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Reminders of past choices bias decisions for reward in humans

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We provide evidence that decisions are made by consulting memories for individual past experiences, and that this process can be biased in favour of past choices using incidental reminders. First, in a standard rewarded choice task, we show that a model that estimates value at decision-time using individual samples of past outcomes fits choices and decision-related neural activity better than a canonical incremental learning model. In a second experiment, we bias this sampling process by incidentally reminding participants of individual past decisions. The next decision after a reminder shows a strong influence of the action taken and value received on the reminded trial. These results provide new empirical support for a decision architecture that relies on samples of individual past choice episodes rather than incrementally averaged rewards in evaluating options and has suggestive implications for the underlying cognitive and neural mechanisms.

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ow do we use experience to guide our choices? One approach is to use a simple heuristic: do what has worked best recently. This idea is captured in prominent reinforcement learning (RL) models, which build a running average of the rewards received for each action taken, updating this average incrementally as new rewards are experienced. Such averages can be updated by error-driven learning, tying this approach to observations of reward prediction errors (RPEs) in the brain^{1,2}. This approach is effective and efficient in the sorts of tasks typically used to study reward learning, which usually involve many repeated choices among a small number of options whose only salient difference is how much or how often they are rewarded²⁻⁵. A running average falls short, however, in more naturalistic settings, where experience is sparse and relevant evidence might concern only a single, or a small number, of individual experiences⁶⁻¹².

Here we provide evidence in favour of a more flexible approach, in which choosers draw on memories for individual instances of relevant previous choices and use them to predict how the current decision might turn out. For example, when deciding where to eat dinner, I might recall a particular evening dining at a particular restaurant and make my decision based on how much I enjoyed the meal I had that night. Making decisions this way allows us to pick and choose the most relevant of previous experiences, rather than relying on static summary representations. The latter would be useless in considering a brand new restaurant, whereas a richer memory representation of each past experience could be used to develop an informed guess, by generalizing from experiences at similar establishments. Along these lines, Plonsky et al. 13 showed that a similarity-based approach to selecting past experiences capture a complex pattern of dependencies of choice behaviour on rare, but impactful, past experiences, leading to near-optimal performance in a wide range of dynamic choice tasks. In this way, evaluating options by sampling the most similar previous experiences resembles non-parametric and kernel-based methods long employed in statistics and machine learning, which draw on individual samples of raw experience 14,15. The effectiveness of these estimation algorithms reinforces recent interest in the idea that humans and animals draw on memory to flexibly compute decision variables at choice time^{11,16-22} rather than relying on precomputed averages. Indeed, recent work has shown that associations formed during even a single experience can affect later choice behaviour^{23,24}.

We present results from two experiments in support of this hypothesis. First, we develop and test a computational model of sequential choice that estimates values by selecting and evaluating individual memories of past trials. We show that this sampling model is a better fit to trial-by-trial choice sequences than is a standard incremental learning algorithm, even in the sort of repeated choice task previously studied using RL models. In a second study, we directly test the prediction that value estimation draws on memories of individual trials, by demonstrating a causal effect on choices of specific, selected experiences, brought to mind by refreshing memories of individual past decision trials.

Results

Experiment 1. We first formally investigated whether a sampling model could provide a better trial-by-trial fit to human choices than standard incremental learning. To test this, we re-analysed choice and neuroimaging data from a previously published learning study⁴. Twenty participants (14 presented with neuroimaging data previously and 6 additional behaviour-only participants included here) performed a series of choices between four virtual slot machines with time-varying payoffs (Fig. 1a,b).

Here we compared a standard incremental learning rule (of the form tested in the original study) with a sampling model that evaluates each option by retrieving the rewards received at individual past choices of that option, stochastically sampled with probabilities given by their temporal recency (see Methods). These sampled rewards are averaged to compute a net value for each option.

In the limit, when a potentially infinite number of samples is considered, these sample-based averages would be identical to those computed by the standard model, as the incremental learning rule recursively computes a recency-weighted running average of the experienced outcomes. When fewer samples are considered, the predicted statistics of resulting choices differ systematically. This is because although the value of the sample is, in expectation, the same as the weighted average of the quantities from which it was drawn, individual samples will fluctuate around this mean. For this reason, choices arising from a nonlinear (for example, max or softmax) comparison of these sampled values across options, do not, in general, have the same statistics as the max of the averages. For instance, the running average approach predicts a fixed, vanishingly small influence of long-past rewards, whereas in the sampling model events from the far past could still have occasional, but sizable, influence on choices. Furthermore, whereas standard incremental learning models imply that the predicted choice distribution is invariant to the particular sequence of outcomes that gave rise to the options' values, given only their means, for sampling, even holding constant the average outcomes, the choice distributions will differ depending on the particular sequence (for example, variance) of outcomes experienced. Our primary model of interest was a sampling model that drew only one sample of past experience. (We also considered variants that drew a greater number of samples; none of these were superior to the one-sample version.) Critically, the two models were carefully set up so as to be matched in all respects other than the key feature of interest (sampling versus averaging): in addition to recency weighting, the choice rules and free parameters were identical.

We computed, for each trial and each participant individually, the probability that each model would produce the choice observed and estimated the parameter values that maximized these choice likelihoods. Comparing these maximum-likelihood estimates we found that, across the population and for all 20 participants individually, choices were better fit by our sampling model than by the learning model (mean log Bayes factor against temporal difference (TD) 8.8867, s.e.m. 1.0811, exceedance probability > 0.99 and Fig. 1c). Fit parameters for each model are shown in Table 1.

We repeated these analyses on simulated data for which the ground truth model was known to verify that these two models were distinguishable from choice behaviour (Supplementary Notes 'Simulated model fits and simulated regression results', Supplementary Fig. 1 and Supplementary Table 1), and also tested the generality of this conclusion under a different specification of the softmax choice noise rule (Supplementary Note 'Alternative forms of choice noise'). We next examined neural correlates of decision variables from the model (RPE and chosen value (CV)), recorded using functional magnetic resonance imaging (fMRI) collected on a subset of participants and verified that, similar to the choices, both were better accounted for by decision variable time series derived from the Sampler model rather than the standard one (Supplementary Note 'Neuroimaging reanalysis' and Supplementary Fig. 2).

Experiment 2. The above results concord with our hypothesis that human decisions are guided by samples of individual past

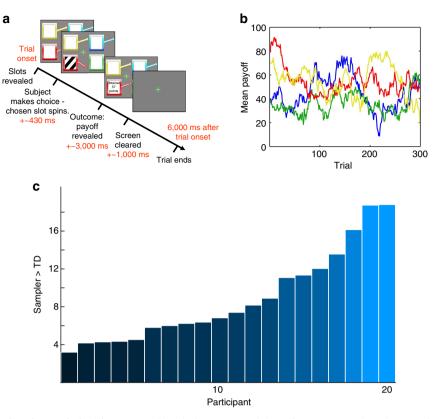


Figure 1 | Restless bandit task and re-analysis. (a) Four-armed bandit from Daw et al. (2006). Participants chose between four slot machines to receive points. (b) Payoffs. The mean amount of points paid out by each machine varied slowly over the course of the experiment. (c) Model comparison. Log Bayes factors favouring sampling over the TD model.

Model	α	β	$oldsymbol{eta^c}$	Log Bayes
TD	0.7754 (0.0472)	8.7069 (0.9564)	0.6498 (0.0193)	8.8867 (1.0811)
Sampler	0.7277 (0.0447)	9.0130 (1.1012)	0.3046 (0.1288)	-

choices, even in a simple repeated choice task. However, one disadvantage of applying sampling models to this type of task is that trials are essentially similar; we cannot directly observe (and must instead marginalize when computing model fit) which individual trials participants have sampled. For this reason, our conclusions are based on an overall comparison of model fit rather than a test of clear qualitative features of either model. Moreover, the experiment was correlational, rather than causal. To address this, we performed a second behavioural experiment to provide more direct support for the hypothesis by bringing the sampling process under experimental control, so as to measure the impact of a single, selected experience on choices. We leveraged the fact that multiple aspects of the choice experience can be bound together as a single representation. Therefore, we used choice-incidental—but still trial-unique—information to tag each choice as a unique event. Specifically, we modified the bandit task from Experiment 1 so that each trial involved a uniquely identifying photograph of an everyday object—a 'ticket' emitted by the chosen slot machine (see also ref. 25). To simplify the task and analysis, the number of choice options was reduced to two from four and outcome values were limited to wins or losses of \$5 (Fig. 2a). The probability that each machine would pay a winning ticket varied from trial to trial (Fig. 2b). Other than the ticket presentation, this task matches the sort of 'two-armed bandit' traditionally employed in RL studies.

Interspersed among the 130 choices were 32 recognition probe trials on which participants were asked whether or not they recognized a given 'ticket' image (Fig. 2c). These probes were intended to bring to mind the specific trial on which the probed image was first experienced—including the bandit chosen before the ticket appeared and the outcome received after. We hypothesized that this reminder would make the original experience more likely to be sampled during the ensuing choice. In contrast, standard running average models predict no such effect, as they base choices only on the summary statistics. For this reason, this experiment (unlike the previous one) exercises a clear, qualitative prediction between the models: sensitivity to probes evoking individual choices. Probes evoked choices that were, on average, 39 trials in the past—a temporal horizon minimizing both the influence of that reward on a recency weighted running average, as well as the likelihood that the reminded trial would still be present in working memory²⁶. After each probe, the choice task continued as before. To incorporate reminded trials into the Sampler model, we added a free parameter α^{evoked} , specifying the likelihood of reminded trials being incorporated into the sample-based value estimation

process. Matching the first analysis, this Sampler model proved a superior fit to choices than did an incremental learning model, for 19 out of 21 participants individually and across the population as a whole, despite being penalized for the additional free parameter (mean log Bayes factor against TD 6.9182, s.e.m. 1.3227, exceedance probability >0.99; Fig. 3a and model parameters in Table 2).

However, the key qualitative test of our hypothesis is whether choices following a recognition probe show an effect of the cued trial. For example, if a given recognition memory probe evoked a trial on which the participant chose the blue bandit and was

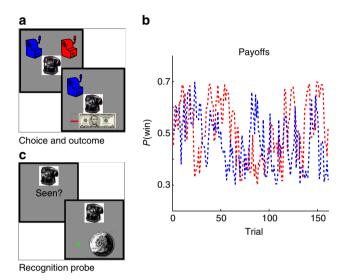


Figure 2 | Ticket bandit task. (a) The ticket-bandit task. Each slot machine ('bandit') delivered tickets—trial-unique photographs—associated with a dollar value—either - \$5 or \$5. (**b**) Payoff probabilities. The probability of each bandit paying out a winning ticket varied slowly over the course of the experiment. Participants were told that their total payout would be contingent both on the number of winning tickets they accrued and their ability to correctly respond on a post-task memory test asking them to recall the reward value and slot machine associated with each ticket. (c) Memory probes. Participants encountered 32 recognition memory probes. On 26 of these probe trials, participants were shown objects that were either received on a previous choice trial ('valid'), whereas on others they were shown new objects that were not part of any previous trial ('invalid'). Participants were asked only to perform a simple old/new recognition judgement—to press 'yes' if they had seen the image previously in this task and 'no' if they had not. After each recognition probe, the sequence of slot machine choices continued as before.

rewarded, then the participant should be more likely to choose the blue bandit on the subsequent choice trial. Conversely, if the choice had resulted in a loss, then they should be more likely to choose the alternative bandit (in this example, red). Incremental RL would be unaffected by the memory probes, because it does not maintain memories for individual trials and because the memory probe trials do not provide relevant direct reward experience that could be used to update running average values for the slot machines. Consistent with the sampling model, we observed that choices following a memory probe were also significantly influenced by the much older experience evoked by the probed ticket (t(20) = 3.8749, P = 0.0009; Fig. 3b). The magnitude of the increase in choice probability was comparable to that of a reward directly received between two and three trials in the past, suggesting that the reminder probe was effective in bringing to mind the associated bandit-reward link, and that reward information was then incorporated into choice, on a large fraction of post-reminder trials. Simulations demonstrated that the Sampler model does indeed capture this qualitative pattern of data (Supplementary Fig. 1).

Discussion

These results establish that choices can be made by sampling from individual trial memories, an architecture wholly different from those traditionally envisioned to explain reward learning. We showed that this model can capture the characteristic—and behaviourally observed²⁷—recency dependence of incremental averaging approaches, yet provides a superior fit to behaviour and decision-related neural signals even in a task originally designed to induce such incremental value updating. This finding goes beyond earlier work that showed that aggregate features of choices—such as a chooser's variable sensitivity to risky options are also consistent with a similar sampling model^{28,29}. Critically, the results of Experiment 1 imply that episodic sampling obtains as an influence on choices even when participants are not explicitly encouraged to encode memories of individual trials. However, the results do not directly test the other part of our hypothesis that choices are driven by memories of individual trials. We therefore tested this idea more directly in a second experiment. In Experiment 2, we strongly encouraged participants to form memories of each trial, so that we could test the influence these memories had on later choices. We brought these past trials to mind by the use of incidental stimulusstimulus associations present at the time of the original decision and showed that reward received on that individual trial had a reliable effect on choices following the reminder.

The fact that there was such a strong incentive to encode individual trial memories in Experiment 2 could, in principle,

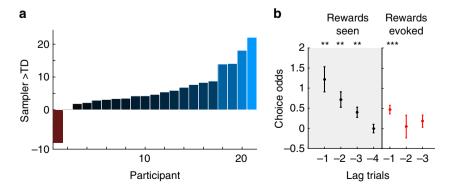


Figure 3 | Ticket bandit results. (a) Model comparison. Log Bayes factors favouring sampling over the TD model. **(b)** Impact of probes. As in standard RL models, choices are affected by previously observed rewards (black points). Here, memory probes evoking past decisions (red) also modulate choices on the subsequent choice trial. Data points are log odds of choosing the righthand option. (*P<0.05, *P<0.01 and **P<0.001).

Table 2 Fit model parameters for Experiment 2.							
Model	α	$lpha^{ ext{evoked}}$	β	eta^{c}	Log Bayes		
TD	0.5552 (0.0862)	-	1.7551 (0.6845)	- 0.0930 (0.2354)	6.9182 (1.3227)		
Sampler	0.5393 (0.0583)	0.4386 (0.0990)	2.2869 (0.4943)	0.5855 (0.3215)	-		

affect the decisions that subjects made. However, just as Experiment 2 extends the results of Experiment 1, Experiment 1 supports the findings of Experiment 2, by showing that the same model obtains on choices in a standard sequential choice experiment (indeed, one that was carried out without reference to the episodic sampling model). Episodic memory is characterized by the formation of stimulus–stimulus associations during a single experience. Therefore, this result suggests a key role in sampling decisions for the episodic memory system.

Episodic sampling is a potentially powerful mechanism for choice, as it allows generalizing past experience to apply to novel situations in which decision makers can rely on neither learned action values nor structured, semantic knowledge. In these situations, the episodic memory system could use what features partially overlap with past experience to pattern complete and make predictions for entirely new outcomes using elements of old ones^{10,12}. Importantly, episodic sampling allows these sorts of generalizations to be made at the time of choice, rather than needing to rely on links inferred before the decision at hand³⁰. Although the latter sort of generalization during encoding can account for many putatively 'model-based' generalization phenomena (for example, ref. 10), retrieval of episodes or semantic information at the time of a choice is necessary for evaluating unanticipated options¹² and to make use of additional information learned between initial encoding and choice²². Such an ability to flexibly generalize—as in latent learning and other revaluation tasks—has previously been associated with a 'model-based' learning system, distinct from the 'model-free' reward averaging approach⁶. However, although a common assumption is that such behaviours arise from more semantic representations—similar to cognitive maps—learned over multiple experiences⁸, it is possible that samples of individual episodes (for example, spatial trajectories) might play an analogous role. Thus, it remains unclear whether episodic sampling similar to that considered here supports model-based learning in other tasks, or if it represents yet a third way of deciding 18,22. Suggestively, we have previously shown that BOLD signal in the hippocampus at the time of choice scales with the difficulty of making these sorts of 'latent learning' decisions consistent with a sampling process that draws on a wider range of memories when associative information is less decisive about the right action to take¹². Relatedly, we have shown that enhanced hippocampal-striatal functional connectivity in a bandit task is associated with reduced influence of incremental reward learning on striatal RPEs and choices, and with enhanced episodic encoding for choice options²⁵.

One open question is the similarity or priority function by which samples are selected from episodic memory. In the terms of non-parametric statistical models, this function corresponds to the kernel and may itself be modifiable by experience²². Here, for simplicity and because of the relative lack of structure in the task, we assumed that samples were drawn according to the recency of the experience. Hertwig and Erev³¹ have shown this is a reasonable approximation when the underlying similarity structure of experiences is unknown. However, this recency may only be an approximation and, in environments with more

definite structure than the one we use here, different functions or additional dimensions of similarity may obtain. (Indeed, the selection function shown here is likely to be an ineffective learner of tasks in which simple temporal similarity is violated, such as those with payoff rules that alternate from trial to trial.) For instance, Plonsky *et al.*¹³ showed that across several kinds of sequential, binary choice tasks, a model that used the recent sequence of outcomes as a template for selecting past experiences had superior performance to one that selected on temporal recency alone. Connecting these sampling procedures to episodic memory opens up the possibility that these effects, and others, can be tied to well-known features of that memory system.

Indeed, the link between reward-guided choice and episodic memory brings into contact two areas of study with welldeveloped bodies of computational theory and widespread impacts for cognition more broadly³² and opens the door for leveraging many other features of episodic memory to explain and perhaps alter—decisions. Although in the present study we focus exclusively on the direct association between action and reward outcome, as the choice trials contain little other useful information, decisions in the real world take place in contexts laden with associative information that could be of relevance to decisions. It seems reasonable to suspect that adaptive decisionmakers can leverage this information to support value estimation. Along these lines, in a follow-up study³³, we provide behavioural and neuroimaging evidence that the context in which a sampled experience was first learned predicts the identity of the next experience to be sampled. Further work will be necessary to understand the way in which the episodic nature of memory samples affects the process by which they are sampled.

A related question is to what extent episodic sampling is separate from 'model-free' prediction error mechanisms associated with dopamine and its targets in striatum^{1,2}. The finding here that neural RPE signals are captured by expectations formed using associative memories concords with our previous observation that RPE signals during goal-directed decision tasks were better explained by expectations derived from hippocampally linked associative information¹¹ and fit with a series of observations that the nucleus accumbens (NAcc) RPE signal reflects a mixture of value signals beyond simple model-free TD learning^{9,11,34}. One possible unifying explanation for these and the previous findings of model-free RPE is that the systems partly overlap, with samples from memory used to train an average-value representation that guides action selection³⁵. In this way, rather than computing sampled values separately, direct and sampled experiences might be mixed freely, buffered through a single prediction error signal and net value store^{36,37}. Notably, RPE signals in striatum are influenced by 'model-based' and hippocampally linked stimulus-stimulus information 11,34 and are disrupted when participants successfully form episodic memories for task-irrelevant incidental material²⁵, suggesting an ongoing, trial-by-trial interaction. (We fit this sort of hybrid model to the data collected in Experiment 2, but the results were equivocal; see Supplementary Notes 'Adding evoked trials to the TD model and hybrid sampler and TD model', and Supplementary Tables 2 and 3).

Altogether, although we show here that episodic sampling can explain choice behaviour and neural signals better than incremental learning alone, normative frameworks support the idea that multiple forms of value learning coexist and their influence on each given choice fluctuates according to momentary demands^{6,38,39}. This finding concords with those cooperative frameworks by showing that the sampling mechanism may contribute to decision-making continually, not just in situations of low experience or uncertain associative structure (*c.f.* ref. 18). A superposition of two recency-weighted processes may be one reason why several studies have observed that actions exhibit a double-exponential form of dependence on past outcomes^{27,40}—with components that have been linked to separate activity in striatum and hippocampus^{11,41}. The results also connect to a known role for episodic memory in ongoing elaborative prospection about future situations⁴².

More broadly, an explicit link between episodic memory and adaptive instrumental control may serve as one rationalization for why it is that organisms encode rich, associatively dense memories of individual events in the first place⁴³. An open question is to what degree the content and persistence of certain memories, rather than others, can be attributed to the potential usefulness of those memories for later decision-making.

Methods

Experiment 1: Four-armed bandit from Daw et al. (2006). Participants completed a four-choice bandit task, choosing in each of 300 trials between four different slot machines and receiving a payoff (between 0 and 100 points) for their choice. The data analysed comprise choice timeseries for the 14 fMRI participants reported previously⁴ and for behavioural analyses are augmented with 6 additional behavioural-only pilot participants collected for (but not reported with) that study. Over the experiment, the mean payoffs for the machines diffused according to independent Gaussian random walks.

For a detailed description of the experimental methods, materials and previous analyses, see the prior report⁴. This section describes the new analyses we performed.

Choice behaviour. Two distinct types of models were compared in their effectiveness at explaining each participant's time series of choices. The first implemented a TD learning approach that kept a running average estimate of action values. The second followed a strategy of sampling from previous experiences to estimate these values at the time of decision.

The standard TD model maintains a value $Q_t^{\text{TD}}(a)$ for each option a, updating it following each choice of a according to the difference, $r - Q_t^{\text{TD}}(a)$, between obtained and expected rewards. The amount of update is controlled by a step-size (learning rate) parameter α^{TD} . Formally, on each timestep we updated the value estimate of each action according to equation (1):

$$Q_t^{\text{TD}}(a_c) = Q_{t-1}^{\text{TD}}(a_c) + \alpha^{\text{TD}}[r_t - Q_{t-1}^{\text{TD}}(a_c)]$$
 (1)

where r_t is the reward received at trial t, a_c is the chosen action and a_u is the unchosen action.

Indexing the previous choices of a as i=1, 2...t-1 choices into the past, this rule can easily be shown⁴⁴ at each step to make $Q_t^{\mathrm{TD}}(a)$ a weighted average of the rewards previously received for that choice:

$$Q_i^{\text{TD}}(a_c) = \alpha^{\text{TD}} \cdot \sum_i (1 - \alpha^{\text{TD}})^i r_{i-1}$$
 (2)

with weights decaying exponentially (by in the choice lag *i*. The resulting decision variable $Q^{\rm TD}$ is linked to choice probabilities according to the standard softmax function with inverse temperature parameter $\beta^{\rm TD}$. A third parameter, β^c , modelled any perseverative effect of the previous trial's choice, irrespective of reward received ^{41,45}. Choice probabilities were therefore estimated via the combined softmax as:

$$p_t(a = A) = \frac{e^{\beta^c I_t^c + \beta^{TD} Q_t^{TD}(A)}}{\sum_i e^{\beta^c I_t^c + \beta^{TD} Q_t^{TD}(a_i)}}$$
(3)

where I_t^c is an indicator function returning 1 if the previous choice (at trial t-1) was identical to that on the current trial (t) and 0 otherwise.

A similar sort of temporally decaying dependence on past rewards can also arise from a strategy of recency-based sampling. Instead of maintaining a running average, the sampling model stochastically samples one previous reward for each option with probability given by the same form of weighting:

$$P(Q_i^{\text{sample}}(a) = r_i) = \alpha^{\text{sample}} (1 - \alpha^{\text{sample}})^{t-i}$$
(4)

In this model, the most recent experience is most likely to be sampled and previous trials are successively exponentially less likely to be sampled. (We also considered a version of the model that samples k rewards, with replacement, and averages them to produce $Q_t^{\text{sample}}(a)$; this model limits to the standard TD model as $k \to \infty$). The models were equated in all aspects other than the functions they used to aggregate past experience.

For the sampling model, this probability is computed as the likelihood-weighted expectation of equation (3). One sample is drawn for each bandit at each trial, independently according to equation (4) within that bandit's reward history. This expectation is taken over every possible individual sample r_b with weighting $P\left(Q_i^{\text{sample}}(a) = r_i\right)$, as given by equation (4). One sample is drawn for each bandit at each trial and so the full choice probability is computed as a vector across bandits, over all possible combinations of samples across bandits (four bandits in Experiment 1, and two bandits in Experiment 2). For illustration, in Experiment 2, the computation of $p_i(a = A_i)$ is written as equation (5):

$$\sum_{j=1}^{t-1} P(Q_t^{\text{sample}}(A_i) = r_j) \cdot \sum_{k=1}^{t-1} P\left(Q_t^{\text{sample}}(A_{\neq i}) = r_k\right) \frac{e^{\beta^c I_t^c + \beta^{\text{sample}} r_j}}{e^{\beta^c I_t^c + \beta^{\text{sample}} r_j} + e^{\beta^c I_t^c + \beta^{\text{sample}} r_k}}$$
(5

For Experiment 1, the expression is analogous but contains four (rather than two) nested sums, corresponding to the reward sampled for each bandit. In both models, initial conditions $(Q^{TD}(\cdot), r_0)$ were set to the median payout value (50 for Experiment 1 and 0 for Experiment 2). In the sampling model, any sample probability not assigned to trials t=1, 2... was assigned to r_0 .

Model comparison. The likelihood of the sequence of obtained choices (each conditional on the rewards and choices up to that point) is then computed as the product over trials of the softmax probabilities, averaging over all combinations of samples in the sampling model and optimizing the free parameters (α^x , β^x , β^c , with the number of samples k = 1 for the sampling models) separately for each participant and model to maximize this likelihood (minimize the negative log likelihood). We compared the fit of candidate models to the choice data using Bayes factors (ref. 46; the ratio of posterior probabilities of the model given the data). We approximated the log Bayes factor using the difference between scores assigned to each model via the Laplace approximation to the model evidence⁴⁷. In participants for whom the Laplace approximation was not estimable for any model (due to a non-positive definite value of the Hessian of the likelihood function with respect to parameters), we approximated the log Bayes factor using the difference in likelihoods penalized using the Bayesian Information Criterion 48. Model comparisons are reported both on the log Bayes factors as estimated for each individual and as aggregated across the population. Parameters were estimated using a maximum a posteriori method, accounting for priors over the parameters⁴⁷. The posterior evidence calculations assumed the following prior distributions, chosen to be unbiased over the parameter ranges seen in previous studies³⁵, and to roll off smoothly at parameter boundaries: for the learning rate parameters, we employed a prior of Beta(1.1, 1.1); for the softmax temperatures and perseveration parameters, we employed a prior of Normal(0, 10). We also report the 'exceedance probability'-the posterior probability that one model is the most prevalent among a set, across a population, as computed using the spm_BMS function included in SPM8 (ref. 49).

fMRI. Region of interest definition. Based on prior studies, we identified regions of interest (ROIs) for each neural decision variable of interest: CV, in the ventromedial prefrontal cortex and RPE, in the NAcc. The NAcc ROI was defined anatomically according to known structural boundaries using the procedure outlined by Breiter $et~al.^{50}$ and used in our previous studies of reward-guided choice; specifically, defining the nucleus' superior border by a line connecting the most ventral point of the lateral ventricle to the most ventral point of the internal capsule at the level of the putamen 11,35,50 . In the case of ventromedial prefrontal cortex, functional and anatomical boundaries are less well-defined; we defined the current ROI as a 10 mm sphere centred on the peak coordinate identified as uniquely responding to 'Goal Value' (defined identically to CV here) by Hare $et~al.^{51}$.

Regression analysis. Candidate timeseries for each variable of interest (CV and RPE) were generated according to each of the two models: TD learning and Sampler. The trial-by-trial predictors were first convolved with a haemodynamic response function for each bandit and then (to account for the hemodynamic lag) we selected the volume two TRs (approximately six and a half seconds) following the relevant event of interest—at decision onset for CV, at outcome receipt for RPE. To account for correlation between the two variables, the time series derived from the Sampler model were then orthogonalized against their TD counterparts. Both variables were then entered into a simultaneous linear regression, which computed their relative contribution to explaining the time series extracted from each of the ROIs. The design matrix also contained twelve participant motion

regressors of no interest. The resulting regression weights were treated as random effects and tested against zero across the population by two-tailed t-test.

Experiment 2. Participants. Thirty individuals (1 left-handed; 14 female; ages 18–38 years, mean 23.0) participated in the study. Participants were recruited from the New York University community as well as the surrounding area, and gave informed consent in accordance with procedures approved by the New York University Committee on Activities Involving Human participants. All had normal or corrected-to-normal vision. All participants received a fixed fee of \$5, unrelated to performance, for their participation in the experiment. Additional compensation of between \$0 and \$18 depended on their performance on the in-task recognition memory probes (\$0.25 each of 32 probes), two pseudorandomly selected choice trials (\$5 each) and a post-task recall test. Average total compensation was \$12.

Exclusion criteria. Data from nine participants were excluded from analysis due to their being unusable, leaving 21 participants analysed here. Seven participants were excluded due to response biases that indicated they did not attempt to learn the rewards associated with each bandit—specifically, that their choices were more than 90% to either option in either block of the experiment. Two participants were excluded due to below-threshold performance on the in-task memory probes (below the 90% confidence level, or four or more incorrect).

Task design. Participants performed a two-armed restless bandit task—a series of 130 choices between two slot machines—with 32 interspersed memory probe trials, for a total of 162 trials. A compulsory rest period of participant-controlled length was inserted after the 81st trial. The experiment was controlled by a script written in Matlab (Mathworks, Natick, MA, USA), using the Psychophysics Toolbox [53]. Before the experiment, participants were given written and verbal instructions as to the types of trials, the buttonpresses required of them, the slot machine payoff probabilities and the rules for determining final payout. Instructions emphasized that there was no pattern linking the content of the ticket photographs to their dollar value or slot machine. Participants were not told that the memory probe trials should have an effect on their choices, nor was any effect implied. They were, however, told that memorizing the content, source (which slot machine the ticket came from) and dollar value of the tickets would impact their total payout after a post-task memory test.

To aid memory, participants were offered a mnemonic strategy. Specifically, they were told that the photographs could be treated as tickets, which could be placed in one of four imaginary 'pockets': left or right, depending on the slot machine, and front or back, depending on the dollar value. During practice, the keypresses required to advance each trial were instructed in the context of this mnemonic strategy (for example, 'Press 'a' to put the ticket in your left hand, now press 'a' again to put the ticket in your front pocket.'). Participants completed two practice trials before beginning the main experiment.

Choice trials. On each choice trial, participants were presented with two slot machines, each on either side of the upper third of the screen (Fig. 2a). The slot machines paid out tickets worth either \$5 or -\$5. Participants were instructed to press a key corresponding to the slot machine they felt had the best chance of paying out a winning (+\$5) ticket rather than a losing (-\$5) ticket on that trial. They could choose either the lefthand (key 'a') or righthand (key 'b') slot machine. The slot machines were visually identical, except for their colour (blue and red) and side of the screen, both of which remained fixed throughout the task. The probability of each machine paying a winning ticket changed independently on each trial according to a diffusing Gaussian random walk with reflecting bounds at 30 and 70% (Fig. 2b). At each timestep t, π_i^t —the probability that machine i would pay out a winning ticket—diffused according to: $\pi_{t+1}^i = \pi_t^i + \nu$ for each i. The diffusion noise ν was selected from a zero-mean Gaussian with standard deviation $\sigma_d=0.1$. The initial payoff probabilities were set to 60% and 40%, with the identity (side, colour) of the superior starting bandit psuedorandomly assigned for each participant.

After selection, the unchosen slot machine was covered and the chosen machine remained alone on the screen for 0.25 s. Then, a trial-unique photograph appeared and remained on the screen until participants again pressed the key corresponding to their chosen slot machine—in the instructions, this corresponded to the mnemonic memory strategy of selecting the left or righthand side 'pockets' for your ticket. When the correct key was pressed, a grey box appeared around the photograph and remained there for at least 500 ms or additional time up to two seconds depending on when the participants pushed the correct key.

At the end of the timeout, the box disappeared and the dollar value associated with the ticket—either — \$5 or \$5—was displayed, as a photograph of a \$5 bill, with a green '+' or a red '-' to the left of the bill image. The bill photograph remained on the screen—along with the chosen slot machine and the trial-unique 'ticket' photograph—until the participant pressed a key corresponding to the value: either 'a' for — \$5 or 'b' for \$5. In the instructions, this was referred to as 'putting the ticket in your front or back pocket', on the side indicated by the slot machine identity. Once the correct key was pressed, the bill photograph was surrounded by a grey box and remained on the screen for two seconds. After each trial, a blank screen was displayed for an inter-trial interval of two seconds.

In-task recognition memory probes. Beginning after the tenth choice trial, 32 memory probe trials were interspersed at pseudorandom intervals throughout the task (Fig. 3c). Each probe trial consisted of a single photograph and the question: 'Is this your ticket? (yes/no)'. Twenty six of these photographs were

Table 3 | Ticket values are modified by performance on a post-task recall memory probe.

Probe outcome	\$5 ticket	- \$5 ticket
Correct on both questions	\$5	- \$2.5
Incorrect on one question	\$0	- \$5

After the main slot machine task, 'tickets' paid out by the machines were presented to the participant again. The participant was asked to recall two specific details associated with the ticket: the machine that paid it out (left or right) and the value of the ticket (—\$5 or \$5). To encourage participants to encode the ticket-machine-value triplet, they were told that the final value of the tickets would depend on both the original value of the ticket and the participant's performance on two post-task recall questions. If they answered either question incorrectly, \$5 tickets were modified to be worth \$0. If they answered both questions correctly, —\$5 tickets were modified to be worth —\$2.5. The payout values of each ticket after the memory tests are described in this table. Values altered by the results of the memory tests are highlighted in bold.

chosen pseudorandomly without replacement from the list of previously seen images; these trials are referred to as valid memory probe trials. The remaining six photographs were novel; these trials are referred to as invalid memory probe trials. Participants were instructed to press 'a' if they remembered seeing that image and 'b' if they did not remember seeing that exact image before.

Correct responses—'yes' for previously seen images and 'no' for images that were not displayed on a previous bandit trial—were rewarded with \$0.25 added to the participant's total payout. This additional reward was indicated by a photograph of a US quarter with a green '+' to the left. Incorrect responses resulted in \$0.25 being deducted from the participant's total payout, indicated by a red '-' to the left of an image of a US quarter. Memory probe rewards were displayed for two seconds.

Rewards for memory probes accumulated over the course of the entire task, rather than for randomly selected rounds—so the total payout could be reduced or increased by as much as \$8.00. Probe images remained on the screen for up to four seconds—if no answer was entered in that time, the trial was scored as incorrect.

Post-task recall memory probes Before the experiment began, participants were instructed to remember as many complete bandit-outcome-ticket triplets as they could. Their memory for these triplets was tested in 21 post-task memory probes

Post-task memory probes were drawn only from the images tested during the experiment as valid in-task memory probes. This time, participants were queried as to their recall of each detail associated with the presented probe image: which slot machine it came from and which dollar value it was associated with. Participants were incentivized to answer correctly by the fact that their total payout in the experiment was predicated on their performance in this memory task. Specifically, payouts for winning (\$5) tickets were reduced, to \$0, if either post-task recall question was answered incorrectly. Payouts for losing (- \$5) tickets were increased, to -\$2.50, if both post-task recall questions were answered correctly (Table 3). (The asymmetry in the final values of the winning and losing tickets was intended to encourage participants to attend to both tasks; if losing tickets could be improved to \$0 by correct memory task responses, then participants would have less incentive to track the payout values of the bandits-in this case, simply choosing the same option on every trial would have a net positive EV; similarly, if incorrect memory responses only reduced winning tickets to some positive value, then there would be less incentive to remember the ticket images.)

Analysis. Our analysis of choice behaviour in this task addressed the hypothesis that memory probe images affect choices in subsequent bandit trials, by evoking the rewards received by choices on past bandit trials.

Choice kernel regression. To test this hypothesis, we first examined the effect of past trials—both bandit trials directly experienced and those evoked by valid memory probes—on choices. Specifically, we entered into a logistic regression one regressor for the rewards received on each of the previous ten trials. If a positive reward was received after choosing the right bandit on trial $t-\tau$, this was coded as a 1 in regressor τ , element t. If a negative reward was received after choosing the right bandit, this was coded as a -1. These values were flipped for lefthand bandits: -1 for positive rewards and 1 for negative rewards.

Next we included ten regressors coding rewards from past choices that were evoked by valid memory probes over the past ten trials. Specifically, if trial $t-\tau$ was a valid memory probe that evoked a trial on which the participant received a positive reward for the right bandit, this was coded as a 1 in the regressor τ , element t, and so forth, following the same coding scheme as directly experienced rewards.

We also entered into the regression matching sets of ten regressors each specifying the identity of the bandit chosen on each of the previous ten trials: $-\,1$ for left, 1 for right. In addition, similarly, for the identity of the bandits evoked by any valid memory probes during the previous ten trials.

Choices of the current bandit were entered as the dependent variable, and coded as -1 (for left) and 1 (for right). The resulting regression weights—indicating the degree to which the current choice was influenced by choices and rewards on a given evoked or directly experienced trial in the past—were tested against zero across the population by two-tailed t-test.

The final regression was thus in the following form:

$$\log \frac{p_{\text{right},t}}{p_{\text{left},t}} \approx \sum_{i=1}^{10} \beta_i^{\text{DR}} \text{DR}_{t-i} + \sum_{i=1}^{10} \beta_i^{\text{ER}} \text{ER}_{t-i} + \sum_{i=1}^{10} \beta_i^{\text{DI}} \text{DI}_{t-i} + \sum_{i=1}^{10} \beta_i^{\text{EI}} \text{EI}_{t-i} \quad (6)$$

where DR is 'directly experienced reward', ER is 'evoked reward', DI is 'directly experienced identity' and EI is 'evoked identity'—each specified over the immediately preceding ten trials.

Models. As in Experiment 1, we tested the fit to behaviour of computational models capturing our hypothesis and the competing, incremental learning hypothesis.

In the TD model, estimated action values were updated only on bandit choice trials; memory probe trials were not considered to have any impact on the values of the slot machines. This model is identical to equation (2) from Experiment 1, with the exception that the initial value $Q^{\text{TD}}(\cdot)$ is set to zero, the mean payout on this experiment. In the second model, estimated values were based on one sample drawn from past experience. In this experiment, the sampling model was augmented with an additional parameter, α^{evoked} , specifying the likelihood that a reminded trial is drawn as a sample. Formally, we distinguish between direct and evoked rewards r_i according to:

$$P\left(Q_{i}^{\text{sample}}(a) = r_{i,\text{direct}}\right) = \alpha^{\text{sample}} \left(1 - \alpha^{\text{sample}}\right)^{t-i}$$

$$P\left(Q_{i}^{\text{sample}}(a) = r_{i,\text{evoked}}\right) = \alpha^{\text{evoked}} (1 - \alpha^{\text{evoked}})^{t-i}$$
(7)

The original Sampling model of Experiment 1, with no impact of reminded trials, is nested within the sampling model and realized when $\alpha^{\rm evoked}$ equals zero.)

This formulation was used in computing choice probabilities for all models. These models were otherwise identical to those used in the analysis of Experiment 1.

Models were compared following an identical procedure as used in Experiment 1.

Data availability. Data for Evnoriment 2 are available upon reasonable request to

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References

- Barto, A. C. in Models of Information Processing in the Basal Ganglia (eds Houk, J. C., Davis, J. L. & Beiser, D. G.) 215–232 (MIT Press, 1995).
- Schultz, W., Montague, P. R. & Dayan, P. A neural substrate of prediction and reward. Science 275, 1593–1599 (1997).
- Sugrue, L. P., Corrado, G. S. & Newsome, W. T. Matching behavior and the representation of value in the parietal cortex. Science 304, 1782–1787 (2004).
- 4. Daw, N. D., O'Doherty, J. P., Dayan, P., Seymour, B. & Dolan, R. J. Cortical substrates for exploratory decisions in humans. *Nature* **441**, 876–879 (2006).
- Behrens, T. E. J., Woolrich, M. W., Walton, M. E. & Rushworth, M. F. S. Learning the value of information in an uncertain world. *Nat. Neurosci.* 10, 1214–1221 (2007).
- Daw, N. D., Niv, Y. & Dayan, P. Uncertainty-based competition between prefrontal and dorsolateral striatal systems for behavioral control. *Nat. Neurosci.* 8, 1704–1711 (2005).
- Hampton, A. N., Bossaerts, P. & O'Doherty, J. P. Neural correlates of mentalizing-related computations during strategic interactions in humans. *Proc. Natl Acad. Sci. USA* 105, 6741–6746 (2008).
- Gläscher, J., Daw, N., Dayan, P. & O'Doherty, J. P. States versus rewards: dissociable neural prediction error signals underlying model-based and modelfree reinforcement learning. *Neuron* 66, 585–595 (2010).
- Simon, D. A. & Daw, N. D. Neural correlates of forward planning in a spatial decision task in humans. J. Neurosci. 31, 5526–5539 (2011).
- Wimmer, G. E. & Shohamy, D. Preference by association: how memory mechanisms in the hippocampus bias decisions. Science 338, 270–273 (2012).
- Bornstein, A. M. & Daw, N. D. Cortical and hippocampal correlates of deliberation during model-based decisions for rewards in humans. *PLoS Comput. Biol.* 9, e1003387 (2013).
- Barron, H. C., Dolan, R. J. & Behrens, T. E. J. Online evaluation of novel choices by simultaneous representation of multiple memories. *Nat. Neurosci.* 16, 1492–1498 (2013).
- Plonsky, O., Teodorescu, K. & Erev, I. Reliance on small samples, the wavy recency effect, and similarity-based learning. *Psychol. Rev.* 122, 621–647 (2015).
- 14. Fan, J. & Gijbels, I. Local Polynomial Modelling and its Applications: Monographs on Statistics and Applied Probability (CRC Press, 1996).
- Ormoneit, D. Kernel-based reinforcement learning. Machine Learn. 49, 161–178 (2002).
- Erev, I. & Barron, G. On adaptation, maximization, and reinforcement learning among cognitive strategies. *Psychol. Rev.* 112, 912–931 (2005).
- 17. Stewart, N., Chater, N. & Brown, G. D. A. Decision by sampling. *Cogn. Psychol.gy* **53**, 1–26 (2006).
- Lengyel, M. & Dayan, P. Hippocampal contributions to control: the third way. *Adv. Neural Inform. Process. Syst.* 20, 889–896 (2008).
- Krajbich, I., Armel, C. & Rangel, A. Visual fixations and the computation and comparison of value in simple choice. Nat. Neurosci. 13, 1292–1298 (2010).

- Giguere, G. & Love, B. C. Limits in decision making arise from limits in memory retrieval. Proc. Natl Acad. Sci. USA 110, 7613–7618 (2013).
- Woodford, M. Stochastic choice: an optimizing neuroeconomic model. Am. Econ. Rev. 104, 495–500 (2014).
- Gershman, S. J. & Daw, N. D. Reinforcement learning and episodic memory in humans and animals: an integrative framework. *Annu. Rev. Psychol.* 68, 101–128 (2017).
- Duncan, K. D. & Shohamy, D. Memory states influence value-based decisions. J. Exp. Psychol. Gen. 145, 1420–1426 (2016).
- Murty, V. P., FeldmanHall, O., Hunter, L. E., Phelps, E. A. & Davachi, L. Episodic memories predict adaptive value-based decision-making. *J. Exp. Psychol. Gen.* 145, 548–558 (2016).
- Wimmer, G. E., Braun, E. K., Daw, N. D. & Shohamy, D. Episodic memory encoding interferes with reward learning and decreases striatal prediction errors. J. Neurosci. 34, 14901–14912 (2014).
- Collins, A. G. E. & Frank, M. J. How much of reinforcement learning is working memory, not reinforcement learning? A behavioral, computational, and neurogenetic analysis. *Eur. J. Neurosci.* 35, 1024–1035 (2012).
- 27. Lau, B. & Glimcher, P. W. Dynamic response-by-response models of matching behavior in rhesus monkeys. *J. Exp. Anal. Behav.* **84**, 555–579 (2005).
- Denrell, J. & March, J. G. Adaptation as Information Restriction. Organization Science 12, 523–538 (2001).
- Erev, I., Ert, E. & Yechiam., E. Loss aversion, diminishing sensitivity, and the effect of experience on repeated decisions. *J. Behav. Decision Making* 21, 575–597 (2008).
- 30. Berens, S. C. & Bird, C. M. The role of the hippocampus in generalizing configural relationships. *Hippocampus* **35**, 591–598 (2017).
- Hertwig, R. & Erev, I. The description-experience gap in risky choice. Trends Cogn. Sci. 13, 517–523 (2009).
- Shohamy, D. & Turk-Browne, N. B. Mechanisms for widespread hippocampal involvement in cognition. J. Exp. Psychol. Gen. 142, 1159–1170 (2013).
- Bornstein, A. M. & Norman, K. A. Reinstated episodic context guides samplingbased decisions for reward. *Nat. Neurosci.*, doi: 101038/nn.4573.
- Daw, N. D., Gershman, S. J., Seymour, B., Dayan, P. & Dolan, R. J. Model-based influences on humans' choices and striatal prediction errors. *Neuron* 69, 1204–1215 (2011).
- Bornstein, A. M. & Daw, N. D. Multiplicity of control in the basal ganglia: computational roles of striatal subregions. *Curr. Opin. Neurobiol.* 21, 374–380 (2011).
- 36. Sutton, R. S. Dyna, an integrated architecture for learning, planning, and reacting. *ACM SIGART Bull* **2**, 160–163 (1991).
- Gershman, S. J., Markman, A. B. & Otto, A. R. Retrospective revaluation in sequential decision making: a tale of two systems. Journal of experimental psychology. *General* 143, 182–194 (2014).
- Keramati, M., Dezfouli, A. & Piray, P. Speed/accuracy trade-off between the habitual and the goal-directed processes. *PLoS Comput. Biol.* 7, e1002055 (2011).
- Simon, D. A. & Daw, N. D. in Advances in Neural Information Processing Systems Vol. 24 (eds Shawe-Taylor, R. S. et al.) Curran Associates, 127–135 (2011).
- Corrado, G. S., Sugrue, L. P., Seung, H. S. & Newsome., W. T. Linear-nonlinear-poisson models of primate choice dynamics. *J. Exp. Anal. Behav.* 84, 581–617 (2005)
- Bornstein, A. M. & Daw, N. D. Dissociating hippocampal and striatal contributions to sequential prediction learning. Eur. J. Neurosci. 35, 1011–1023 (2012).
- Buckner, R. L. & Carroll, D. C. Self-projection and the brain. Trends Cogn. Sci. 11, 49–57 (2006).
- Anderson, J. R. in Varieties of Memory and Consciousness: Essays in Honor of Endel Tulving (eds Roediger III, H. L. & Craik, F. I. M.) 195–210 (Erlbaum, 1989)
- Bayer, H. M. & Glimcher, P. W. Midbrain dopamine neurons encode a quantitative reward prediction error signal. *Neuron* 47, 129–141 (2005).
- 45. Kass, R. E. & Raftery, A. E. Bayes factors. J. Am. Stat. Assoc. 90, 773-795 (1995).
- Mackay, D. J. C. Information Theory, Inference, and Learning Algorithms (Cambridge Univ. Press, 2003).
- Schwarz, G. Estimating the dimension of a model. Ann. Stat. 6, 461–464 (1978).
- Stephan, K. E., Penny, W. D., Daunizeau, J., Moran, R. J. & Karl, J. Bayesian model selection for group studies. *NeuroImage* 46, 1004–1017 (2009).
- 49. Breiter, H. C. et al. Acute effects of cocaine on human brain activity and emotion. Neuron 19, 591-611 (1997).
- Hare, T. A., O'Doherty, J. P., Camerer, C. F., Schultz, W. & Rangel, A. Dissociating the role of the orbitofrontal cortex and the striatum in the computation of goal values and prediction errors. *J. Neurosci.* 28, 5623–5630 (2008).
- 51. Brainard, D. H. The psychophysics toolbox. *Spatial Vision* **10**, 433–436

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Author contributions

N.D.D. contributed data from Experiment 1. M.W.K. and A.M.B. performed the reanalysis of Experiment 1. A.M.B., N.D.D. and D.S. contributed to conceptualizing a role for episodic memory in decision-making. A.M.B. and N.D.D. designed Experiment 2. A.M.B. ran Experiment 2. A.M.B. analysed the data from Experiment 2. A.M.B. and N.D.D. wrote the paper, with input from M.W.K. and D.S.

Additional information

Supplementary Information accompanies this paper at http://www.nature.com/naturecommunications

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