward prediction error hypothesis for phasic dopamine to account also for the observed properties of tonic dopamine 1 levels [3]. It has since been used to emphasize the relative nature of reward-based decisionmaking [4] in explanations of human and animal behaviour in foraging [5], free-operant conditioning [6], perceptual decision-making [7, 8], cognitive effort/control [8, 9], and even economic exchange [10].

Unlike the alternative discount-reward approach, AR-RL is a theoretically well-defined 35 ₃₆ and numerically stable formulation for long horizon decision problems [11], such as those 37 in *continuing environments* in which there is no definite end [12]. Solutions to AR-RL ³⁸ problems maximize average reward, in contrast to traditional fixed accuracy criteria in ³⁹ perceptual decision-making tasks that focus on maximizing trial reward alone [13]. The 40 solutions to AR-RL formulations of tasks of long sequence of trials are decision boundaries ⁴¹ in the state space of a trial. Determining this decision boundary requires maximizing the ⁴² relative value, defined using the opportunity cost of time. The resulting optimal decision 43 boundaries typically 'collapse' over a trial: they cut deliberation short, e.g. in tasks where ⁴⁴ trial difficulty is variable [7, 14]. Up to now, however, AR-RL and most of its applications 45 have focused on fixed context and have used the stationary average reward as the fixed ⁴⁶ opportunity cost of time, which ignores context-dependent performance variation. This is 47 perhaps not surprising given that in psychological and neuroscientific studies of decision-48 making, we usually eliminate such contextual factors from the experimental design such ⁴⁹ that our models describe stationary behaviour. However, the brain mechanisms under study 50 are adapted to a more diverse natural world in which changing environmental factors are ⁵¹ often relevant, hard to infer and vary over time [4].

We pursue a theory of approximate relative-value decision-making under uncertainty in a 52 ⁵³ setting relevant to decision-making neuroscience. We start by showing that value in AR-RL ⁵⁴ can be expressed using the opportunity costs of deliberation and commitment. Here, the ⁵⁵ commitment cost is the shortfall in reward (relative to the maximum possible in a trial) ⁵⁶ that is expected to be lost when committing to a decision at a given time. Highlighting the 57 risk of value representations in non-stationary environments, we propose an approximation ⁵⁸ to the AR-RL value-optimal solution, Performance-Gated Deliberation (PGD), that uses ⁵⁹ the increasing opportunity cost of time in a trial to collapse the decision boundary directly, ⁶⁰ by-passing the need to maximize relative value. PGD thus reduces decision-making to 61 estimating two opportunity costs: a commitment cost learned from the statistics of the 62 environment and a deliberation cost estimated from tracking one's own performance in that ⁶³ environment. It explains how an agent, without explicitly tracking context parameters or 64 storing a value function, can trade-off speed and accuracy according to performance at ⁶⁵ the typically longer timescales over which context changes. We propose that deliberation 66 cost is then directly encoded as "urgency" in the neural dynamics underlying decision-⁶⁷ making [7, 15–17]. The theory is thus directly testable using both behaviour and neural 68 recordings.

To illustrate how PGD applies in a specific continuing decision-making task, and to make the links to a neural implementation explicit, we analyze behavior and neural recordings

4

⁷¹ collected over eight years from two non-human primates (NHPs) [18, 19]. They performed ⁷² successive trials of the "tokens task", a probabilistic guessing task in which information ⁷³ about the correct choice is continuously changing within each trial, and a task parame-⁷⁴ ter controlling the incentive to decide early (the context) is varied over longer timescales. ⁷⁵ Behavior in the task, in both humans [16] and monkeys [19], provides additional support 76 to an existing hypothesis about how neural dynamics implements time-sensitive decision- τ making [15]. Specifically, neural recordings in monkeys suggest that the evidence needed ⁷⁸ to make the decision predominates in dorsolateral prefrontal cortex [20]; a growing context-⁷⁹ dependent urgency signal is provided by the basal ganglia [21]; and the two are combined to ⁸⁰ bias and time, respectively, a competition between potential actions that unfolds in dorsal ⁸¹ premotor and primary motor cortex [18]. Similar findings have been reported in other tasks -⁸² for example, in the frontal eve fields during decisions about eve-movements [17]. We propose ⁸³ PGD as a theoretical explanation for why decision-making mechanisms are organized in this ⁸⁴ way. As an algorithm, it serves as a robust means to balance immediate rewards and the cost ⁸⁵ of time across multiple timescales. As a quantitative model, it serves to explain concurrently ⁸⁶ recorded behaviour and neural urgency in continuing decision-making tasks. From neural ⁸⁷ recordings in non-human primates and and behaviour in human and non-human primates, ⁸⁸ we show that it does so more accurately than AR-RL solutions. Adapting PGD to the ³⁹ random dot motion task in which urgency was first characterized [17], we make quantitative ⁹⁰ predictions about neural urgency is such tasks, which we validate on their data within error 91 bounds.

RESULTS

A. Theory of performance-gated deliberation

Opportunity costs of deliberation and commitment, and drawbacks of average-reward reinforcement learning

We consider a class of tasks consisting of a long sequence of trials indexed by k =97 1, 2, ... (see fig. 1a), each of which provides the opportunity to obtain some reward by choos-99 ing correctly. In each trial, a finite sequence of states, S_t , $t = 0, ..., t_{\text{max}}$, is observed that 100 provide evidence for an evolving belief about the correct choice among a fixed set of options. 101 To keep notation simple, we suppress denoting the trial index, k, on quantities such as trial 102 state, S_t , that also depend on trial time, t. The time of decision, t_k^{dec} , and the chosen option 103 determine both the reward received, R_k , and the trial duration, $T_k \geq t_k^{\text{dec}}$. Importantly, 104 decision timing can affect performance because earlier decisions typically lead to shorter 105 trials (and thus more trials in a given time window), while later decisions lead to higher 106 accuracy. Effectively balancing such speed-accuracy trade-offs is central to performing well 107 in continuing episodic task settings. For a fixed strategy, the *stationary reward rate* (see 108 slope of dashed line in fig. 1a(right)) is

$$\rho := \lim_{k \to \infty} \sum_{k} R_k / \sum_{k} T_k .$$
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¹⁰⁹ For a stochastic environment, the definition of ρ includes an ensemble average. Free-operant ¹¹⁰ conditioning, foraging, and several perceptual decision-making tasks often fall into this class.

92

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5



Figure 1. AR-RL and Performance-Gated Deliberation. (a) Task setting. Left: Within trial state, S_t evolves over trial time t in successive trials indexed by k. The decision 'A' is reported at the decision time t_k^{dec} (red cross), determining trial reward, R_k , and trial duration, T_k . Right: Sketch of cumulative reward versus cumulative duration. Context-conditioned reward rate (slope of red line), varies with alternating context (labelled 1 and 2) around average reward, ρ (dashed line). (b) Decision rules based on opportunity costs of commitment, C_t^{com} , and deliberation, C_t^{del} . The AR-RL rule (black 'x') finds t that minimizes $C_t^{\text{del}} + C_t^{\text{com}}$. The PGD rule (black cross) finds t^{dec} at which they intersect, $C_t^{\text{del}} = C_t^{\text{com}}$. (c) Schematic diagram of each algorithm's dependency. PGD computes a decision time directly from the two opportunity costs, while AR-RL uses both to first estimate a value function, whose maximum specifies the decision time. (d) Loss (error in performance with respect to the optimal policy, $(\rho^* - \rho)/\rho^*$) over learning time in a patch-leaving task (AR-RL: brown, PGD: black). The arrow indicates when the state labels were randomly permuted.

¹¹¹ Previous work [7, 22] has studied the belief of correct report for binary rewards, $b_t = P(R_k = 12 \ 1 | \mathbf{S}_t, t^{\text{dec}} = t)$, which also gives the expected trial reward, $\bar{r}_t = b_t \cdot 1 + (1 - b_t) \cdot 0 = b_t$ [7] ¹¹³ (see [23] for more about the relationship between value-based and perceptual decisions). \mathbf{S}_t ¹¹⁴ denotes the state sequence observed so far, (S_0, \ldots, S_t) . We consider greedy strategies that ¹¹⁵ report the choice with the largest belief at decision time. The decision problem is then about ¹¹⁶ when to decide.

Average-reward reinforcement learning (AR-RL), first proposed in artificial intelli-

6

¹¹⁸ gence [24], was later incorporated into reward prediction error theories of dopamine sig-¹¹⁹ nalling [3] and employed to account for the opportunity cost of time [6]. AR-RL was ¹²⁰ subsequently used to study reward-based decision-making in neuroscience and psychol-¹²¹ ogy [7, 8, 25, 26]. AR-RL centers around the average-adjusted future return, which penalizes ¹²² the passage of time according the average reward. A reporting decision is associated with a ¹²³ return that for trial-based tasks combines the remainder of the current trial and all future ¹²⁴ trials, $\bar{r}_t - \rho(T_k - t) + \sum_{k'>k} (R_{k'} - \rho T_{k'})$, where ρ (*c.f.* eq. (1)) is either estimated online ¹²⁵ or obtained self-consistently (see Methods for details). Value is defined as the future return ¹²⁶ averaged over trial sequence realizations. This average of a sum of reward deviations into ¹²⁷ the future converges on account of the decaying effects of the state at which the decision is ¹²⁸ made. The AR-RL algorithms we consider aim to achieve the highest ρ by also maximizing ¹²⁹ the average-adjusted value. We now provide an alternative, but equivalent definition of ¹³⁰ average-adjusted trial return in terms of opportunity costs incurred by the agent.

We denote the opportunity cost of committing at time t within a trial as C_t^{com} , defined as the difference

$$\mathcal{C}_t^{\rm com} = r_{\rm max} - \bar{r}_t \;, \tag{2}$$

¹³³ where r_{max} is the maximum trial reward possible *a priori*. Within a trial, an agent lowers ¹³⁴ its commitment cost towards zero by accumulating more evidence, i.e. by waiting. Waiting, ¹³⁵ however, incurs another opportunity cost: the reward lost by not acting. We denote this ¹³⁶ opportunity cost of deliberation incurred up to a time t in a trial as C_t^{del} . In AR-RL, the ¹³⁷ constant opportunity cost rate of time is integrated so that for $T_k = t_k^{\text{dec}}$,

$$\mathcal{C}_t^{\text{del}} = \rho t \ . \tag{3}$$

¹³⁸ With these definitions, the average-adjusted trial return for deciding at a time t can be ¹³⁹ expressed as $r_{\text{max}} - (C_t^{\text{com}} + C_t^{\text{del}})$. It is maximized by jointly minimizing C_t^{del} and C_t^{com} (fig. 1b), ¹⁴⁰ giving the AR-RL optimal solution (see Methods for a formal statement and solution of the ¹⁴¹ AR-RL problem). Expressed in this way, the average-adjusted trial return emphasizes the ¹⁴² more general perspective that an agent's solution to the speed-accuracy trade-off is about ¹⁴³ how it balances the decaying opportunity cost of commitment and the growing opportunity ¹⁴⁴ cost of deliberation.

Despite their utility, value representations such as the average-adjusted trial return can 145 ¹⁴⁶ be a liability in real world tasks where task statistics are non-stationary. To illustrate this, we consider the following foraging task. An foraging agent feeds among a fixed set of food (e.g. berry) patches. Total berries consumed in a patch saturates with duration t according to a given saturation profile, shared across patches, as the fewer berries left are harder to find. Patches differ in their richness (e.g. berry density), which is randomly sampled and $_{151}$ fixed over the task. Denoting patch identity (serving as context) by s, the food return is $_{152}$ directly observed and deterministic given s. To perform well, the agent needs to decide when to move on from depleting the current patch. Further details about the task and its solution are given in the Methods. For a broad class of online AR-RL algorithms, the agent learns the ¹⁵⁵ average-adjusted trial return as a function of state and time. For a given patch, it then leaves when this return is at its maximum (c.f. fig. 1b). In fig. 1d, we show how the performance (brown line) approaches that of the optimal policy in time as the estimation of the AR-RL ¹⁵⁸ trial return improves with experience (see Methods for implementation details). However, if ¹⁵⁹ the agent's environment undergoes a significant disturbance (e.g. a forest fire due to which ¹⁶⁰ the patch locations are effectively re-sampled), the performance of this AR-RL algorithm can ¹⁶¹ drop back to where it started. We implement such a disturbance via random permutation

7

¹⁶² of the state labels at the time indicated by the arrow in fig. 1d. This is true over a range of ¹⁶³ learning rates and the number of patches (fig. S8). More generally, any approach that relies ¹⁶⁴ on estimating state-value associations shares this drawback, including those approaches that ¹⁶⁵ implicitly learn those associations by directly learning a policy instead [27]. Could context-¹⁶⁶ dependent decision times be obtained without having to associate value or action to state? ¹⁶⁷ A means to do so is presented in the next section.

168

2. Performance-Gated Deliberation

We propose that instead of maximizing value as in AR-RL, which minimizes the sum of the two opportunity costs, $C_t^{\text{del}} + C_t^{\text{com}}$, the agent simply takes as its decision criterion when they intersect (shown as the black cross in fig. 1b).

$$t^{\text{dec}} := \min_{t} \left\{ t \mid \mathcal{C}_{t}^{\text{del}} \ge \mathcal{C}_{t}^{\text{com}} \right\} \quad (\text{PGD decision rule}) \tag{4}$$

¹⁷² We call this heuristic rule at the center of our results *Performance-Gated Deliberation* ¹⁷³ (PGD). Plotted alongside the AR-RL performance in fig. 1d for our example foraging task, ¹⁷⁴ PGD (black line) achieves better performance than AR-RL overall. It is also insensitive to ¹⁷⁵ the applied disturbance since PGD uses C_t^{del} and C_t^{com} directly when deciding, rather than ¹⁷⁶ as input to problem of optimizing average-adjusted value as in AR-RL (fig. 1c).

¹⁷⁷ We constructed the above task so that PGD is the AR-RL optimal solution. In general, ¹⁷⁸ however, PGD is a well-motivated approximation to the optimal strategy, so we call it a ¹⁷⁹ heuristic. In the more general stochastic setting where there is residual uncertainty in trial ¹⁸⁰ reward at decision time, the PGD agent will have to learn the association between state ¹⁸¹ and expected reward, \bar{r}_t . This association is learned from within-trial correlations only. In ¹⁸² contrast, the opportunity cost of time as the basis for the deliberation cost depends on ¹⁸³ across-trial correlations that together determine the overall performance. It is thus more ¹⁸⁴ susceptible to non-stationarity. A typical task setting is when the value of the same low-level ¹⁸⁵ action plan differs across context. From hereon, we will assume the agent has learned the ¹⁸⁶ stationary opportunity cost of commitment and so focus on resolving the remaining problem: ¹⁸⁷ how to learn and use an opportunity cost of deliberation that exhibits non-stationarity on ¹⁸⁸ the longer timescales over which context varies.

189

3. Reward filtering for a dynamic opportunity cost of deliberation

The state disturbance in the toy example above altered task statistics at only a single ¹⁹⁰ time point. In general, however, changes in task statistics over time can occur throughout ¹⁹² the task experience. A broader notion of deliberation cost beyond the static average reward ¹⁹³ is thus needed-one that can account for extended timescales over which performance varies. ¹⁹⁴ Such a cost serves as a dynamic reference in a relative definition of value based on a non-¹⁹⁵ stationary opportunity cost of time. We first address how performance on various timescales ¹⁹⁶ can be estimated.

As a concrete example, we make use of the task that we will present in detail in the following section. This task has a context parameter, α , that can vary in time on characteristic timescales longer than the moment-to-moment and can serve as a source of non-stationarity in performance. Here, the context sequence, α_k , varies on a single timescale, e.g. through petriodic switching between two values. The resulting performance (fig. 2a(top)) varies around

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Figure 2. Non-stationary opportunity cost. (a) Top: Dynamics of trial performance ($\rho_k^{\text{trial}} := R_k/T_k$; blue) with its distribution as well as dynamics of between context-conditioned averages of performance ($\rho_{\alpha} = \langle \rho_k^{\text{trial}} \rangle_{k|\alpha}$; orange), and the effectively stationary average performance ($\rho \sim \langle \rho_k^{\text{trial}} \rangle_k$; purple). Bottom: these are decomposed into a hierarchy by filtering reward history on trial, context, and long timescales, respectively. (b) Two hypothetical forms for context-specific trial opportunity cost. Top: Trial-unaware cost in which context varies the slope around ρ . Bottom: Trial-aware cost in which context varies the slope around ρ .

 $_{202}$ the stationary average, ρ (purple), with context variation due to the switching (orange), as ²⁰³ well as context-conditioned trial-to-trial variation (blue). The decomposition of time-varying ²⁰⁴ performance into these multiple, timescale-specific components can be achieved by passing ²⁰⁵ the reward signal through parallel filters, each designed to retain the signal variation specific ²⁰⁶ to that timescale (fig. 2a(bottom)). There are multiple approaches to this decomposition. We chose a heuristic approach in which the performance over a finite memory timescale can $_{208}$ be estimated by filtering the sequence of rewards through a simple low-pass filter [8, 28]. This filter is defined by an integration time, τ , tuned to trade off the bias and variance 209 ²¹⁰ of the estimate in order to best capture the variation on the desired timescale (e.g. how ²¹¹ performance varies over different contexts). We denote such an estimate $\hat{\rho}_k^{\tau}$, and show in $_{212}$ the Methods that it approximates the average reward over the last τ time units. We discuss ²¹³ the question of biological implementation in the discussion, but note here that the number $_{214}$ and values of τ needed to represent performance variation in a given task could be learned ²¹⁵ or selected from a more complete set in an online fashion during task learning. In an exper-²¹⁶ imental setting, these learned values can in principle be inferred from observed behaviour ²¹⁷ and we developed such an approach in the analysis of data that we present in the following 218 section.

Applying this heuristic decomposition here, the stationary reward rate, ρ , can be esti-220 mated to high precision by using a long integration time, τ_{long} , to the reward sequence R_k ,

9

²²¹ producing the estimate $\hat{\rho}_k^{\tau_{\text{long}}}$. If $\boldsymbol{\alpha}_k$ were a constant sequence, $\mathcal{C}_t^{\text{del}} = \hat{\rho}_k^{\tau_{\text{long}}} t$, the station-²²² ary opportunity cost of deliberation eq. (3) of AR-RL. However, in this example context ²²³ varies on a specific timescale, to which the former is insensitive. Thus, a second filtered ²²⁴ estimate $\hat{\rho}_k^{\tau_{\text{context}}}$ is needed to estimate performance on this timescale. Unlike $\hat{\rho}_k^{\tau_{\text{long}}}$, this es-²²⁵ timate tracks the effective instantaneous, context-specific performance, ρ_{α_k} . Its estimation ²²⁶ error arises from a trade-off, controlled by the integration time, τ_{context} , between its speed ²²⁷ of adaptation and its finite memory.

We consider two distinct hypotheses for how to extend AR-RL to settings where performance varies over context. The first hypothesis, $C_t^{\text{del}} = \rho_{\alpha} t$, is the straightforward, *trialunaware* extension of eq. (3), shown in fig. 2b(top). Here, performance is tracked only on a timescale sufficient to capture context variation and the corresponding cost estimate, $\hat{\rho}_{k-1}^{\text{rcontext}}$, is incurred moment-to-moment, neglecting the trial-based task structure. However, this incorrectly lumps together two distinct opportunity costs: those incurred by momentby-moment decisions and those incurred as a result of the effective planning implied by performance that varies over context. In particular, context is defined over trials not moments, and thus the context-specific component of opportunity cost of a trial is a sunken the outset of a trial. This inspires a second *trial-aware* hypothesis

$$\mathcal{C}_t^{\text{del}} = \rho t + (\rho_\alpha - \rho) T_\alpha \,. \quad \text{(trial-aware opportunity cost)} \tag{5}$$

238 Equation (5) is plotted over trial time t in fig. 2b(bottom). Its first term is the AR-RL ²³⁹ contribution from the stationary opportunity cost of moment-to-moment decisions using 240 the stationary reward rate, ρ estimated with $\hat{\rho}_k^{\tau_{\text{long}}}$. The second, novel term in eq. (5) is a ²⁴¹ context-specific trial cost deviation incurred at the beginning of each trial and computed as ²⁴² the average deviation in opportunity cost accumulated over a trial from that context (T_{α} 243 is the average duration of a trial in context α). This deviation fills the cost gap made by ²⁴⁴ using the stationary reward rate ρ in the moment-to-moment opportunity cost instead of ²⁴⁵ the context-specific average reward, ρ_{α} . This baseline cost derived from the orange time 246 series in fig. 2a(bottom) vanishes in expectation, as verified through the mixed-context ²⁴⁷ ensemble average reward (e.g. $\rho \equiv \sum_{\alpha} \rho_{\alpha} T_{\alpha} / \sum_{\alpha} T_{\alpha}$ when the context is distributed evenly ²⁴⁸ among trials such that $\sum_{\alpha} (\rho_{\alpha} - \rho) T_{\alpha} = 0$). Thus, this opportunity cost reduces to that 249 used in AR-RL when ignoring context, and suggests a generalization of average-adjusted ²⁵⁰ value functions to account for non-stationary context. We estimate this baseline cost using $_{251}$ $(\hat{\rho}_{k-1}^{\tau_{\text{context}}} - \hat{\rho}_{k-1}^{\tau_{\text{long}}})T_{k-1}$, where we have used the sample T_{k-1} in lieu of the average T_{α} . See fig. S1 $_{252}$ for a signal filtering diagram that produces this estimate of eq. (5) from reward history. A ²⁵³ main difference between the cost profiles from the two hypotheses is the cost at early times. ²⁵⁴ Both the behaviour and neural recordings we analyze below seem to favor the second, trial-²⁵⁵ aware hypothesis eq. (5). We hereon employ that version in the main text, and show the ²⁵⁶ results for the trial-unaware hypothesis in fig. S7.

257

B. Neuroscience application: PGD in the tokens task

In this section, we apply the PGD algorithm to the "tokens task" [16]. We first give a simulated example with periodic context dynamics. We then present an application to a set of non-human primate experiments in which context variation was non-stationary [19]. For the latter, we used the decision time dynamics over trials to fit a model for each of the used the validated the models by assessing their ability to explain (1) the

10

²⁶³ concurrently recorded behaviour via their context-specific behavioural strategies and (2) the ²⁶⁴ neural activity in premotor cortex (PMd) via the temporal profile of the underlying neural ²⁶⁵ urgency signals.

In the tokens task, the subject must guess as to which of two peripheral reaching targets will receive the majority of tokens that randomly jump, one by one every 200ms, from a central pool initialized with a fixed number of tokens. Importantly, after the subject reports, the interval between remaining jumps contracts to once every 150ms (the "slow" condition) or once every 50ms (the "fast" condition), giving the subject the possibility to save time by taking an early guess. The interval contraction factor, $1 - \alpha$, for slow ($\alpha = 1/4$) and fast $\alpha = 3/4$) condition is parametrized $\alpha \in [0, 1]$, the incentive strength to decide early, which are serves as the task context.

In contrast to the patch leaving task example from Section A, the tokens task has many 275 within-trial states and the state dynamics is stochastic. With the t^{th} jump labelled $S_t \in$ 276 $\{-1,1\}$ serving as the state, for the purposes of prediction, the history of states can be 277 compressed into the tokens difference, $N_t = \sum_{i=1}^t S_i$, between the two peripheral targets 278 with $N_0 = 0$. The dynamics of N_t is an unbiased random walk (see fig. 3a), with its current 279 value sufficient to determine the belief of a correct report, b_t (computed in Methods). Since 280 for binary rewards, b_t is also the expected reward, N_t is also sufficient for determining the 281 opportunity cost of commitment, C_t^{com} (eq. (2)). We display this commitment cost dynamics 282 in fig. 3b. It evolves on a lattice (gray), always starting at 0.5 (for p = 1/2) and ending at 0 283 for all p. We assume the agent has learned to track this commitment cost. The PGD agent 284 uses this commitment cost, along with the estimate of the trial-aware deliberation cost, to 285 determine when to stop deliberating and report its guess.

1. A simulated example for a regularly alternating context sequence

287

We first show the behaviour of the PGD algorithm in the simple case where α switches back and forth every 300 trials (see fig. 3). We call such segments of constant α 'trial blocks', with context alternating between slow ($\alpha = 1/4$) and fast ($\alpha = 3/4$) blocks. The decision space in PGD is a space of opportunity costs, equivalent to the alternative decision space formulated using beliefs [7]. In particular, one can think of the deliberation cost as the decision boundary (fig. 3b). This boundary is dynamic (see Supplemental video), depending on performance history via the estimates, $\hat{\rho}_k^{\tau_{context}}$ and $\hat{\rho}_k^{\tau_{long}}$, of the context-conditioned and stationary average reward, respectively. The result of these dynamics is effective context planning: the PGD algorithm sacrifices accuracy to achieve shorter trial duration in trials of the fast block, achieving a higher context-conditioned reward rate compared to decisions understood by analyzing the dynamics of $\hat{\rho}_k^{\tau_{context}}$ and $\hat{\rho}_k^{\tau_{long}}$, and their effect on the dynamics of the decision time ensemble.

The two performance estimates behave differently from one another solely because of their distinct integration times. Ideally, an agent would choose τ_{context} to be large enough that it serves to average over trial-to-trial fluctuations in a context, but short enough to not average over context fluctuations. In contrast, the value of τ_{long} would be chosen large enough to average over context fluctuations. We apply those choices in this simulated example, with rounded values chosen squarely in the range in which the values inferred from the behaviour in the following application will lie. As a result of this chosen values, the context estimate $\hat{\rho}_k^{\tau_{\text{context}}}$ relaxes relatively quickly after context switches to the context-



Figure 3. PGD agent performs the tokens task for periodic context switching. (a) A tokens task trial. Left: Tokens jump from a center to a peripheral region (gray circles). Right: The tokens difference, N_t , evolves as a random walk that accelerates according to α (here 3/4) post-decision time, t^{dec} . The trial duration is T, which includes an inter-trial interval. (b) Decision dynamics in cost space obtained from evidence dynamics in (a). Commitment cost trajectories (gray lattice; thick gray: trial-averaged) start at $C_{t=0}^{\text{com}}$ and end at 0. Trajectory from (a) shown in black. t^{dec} (black cross) is determined by the crossing of the commitment and deliberation cost. (c) Incentive strength switches between two values every 300 trials. (d) Expected rewards filtered on τ_{long} ($\hat{\rho}_k^{\tau_{\text{long}}}$, purple) and τ_{context} ($\hat{\rho}_k^{\tau_{\text{context}}}$, green). Black dashed lines from bottom to top are $\rho_{\alpha=1/4}$, ρ , and $\rho_{\alpha=3/4}$.

conditioned stationary average performance (dashed lines in fig. 3d), but exhibits stronger fluctuations as a result. The estimate of the stationary reward, $\hat{\rho}_k^{\tau_{\text{long}}}$, on the other hand has relatively smaller variance. This variance results from the residual zigzag relaxation over the period of the limit cycle. Given the characteristic block duration, T_{block} , we can be more precise. In particular, when T_{block} is much less than τ_{long} ($T_{\text{block}}/\tau_{\text{long}} \ll 1$), the within-block are exponential relaxation is roughly linear. Thus, the average unsigned deviation between $\hat{\rho}_k^{\tau_{\text{long}}}$ and the actual stationary reward, ρ , can be approximated using $1 - \exp\left[-T_{\text{block}}/\tau_{\text{long}}\right] \approx$ $T_{\text{block}}/\tau_{\text{long}} \ll 1$. This scaling fits the simulated data well (fig. S2d: inset).

The dynamics of these two performance estimates drives the dynamics of the k-conditioned are decision time ensemble via how they together determine the deliberation cost (eq. (5); Supare plemental video). For example, the mean component of this ensemble relaxes after a context

12

³²⁰ switch to the context-conditioned average, while the fluctuating component remains strong ³²¹ due to the sequence of random walk realizations (fig. S2c). In the case of periodic context, ³²² the performance estimates and thus also the decision time ensemble relax into a noisy peri-³²³ odic trajectory over the period of a pair of fast and slow blocks (fig. 3d). Over this period, ³²⁴ they exhibit some stationary bias and variance relative to their corresponding stationary ³²⁵ averages (distributions shown in fig. S2e).

2. Fit to behavioural data from non-human primates and model validation

326

Next, we fit a PGD agent to each of the two non-human primates' behaviour in the to kens task experiments reported in [19] and compare to AR-RL solutions. As with the above example (*c.f.* fig. 3), trials were structured in alternating blocks of $\alpha = 1/4$ and $\alpha = 3/4$. Figure 4a shows context-switching α -sequence from these experiments, which, in are contrast to the above example exhibits large, irregular fluctuations in block size [29].

So far, PGD has only two free parameters: the two filtering time constants, τ_{long} and 333 ³³⁴ τ_{context} . We anticipated only a weak dependence of the fit on the τ_{long} , so long as it exceeded ³³⁵ the average duration of a handful of trial blocks enabling a sufficiently precise estimate of 336 ρ . In contrast, the context filtering timescale, τ_{context} , is a crucial parameter as it dictates ³³⁷ where the PGD agent lies on a bias-variance trade-off in estimating ρ_{α_k} , the value of which $_{338}$ determines the context-specific contribution to the deliberation cost (eq. (2)). To facilitate ³³⁹ the model's ability to fit individual differences, we introduce a subjective reward bias factor, 340 K, that scales the rewards fed into the performance filters. We also add a tracking-cost sen-341 sitivity parameter, ν , that controls τ_{context} to avoid wasting adaptation speed (see Methods ³⁴² for details). The latter made it possible to fit the asymmetric switching behaviour observed ³⁴³ in the average decision time dynamics. With these four parameters, we quantitatively match the baselines and exponential-like relaxation of the average decision time dynamics around 344 the two context switches (fig. 4b,c; see Methods for fitting details). 345

A comparison of the best-fitting parameter values over the two monkeys (fig. 4d-f) sug-347 gests that the larger the reward bias, K (fig. 4e), the more hasty the context-conditioned 348 performance estimate (the smaller τ_{context}), and the lower the sensitivity to the tracking cost 349 (fig. 4f). This is consistent with the hypothesis that subjects withhold cognitive effort in 350 contexts of higher perceived reward [8]. Along with the correspondence in temporal statistics 351 of the behaviour (e.g. fig. S6), the fitted model parameters for the two subjects provides a 352 basis on which to interpret the subject differences in the results of the next section, in par-353 ticular their separation on a speed-accuracy trade-off, as originating in the distinct reward 355 sensitivity shown here.

To better understand where both the data and the learned PGD agent lie in the space of strategies for the tokens task, we computed reward-rate (AR-RL) optimal solutions for a given fixed context, α (here $\alpha \in [0, 1]$), using the same approach as [7] (conventional discount-reward value iteration achieved the same solution in the limit of the undiscounted the same solution in the limit of the undiscounted dynamic programming approach involves iterating Bellman's equation to obtain the optimal value functions from which the optimal policy and its reward rate can be obtained (see Methods for details). The optimal reward rate as a function α is shown in fig. 5a. The strategies generating these reward rates interpolate from the wait-for-certainty strategy at to the one-and-done strategy [30] at high α . The α -conditioned reward rates achieved by the two primates with their corresponding PGD model, and a reference human [31]

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13



Figure 4. PGD model fit to NHP behaviour for non-stationary α -dynamics reported in Ref. [19]. (a) Block length sequence used in the experiment. (b,c) decision times (dots) aligned on the contextswitching event type (fast-to-slow in gray; slow-to-fast in color) and averaged. Shaded regions are the standard error bounds of the models' average decision times. (d) Error evaluated on a $(\hat{\tau}_{\text{context}}, \hat{\tau}_{\text{long}})$ -plane cut through the parameter space at the best-fitting $\nu = \hat{\nu}^*$ and $K = \hat{K}^*$ (gray area indicates timescales within an order of magnitude of the end of the experiment). Contours show the first 10 contours incrementing by 0.01 error from the minimum (shown as a circle marker). Colors refer to subject, as in (b) and (c). (e) Same for $(\hat{\tau}_{\text{context}}, \hat{K})$ at $\hat{\tau}_{\text{long}} = \hat{\tau}^*_{\text{long}}$ and $\nu = \hat{\nu}^*$. (f) Same for $(\hat{\tau}_{\text{context}}, \hat{\nu})$ at $\hat{\tau}_{\text{long}} = \hat{\tau}^*_{\text{long}}$ and $K = \hat{K}^*$.

367 are also shown in fig. 5a. They clearly fall below the optimal strategy, and, as expected, 368 above the strategy that picks one of the three actions (report left, report right, and wait) at 369 random.

To confirm that this similarity in performance between PGD and the data arises from and a better fit to the behaviour than AR-RL, we plotted the distribution of the differences between model and data decision times, $|\Delta t_{dec}|$, conditioned on the context (fig. 5b,c). For around comparison with previous work [7] and to account for deliberation cost in AR-RL, we added to the AR-RL reward objective a constant auxiliary deliberation cost rate, c, incurred up to the decision time in each trial, and chose the cost rate, c^* , that gave the lowest mean difference. In both contexts, PGD exhibits lower error than this c^* AR-RL solution.

³⁷⁷ To reveal the source of this discrepancy in both performance and behaviour, we turned





Figure 5. Context-conditioned analysis of PGD and comparison to AR-RL models. (a) Shown is the reward rate as a function of incentive strength, α (wait-for-certainty strategy shown in brown; one-and-done strategy shown in red). We additionally show the slow and fast contextconditioned reward rates for the two primates and the PGD model fitted to them, as well as a reference expert human. Reward rates for the human and non-human primates are squarely in between the best (black dashed) and uniformily random (gray) strategy. (b,c) The distribution over trials of differences in decision times between model and data, $|\Delta t_{\rm dec}| = |t_{\rm dec,data} - t_{\rm dec,model}|$ conditioned on slow and fast block contexts. Solid lines are for PGD. Dotted lines are for the AR-RL solution using the cost rate, c^* , with the lowest mean error. The residual sum of squares (RRS) for each model/block combination is displayed. (d-g) Interpolated state-conditioned survival probabilities, $P(t^{\text{dec}} = t | N_t, t)$, over slow (d,f) and fast (e,g) blocks. White dotted lines show the $P(t^{\text{dec}} = t | N_t, t) = 0.5$ contour. (h,i) State-conditioned decision time frequencies (cross size) from AR-RL optimal decision boundaries across different values of the cost rate, c (colored crosses) for slow (h) and fast (i) conditions. Only samples with $N_t < 0$ and $N_t > 0$, respectively, are shown. For comparison, the reflected axes shows as gray crosses the state-conditioned decision time frequencies of the data.

15

³⁷⁸ to analyzing the corresponding policies of PGD and c-based AR-RL agents. A robust and ³⁷⁹ rich representation of the behavioural statistics is the state and time-conditioned survival ³⁸⁰ probability that a decision has not yet occurred. It serves as a summary of the action policy ³⁸¹ associated with a stationary strategy (see Methods for its calculation from response times). ₃₈₂ Applied equally to the decision times of both model and data, it can provide a means of ³⁸³ comparison even in this non-stationary setting. We give this conditional probability for ₃₈₄ each of the two contexts for subject 1 and its fitted PGD model in fig. 5d-g. We left the ³⁸⁵ many possible noise sources underlying the behaviour out of the model in order to more ³⁸⁶ clearly demonstrate the PGD algorithm. However, such noise sources would be necessary ³⁸⁷ to quantitatively match the variability in the data (e.g. added noise in the performance ₃₈₈ estimates leads to larger variability in the location of the decision boundary and thus also to larger spread in these survival probability functions (not shown)). In the absence of these noise sources, we see the model underestimates the spread of probability over time ³⁹¹ and tokens state. Nevertheless, the remarkably smooth average strategy is well captured by ³⁹² the model (white dashed lines in fig. 5d-g). Specifically, policies approximately decide once ³⁹³ either of the peripheral targets receive a certain number of tokens. Comparing results across ³⁹⁴ context, we find that fast block strategies (fig. 5e,g) exhibit earlier decision times relative $_{395}$ to slow block strategies (fig. 5d,f) in both model and data. The strategies for subject 2 ³⁹⁶ are qualitatively similar, but shifted to earlier times relative to subject 1 (fig. S3). Our ³⁹⁷ model explains this inter-individual difference as resulting from subject 2's larger reward $_{398}$ bias and faster context integration (c.f. fig. 4e). The correspondence between the PGD ³⁹⁹ model and data over the many token states in fig. 5d-g explains their similar performance (c.f. fig. 5a). This similarity in policy is remarkable given that the model has essentially 401 only a single, crucial degree of freedom (τ_{context}), a priori unrelated to how decision times 402 depend on token state. Note that in both the fitted PGD model and the primate behaviour, residual ambiguity $(N_t \approx 0)$ is resolved at intermediate trial times (fig. 5b-e). 403 The AR-RL strategies are plotted across c in fig. 5g,h. In contrast, they give no interme-404 405 diate decision times at ambiguous $(N_t \approx 0)$ states, invariably waiting until the ambiguity ⁴⁰⁶ resolves. This in fact holds over the entire (α, c) -plane (see fig. S9 for the complete depen-⁴⁰⁷ dence), and also under the addition of a movement cost, i.e. a constant cost incurred by

⁴⁰⁸ either of the reporting actions (data not shown). Thus, whereas AR-RL policies shift around ⁴⁰⁹ the edges of the relevant decision space as α or c is varied, the PGD policy lies squarely ⁴¹⁰ in the bulk, tightly overlaying the policy extracted from the data. We conclude that the ⁴¹¹ context-conditioned strategies of the non-human primates in this task are well-captured by ⁴¹² PGD, while having little resemblance to the behaviour that would maximize reward rate ⁴¹³ with or without a fixed deliberation cost rate. We address the additional freedom of a ⁴¹⁴ time-varying cost rate in the discussion.

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3. Neural urgency and context-dependent opportunity cost

So far, we have fit and analyzed the PGD model with respect to recorded behaviour. Here, 417 we take a step in the important direction of confronting the above theory of behaviour with 418 the neural dynamics that we propose drive it. The proposal for the tokens task mentioned 419 at the end of the introduction has evidence strength and urgency combining in PMd, whose 420 neural dynamics implements the decision process. In fig. 6a, we restate in a schematic 421 diagram an implementation of this dynamics that includes a collapsing decision boundary. 422 In the one-dimensional belief space for the choice (fig. 6a(top)) [7, 32], the rising belief

16

⁴²³ collides with the collapsing boundary to determine the decision time. In the equivalent ⁴²⁴ commitment and deliberation cost formulation developed here (fig. 6a(middle)), the falling ⁴²⁵ commitment cost collides with the rising deliberation cost. The collapsing boundary in ⁴²⁶ belief space can be parametrized as $C - u_t$, where C is the initial strength of belief, e.g. ⁴²⁷ some desired confidence, that is lowered by a growing function of trial time $u_t > 0$. The ⁴²⁸ decision criterion is then $b_t > C - u_t$, where b_t is the belief, i.e. the probability of a correct ⁴²⁹ report. For AR-RL optimal policies, u_t emerges from value maximization and thus has a ⁴³⁰ complicated dependence on the opportunity cost sequence, C_t^{del} . For PGD, in contrast, C⁴³¹ is interpreted as the maximum reward r_{max} and u_t is identically C_t^{del} . For a linear neural ⁴³² encoding model in which belief, rather than evidence, is encoded in neural activity, the sum ⁴³³ of the encoded belief \tilde{b}_t and the encoded collapsing boundary, \tilde{u}_t , evolve on a one-dimensional ⁴³⁴ choice manifold. According to the proposal, when this sum becomes sufficiently large (e.g. ⁴³⁵ $\tilde{b}_t + \tilde{u}_t > \tilde{C}$ for some threshold \tilde{C}), PMd begins to drive the activity in downstream motor ⁴³⁶ areas towards the associated response.

Neural urgency was computed from the PMd recordings of [19] in [33]. This computation 437 438 relies on the assumption that while a single neuron's contribution to b_t will depend on 439 its selectivity for choice (left or right report), the urgency \tilde{u}_t is a signal arising from a population-level drive to all PMd neurons, irrespective of their selectivity. Thus, \tilde{u}_t can 441 be extracted from neural recordings by conditioning on zero-evidence states $(b_t = 0)$ and ⁴⁴² averaging over cells. In [33], error bars were computed at odd times via bootstrapping; data 443 at even times was obtained by interpolating between $N_t = \pm 1$; and data was pooled from 444 both subjects. We have excluded times at which firing rate error bars exceed the range 445 containing predictions from both blocks. To assess the correspondence of the components 446 of the deliberation cost developed here and neural urgency, in fig. 6b we replot their result $_{447}$ (c.f. fig.8b of [33]). We overlay the mean (+/- standard deviation) of the opportunity cost ⁴⁴⁸ sequence, $\mathcal{C}_t^{\text{del}}$ (shaded area in fig. 4; averaged over all trials produced by applying the two 449 fitted PGD models on the data sequence and conditioning the resulting average within-450 trial deliberation cost on context). To facilitate our qualitative comparison, we convert ⁴⁵¹ cost to spikes/step simply by adjusting the y-axis of the deliberation cost. The observed ⁴⁵² urgency signals then lie within the uncertainty of the context-conditioned deliberation cost ⁴⁵³ signals computed from the fitted PGD models. There are multiple features of the qualitative ⁴⁵⁴ correspondence exhibited in fig. 6b: (1) the linear rise in time; (2) the same slope across 455 both fast and slow conditions; and (3) the baseline offset between conditions, where the fast ⁴⁵⁶ condition is offset to higher values than the slow condition. Such features would remain ⁴⁵⁷ descriptive in the absence of a theory. With the theory we have presented here, however, ⁴⁵⁸ each has their respective explanations via the interpretation of urgency as the opportunity 459 cost of deliberation: (1) the subject uses a constant cost per token jump, (2) this cost rate 460 refers to moment-to-moment decisions, irrespective of context, that is reflective of the use 461 of the context-agnostic stationary reward, and (3) trial-aware planning over contexts leads ⁴⁶² to an opportunity cost baseline offset with a sign given by the reward rate deviation $\rho_{\alpha} - \rho$ with respect to the stationary average, ρ . 463

⁴⁶⁴ Up to now, the computational and neural basis for urgency has remained largely un-⁴⁶⁵ explored in normative approaches, which also typically say little about adaptation effects ⁴⁶⁶ (see [34] for a notable exception). In summary, we exploited the adaptation across context ⁴⁶⁷ switches to learn the model and explained earlier responses in high reward rate contexts ⁴⁶⁸ as the result of a higher opportunity cost of deliberation. While this qualitative effect is ⁴⁶⁹ expected, we go beyond existing work by quantitatively predicting the average dependence

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Figure 6. Comparing neural urgency and collapsing decision boundaries. (a) Top: Rising belief (blue) meets collapsing decision boundary (black dashed) in belief space. Middle: Falling commitment cost (blue) meets rising deliberation cost (black-dashed) in cost space. Bottom: Belief/commitment cost is encoded (blue) into a low-dimensional neural manifold, with the addition of an urgency signal (orange) (c.f. fig.8 in [7]). The decision (red circle) is taken when the sum passes a fixed threshold (black-dashed). (b) Deliberation cost maps onto the urgency signal extracted from zero-evidence conditioned cell-averaged firing rate in PMd (200ms time steps).

⁴⁷⁰ on both time and state (fig. 5b-e) as well as the qualitative form of urgency signal (fig. 6b). ⁴⁷¹ Taken together, the data is thus consistent with our interpretation that neural activity un-⁴⁷² derlying context-conditioned decisions is gated by opportunity costs reflective of a trial-aware ⁴⁷³ timescale hierarchy computed using performance estimation on multiple timescales.

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DISCUSSION

We introduced PGD, a heuristic decision-making algorithm for continuing tasks that 475 gates deliberation based on performance. We constructed a foraging example for which 476 PGD is the optimal strategy with respect to the average-adjusted value function of averagereward reinforcement learning (AR-RL). While this will not be true in general, PGD does strike a balance between strategy complexity and return. The PGD decision rule does not 479 depend on task specifics and exploits the stationarity of the environment statistics while simultaneously hedging against longer term non-stationarity in reward context. It does so 481 by splitting the problem into two fundamental components—learning the statistics of the ⁴⁸³ environment in order to compute the opportunity cost of commitment, and tracking one's ⁴⁸⁴ own performance in that environment with which to compute the opportunity cost of deliberation. This splitting is not only crucial to making efficient use of the opportunity cost of time in non-stationary settings. Building on the field's current understanding of how the cortico-basal ganglia system supports higher-level decision-making [35], we propose that the 487 cost of deliberation arises from performance estimated on multiple, behaviourally-relevant timescales and is broadcast to multiple, lower-level decision-making areas to gate the speed ⁴⁹⁰ of their respective evidence-driven attractor dynamics. Incorporating this cost into existing

18

⁴⁹¹ models of such dynamics [32, 36, 37] is an interesting direction for future work. Consistent ⁴⁹² with this picture, PGD's explanatory power was borne out at both the behavioural and ⁴⁹³ neural levels for the tokens task data we analyzed. In particular, a deliberation cost con-⁴⁹⁴ structed from trial-aware planning was supported independently by both these data sources. ⁴⁹⁵ We used behavioural data to fit and validate the theory, and neural recordings to provide ⁴⁹⁶ evidence of one of the neural correlates it proposes: the temporal profile of neural urgency.

Scientific and clinical implications In our proposal, we have linked two important and 497 ⁴⁹⁸ related, but often disconnected fields: the systems neuroscience of the neural dynamics of decision-making and the cognitive neuroscience of opportunity cost and reward sensitivity. 400 The view that tonic dopamine encodes average reward is two decades old [3]. However, the 500 existence of a reward representation decomposed by timescale has received increasing empirical support only in recent years, from cognitive results [38-40] to a recent unified view of 502 $_{503}$ how dopamine encodes reward prediction errors using multiple discount factors [41, 42] and $_{504}$ of dopamine as encoding both value and uncertainty [43]. Dopamine's effect on time perception has been proposed [44] and has empirical support [45], but the mechanism by which its putative effect on decision speed is implicated in the neural dynamics of the decision-506 ⁵⁰⁷ making areas driving motor responses was unknown. Our theory fills this explanatory gap by considering dynamic evidence tasks and parametrizing urgency using a multiple-timescale 508 representation of performance. One candidate for the latter's neural implementation is in ⁵¹⁰ the complex spatio-temporal filtering of dopamine via release-driven tissue diffusion and ⁵¹¹ integration via DR1 and DR2 binding kinetics [46]. Subsequent neural filtering and compu-⁵¹² tation by striatal network activity could also play a role [47]. The study of spatiotemporal ⁵¹³ filtering of dopamine is increasingly accessible experimentally [48, 49] and provides an excit-⁵¹⁴ ing direction for multiscale analysis of behaviour. Our proposal that urgency is the means ⁵¹⁵ by which the neural representation of reward ultimately affects neural dynamics in decision-⁵¹⁶ making areas frames a timely research question on which these experimental methods could 517 shed light.

We applied PGD to decisions playing out in PMd, a decision-making area relevant to arm 518 ⁵¹⁹ movements. PGD appears to be relevant to other kinds of decisions, however. For instance, ⁵²⁰ a large body of work has studied decisions through recordings in lateral intraparietal cortex ⁵²¹ in random dot motion tasks whose environment is formally similar to that of the tokens task. One seminal study identified an urgency signal with the same properties as those exhibited by the tokens task: a linear rise at early trial times that is independent of trial evidence and an offset with sign given by the reward rate deviation of the current context, here two and four-choice trials [17]. While decision boundaries obtained using AR-RL are evidenceindependent, these models require tailored cost functions that are fit to those experiments $_{527}$ in a procedure that assumes optimality *a piori* [7]. Here, we offer an alternative explanation that behaviour is in fact suboptimal, with the decision boundary determined directly by the 528 estimated opportunity cost only. PGD decision boundaries are thus independent of evidence ⁵³⁰ by construction. In contrast to the tokens task, however, context in these random dot task ⁵³¹ experiments was sampled randomly and thus its dynamics lacked temporal correlation [17]. ⁵³² In this case, a natural hypothesis from our approach is that a pair of performance filters, one for each context, tracks the reward history in two parallel streams. In this case, our 534 theory would predict that the ratio of slopes of urgency across the two contexts reflects ⁵³⁵ the ratio of context-conditioned reward rates. An estimation procedure described in the Methods for this data [17] agrees to within 20% error, providing support for the hypothesis ⁵³⁷ that PGD underlies non-human primate behaviour on this widely-studied task. Within the

19

⁵³⁸ context of the drift-diffusion models typically used to understand neural activity for that ⁵³⁹ task, PGD provides a principled mechanism that implements collapsing decision boundary. ⁵⁴⁰ PGD is thus easily incorporated into such models and testing the generality of our theory ⁵⁴¹ using tailored experiments in this setting is an important next step.

⁵⁴² Urgency may play a role in both decision and action processes, potentially providing a ⁵⁴³ transdiagnostic indicator of a wide range of cognitive and motor impairments in Parkinson's ⁵⁴⁴ disease and depression [50]. Our theory offers a means to ground these diverse results in ⁵⁴⁵ neural dynamics by formulating opportunity cost estimation as the underlying causal factor ⁵⁴⁶ linking vigor impairments (e.g. in Parkinson's disease) and dysregulated dopamine signalling ⁵⁴⁷ in the reward system [50–52]. We provide a concrete proposal for a signal filtering system ⁵⁴⁸ that extracts a context-sensitive opportunity cost from a reward prediction error sequence ⁵⁴⁹ putatively encoded by dopamine. Neural recordings of basal ganglia provide a means to ⁵⁵⁰ identify the neural substrate for this system.

⁵⁵¹ Commitment cost estimation Beyond the estimation of the opportunity cost of deliber-⁵⁵² ation, we assumed that the agent had a precise estimate of the expected reward, which it ⁵⁵³ used to compute the within-trial commitment cost. For the tokens task, a recorded signal in ⁵⁵⁴ dorsal lateral prefrontal cortex of non-human primates correlates strongly with belief [20], ⁵⁵⁵ equivalent to the expected reward for binary rewards). How this quantity is computed by ⁵⁵⁶ neural systems is not currently known. However, for a general class of tasks, a generic, ⁵⁵⁷ neurally plausible means to learn the expected reward is via distributional value codes [43]. ⁵⁵⁸ For example, the Laplace code is a distributional value representation that uses an ensemble ⁵⁵⁹ of units over a range of temporal discount factors and reward sensitivities [53]. The authors ⁵⁶⁰ show that expected reward is linearly decodeable from this representation.

Experimental predictions A feature of our decision-making theory is that it is highly 561 ⁵⁶² vulnerable to falsification. First, with regards to behaviour via the shape of the action ⁵⁶³ policy using our survival probability representation (*c.f.* fig. 5b-e,g,h), PGD varies markedly ⁵⁶⁴ with reward structure and thus provides a wealth of predictions for how observed behaviour ⁵⁶⁵ should be altered by it. For example, a salient feature of the standard tokens task is its ⁵⁶⁶ reflection symmetry in the tokens difference, N_t . We can break this symmetry for which the ⁵⁶⁷ theory predicts a distinctly asymmetric shape (fig. S10; for details see Methods). Our theory 568 is also prescriptive for neural activity via the temporal profile of neural urgency. The slope of $_{569} \mathcal{C}_t^{\text{del}}$ remained fixed across blocks for relatively short block lengths used in the data analyzed 570 here. In the opposite limit, $T_{\rm block}/\tau_{\rm long} \gg 1$, $\rho_k^{\tau_{\rm long}}$ approaches ρ_{α} except when undergoing 571 large, transient excursions after context switches. Thus, the deliberation cost is given by $_{572}$ the first component in eq. (5) most of the time, with the context specific reward rate as the ⁵⁷³ slope. One simple prediction is that the slope of urgency should exhibit increasing variation as the duration of the blocks increases. 574

Reinforcement learning theory We suggest how to generalize average-adjusted value from functions to context-varying opportunity cost of time in a way that reduces to AR-RL from known of the fixed or not tracked. This adds a continuing task perspective to episodic AR-RL, in line with recent work in machine learning, which is arguably the more appropriate from perspective entailed in the estimation of these costs parallels a recent epistemic interpretation of the discount-reward formulation as encoding knowledge about the volatility for the environment [54].

Our work also suggests a new class of reinforcement learning algorithms between model-⁵⁸⁴ based and model-free: only parts of the algorithm need adjustment upon task structure

20

⁵⁸⁵ variation. This is reminiscent of how the effects of complex state dynamics are decoupled ⁵⁸⁶ from reward when using a successor representation [55], but tailored for the average-reward ⁵⁸⁷ rather than the discount-reward formulation. We have left analysis of the algorithmic com-⁵⁸⁸ plexity of PGD to future work, but expect performance improvements, as with successor ⁵⁸⁹ representations, in settings where decoupling the learning of environment statistics from the ⁵⁹⁰ learning of reward structure is beneficial.

Comparison with humans In the space of strategies, PGD lies in a regime between fully 591 exploiting assumed task knowledge (average-case optimal) and assumption-free adaptation 592 (worst-case optimal). Highly incentivized human behaviour is likely to be more structured 593 than PGD because of access to more sophisticated learning. While some humans land on 594 the optimal one-and-done policy in the fast condition when playing the tokens task [56]. most do not. The human brain likely has all the components needed to implement PGD. Nevertheless, the situations in which we actually exploit PGD, if any, are as yet unclear. In 597 particular, how PGD and AR-RL relate to existing behavioural models tailored to explain ⁵⁹⁹ relative-value, context-dependent decision-making in humans [4], such as scale and shift adaptation [57], is an open question. Whether or not PGD is built into our decision-making, the question remains if PGD is optimal with respect to some bounded rational objective. ₆₀₂ In spite of the many issues with the latter approach [58], using it to further understand the ⁶⁰³ computational advantages of PGD is an interesting direction for future work.

Despite our putative access to sophisticated computation, humans still exhibit measurable bias in how we incorporate past experience [59]. One simple example is the win-stay/loseshift strategy, a more rudimentary kind of performance-gated decision-making than PGD, which explains how humans approach the rock-paper-scissors game [60]. In that work, numerical experiments demonstrated that this strategy outperforms at a population level the optimal Nash equilibrium for this game, demonstrating that the use of such seemingly suboptimal strategies can confer a surprising evolutionary advantage. This example supports the claim that relatively simple and nimble strategies such as PGD make for attractive candidates when acknowledging that a combination of knowledge and resource limitations over task, development, and evolutionary timescales have shaped decision-making in nonstationary environments.

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METHODS

⁶¹⁶ Code for simulations and main figure generation (written in Python 3) is publicly acces-⁶¹⁷ sible as a online repository: https://github.com/mptouzel/dyn_opp_cost/.

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Patch leaving task

⁶¹⁹ We devised a mathematically tractable patch leaving task for which PGD learning is ⁶²⁰ optimal with respect to the average-adjusted value function. Here the value is simply the ⁶²¹ return from the patch. This value function is related, but not equivalent to the marginal ⁶²² value of optimal foraging, for which the decision rule is $C_t^{\text{del}} > r_{\text{max}} - C_t^{\text{com}} = \bar{r}_t$ [5]). This ⁶²³ choice of task allowed us to compare PGD's convergence properties relative to conventional ⁶²⁴ AR-RL algorithms that make use of value functions. In contrast to PGD, the latter requires ⁶²⁵ exploration. For a comparison generous to the AR-RL algorithm, we allowed it to circumvent ⁶²⁶ exploration by estimating the value function from off-policy decisions obtained from the

21

⁶²⁷ PGD algorithm using the same learning rate. We then compared them to PGD using their ⁶²⁸ on-policy, patched-averaged reward. This made for a comparison based solely between the ⁶²⁹ parameters of the respective models. If we did not allow for this, the AR-RL algorithms ⁶³⁰ would have to find good learning signals by exploring. In any form, this exploration would ⁶³¹ lead them converge substantially slower. This setting thus provides a lower bound on the ⁶³² convergence times of the AR-RL algorithm.

In this task, the subject randomly samples (with replacement) d patches, each of a distinct, fixed, and renewable richness defined by the maximum return conferred. These maximum returns are sampled before the task from a richness distribution, $p(r_{\text{max}})$, with $r_{\text{max}} > 0$ and are fixed throughout the experiment. The trials of the task are temporally extended periods during which the subject consumes the current patch. After a time t in a patch, the return is defined $r(t) = r_{\text{max}}(1 - (\lambda t)^{-1})$. This patch return profile, $1 - (\lambda t)^{-1}$, is shared across all patches and saturates in time with rate λ , a parameter of the environment that for mathematical convenience, but also evokes situations where leaving a patch soon after arriving is prohibitively costly (e.g. when transit times are long). A stationary policy is then a leaving time, t_s , for each of d patches, where the s-subscript indexes the patch. Given any policy, the stationary reward rate for uniformly random sampling of patches is then defined as

$$\rho = \sum_{s=1}^{d} r_s(t_s) \bigg/ \sum_{s=1}^{d} t_s .$$
 (6)

⁶⁴⁶ We designed this task to (1) emphasize the speed-return trade-off typical in many delibera-⁶⁴⁷ tion tasks, and (2) have a tractable solution with which to compare convergence properties ⁶⁴⁸ of PGD and AR-RL value function learning algorithms.

A natural optimal policy is the one that maximizes the average-adjusted trial return, 649 ₆₅₀ $Q(r,t) = r - \rho t$. Given the return profile we have chosen, the corresponding optimal decision ⁶⁵¹ time, t_s^* , in the sth patch obtained by maximizing $r - \rho t$ is $t_s^* = \sqrt{r_{\max,s}/(\lambda \rho)}$, which scales ₆₅₂ inversely with the reward rate so that decision times are earlier for larger reward rates, because consumption (or more generally deliberation) at larger reward rates costs more. We 653 ⁶⁵⁴ chose this return profile such that stationary PGD learning gives exactly the same decision 655 times: the condition $C_t^{\text{del}} = C_t^{\text{com}}$ for patch *s* here takes the form $\rho t_s = r_{\max,s}/(\lambda t_s)$. Thus, 656 they share the same optimal reward rate, ρ^* . Using t_s^* for each patch in eq. (6) gives a 657 self-consistency equation for ρ with solution $\rho^* = \lambda \mu_1^2/4\mu_{1/2}^2$, where $\mu_n = \langle r_{\max}^n \rangle_{p(r_{\max})}$ (we $_{658}$ have assumed d is large here to remove dependence on s). Described so far in continuous 659 time, the value function was implemented in discrete time such that the action space is a finite set of decision times selected using the greedy policy, $t^* = \operatorname{argmax}_t Q(r, t)$, where $\hat{Q}(r,t)$ is the estimated trial return. As a result, there is a finite lower bound on the ₆₆₂ performance gap, i.e. the relative error, $\epsilon = (\rho^* - \rho)/\rho^* > 0$ for the AR-RL algorithm. ⁶⁶³ Approaching this bound, convergence time for both PGD and AR-RL learning is limited ₆₆₄ by the integration time τ of the estimate $\hat{\rho}_k^{\tau}$ (c.f. eq. (8)) of ρ . We note that PGD learns ⁶⁶⁵ faster in all parameter combinations tested. To demonstrate the insensitivity of PGD to the state space representation, at 5×10^5 time steps into the experiment we shuffled the labels 667 of the states. PGD is unaffected, while the value function-based AR-RL algorithm is forced 668 to relearn and in fact does so slower than in the initial learning phase, due to the much 669 larger distance between two random samples, than between the initial values (chosen near 670 the mean) and the target sample.

22

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Filtering performance history

For unit steps of discrete time, the step-wise update of the performance estimate, $\hat{\rho}_t^{\tau}$, is

$$\hat{\rho}_t^{\tau} = (1 - \beta)\hat{\rho}_{t-1}^{\tau} + \beta R_t , \qquad (7)$$

⁶⁷³ with $\beta = 1/(1+\tau)$ called the learning rate, and τ the characteristic width of the exponential ⁶⁷⁴ window of the corresponding continuous time filter over which the history is averaged. We ⁶⁷⁵ add τ as a superscript when denoting the estimate to indicate this. Exceptionally, here t ⁶⁷⁶ indexes absolute time rather than trial time. Note that a continuous-time formulation of ⁶⁷⁷ the update is possible via an event-based map given the decision times in which the reward ⁶⁷⁸ event sequence is given as a sum of delta functions. In either case, to leading order in β , ⁶⁷⁹ $\hat{\rho}_t^{\tau} \approx \beta \sum_i^t R_i$, i.e. the filter sums past rewards. Thus, when $\tau \sim \mathcal{O}(t) \gg 1$, $\beta \sim \mathcal{O}(1/t) \ll 1$ ⁶⁸⁰ and so $\hat{\rho}_t^{\tau} \approx \beta \sum_i^t R_i \to \rho$ when t is large.

The rewards in this task are sparse: $R_t = 0$ except when a trial ends and the binary trial reward R_k (1 or 0) is received. A cumulative update of eq. (7) that smooths the reward uniformly over the trial duration and is applied once at the end of each trial is thus more computationally efficient. Resolving a geometric series leads to the cumulative update [8, 28]

$$\hat{\rho}_{k}^{\tau} = (1 - \beta)^{T_{k}} \hat{\rho}_{k-1}^{\tau} + (1 - (1 - \beta)^{T_{k}}) \rho_{k}^{\text{trial}} , \qquad (8)$$

where the smoothed reward, $\rho_k^{\text{trial}} = R_k/T_k$, can be interpreted as a trial-specific reward rate. The initial estimate, $\hat{\rho}_0^{\tau}$, is set to 0. Exceptionally, $\hat{\rho}_1^{\tau} = R_1/T_1$, after which eq. (8) is used. Using the first finite sample as the first finite estimate is both more natural and robust than having to adapt from zero. We will reuse this filter for different τ and denote the filtered estimate from its application with a τ -superscript, $\hat{\rho}_k^{\tau}$. For example, the precision of $\hat{\rho}_k^{\tau_{\text{long}}}$ as an estimate of a stationary reward rate ρ is set by how many samples it averages over, which is determined by the effective length of its memory given by τ_{long} . Since we assume the subject has learned the expected reward, \bar{r}_t , we use it instead of R_k when computing ρ_k^{trial} .

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Tokens task: a random walk formulation

The tokens task is a continuing task of episodes (here trials), which can be formulated ⁶⁹⁷ using the token difference, N_t . Each trial effectively presents to the agent a realization ⁶⁹⁸ of a finite-length, unbiased random walk, $N_{t_{\text{max}}} = (N_0, \ldots, N_{t_{\text{max}}})$ with $N_t = \{-t, \ldots, t\}$ ⁶⁹⁹ and $N_0 = 0$. We express time in units of these steps. The agent observes the walk and ⁷⁰⁰ reports its prediction of the sign of the final state, $\operatorname{sign}(N_{t_{\text{max}}}) = \pm 1$ (t_{max} is odd to exclude ⁷⁰¹ the case it has no sign). The time at which the agent reports is called the decision time, ⁷⁰² $t^{\text{dec}} \in \{0, 1, \ldots, t_{\text{max}}\}$. For a greedy policy, $\operatorname{sign}(N_t)$ can be used as the prediction (and ⁷⁰³ the reporting action selected randomly if $N_{t^{\text{dec}}} = 0$). The decision-making task then only ⁷⁰⁴ involves choosing when to decide. In this case, the subject receives reward $R = \Theta(N_{t_{\text{max}}}N_{t^{\text{dec}}})$ ⁷⁰⁵ at the end of the random walk, i.e. a unit reward for a correct prediction, otherwise nothing ⁷⁰⁶ (Θ is the Heaviside function: $\Theta(x) = 1$ if x > 0, zero otherwise).

⁷⁰⁷ An explicit action space beyond decision time is not necessary for the case of greedy ⁷⁰⁸ actions. It can nevertheless be specified for illustration in an Markov decision process (MDP) ⁷⁰⁹ formulation: the agent waits ($a_t = 0$ for $t < t^{\text{dec}}$) until it reports its prediction, $a_{t^{\text{dec}}} = \pm$, ⁷¹⁰ after which actions are disabled and the prediction is stored in an auxiliary state variable

23

⁷¹¹ used to determine the reward at the end of the trial. A MDP formulation for a general class ⁷¹² of perceptual decision-making tasks, including the tokens and random dots task, is given in ⁷¹³ Methods).

Perfect accuracy in this task is possible if the agent reports at t_{max} since $R = \Theta(N_{t_{\text{max}}}^2) =$ ⁷¹⁵ 1. The task was designed to study reward rate maximizing policies. In particular, the task ⁷¹⁶ has additional structure that allows for controlling what this optimal policy is through the ⁷¹⁷ incentive to decide early, α , incorporated into the trial duration for deciding at time t in the ⁷¹⁸ trial,

$$T(t) = t + (1 - \alpha)(t_{\max} - t) + T_{\text{ITI}}.$$
(9)

⁷¹⁹ Here, a dead time between episodes is added via the inter-trial interval, $T_{\rm ITI}$, to make ⁷²⁰ suboptimal the strategy of predicting randomly at the trial's beginning. We emphasize that ⁷²¹ it is through the trial duration that α serves as a task parameter controlling the strength ⁷²² of the incentive to decide early. When α is fixed, we denote the corresponding optimal ⁷²³ stationary reward rate, ρ_{α} , obtained from the reward rate maximizing policy. This policy ⁷²⁴ shifts from deciding late to deciding early as α is varied from 0 to 1 (*c.f.* fig. S9f,g).

We consider a version of the task where α is variable across two episode types, a slow reference ($\alpha = 1/4$) and fast ($\alpha = 3/4$) type. The agent is aware that the across-trial α dynamics reference (maybe even adversarial), whereas the within-trial random walk dynamics reference (controlled by the positive jump probability, here p = 1/2) can be assumed fixed (see the reference the response of the expression for the expected reward, \bar{r}_t .

Expected trial reward for the tokens task

We derived and used an exact expression for the expected reward in a trial of the tokens task. We derive that expression here as well as a simple approximation. The state sequence task is formulated as a t_{max} -length sequence of random binary variables, $S_{t_{\text{max}}} = (S_1, \ldots, S_{t_{\text{max}}})$, $T_{34} S_t = \pm 1, i = 1, 2, \ldots, t_{\text{max}}$. Consider a simple case in which each is an independent and tast identically distributed Bernoulli sample, $P(s) = p^{\frac{1+s}{2}}(1-p)^{\frac{1-s}{2}}$, for jump probability $p \ge 1/2$. The distribution of $S_{t_{\text{max}}}$ is then

$$P(\boldsymbol{s}_{t_{\max}}) = \prod_{i=1}^{t_{\max}} P(s_i) .$$
(10)

⁷³⁷ We will use this distribution to compute expectations of quantities over this space of trajec-⁷³⁸ tories, namely the sign of $N_t = \sum_{i=1}^t S_i$, for some $0 \le t \le t_{\max}$ and in particular the sign of ⁷³⁹ the final state, $\xi := \operatorname{sgn}(N_{t_{\max}}) \in \{+, -\}$ given $N_t = n$. Note that N_t is even if t is even and ⁷⁴⁰ same with odd values. We remove the case of no sign in $N_{t_{\max}}$ by choosing t_{\max} to be odd. ⁷⁴¹ First, consider predicting $\operatorname{sgn}(N_t)$ with no prior information. The token difference, $-t \le$ ⁷⁴² $N_t \le t$, appears directly in $P(\boldsymbol{s}_{t_{\max}})$. Marginalizing (here just integrating out) the additional ⁷⁴³ degrees of freedom leads to a binomial distribution in the number of S_i for $i \le t$ for which ⁷⁴⁴ $S_i = +1$, $N_t^+ = \sum_{i=1}^t \Theta(s_i) = (t + N_t)/2$,

$$P(N_t^+ = n) = {t \choose n} p^n (1-p)^{t-n} , \qquad (11)$$

24

745 with $n \in \{0, \ldots, t\}$ and $N_t = 2N_t^+ - t$. Thus, the probability that $N_t > 0$, i.e. $N_t^+ > t/2$, is

$$P(N_t > 0) = \sum_{n=0}^{t} {t \choose n} p^n (1-p)^{t-n} \Theta(n-t/2) .$$
(12)

Now consider predicting $\xi = \operatorname{sgn}(N_{t_{\max}})$, given the observation $N_t = n$. Define $t' = t_{\max} - t$ 747 as the remaining time steps to the predicted time and $N_{t'} = \sum_{i=t+1}^{t_{\max}} s_i$, i.e. the total count 748 in the remaining part of the realization. Then the probability of $\xi = +$ conditioned on the 749 state $N_t = n$, denoted $p_{n,t}$, is defined in the same way as $P(N_t > 0)$,

$$p_{n,t}^{+} := P(\xi = +|N_t = n) = \sum_{n'=0}^{t'} {t' \choose n'} p^{n'} (1-p)^{t'-n'} \Theta(n' - (t'-n)/2) .$$
(13)

⁷⁵⁰ where $N_{t'}^+ = n'$ is the number of positive jumps in the remaining $t' = t_{\text{max}} - t$ steps and we ⁷⁵¹ have used $N_{t_{\text{max}}} = N_t + N_{t'} = N_{t'}^+ - (t' - N_t)/2$. The $\Theta(n' - (t' - n)/2)$ factor effectively changes ⁷⁵² the lower bound of the sum to max $\{0, \lceil (t' - n)/2 \rceil\}$, where $\lceil \cdot \rceil$ rounds up. If $\lceil (t' - n)/2 \rceil \le 0$ ⁷⁵³ then $p_{n,t}^+ = 1$ since the sum is over the domain of the distribution, which is normalized. ⁷⁵⁴ Otherwise, the lower bound is $\lceil (t' - n)/2 \rceil$, and the probability of $\xi = +1$ is

$$p_{n,t}^{+} = \sum_{n' = \lceil (t'-n)/2 \rceil}^{t'} {t' \choose n'} p^{n'} (1-p)^{t'-n'} .$$
(14)

⁷⁵⁵ For odd t_{max} , the probability that $\xi = -$ is denoted $p_{n,t}^- = 1 - p_{n,t}^+$. For the symmetric case, ⁷⁵⁶ p = 1/2,

$$p_{n,t}^{+} = \frac{1}{2^{t'}} \sum_{n' = \lceil (t'-n)/2 \rceil}^{t'} {t' \choose n'} , \qquad (15)$$

⁷⁵⁷ when $\lceil (t'-n)/2 \rceil > 0$ and 1 otherwise. This expression is equivalent to equation 5 in [16], ⁷⁵⁸ which was instead expressed using $N_{t'}^-$.

The space of trajectories, i.e. of $\dot{s}_{t_{\text{max}}}$, maps to a space of trajectories for $p_{n,t}^+$ defined on resonant evolving lattice in belief space. The expected reward in this case is,

$$\bar{r}_t := \langle r | N_t = n \rangle = \mathbb{E}\left[\Theta(N_{t_{\max}}N_t) | N_t = n\right]$$
(16)

$$= \max\{p_{n,t}^+, 1 - p_{n,t}^+\}$$
(17)

$$=b_t , \qquad (18)$$

⁷⁶¹ where the belief of correct report $b_t := \max\{p_{n,t}^+, 1 - p_{n,t}^+\}$. The commitment cost $C_t^{\text{com}} = r_{62} r_{\text{max}} - \bar{r}_t$, then also evolves on a lattice (see fig. 3(b)). More generally, $\bar{r}_t = \Delta r b_t + r_{\text{incorrect}}$ ⁷⁶³ for Δr the difference of correct r_{correct} (here 1) and incorrect $r_{\text{incorrect}}$ (here 0) rewards. Since ⁷⁶⁴ $r_{\text{max}} = r_{\text{correct}}$, we have $C_t^{\text{com}} = \Delta r(1 - b_t)$. For p = 1/2 and $\Delta r = 1$, $C_{t=0}^{\text{com}} = 1/2$. ⁷⁶⁵ The shape of $p_{n,t}^+$ is roughly sigmoidal, admitting the approximation,

$$p_{n,t}^+ \approx \frac{1}{1 + \exp\left[-(at+b)n\right]}$$
 (19)

where fitting constants a and b depend on t_{max} . For $t_{\text{max}} = 15$, a = 0.03725 and b = 0.3557. We demonstrate the quality of this approximation in fig. S5. Approximation error is worse to near t_{max} . More than 95% of decisions times in the data we analyzed occur before 12 time steps, where the approximation error in probability is less than 0.05. A similar approximation without time dependence was presented in [16]. We nevertheless used the τ_{11} exact expression eq. (15) in all calculations.

PGD implementation and fitting to relaxation after context switches

⁷⁷³ We identified the times of the context switches in the data and their type (slow-to-fast ⁷⁷⁴ and fast-to-slow). Taking a fixed number of trials before and after each event, we averaged ⁷⁷⁵ the decision times over the events to create two sequences of average decision times around ⁷⁷⁶ context switches (the result is shown in fig. 4a,b). We used a uniformly weighted squared-⁷⁷⁷ error objective, minimized with the standard (Nelder-Mead) simplex routine in python's ⁷⁷⁸ scientific computing library's optimization package.

Survival probabilities over the action policy

779

Behavioural analyses typically focus on response time distributions. From the perspective 780 of reinforcement learning, this is insufficient to fully characterize the behaviour of an agent. 781 782 Instead, the full behaviour is given by the action policy. In this setting, a natural represen-783 tation of the policy is the probability to report as a function of both the decision time and the environmental state (see fig. 5). These are computed from the histograms of $(N_{t^{\text{dec}}}, t^{\text{dec}})$, over trials. However, the histograms themselves do not reflect the preference of the agent to decide at a particular state and time because they are biased by the different frequencies with which the set of trajectories visit each state and time combination. While there are 788 obviously the same number of trajectories at early and late times, they distribute over many 789 more states at later times and so each state at later times is visited less on average than states 790 at earlier times. We can remove this bias by transforming the data ensemble to the ensemble ⁷⁹¹ of two random variables: the state conditioned on time $(N_t|t)$, and the event that $t = t^{\text{dec}}$. ⁷⁹² Conditioning this ensemble on the state gives $P(t = t^{\text{dec}} | N_t, t) = p(N_t, t = t^{\text{dec}} | t) / p(N_t | t)$. To reduce estimator variance, we focus on the corresponding survival function, $P(t < t^{\text{dec}} | N_t, t)$. ⁷⁹⁴ So, $P(t < t^{\text{dec}}|N_t, t) = 1$ when t = 0 and decays to 0 as t and $|N_t|$ increase. Unlike the ⁷⁹⁵ unconditioned histograms, these survival probabilities vary much more smoothly over state ⁷⁹⁶ and time. This justifies the use of the interpolated representations displayed in fig. 5b-e. ⁷⁹⁷ Note that to simplify the analysis, we have binned decision times by the 200 ms time step ⁷⁹⁸ between token jumps. This is justified by the small deviations from uniformity of decision times modulo the time step shown in fig. S11. 799

⁸⁰⁰ Episodic decision-making and dynamic programming solutions of value iteration

We generalize the mathematical notation and description of an existing AR-RL formulation and dynamic programming solution of the random dots task [7], a binary perceptual evidence accumulation task extensively studied in neuroscience. To align notation with convention in reinforcement learning theory, exceptionally here s denotes the belief state variable, i.e. a representation of the task state sufficient to make the decision (e.g. the tokens difference, N_t , in the case of the tokens task). We connect this extended formulation to account for a dynamic deliberation cost. We write it in discrete time, though the continuous time version is equally tractable.

The problem is defined by a recursive optimality equation for the value function V(s|t)⁸¹⁰ in which the highest of the action values, Q(s, a|t), is selected. We formalize the non-⁸¹¹ stationarity within episodes by conditioning on the trial time, t, where t = 0 is the trial start ⁸¹² time. Q(s, a|t) is the action-value function of average-reward reinforcement learning [11], i.e.

26

⁸¹³ the expected sum of future reward deviations from the average when selecting action a when ⁸¹⁴ in state s, at possible decision time t within a trial, and then following a given action policy ⁸¹⁵ π thereafter. The action set for these binary decision tasks consists of *report left* (-), *report* ⁸¹⁶ *right* (+), and *wait*. When *wait* is selected, time increments and beliefs are updated with ⁸¹⁷ new evidence. We use a decision-time conditioned, expected trial reward function, r(s, a|t)⁸¹⁸ with $a = \pm$, that denotes the reward expected to be received at the end of the trial after ⁸¹⁹ having reported \pm in state s at time t during the trial. Note that r(s, a|t) can be defined ⁸²⁰ in terms of a conventional reward function r(s, a) if the reported action, decision time, and ⁸²¹ current time are stored as an auxiliary state variable so they can be used to determine the ⁸²² non-zero reward entries at the end of the trial.

The average-reward formulation of Q(s, a|t) naturally narrows the problem onto determining decisions within only a single episode of the task. To see this, we pull out the contribution of the current trial,

$$Q(s,a|t) = \mathbb{E}^{\pi} \left[\sum_{t'=t}^{T} R_t - \rho \, \left| S_t = s, A_t = a \right] + V(s|T+1) \right]$$
(20)

where T is the (possibly stochastic) trial end time and V(s|T+1) is the state value at the start of the following trial, which does not depend on s_t and a_t for independently sampled trials. Following conventional reinforcement learning notation, the expectation \mathbb{E}^{π} is over all randomness conditioned on following the policy, π , which itself could be stochastic [11]. When trials are identically and independently sampled, the state at the trial start is the same for all trials and denoted s_0 with value V_0 . Thus, the value at the start of the trial $V(s|t=0) = V(s|T+1) = V_0$ equals that at the start of the next trial and so, by construction, the expected trial return (total trial rewards minus trial costs) must vanish (we will show this explicitly below). Note that the value shift invariance of eq. (20) can be fixed so that $V_0 = 0$.

The optimality equation for V(s|t) arises from a greedy action policy over Q(s, a|t): it selects the action of the largest of Q(s, -|t), Q(s, +|t), and Q(s, wait|t). The value expression for the wait-action is incremental, and so depends on the value at the next time step. In selects the action for the two reporting actions integrates over the remainder of the trial selection is made and so depends on the value at the start of the following trial. The resulting optimality equation for the value function V(s|t) is then

$$V(s|t) = \max_{a} Q(s, a|t) ,$$

$$Q(s, \pm|t) = r(s, \pm|t) - \sum_{t'=t+1}^{T} c_{t'} + V(s|t = T+1) ,$$

$$Q(s, wait|t) = -c_t + \mathbb{E}_{s_{t+1}|s} \left[V(s_{t+1}|t+1) \right] ,$$

$$V(s|t = 0) = V(s|t = T+1) .$$
(21)

⁸⁴² Here, $t = 0, 1, \ldots, t_{\text{max}}$ within the current trial and $t = T + 1, T + 2 \ldots$ in the following ⁸⁴³ trial, with t_{max} the latest possible decision time in a trial, and T = T(t) the decision-time ⁸⁴⁴ dependent trial duration. For inter-trial interval T_{ITI} , T satisfies $T_{\text{ITI}} \leq T \leq t_{\text{max}} + T_{\text{ITI}}$. ⁸⁴⁵ c_t is the cost rate at time t. The second term in Q(s, wait|t) uses the notation $\mathbb{E}_{x|y}[z]$, i.e. ⁸⁴⁶ the expectation of z with respect to p(x|y). The last line in eq. (21) is the self-consistency ⁸⁴⁷ criterion imposed by the AR-RL formulation, which demands that the expected value at

27

the beginning of the trial be the expected value at the beginning of the following trial. The greedy policy then gives a single decision time for each state trajectory as the first time when Q(s, -|t) > Q(s, wait|t) or Q(s, +|t) > Q(s, wait|t), with the reporting action determined by which of Q(s, -|t) and Q(s, +|t) is larger. For given c_t , dynamic programming provides a solution to eq. (21) [7] by recursively solving for V(s|t) by back-iterating in time from the est at $t = t_{\text{max}}$ is set by the best of the two reporting (±) actions, which do not have a recursive dependence on the value and so can seed the recursion.

We now interpret this general formulation in terms of opportunity costs. For the choice ⁸⁵⁷ of a static opportunity cost rate of time, $c_t = \rho$. This is the AR-RL case. As in [7], a ⁸⁵⁸ constant auxiliary deliberation cost rate, c, incurred only up to decision time can be added, ⁸⁵⁹ $c_t = \rho + c\Theta(t^{dec} - t)$. Of course, ρ is unknown *a priori*. For this solution method, its value ⁸⁶⁰ can be found by exploiting the self-consistency constraint, V(s|t=0) = V(s|t=T+1). This ⁸⁶¹ dependence can be seen formally by taking the action value eq. (20), choosing *a* according ⁸⁶² to π to obtain the state value, V(s|t), and evaluating it for t = 0,

$$V(s|t=0) = \mathbb{E}_{t^{\text{dec}}}\left[\sum_{t=0}^{T} R_t - \rho\right] + V(s|t=T+1)$$
(22)

$$= \mathbb{E}_{t^{\text{dec}}} \left[r(t^{\text{dec}}) - \rho T(t^{\text{dec}}) \right] + V(s|t = T + 1)$$
(23)

$$= \bar{R} - \rho \bar{T} + V(s|t = T + 1) .$$
(24)

⁸⁶³ Here, $\bar{R} = \mathbb{E}_{t^{\text{dec}}} \left[r(t^{\text{dec}}) \right]$ and $\bar{T} = \mathbb{E}_{t^{\text{dec}}} \left[T(t^{\text{dec}}) \right]$ denotes the expectations over the trial en-⁸⁶⁴ semble that, when given the state sequence, transforms to an average over t^{dec} , the trial deci-⁸⁶⁵ sion time, defined as when V(s|t) achieves its maximum on the state sequence, $(s_0, \ldots, s_{t_{\max}})$. ⁸⁶⁶ The expected trial reward function, $r(t) := \max_{a \in \{-,+\}} r(s, a|t)$ is the expected trial reward ⁸⁶⁷ for deciding at t. Imposing the self-consistency constraint on eq. (24) recovers the definition ⁸⁶⁸ $\rho = \bar{R}/\bar{T}$.

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Asymmetric switching cost model

Here, we present the model component that accounts for the asymmetric relaxation timescales after context switches. The basic assumption is that tracking a signal at a higher temporal resolution should be more cognitively costly, so that adapting from faster to slower ervironments should happen more quickly than the reverse, so as to not pay this cost unare necessarily. We now develop this idea formally (see fig. S4).

Let T_{track} and T_{sys} be the timescale of tracking and of the tracked system, respectively. ⁸⁷⁵ One way to interpret the mismatch ratio, $T_{\text{sys}}/T_{\text{track}}$, is via an attentional cost rate, q. ⁸⁷⁷ This rate should decay with T_{track} : the slower the timescale of tracking, the lower the ⁸⁷⁸ cognitive cost. For simplicity, we set $q = 1/T_{\text{track}}$ (fig. S4a). Integrating this cost rate over a ⁸⁷⁹ characteristic time of the system is then the tracking cost, $Q = qT_{\text{sys}} = T_{\text{sys}}/T_{\text{track}}$, which is ⁸⁸⁰ also the mismatch ratio. We propose that Q enters the algorithm via a scale factor on the ⁸⁸¹ integration time of the reward filter for $\hat{\rho}_k^{\tau_{\text{context}}}$, τ_{context} . We redefine τ_{context} as

$$\tau_{\rm context} \leftarrow \frac{\tau_{\rm context}}{1+Q^{\nu}} ,$$
(25)

where ν is a sensitivity parameter that captures the strength of the nonlinear sensitivity of the speed up (for $\nu > 1$) or slow down (for $\nu < 1$) in adaptation with the tracking cost,

⁸⁸⁴ Q (fig. S4a shows how this timescale varies over Q for three values of ν). A natural choice ⁸⁸⁵ for T_{sys} is T_k , the trial duration. For T_{track} , we introduce the filtered estimate of the trial ⁸⁸⁶ duration, $\hat{T}_k^{\tau_{\text{context}}}$ (computed using the same simple low-pass filter *c.f.* eq. (8)). Thus, the ⁸⁸⁷ tracking timescale adapts to the system timescale. As a result of how τ_{context} is lowered by Q⁸⁸⁸ for $\nu > 1$, this adaptation is faster in the fast-to-slow transition relative to the slow-to-fast ⁸⁸⁹ transition.

Prediction for asymmetric rewards

Given a payoff matrix, $\mathbf{R} = (r_{s,a})$, where $r_{s,a}$ is the reward for reporting $a \in \{-, +\}$ in the trial realization leading to s, here the sign of $N_{t_{\max}}$, and the probability that the rightward choice is correct, $p_{n,t}^+$, the expected reward for the two reporting actions in a trial is given by the matrix equation

$$\left[\langle r|a = +, n, t \rangle \ \langle r|a = -, n, t \rangle \right] = \left[p_{n,t}^+ \ 1 - p_{n,t}^+ \right] \left[\begin{matrix} r_{++} & r_{+-} \\ r_{-+} & r_{--} \end{matrix} \right]$$

⁸⁹⁵ Here, the corresponding reported choice is $a^* = \operatorname{argmax}_{a \in \{-,+\}} \langle r | a, n, t \rangle$. In this paper and ⁸⁹⁶ in all existing tokens tasks, \mathbf{R} was the identity matrix. In this case, and for all cases where ⁸⁹⁷ \mathbf{R} is a symmetric matrix, $\mathbf{R} = \mathbf{R}^{\top}$, an equivalent decision rule is to decide based on the sign ⁸⁹⁸ of N_t . When \mathbf{R} is not symmetric, however, this is no longer a valid substitute. Asymmetry ⁸⁹⁹ can be introduced through the actions and the states.

Using an additional parameter γ , we introduce asymmetry via a bias for + actions that ⁹⁰⁰ leaves the total reward unchanged by replacing the payoff matrix with

$$\boldsymbol{R}_{\text{asym}} = \begin{bmatrix} r_{++}(1+\gamma) & r_{+-}(1-\gamma) \\ r_{-+}(1+\gamma) & r_{--}(1-\gamma) \end{bmatrix}$$

⁹⁰² The result for $\gamma = -0.6, 0$, and 0.6 is shown in fig. S10. For $\gamma > 0$ the decision boundary for ⁹⁰³ a = + shifts up proportional to γ . For $\gamma < 0$ the decision boundary for a = - shifts down ⁹⁰⁴ proportional to $-\gamma$. The explanation is that the components are set and exchange where ⁹⁰⁵ the decision is exchanged, $N_t = 0$ for the symmetric case. This changes to $N_t \propto \pm \gamma$ for the ⁹⁰⁶ asymmetric $\gamma \neq 0$ case.

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Comparing reward rates and slopes of urgency

Reference [17] parametrize urgency with the saturation value, u_{∞} , and the half-maximum, $\tau_{1/2}$. The initial slope is given by their ratio. We used the context-conditioned values published in Table 1 in [17] for the n = 70 (no 90° control) dataset. The context-conditioned τ_{11} reward rates, ρ_{α} , are computed as the accuracy $\langle R \rangle_{|\alpha}$ divided by the average trial time, $\langle T \rangle_{|\alpha}$ τ_{12} for choice number $\alpha \in \{2, 4\}$ as context. We computed $\langle R \rangle_{|\alpha=2} = 0.71$ and $\langle R \rangle_{|\alpha=4} = 0.49$. τ_{12} for choice number $\alpha \in \{2, 4\}$ as context. We computed $\langle R \rangle_{|\alpha=2} = 0.71$ and $\langle R \rangle_{|\alpha=4} = 0.49$. τ_{12} for the trial time is the sum of the response time, the added time penalty if incorrect, and the τ_{14} inter-trial interval. We computed the response times $t_{response,\alpha=2} = 0.527$ and $t_{response,\alpha=4} = 0.725$. While the dataset contains the response times, it does not have the latter two. The τ_{14} time penalty was on the order of 1 second, as was the time penalty [61]. Under those τ_{17} estimates, the reward rates are $\rho_{\alpha=2} = 0.40$ and $\rho_{\alpha=4} = 0.22$. The ratio between slopes is τ_{18} and the ratio of reward rates was 2.3 giving an error of about 20%.

919

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- ⁹²⁶ [1] David I Green, "Pain-Cost and Opportunity-Cost," The Quarterly Journal of Economics 8,
 ⁹²⁷ 218–229 (1894).
- Vektor Dewanto, George Dunn, Ali Eshragh, Marcus Gallagher, and Fred Roosta, "Average reward model-free reinforcement learning: a systematic review and literature mapping,"
 arXiv:2010.08920 [cs.LG].
- [3] Nathaniel D Daw and David S Touretzky, "Long-term reward prediction in TD models of the
 dopamine system," Neural computation 14, 2567–2583 (2002).
- [4] Lindsay E Hunter and Nathaniel D Daw, "Context-sensitive valuation and learning," Current
 Opinion in Behavioral Sciences 41, 122–127 (2021).
- [5] Nils Kolling and Thomas Akam, "(Reinforcement?) Learning to forage optimally," Current
 Opinion in Neurobiology 46, 162–169 (2017).
- ⁹³⁷ [6] Yael Niv, Nathaniel D Daw, and Peter Dayan, "How fast to work : Response vigor, motivation
 ⁹³⁸ and tonic dopamine," in *Neural Information Processing Systems* (2005).
- [7] Jan Drugowitsch, Rubén Moreno-Bote, Anne K Churchland, Michael N Shadlen, and Alexandre Pouget, "The Cost of Accumulating Evidence in Perceptual Decision Making," The Journal
 of Neuroscience 32, 3612 LP 3628 (2012).
- [8] A Ross Otto and Nathaniel D Daw, "The opportunity cost of time modulates cognitive effort,"
 Neuropsychologia 123, 92–105 (2019).
- [9] A Ross Otto and Eliana Vassena, "It's all relative: Reward-induced cognitive control modulation depends on context." Journal of Experimental Psychology: General 150, 306–313 (2021).
- Germain Lefebvre, Aurélien Nioche, Sacha Bourgeois-gironde, and Stefano Palminteri, "Contrasting temporal difference and opportunity cost reinforcement learning in an empirical money-emergence paradigm," Proceedings of the National Academy of Sciences 115, E11446
 LP – E11454 (2018).
- ⁹⁵¹ [11] Richard S Sutton and Andrew G Barto, *Reinforcement learning: An introduction, 2nd ed.*,
 ⁹⁵² Adaptive computation and machine learning. (The MIT Press, Cambridge, MA, US, 2018)
 ⁹⁵³ pp. xxii, 526-xxii, 526.
- ⁹⁵⁴ [12] Khimya Khetarpal, Matthew Riemer, Irina Rish, and Doina Precup, "Towards Continual
 ⁹⁵⁵ Reinforcement Learning: A Review and Perspectives," arXiv:2012.13490 [cs.LG].
- 956 [13] Roger Ratcliff, "A theory of memory retrieval." Psychological Review 85, 59–108 (1978).
- ⁹⁵⁷ [14] Gaurav Malhotra, David S Leslie, Casimir J H Ludwig, and Rafal Bogacz, "Time-varying
 ⁹⁵⁸ decision boundaries : insights from optimality analysis," Psychon Bull Rev 25, 971–996 (2018).
- Jochen Ditterich, "Evidence for time-variant decision making," European Journal of Neuro science 24, 3628–3641 (2006).

30

- [16] Paul Cisek, Geneviève Aude Puskas, and Stephany El-Murr, "Decisions in Changing Con-961 ditions: The Urgency-Gating Model," The Journal of Neuroscience 29, 11560 LP – 11571 962 (2009).
- [17] Anne K Churchland, Roozbeh Kiani, and Michael N Shadlen, "Decision-making with multiple 964 alternatives," Nature Neuroscience 11, 693-702 (2008). 965
- [18] David Thura and Paul Cisek, "Deliberation and Commitment in the Premotor and Primary 966 Motor Cortex during Dynamic Decision Making," Neuron 81, 1401–1416 (2014). 967
- [19] David Thura, Ignasi Cos, Jessica Trung, and Paul Cisek, "Context-Dependent Urgency Influ-968
- ences Speed–Accuracy Trade-Offs in Decision-Making and Movement Execution," The Journal 969 of Neuroscience **34**, 16442 LP – 16454 (2014). 970
- [20] David Thura, Jean-François Cabana, Albert Feghaly, and Paul Cisek, "Unified neural dy-971 namics of decisions and actions in the cerebral cortex and basal ganglia," bioRxiv (2020), 972 10.1101/2020.10.22.350280.973
- [21] David Thura and Paul Cisek, "The Basal Ganglia Do Not Select Reach Targets but Control 974 the Urgency of Commitment," Neuron 95, 1160–1170.e5 (2017). 975
- [22] Peter Janssen and Michael N Shadlen, "A representation of the hazard rate of elapsed time 976 in macaque area LIP," Nature Neuroscience 8, 234–241 (2005). 977
- Satohiro Tajima, Jan Drugowitsch, and Alexandre Pouget, "Optimal policy for value-based [23]978 decision-making," Nature Communications 7, 12400 (2016). 979
- [24] Anton Schwartz, "A Reinforcement Learning Method for Maximizing Undiscounted Rewards," 980 in International Conference on Machine Learning, Vol. 0 (1993). 981
- Yael Niv, Nathaniel D Daw, Daphna Joel, and Peter Dayan, "Tonic dopamine: opportunity |25|982 costs and the control of response vigor," Psychopharmacology 191, 507–520 (2007). 983
- Sara M Constantino and Nathaniel D Daw, "Learning the opportunity cost of time in a patch-984 [26]foraging task," Cogn Affect Behav Neurosci. 15, 837 (2015). 985
- [27] Benjamin Y Hayden and Yael Niv, "The case against economic values in the orbitofrontal 986 cortex (or anywhere else in the brain)," PsyArXiv 10.31234/osf.io/7hgup. 987
- Nathaniel D Daw, "Advanced Reinforcement Learning," in *Neuroeconomics*, edited by Paul W [28]988 Glimcher and Ernst B T Neuroeconomics (Second Edition) Fehr (Academic Press, San Diego, 989
- 2014) 2nd ed., Chap. 16, pp. 299–320. 990

- These were primarily as as result of the experimenter adapting to fluctuations in motivation [29]991
- of the subject. D. Thura. Personal communication. 992
- [30] Edward Vul, Noah Goodman, Thomas L Griffiths, and Joshua B Tenenbaum, "One and 993 Done? Optimal Decisions From Very Few Samples," Cognitive Science 38, 599-637 (2014). 994
- [31] Single subject behavioural data shared by Thomas Thierry. 995
- [32] Surva Ganguli, James W Bisley, Jamie D Roitman, Michael N Shadlen, Michael E Goldberg, 996
- and Kenneth D Miller, "One-Dimensional Dynamics of Attention and Decision Making in 997 LIP," Neuron 58, 15–25 (2008). 998
- [33] David Thura and Paul Cisek, "Modulation of Premotor and Primary Motor Cortical Activity 999 during Volitional Adjustments of Speed-Accuracy Trade-Offs," The Journal of Neuroscience 1000 **36**, 938 – 956 (2016). 1001
- [34]Kiyohito Iigaya, Yashar Ahmadian, Leo P Sugrue, Greg S Corrado, Yonatan Loewenstein, 1002 William T Newsome, and Stefano Fusi, "Deviation from the matching law reflects an optimal 1003
- strategy involving learning over multiple timescales," Nature Communications 10, 1466 (2019). 1004
- Long Ding and Joshua I. Gold, "The Basal Ganglia's Contributions to Perceptual Decision [35]1005
- Making," Neuron **79**, 640–649 (2013). 1006

- 31
- 1007 [36] Kong-Fatt Wong and Xiao-Jing Wang, "A Recurrent Network Mechanism of Time Integration
 in Perceptual Decisions," The Journal of Neuroscience 26, 1314 1328 (2006).
- 1009 [37] Alex Roxin and Anders Ledberg, "Neurobiological Models of Two-Choice Decision Making
- Can Be Reduced to a One-Dimensional Nonlinear Diffusion Equation," PLOS Computational
 Biology 4, e1000046 (2008).
- 1012 [38] David Meder, Nils Kolling, Lennart Verhagen, Marco K Wittmann, Jacqueline Scholl, Kristof-
- ¹⁰¹³ fer H Madsen, Oliver J Hulme, Timothy E J Behrens, and Matthew F S Rushworth, "Simul-¹⁰¹⁴ taneous representation of a spectrum of dynamically changing value estimates during decision
- ¹⁰¹⁵ making," Nature Communications 8 (2017), 10.1038/s41467-017-02169-w.
- ¹⁰¹⁶ [39] "Predictive Representations in Hippocampal and Prefrontal Hierarchies,".
- ¹⁰¹⁷ [40] Jan Zimmermann, Paul W Glimcher, and Kenway Louie, "Multiple timescales of normalized ¹⁰¹⁸ value coding underlie adaptive choice behavior," Nature Communications **9**, 3206 (2018).
- 1019 [41] HyungGoo R Kim, Athar N Malik, John G Mikhael, Pol Bech, Iku Tsutsui-Kimura, Fangmiao
- Sun, Yajun Zhang, Yulong Li, Mitsuko Watabe-Uchida, Samuel J Gershman, and Naoshige
 Uchida, "A Unified Framework for Dopamine Signals across Timescales," Cell 183, 1600–
 1616.e25 (2020).
- Paul Masset, Athar N. Malik, HyungGoo R. Kim, Pol Bech, and Naoshige Uchida, "A diversity of discounting horizons explains ramping diversity in dopaminergic neurons," in COSYNE Abstracts (2021).
- ¹⁰²⁶ [43] Angela J Langdon and Nathaniel D Daw, "Beyond the Average View of Dopamine," Trends ¹⁰²⁷ in Cognitive Sciences **24**, 499–501 (2020).
- ¹⁰²⁸ [44] John G Mikhael and Samuel J Gershman, "Adapting the flow of time with dopamine," Journal ¹⁰²⁹ of Neurophysiology **121**, 1748–1760 (2019).
- ¹⁰³⁰ [45] Ido Toren, Kristoffer C Aberg, and Rony Paz, "Prediction errors bidirectionally bias time ¹⁰³¹ perception," Nature Neuroscience **23**, 1198–1202 (2020).
- ¹⁰³² [46] Lars Hunger, X Arvind Kumar, and X Robert Schmidt, "Abundance Compensates Kinet¹⁰³³ ics : Similar Effect of Dopamine Signals on D1 and D2 Receptor Populations," Journal of
 ¹⁰³⁴ Neuroscience 40, 2868–2881.
- ¹⁰³⁵ [47] Julia Cox and Ilana B Witten, "Striatal circuits for reward learning and decision-making,"
 ¹⁰³⁶ Nature Reviews Neuroscience 20, 482–494 (2019).
- 1037 [48] Helen N Schwerdt, Hideki Shimazu, Ken-ichi Amemori, Satoko Amemori, Patrick L Tierney,
- Daniel J Gibson, Simon Hong, Tomoko Yoshida, Robert Langer, Michael J Cima, and Ann M
 Graybiel, "Long-term dopamine neurochemical monitoring in primates," Proceedings of the
 National Academy of Sciences 114, 13260 LP 13265 (2017).
- 1041 [49] Tommaso Patriarchi, Jounhong Ryan Cho, Katharina Merten, Mark W Howe, Aaron Marley,
- ¹⁰⁴² Wei-hong Xiong, Robert W Folk, Gerard Joey Broussard, Ruqiang Liang, Min Jee Jang,
- Haining Zhong, Daniel Dombeck, Mark Von Zastrow, Axel Nimmerjahn, Viviana Gradinaru,
 John T Williams, and Lin Tian, "Ultrafast neuronal imaging of dopamine dynamics with
- designed genetically encoded sensors," Science **4422** (2018), 10.1126/science.aat4422.
- 1046 [50] Matthew A Carland, David Thura, and Paul Cisek, "The Urge to Decide and Act: Implica-
- tions for Brain Function and Dysfunction," The Neuroscientist **25**, 491–511 (2019).
- ¹⁰⁴⁸ [51] Samuel J Gershman and Naoshige Uchida, "Believing in dopamine," Nature Reviews Neuro-¹⁰⁴⁹ science **20**, 703–714 (2019).
- 1050 [52] Andrew Westbrook and Todd S Braver, "Dopamine Does Double Duty in Motivating Cognitive
- 1051 Effort," Neuron **91**, 708 (2016).

- 1052 [53] Pablo Tano, Peter Dayan, and Alexandre Pouget, "A Local Temporal Difference Code for
- 1053 Distributional Reinforcement Learning," in Advances in Neural Information Processing Sys-
- *tems*, Vol. 33, edited by H Larochelle, M Ranzato, R Hadsell, M F Balcan, and H Lin (Curran
 Associates, Inc., 2020) pp. 13662–13673.
- ¹⁰⁵⁶ [54] William Fedus, Carles Gelada, Yoshua Bengio, Marc G Bellemare, and Hugo Larochelle, ¹⁰⁵⁷ "Hyperbolic Discounting and Learning over Multiple Horizons," arXiv:1902.06865 [stat.ML].
- ¹⁰⁵⁸ [55] I Momennejad, E M Russek, J H Cheong, M M Botvinick, N D Daw, and S J Gershman,
- ¹⁰⁵⁹ "The successor representation in human reinforcement learning," Nature Human Behaviour 1, 680–692 (2017).
- 1061 [56] Personal communication, Thomas Thierry.
- ¹⁰⁶² [57] Stefano Palminteri and Maël Lebreton, "Context-dependent outcome encoding in human re-¹⁰⁶³ inforcement learning," Current Opinion in Behavioral Sciences **41**, 144–151 (2021).
- ¹⁰⁶⁴ [58] Ernest S Davis and Gary F Marcus, "Computational limits don't fully explain human cognitive
 ¹⁰⁶⁵ limitations," Behavioral and Brain Sciences 43, e7 (2020).
- $\begin{bmatrix} 1005 \\ 1005 \end{bmatrix} = \begin{bmatrix} 1001 \\ 1$
- ¹⁰⁶⁶ [59] Arman Abrahamyan, Laura Luz Silva, Steven C Dakin, Matteo Carandini, and Justin L Gard ¹⁰⁶⁷ ner, "Adaptable history biases in human perceptual decisions," Proceedings of the National
 ¹⁰⁶⁷ A human f G is a state LD = E2557 (2016)
- 1068 Academy of Sciences **113**, E3548 LP E3557 (2016).
- ¹⁰⁶⁹ [60] Zhijian Wang, Bin Xu, and Hai-Jun Zhou, "Social cycling and conditional responses in the ¹⁰⁷⁰ Rock-Paper-Scissors game," Scientific Reports 4, 5830 (2014).
- 1071 [61] A. Churchland. Personal communication.



Figure S1. Reward filtering scheme for online computation of within-trial opportunity cost. With t denoting absolute time, the reward sequence, R_t , is integrated on both a stationary (τ_{long}) and context (τ_{context}) filtering timescale to produce estimates of the stationary and context-specific reward rates, respectively. These are large and small, respectively, relative to the average context switching timescale, T_{block} . The estimate of the context-specific offset, o_t is computed by time-integrating the difference of these two estimates. In this filtering, when a trial terminates, the effective operation is that C_t^{del} is set to o_t , and the latter is zeroed. Thus, the opportunity cost starts at this offset and then integrates ρ_{long} , $C_{t,k}^{\text{del}} = o_{T_{k-1},k-1} + \rho_{\text{long},k-1}t$, where $o_{T_{k-1},k-1} = (\rho_{\text{context},k-1} - \rho_{\text{long},k-1})T_{k-1}$. Notes on the computational graph: Arrows pass the value at each time step (dashed arrows only pass the value when a trial terminates). Links annotated with '-' multiply the passed quantity by -1.

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Figure S2. *PGD agent plays the tokens task with periodic* α -dynamics. (a) Trials are grouped into alternating trial blocks of constant α (fast (orange) and slow (blue) conditions). (b) Here, trial block durations are constant over the experiment. (c) Decision times over the trials from (a) distribute widely, but relax after context switches. (d) Block-averaged decision times remain stationary. Inset shows the context-conditioned trial-averaged reward $\langle R_k \rangle$ and trial duration $\langle T_k \rangle$ (orange and blue dots; black is unconditioned average; $\langle \cdot \rangle$ denotes the trial ensemble average). Lines pass through the origin (slope given by the respective reward rate). (e) Distribution of estimates have lower variance than the trial reward rates, ρ^{trial} (gray). The conditioned averages of $\hat{\rho}_k^{\tau_{\text{context}}}$ shown as blue and orange. (f) The relative error in estimating ρ , $E_t = \frac{1}{t} \sum_k^t |\hat{\rho}_k^{\tau_{\text{long}}} - \rho|/\rho$, for $\tau_{\text{long}} = 10^3(\text{circle})$, $10^4(\text{square})$, $10^5(\text{triangle})$. Inset shows that $E_{T_{\text{exp}}} \propto (\tau_{\text{long}}/T_{\text{block}})^{-1}$ over a grid of τ_{long} and T_{block} as expected (black line).



Figure S3. Comparison of PGD and NHP in non-stationary α dynamics from [19]: Subject 2. Same as fig. 5.



Figure S4. Asymmetric switching cost model. (a) Attentional cost rate, q, is set to be inversely proportional to tracking timescale, T_{track} . (b) Filtering timescale τ_{context} is scaled down with tracking cost, $Q = T_{\text{sys}}/T_{\text{track}}$ from a base timescale, here denoted τ_0 (shown for three values of sensitivity $\nu = 2, 4, 8$).



Figure S5. sigmoidal approximation to expected reward. (a) the approximation explained in Methods: State-conditioned expected trial reward, for different dec,[p]ision times. (b) The error in the approximation for different decision times.

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Figure S6. Model validation on behavioural statistics from [19]. (a,b) Running average (last 1000 trial) of trial reward rate ρ_k^{trial} . (c,d) Histograms of trial reward rate, ρ_k^{trial} (c) and trial duration, T_k (d). (e) Auto-correlation function of trial duration. (f) Data vs. model decision time (gray-scale is count; white dashed line is perfect correlation; actual Pearson correlation is shown)

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Figure S7. Comparison of trial-aware and trial-unaware results. (a,b) 1/2-Survival probability contours for subject 1 (dashed), trial-aware PGD (blue), and trial-unaware PGD (red) for slow (a) and fast (b) context-conditioned data. (c) Opportunity cost for trial-unaware PGD (compare with fig. 2b). Opportunity cost range adjusted here such that data within standard error of trial-unaware PGD model prediction for slow block (blue).

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Figure S8. Comparison of PGD and AR-RL learning on a patch leaving task. Performance is defined as relative regret rate, $(\hat{\rho} - \rho^*)/\rho^*$ (PGD (dots); AR-RL (lines)). (a) Performance over different sizes of the state vector (d = 100 (blue), 200 (orange), 300 (green)). (b) Performance over different learning rates (parametrized by integration time constant, $\tau = 1 \times 10^4$ (blue), 2×10^4 (orange), 3×10^4 (green)). (parameters: $\lambda = 1/5$; r_{max} sampled uniformily on [0,1]). A random state label permutation is made at the time indicated by the black arrow. Values were initialized at -1.



Figure S9. Reward rate optimal strategies in (α, c) plane. (a) The reward-rate maximizing policy interpolates from the wait-for-certainty strategy at weak incentive (low α) and low deliberation cost (low c), to the one-and-done strategy at strong incentive (high α) and high deliberation cost (high c). Dashed lines bound a transition regime between the two extreme strategies. Red line denotes where they have equal performance. (b-e) Slices of the (α, c) -plane. Shown are the reward rate as a function of α (b,c) and c (d,e) (wait-for-certainty strategy is shown in blue; one-and-done strategy is shown in orange). N is the magnitude of the token difference



Figure S10. Asymmetric action rewards skew survival probability. Here, we plot the half-maximum of the PGD survival probability for three values of the action reward bias, $\gamma = -0.6, 0, 0.6$ (blue, black and orange, respectively). Other model parameters same as in fitted model.



Figure S11. Decision times relative to token jumps. Here, we plot the histograms of decision times using their position between token jumps, the step fraction. The data is separated by α and monkey.