A flexible and generalizable model of online latent-state learning

Amy L Cochran[1*], Josh M Cisler[2]

1 Department of Biostatistics and Medical Informatics, University of Wisconsin, Madison, WI, USA

2 Department of Psychiatry, University of Wisconsin, Madison, WI, USA

* cochran4@wisc.edu

Abstract

Traditional learning models fail to describe important phenomena, notably the rapid return of fear after extinction. To address this shortfall, evidence converged on the idea that learning agents rely on latent-state inferences, i.e. an ability to index disparate associations between observations and infer which index (i.e. latent state) is presently active. Our goal was to develop a model of latent-state inferences that uses *online* (also known as recursive or adaptive) updates and that can describe behavior in a diverse set of experiments. The resulting model combines a Rescorla-Wagner rule, for which updates to associations are proportional to prediction error, with an approximate Bayesian rule, for which beliefs in latent states are proportional to prior beliefs and an approximate likelihood based on current associations. In simulation, we demonstrate the model's ability to reproduce learning effects both famously explained and not explained by the Rescorla-Wagner model, including rapid return of fear after extinction, the Hall-Pearce effect, partial reinforcement extinction effect, backwards blocking, and memory modification. We find this feat is not trivial, as other models are unable to reproduce all these effects. We then show our model is practical for explaining behavioral data using a novel dataset (n=55) of a bandit task with switches in arm-reward contingencies. Lastly, we derive our model as an online algorithm to maximum likelihood estimation, demonstrating it is an efficient approach to outcome prediction. Establishing such a framework is a key step towards quantifying normative and pathological ranges of latent-state inferences in various contexts.

Author summary

Computational researchers are increasingly interested in a structured form of learning known as latent-state inferences. Latent-state inferences is a type of learning that involves categorizing, generalizing, and recalling disparate associations between observations in one's environment and is used in situations when the correct association is latent or unknown. This type of learning has been used to explain overgeneralization of a fear memory and the cognitive role of certain brain regions important to cognitive neuroscience and psychiatry. Accordingly, latent-state inferences are an important area of inquiry. Through simulation, psychological experiment, and theory, we establish a new model of latent-state inferences. Moving forward, we aim to use this framework to measure latent-state inferences in healthy and psychiatric populations.

Introduction

Learning and decision-making are fundamental aspects of day-to-day human life. Indeed, many mental health disorders can be conceptualized from the perspective of biases or errors in learning and decision-making [1]. Accordingly, the study of how humans learn and make decisions is an important topic of inquiry and the past two decades has witnessed a significant surge in the application of computational modeling approaches to the problem of human learning and decision-making [2–5]. One significant insight this literature has made is the differentiation of model-free and model-based learning [6,7]. Model-free learning refers to relatively simple updating of cached values based on trial-and-error experience. An early and influential example of model free learning is the Rescorla-Wagner (RW) model [8], which proposed that associative strength (i.e. the degree to which a cue predicted an outcome) updates in response to new experiences in proportion to the magnitude of a prediction error (i.e., how wrong the current prediction was). The RW model, and similar model-free formulations that it inspired [9,10], powerfully explained many aspects of learning.

Nonetheless, a notable problem for model-free accounts of learning was the phenomenon of rapid recovery of responding following extinction learning [11]. That is, model-free accounts of learning based on trial-by-trial updating predict that the return of responding following extinction should essentially occur at the same rate as initial learning. This prediction is not supported by a wealth of data demonstrating that extinguished responding can rapidly return via reinstatement (i.e., an un-signalled presentation of the outcome), context renewal (i.e., returning to the initial acquisition learning context), or spontaneous recovery (i.e., return of responding following the passage of time) [12]. This rapid return of extinguished responding is an important phenomenon with implications for treatment of clinical disorders such as anxiety disorders and

substance use [13, 14].

One of the first applications of a model-based account of learning that could address rapid renewal of responding following extinction was by Redish and colleagues in 2007 [11]. They argued that an agent performs two concurrent tasks during a learning experiment: 1) updating associative strength between a cue and an outcome, and 2) recognizing the current state of the experiment and storing separate associative strengths for the acquisition and extinction contexts. Rapid recovery of responding can be explained by the agent inferring that the current state of the experiment has changed to the acquisition phase and therefore responding to the cue with the associative strength stored for the acquisition phase. Though this initial formulation had limitations [15], it stimulated subsequent research developing latent-state models of reinforcement learning [16–19]. For example, if an agent is assumed to infer latent states to explain observed relationships between stimuli, actions, and outcomes, and if inferring separate latent states for acquisition and extinction phases of an experiment explains rapid recovery of responding, then it follows that blurring the experimental distinction between acquisition and extinction phases would result in less recovery of responding following extinction. That is, if the contingency between cue and outcome (e.g., 50% contingency) during the acquisition phase slowly transitions to extinction, rather than an abrupt change, then an agent is less likely to infer a separate extinction context and more likely to modify the initial acquisition memory. Because the acquisition associative strength is lower, less subsequent recovery of responding would be expected in a recall test. A carefully designed experiment using an animal model demonstrated exactly this observation, providing strong evidence for a latent-state model of learning [20].

The application and utility of latent-state learning models is not confined to explaining recovery of responding following extinction. Tolman's early theory of cognitive maps [21] posits that an agent forms abstract representations of a task's state space. There has been a recent resurgence of interest in this concept of cognitive maps in the cognitive neuroscience field informed by latent-state computational models [17, 22, 23]. This work, conducted both using animal and human experimental models, suggests that essential functions of the orbitofrontal cortex (OFC) and hippocampus are the encoding and representation of latent task space (i.e., forming cognitive maps of a task's state space) [23]. This theory potentially powerfully describes a superordinate process that explains why the OFC and hippocampus are implicated in such a diverse array of cognitive and emotional functions. This theory also has implications that tie back into extinction learning and clinical disorders and suggest a novel understanding of the role of the OFC in mediating poor extinction learning. However, it is important to recognize that the role of the OFC remains highly debated.

As can be seen, latent-state theories have significant implications for our understanding of normative and disordered mechanisms of learning. Despite the significant progress in this area, a current weakness is that

existing computational models of latent-state learning suffer from several limitations. These include imposing demands on human computation and memory that grow in the number of trials (e.g., storing the complete history of past outcomes), failing to describe more traditional experiments of associative learning, and requiring assumptions that restrict which experiments can be described. An additional limitation is that there is not one model that has been consistently applied to all recent experiments examining latent-state learning [20, 24–26]. These disparate models negate a unified account of latent-state learning. Indeed, one of the strengths of computational modeling applied to the problem of decision-making is that the learning theory is mathematically formalized [27]. This strength is lost when the mathematical formulation differs with each study and task because the underlying learning theory also necessarily differs across tasks. By contrast, having a single model that can characterize latent-state learning across the many different tasks used to probe latent-state learning would have both theoretical and practical significance. Theoretical significance would come from formalizing a theory of latent-state learning that generalizes across tasks, thereby potentially powerfully describing the essential nature of latent-state learning. Practical significance would come from providing the research community with a general and flexible model that can be used to integrate hypotheses and findings across the many research domains that propose latent-state models of learning. The purpose of the current work is to present a model of online latent-state learning and to verify the suitability and utility of this model for a diverse set of experiments.

Results

Model

We present a model of how an agent uses latent states to learn in a psychological experiment. Each experiment is assumed to consist of a sequence of trials, wherein a learning agent is exposed to a collection of cues followed by rewards (and/or penalties). We use the following notation. Cue n on trial t is denoted by $c_n(t)$ and takes a value of 1 when the cue is present and 0 otherwise. Cues are collected in a vector $\vec{c}(t)$. Rewards on trial t are denoted by $R(t)$. After trial t, the strength of the association between cue n and rewards is denoted by $V_n(t)$. An apostrophe denotes transposition, and the notation $X(1{:}t)$ is shorthand for $(X(1), \ldots, X(t))$.

Our model assumes a learning agent has a certain world view about how rewards are generated and wants to predict rewards efficiently based on this world view (Fig 1A). Their world view assumes rewards are a function of which cues are presented, a latent state, and a latent error. A latent state is an index to disparate

associations between cues and rewards. In order to predict rewards, they must invert their view of the world   90

to infer which latent state is active, how cues are associated with rewards for each latent state, and the   91

expected uncertainty in rewards due to the latent error. We proceed to describe our model for how an agent   92

performs these tasks. We leave details to the Methods section about how our model of latent-state learning   93

can be derived formally from this world view.   94

Fig 1. A) A learning agent's world view whereby rewards are generated according to cues, a latent state, and a latent error. In order to predict rewards, they must infer which latent state is active, the relationship between cues and rewards for each latent state, and the expected uncertainty in rewards due to the latent error. B) The proposed model for how a learning agent inverts their world view. They first observe cues to generate expectations or predictions for rewards based on L estimates of associative strengths corresponding to L latent states. Upon observing rewards, they use errors in their predictions to update associative strengths, measures of uncertainty, and beliefs in which state is active. The degree to which associative strengths can be updated depends on both the agent's belief in the corresponding latent state and the corresponding effort matrix, which keeps track of how cues covary.

Building upon the Rescorla-Wagner (RW) model   95

If an agent believed in only one latent state, then a RW model could be used to determine the relationship

between cues and rewards. The RW model proposes a linear relationship between cues and expected rewards.

This relationship is captured by an associative strength for each cue updated according to:

$$V_n(t+1) = \begin{cases} V_n(t) + \alpha_n(t) \left(R(t) - \sum_{n \, : \, c_n(t)=1} V_n(t) \right) & \text{if } c_n(t) = 1 \\ \\ V_n(t) & \text{if } c_n(t) = 0, \end{cases}$$

with initial associative strength $V_n(1) = 0$. Term $\alpha_n(t)$ is referred to as *associability* and is constant in the RW model. Collecting associative strengths in a vector $\vec{V}(t)$, we can express this update concisely as

$$\vec{V}(t+1) = \vec{V}(t) + A(t)\,\vec{c}(t)\left(R(t) - \vec{c}(t)'\vec{V}(t) \right).$$

The term $A(t)$ is now a matrix capturing associability and in the RW model, is diagonal with terms $\alpha_n(t)$ along its diagonal. Changes in associative strength depends on the learning agent's current expectation or best guess $\vec{c}(t)'\vec{V}(t)$ for rewards given observed cues. If their expectation exceeds the actual reward, then associative strength declines for each cue present. If their expectation is below the actual reward, then associative strength increases for each cue present. By updating associative strength in vector form, we see that rewards are captured by a linear regression model of cues. Hence, cues can be continuous or represent interactions, i.e. the presence or absence of a compound of cues. That is, the same update formula applies for continuous cues and interaction terms.

Allowing for latent-state learning

We build upon the RW model by allowing a learning agent to propose L competing (i.e. mutually-exclusive) *associations* between cues and rewards. The index to each association is referred to as a *latent state* and is associated with its own RW model for predicting rewards from cues. The agent learns about associations for each latent state while learning about their belief in which latent state best explains current observations. We capture belief in latent state l on trial t as a positive variable $p_l(t)$ such that beliefs sum to one: $p_1(t) + \ldots + p_L(t) = 1$.

We let $\vec{V}_l(t)$ represent the vector of associative strengths and $A_l(t)$ represent the associability matrix for latent state l. The learning agent updates associative strength as before except for the subscript l:

$$\vec{V}_l(t+1) = \vec{V}_l(t) + A_l(t)\vec{c}(t)E_l(t), \tag{1}$$

where we used the shortened notation $E_l(t)$ to represent the prediction error for latent state l on trial t:

$$E_l(t) = R(t) - \vec{c}(t)'\vec{V}_l(t). \tag{2}$$

An associability process

Associability is constant in the RW model. By contrast, experiments such as those by Hall and Pearce [9, 28] suggest associative strength is path-dependent: associability, i.e. how quickly associative strength changes, depends on the history of observations and changes over time [10]. We describe associability as depending on current beliefs and a matrix $B_l(t)$ which we refer to as an *effort* matrix:

$$A_l(t) = \alpha_0 p_l(t) B_l(t)^{-1}, \tag{3}$$

for a patent-specific parameter $\alpha_0 \in [0, 1]$ controlling how much an individual weights older observations versus newer observations. The effort matrix $B_l(t)$ is updated according to

$$B_l(t+1) = B(t) + \alpha_0 \left(p_l(t)\vec{c}(t)\vec{c}(t)' - B(t) \right). \tag{4}$$

with $B_l(1) = I$. The effort matrix estimates the matrix of cue second moments associated with each latent state. Second moment information of cues has been integrated into other models of learning, but usually in the form of variances and covariances rather than second moments directly [11, 19]. The use of the effort matrix is motivated in our derivation of our model in the Methods section and is similar to matrices found in an online algorithm for solving the machine learning problem of contextual bandits with linear rewards [29].

Updating latent-state beliefs

Beliefs link latent states in our model. The assumed world view of the learning agent proposes that rewards are generated on trial t as follows: a latent random variable $X(t) \in \{1, \ldots, L\}$ is drawn from some distribution and rewards are drawn depending on the current latent state $X(t)$ and cues $\vec{c}(t)$ (Fig 1A). Further, rewards are assumed to be mutually independent between trials conditional on latent states, and latent states are assumed to be Markovian, i.e. $X(t)$ depends on $X(1{:}t-1)$ only through $X(t-1)$. Under these assumptions, Bayes law yields a posterior distribution over latent states from observed rewards given

by:

$$\mathbb{P}\left[X(1{:}t) = k(1{:}t)\,\big|\,R(1{:}t)\right] \propto \mathbb{P}\left[X(1{:}t) = k(1{:}t)\right]\prod_{s=1}^{t}\mathbb{P}\left[R(s)\,\big|\,X(s) = l(s)\right],$$

where we suppress the dependence on cue vectors to shorten notation. The posterior probability $\rho_l(t)$ that latent state $X(t)$ is l based on observations up to trial t is then:

$$\mathbb{P}\left[X(t) = l\big|R(1{:}t)\right] = \sum_{k(1),\dots,k(t-1)=1}^{L}\mathbb{P}\left[X(1{:}t-1) = k(1{:}t-1), X(t) = l\,\big|\,R(1{:}t)\right].$$

From this expression, we see that optimal Bayesian inference requires enumerating over all possible sequences [120] of latent states, the number of which grows exponentially with t. Such computation is generally considered [121] optimistic for human computation [30]. Efforts are made to reduce computation such as with particle [122] methods [17], assuming one transition between states [25], or using a functional form of memory that does [123] not maintain the entire history of observations. [124]

We capture a more general dynamic learning environment similar to [30], by assuming latent states are a Markov chain in which a latent variable transitions to itself from one trial to the next with probability $1 - \gamma(L-1)/L$ and transitions to a new state (all new states being equally-likely) with probability $\gamma(L-1)/L$. This assumption reflects that learning often involve blocks of consecutive trials, or stages, in which rewards are generated in an identical manner. An optimal Bayesian filtering equation could then be used to update beliefs

$$\rho_l(t) \propto \left((1-\gamma)\,\rho_k(t-1) + \frac{\gamma}{L}\right)\mathbb{P}\left[R(t)\big|X(t) = l\right].$$

Unfortunately, this equation requires the learning agent knows the probability distribution of rewards for each latent state. We thus do not think the agent reasons in an optimal Bayesian way. Instead, we initialize beliefs $p_l(0) = 1/L$ and propose that the agent uses an *approximate Bayesian filtering equation* (cf. [31])

$$p_l(t) = \frac{1}{l_0}\left((1-\gamma)\,p_l(t-1) + \frac{\gamma}{L}\right)\phi\left(\frac{E_l(t)}{\sigma(t)}\right), \tag{5}$$

where ϕ is the probability density function of a standard normal random variable; $\sigma(t)$ is the agent's estimate

of the standard deviation of rewards at the start of trial t; and

$$l_0 = \sum_{l=1}^{L} \left((1 - \gamma)\, p_l(t-1) + \frac{\gamma}{L} \right) \phi \left(\frac{E_l(t)}{\sigma(t)} \right) \tag{6}$$

is a normalizing constant to ensure beliefs sum to 1. In other words, we replace the distribution/density   125
function of rewards in the optimal Bayesian filtering equation with a normal density function with mean   126
given by the agent's current estimate $\vec{c}_l(t)'\vec{V}_l(t)$ and standard deviation given by the agent's current estimate   127
$\sigma(t)$. Even though we use a normal density function, rewards do not need to be normally-distributed or even   128
continuous.   129

Measuring expected uncertainty   130

The learning agent may also track the standard deviation $\sigma(t)$. With only one standard deviation, we assume
the learning agent estimates the standard deviation *pooled* over each latent state and each cue vector. We
use the following update:

$$\sigma^2(t+1) = \sigma^2(t) + \beta_0 \left(\overline{E^2}(t) - \sigma^2(t) \right) \tag{7}$$

where $\sigma^2(1) = \sigma_0^2$ is some initial estimate of the variance; β_0 is a constant associability parameter for
variance $\sigma^2(t)$; and $\overline{E^2}(t)$ is the squared prediction error averaged over latent states:

$$\overline{E^2}(t) = \sum_l p_l(t) E_l(t)^2. \tag{8}$$

Variance is also used in other models of latent-state learning, such as the model in Redish et al [11] and the   131
model in Gershman et al [19]. Further, Yu and Dayan [30] proposed a neural mechanism, via acetylcholine   132
signalling, by which a learning agent keeps track of expected uncertainty in changing environments. As a   133
staple to statistical inference, variance is a natural candidate for quantifying expected uncertainty.   134

Measuring unexpected uncertainty   135

Many latent-state learning models assume the number of latent state can grow [11, 15, 17]. Similarly, we use
an *online* mechanism to trigger an addition of new latent states when needed to explain rewards. This
mechanism reflects the need to capture *unexpected uncertainty*, as discussed in Yu an Dayan [30]. It is based
on Page's algorithm [32] for solving a problem known as change point detection (cf. [33]). The idea is to test

whether we can reject our current model in favor of an alternative model with an extra latent state. Specifically, we let L be the number of latent states actively considered on trial t. We keep track of model performance using a statistic $q(t)$:

$$q(t+1) = \max\left(q(t) + \log\left(\frac{\phi(0)}{l_0} \right) - \delta, 0 \right) \tag{9}$$

with a scalar parameter δ and initialized $q(1) = 0$. Importantly, $\phi(0)/l_0$ is the likelihood ratio between a one-state model with expected rewards given by actual rewards $R(t)$ and the current model, where l_0 was the normalizing constant for beliefs at Eq (6). If $q(t)$ exceeds a threshold η, we let

$$q(t+1) = p_{L+1}(t) = 0, \qquad \vec{V}_{L+1}(t) = \frac{R(t)}{\vec{c}(t)'\vec{c}(t)}\vec{c}(t); \qquad \vec{B}_{L+1}(t) = I. \tag{10}$$

Note that $\vec{c}(t)'\vec{V}_{L+1}(t) = R(t)$ so that the current reward is the expected reward for the new state. We then replace L with $L+1$.

Changes in context

Learning often involves changes in context whether it be new surroundings, backdrop, or point in time. Context changes are believed to influence learning in a distinctly different way than a cue influences learning [12]. Our model supposes that these changes in context corrodes current beliefs, whereby beliefs in a latent state on a previous trial are thought to be less informative on the present trial. For a temporal shift in context, our model supposes this corrosion increases with time. While the Gershman (2017) model uses a temporal kernel, we replace beliefs $p_l(t)$ at the end of trial t with

$$(1-\gamma)^{ITI-1}p_l(t) + \left(1 - (1-\gamma)^{ITI-1}\right)/L \tag{11}$$

where ITI is the time between trials within a task phase. This update amounts to repeatedly updating latent state beliefs for $ITI - 1$ iterations according to the Markov chain transition probabilities. Beliefs would become evenly distributed between latent states in the limit as ITI goes to infinity. For a visual or spatial shift in context, we set $ITI = \infty$ in equation above, leading to uniform beliefs $1/L$, i.e. all latent states are believed to be equally likely. As a result, an agent does not carry their beliefs forward to the next trial when there is a spatial or visual change of context.

A temporal shift in context might also alter how associative strengths and beliefs are updated. Gershman et al [19] describes the influence of retrieval on a memory by way of "rumination". This mechanism is

captured in the model of Gershman (2017) [19] and in our model by repeatedly replacing associative strengths $\vec{V}_l(t)$ with the new estimate $\vec{V}_l(t+1)$ performing the updates at the end of trial t again. This process is repeated for a number of trials given by $\min\{\chi, ITI - 1\}$ where χ is a patient-specific parameter.

Putting it all together

Our latent-state model of learning is summarized in Fig 1B and in Algorithm 1. Briefly, a learning agent undergoes the following each trial: they observe cues, generate expectations for rewards, observe a reward, measure error, update their belief in latent states, and update associative strength and measures of uncertainty. In between trials, a learning agent may adjust their beliefs and associative strengths to account for contextual shifts whether they be visual, spatial, or temporal. Further, parameters can be tuned to describe differences in learning between agents: α_0 influences the rate of learning associative strengths, β_0 influences the rate of learning variance, γ influences transitions between latent state, σ_0 influences initial expected uncertainty, ν and δ influence unexpected uncertainty, and χ determines rumination.

Our model is grounded by six major predictions or assumptions about how an agent learns:

1) An agent learns *online*. Model variables are updated on each trial using only current observations and current values of model variables. Consequently, an agent can learn without remembering observations of cues and rewards from prior trials. Their memory and computational requirements grow only in the number of latent states rather than the number of trials, which may be a more realistic reflection of how humans learn. The Rescorla-Wagner model and other models also use online updates for certain model variables [8, 11, 17, 19].

2) An agent uses latent states to *predict rewards*. In deriving our model (see Methods section), we start with the assumption that the agent's goal is to predict rewards. The role of a latent state is to account for different predictions that can be made upon observing cues. By contrast, the role of a latent state could be to discriminate between different sets of cues-reward observations [11, 17], which in turn could be used to predict rewards.

3) Associability, i.e. how quickly associations are updated, depends on beliefs in latent states and cue novelty. Associative strength updates quickly for latent states believed to be active, but slowly for latent states believed to be inactive. This feature is similar to the Mackintosh model [34] in that associability increases when rewards are accurately predicted. However, associative strength also updates quickly when an agent switches their belief to a new latent state to account for unexpected or surprising outcomes, i.e. outcomes in which prediction error is larger relative to the expected uncertainty. This feature is similar to Pearce-Hall model [9], which supposes associability increases in the presence of uncertainty.

Meanwhile, the effort matrix, which tracks how often a cue appears (diagonal entries) and a pair of cues
appear (off-diagonal entries), also determines associability. As a result, associative strength is updated both
more quickly when novel sets of cues are presented.

4) Latent states are relatively stable between trials. Latent states are expected to be the same
from one trial to the next with a certain probability. This assumption causes beliefs in latent states to be
relatively consistent between trials. An alternative is to assume latent states are exchangeable between trials,
allowing beliefs in latent states to shift more sharply between trials [17]. Stability between trials, however,
might degrade with time. Our model assumes that with time, the last trial becomes less and less informative
about the latent state, to the point that the agent eventually believes each latent state is equally likely
regardless of prior beliefs.

5) Beliefs are maintained over multiple latent states. Our model allows an agent to maintain
beliefs over multiple latent states on each trial as opposed to believing in only latent state. For example, an
agent is able to believe that two latent states are equally likely. This allows the agent to explicitly state their
beliefs in competing associations, as is required by some learning tasks [26]. Maintaining beliefs over multiple
states is a common feature for Bayesian models [17, 19, 25, 26]. An alternative is to classify each trial to one
state [11, 35].

```
/* Loop through trials                                              */
for each trial t = 1, . . . , T do
    /* State inference                                              */
    Update latent-state beliefs $p_1(t), \ldots, p_L(t)$ with Eq (5);
    /* Unexpected uncertainty learning                              */
    Update change point statistic $q(t)$ with Eq (9);
    if $q(t+1) > \nu$ then
        /* Add new state                                            */
        Set $\vec{V}_{L+1}(t), \vec{B}_{L+1}(t), \vec{p}_{L+1}(t), q(t+1)$ with Eq (10);
        Update number of latent states $L \leftarrow L + 1$;
    end
    /* Value learning                                               */
    Update associative strengths $\vec{V}_1(t), \ldots, \vec{V}_L(t)$ with Eq (1);
    Update effort matrices $\vec{B}_1, \ldots, \vec{B}_L(t)$ with Eq (4);
    /* Expected uncertainty learning                                */
    Update variance $\sigma^2(t)$ with Eq (7);
    /* Account for context                                          */
    if Context shifts after trial t then
        Repeatedly update associative strength $\vec{V}_l(t)$ with Eq (1) if temporal shift;
        Update latent-state beliefs with Eq (11);
    end
end
```

Algorithm 1: Proposed model of latent-state learning which updates variables *online*, i.e. does not keep track of past cues and rewards.

Overview of simulation

We used simulation to verify that our latent-state model can reproduce a set of historically-important effects observed from experiments on learning (Table 1). Our model was compared to the Rescorla-Wagner (RW) model and two latent-state learning models: the infinite-mixture model of Gershman and Niv in [17], and the Gershman (2017) model in [19]. Other model comparisons are considered in S1 Text. For each model, we computed associative strength per cue and belief in each latent state. For our model, the infinite-mixture model, and the Gershman (2017) model, we defined associative strength as the expected reward conditional

Experiment	Observed effect
Blocking [36]	Associative strength of a neutral cue is blocked when reinforced with a cue already associated with the reward
Overexpectation [37]	Associative strength of two cues weakens when these cues are reinforced together after reinforcing each cue separately
Conditioned inhibition [38]	A cue can inhibit responding to a reward after alternating between reinforcing a separate cue and presenting both cues with a weaker reward or no reward.
Backwards blocking [39]	Associative strength of a cue weakens when a separate cue is reinforced after reinforcing both cues together
Rescorla (2000) [40]	Changes in associative strength can be greater for one cue over another when presented at the same time
Wilson *et al* (1992) [41]	Associative strength of a cue increases more when the cue is a less accurate predictor of a reward (also known as the Hall-Pearce effect [28])
Partial reinforcement [42–44]	Associative strength of a cue is slower to extinguish after partial reinforcement compared to continuous reinforcement (also known as the partial reinforcement extinction effect)
Renewal [12]	Associative strength of cue returns quickly after extinction, particularly when extinction is associated with a different context
Spontaneous recovery [12]	Associative strength of a cue returns more greatly after extinction with a longer interval between extinction and renewal
Memory modification [45, 46]	Associative strength of a cue to a fearful outcome can be weakened if extinction occurs within a certain time window following a single retrieval trial

Table 1. Simulated experiments and main effect observed in each experiment. Models were evaluated based on its ability to reproduce the observed effect for each simulated experiment.

on the cue being presented alone and belief in a latent state as the estimated probability of the latent state given observations. Our model includes pairwise interactions between cues as additional cues and also centers rewards (i.e. $R(t) = -1/2$ and $R(t) = 1/2$ rather than $R(t) = 0$ and $R(t) = 1$). Additional details about the simulation are found in the Methods section.

Blocking and other effects

We first tested whether our model could reproduce learning effects that established the RW model as a powerful model of associative learning (Fig 2). In a blocking experiment, for instance, a learning agent is conditioned to associate a cue (Cue A) with a reward, leading to a high associative strength for Cue A. Afterwards, Cue A is repeatedly presented with another cue (Cue B). The compound is then reinforced with a reward. Even though Cue B is reinforced, Cue B does not acquire the high associative strength as Cue A did. That is, conditioning of Cue A *blocked* conditioning of cue B. The RW model, our latent-state model, and the Gershman (2017) model predict blocking because of the way associative strength is updated in each model: updates depend on prediction error and predictions depend on the sum of associative strengths of

present cues. Consequently, associative strength of Cue B changes slightly, because the compound of an ²¹⁵
excitatory Cue A and neutral Cue B yields small prediction errors. ²¹⁶

Our latent-state model reproduces overexpectation and conditioned inhibition, two other effects explained ²¹⁷
by the RW model. An overexpectation experiment consists of reinforcing two cues separately with a reward ²¹⁸
and then later as a compound. When presented as a compound, associative strengths decrease. A ²¹⁹
conditioned inhibition experiment consists of intermixing the reinforcement of one cue (Cue A) with the ²²⁰
omission of a reward when the cue is presented with another cue (Cue B). Cue A increases in associative ²²¹
strength while Cue B decreases in associative strength. As with blocking, the RW model, our latent-state ²²²
model, and the Gershman (2017) model capture overexpectation and conditioned inhibition because of the ²²³
way associative strength is updated. For overexpectation, expectations are twice actual rewards when the ²²⁴
cues are presented together, yielding negative prediction errors and a decrease in associative strength for each ²²⁵
cue. For conditioned inhibition, the negative associative strength of Cue B negates the positive associative ²²⁶
strength of Cue A so that the sum of their associative strengths predicts the weaker reward. The ²²⁷
infinite-mixture model does not update associative strength in the same way as the other three models, ²²⁸
leading to different predictions for blocking, overexpectation, or conditioned inhibition experiments. ²²⁹

Fig 2. Simulated associative strengths of cues during blocking, overexpectation, and conditioned inhibition experiments, which the Rescorla-Wagner (RW) model famously could explain. Built upon the RW model, our latent-state model and the Gershman (2017) model can also explain these experiments.

Learning effects not explained by the RW model

We also tested whether our latent-state learning model could reproduce historically-important learning effects not predicted by the RW model due to its assumption of constant associability (Fig 2).

Associability depends on beliefs in latent-states

Latent-state learning allows an agent to attribute sudden shifts in associations to new latent states, even when the shift is unsignalled. Associability starts relatively high when learning about a new state, thereby explaining the well-known Pearce-Hall effect [9] in which associability remains high in unpredictable circumstances.

    To illustrate, we find that a shift in latent state beliefs can explain Experiment 1 in [41]. This experiment involved two groups, Group C and Group E. For Group C, the experiment consisted of alternating between reinforcing and not reinforcing two cues (a tone and a light) with a reward (food), followed by reinforcing just the light. Group E had similar experiment conditions, except the tone was omitted in the middle of the experiment on non-reinforced trials. The omission was expected to decrease the associative strength of the light for Group E relative to Group C, which would be observed when the light was paired with food. Surprisingly, Group E responded more favorably to the light than Group C. The leading explanation was that associability of the light was higher for Group E than Group C [10].

Fig 3. Associability depends on latent-state beliefs. Associative strength was simulated for two groups in Experiment 1 from Wilson [41], which pointed to greater associability for the light in Group E than Group C in the final trials of the experiment. Our model predicts that only Group E detects the change in experimental conditions, allowing them to shift their beliefs and have greater associability for the light. Beliefs in the first latent state (solid lines) and the second latent state (dashed lines) are shown for the three models with latent states.

    Our latent-state model offers an explanation: that while indeed the associative strength of the light does decrease for Group E relative to Group C, the pairing of the light solely with food is so surprising to Group

E that they correctly infer the change in experimental conditions thereby shifting to a new latent state to

build new expectations for the light (Fig 3). This shift would be observed as higher associability of the light

in Group E than Group C, agreeing with experimental results. By contrast, the other models for the given

set of parameters do not detect the change in experimental conditions, causing learning agents to respond

more greatly to the light in experimental condition C compared to E.

Other learning effects can also be similarly explained by our model as a shift in latent-state beliefs (Fig 4).

Experiments 1A-B in [40], for example, examined whether a cue (Cue A) would undergo the same change in

associative strength as another cue (Cue B) when presented together, as would be predicted by the RW

model. Rescorla, however, found that Cue A decreased more in associative strength than Cue B. Our

latent-state model explains that differences in associability arises because of a shift in latent states. In

Experiment 1B, for example, the omission of the reward is so surprising to the agent—Cue A was always

reinforced—that learning agents correctly infer a change in conditions and switch their beliefs to a new latent

state in which they do not expect Cues A and B to be reinforced. Since rewards were originally expected to

follow Cue A but not cue B, the associative strength of Cue A experiences the greater change. By contrast,

only the infinite-mixture model captured greater associability of Cue B during stage 2 in experiment 1A.

Otherwise, the other models for the given set of parameters do not detect the change in experimental

conditions, causing cues A and B to change a similar amount during Stage 2.

The partial extinction learning effect (PREE) is also similarly explained by our model as a shift in

latent-state beliefs (Figure 5). Partial reinforcement involves alternating between reinforcing and not

reinforcing a cue. It was found that a response was harder to extinguish after partial reinforcement compared

to continuous reinforcement [42]. Our model explains that an agent is better able to discriminate between

reinforcement and extinction with a continuous reinforcement schedule. This allows the agent to switch their

beliefs to a new latent state and thus experience faster extinction. PREE was also observed even if

continuous reinforcement was administered in between partial reinforcement and extinction [43,44]. In this

case, an agent is still better able to discriminate between reinforcement and extinction with a continuous

reinforcement schedule, but differences between reinforcement schedules are smaller as an agent shifts their

beliefs to a new latent state in both cases. The other three models, however, do not capture greater

extinction for continuous reinforcement compared to partial reinforcement in either of the two PREE

simulation experiments.

Fig 4. Another demonstration of associability depending on beliefs in latent states. Total change in associative strength was simulated during Stage 2 of Experiments 1A-B from Rescorla (2000) [40]. Cue B underwent greater increases in associative strength during Stage 2 than Cue A in both experiments, i.e. associability was higher for Cue B than Cue A. The RW model does not capture these effects since associability is constant, whereas our model captures these effects because latent-state beliefs alters associability. Beliefs in the first latent state (solid lines) and the second latent state (dashed lines) are shown for the three models with latent states.

Associability depends on history of cue presentation

Our model encodes the history of cue presentation in effort matrices $B_l(t)$, endowing associability in our model with unique properties. One example is backwards blocking (Fig 6). A compound of two cues (Cues A and B) is reinforced with a reward followed by only one of the cues (Cue A) being reinforced. In the second part, Cue A gains associative strength while Cue B loses associative strength *even though it is not presented.* The RW model cannot explain backwards blocking, because a cue must be present to change its associative strength. The infinite-mixture and the Gershman (2017) model also did not capture backwards blocking. By contrast, the associative strength of Cue B correctly decreased in the second part for our latent-state model. Our latent-state model first learns the compound predicts a reward. Without additional information, it splits associative strength equally between the two cues. Later, our latent-state model learns Cue A predicts

Fig 5. Partial reinforcement extinction effect. Associative strength of a cue is harder to extinguish after partial reinforcement compared to continuous reinforcement (Experiment 1). This effect is observed even when partial reinforcement is followed by continuous reinforcement prior to extinction (Experiment 2). Our model captures these effects, because an agent is able to better able to discriminate between reinforcement and extinction with continuous reinforcement schedule. The agent can thus shift their beliefs to a new latent state in order to build new associations for extinction. Beliefs in the first latent state are shown for the three models with latent states.

rewards. Reconciling both parts, our latent-state model increases the associative strength of Cue A while decreasing the associative strength of Cue B. In other words, our latent-state model learns about the *difference* in associative strengths between Cues A and B. Mathematically, applying the inverse of $B_l(t)$ to the cue vector $c(t)$ rotates the cue vector from being in the direction of Cue A to be in the direction of Cue A minus Cue B.

Learning with latent-states

To complete our simulation study, we tested if our latent-state model could describe more recent experiments that examine latent-state learning.

Fig 6. Associability depends on history of cue presentation. Associative strength was simulated for a backwards blocking experiment in which associative strength decreases for a cue that is not even presented. In our model, associability depends on the history of cue presentation through effort matrices. After the combined associative strength of Cue A and Cue B is learned, these effort matrices rotate the direction of learning into the direction of the difference of Cue A and Cue B. As a result, associative strength of Cue B decreases even though Cue A is presented alone, thereby allowing our model to capture backwards blocking.

Renewal

Latent-state learning was offered as an explanation of renewal of expectations after extinction and the role that context plays in this renewal [11, 12, 15, 16]. We simulated renewal wherein a cue is reinforced, extinguished, and then reinforced (Fig 7). Two experimental conditions are considered: one in which the same visual/spatial context is provided through each phase and another in which a different visual/spatial context is provided during the extinction phase. Following [8] and [19], context was encoded as a separate cue in all models except our latent-state model. Our latent-state model encodes context as a shift in beliefs. Note for this example, expected rewards are given on each trial to account for the contribution of both the cue and the context rather than associative strength which only accounts for the contribution of the cue.

In both experimental conditions, our latent-state learner slowly detects the shift when arm-reward contingencies first shift and switches their beliefs to a new latent state, effectively consolidating the memory of the first task phase. By detecting this shift, the agent can both construct new associative strengths for each arm and preserve the old associative strengths that were accurate for the first block of trials. These associative strengths are later *recalled* when arm-reward contingencies revert back and the agent switches their beliefs back to the original latent state. Introducing a different context during extinction allowed the learning agent to better detect the underlying structure of the task, recalling the first latent state earlier during reinstatement, resulting in higher associative strength in the final phase. This improvement in detection predicted by the model, however, was very slight and most noticeable in the first few renewal trials. The infinite-mixture model predicts that the learning agent switches beliefs in a new latent state only when a different context is introduced otherwise using one latent state to predict all phases of the task. For the given parameters, the Gershman (2017) model uses only one latent state for all three phases of the task for both the no renewal and renewal conditions. Consequently, other than our model, only the infinite-mixture model

Fig 7. Changes in context influences beliefs. The first experiment examines expected rewards when a response is reinstated after extinction with and without a change in a visual/spatial context. The second experiment examines associative strength of cue when a response is reinstated after extinction with and without a change in a temporal context (i.e. a time delay between trials). In both experiments, our model shows a rapid reinstatement of expectations as the agent switch their beliefs back to the first latent state. Our model also shows that rapid reinstatement is more robust with changes in context, particular in the first few trials of reinstatement. Expected rewards are depicted in the first experiment rather than associative strengths to account for the influence of context on expectations in addition to the cue, since the RW, infinite-mixture, and Gershman (2017) models treat context as an additional cue. Beliefs are shown for the first latent state.

correctly predicts a rapid reinstatement of expectations and only when there is a context shift. 317

Spontaneous recovery 318

Latent-state learning also provides an explanation for how changes in temporal context influences 319

reinstatement. For example, spontaneous recovery is a learning effect wherein the associative strength of a 320

cue is reinstated more strongly after a time delay between extinction and renewal [12]. We simulated 321

spontaneous recovery using the same schedule as renewal, but adding a time delay between extinction and 322

renewal. Our model accounts for time delays by shifting beliefs towards uniform beliefs over latent states. 323

Uniform beliefs reflects that beliefs from a prior trial is less informative on a trial when there is a time delay   324
between trials. As a result, a time delay causes the associative strength of the cue predicted by our model to   325
be about half the associative strength at the end of its initial acquisition (first latent state) and at the end of   326
extinction (second latent state). Without the time delay, the associative strength predicted by our model is   327
simply the associative strength at the end of extinction. Only the Gershman (2017) model adjusts its   328
prediction for context changes due to temporal shifts, but does not capture a more robust return of   329
associative strength due to a temporal shift. The other models also do not capture this more robust return.   330

Memory modification   331

Our last simulation experiment explores the potential role of latent-state learning in memory modification.   332
Monfils et al [45] in a rodent model and later Schiller et al [46] in humans showed that a single *retrieval* trial   333
between acquisition and extinction can reduce the fear response to the cue in latter tests. The time between   334
retrieval and extinction was further shown to modify this effect. Only certain windows of times reduced the   335
fear response. They proposed that the retrieval trial helped to reconsolidate the acquisition memory to be   336
less fearful within this *reconsolidation* window. The Gershman (2017) model [19] includes a computational   337
explanation of this phenomenon. With a similar Bayesian framework to the Gershman (2017) model, our   338
model included similar memory modifications.   339

We simulated the Monfils-Schiller experiment in [19], varying the time between retrieval and extinction   340
(Fig 8). Our model reproduces a qualitatively similar result as the Gershman (2017) model. There is a   341
reconsolidation window which can help reduce the fear memory. For example in both models, associative   342
strength of the cue during testing is smaller for a time of 5 from retrieval to extinction relative to time of 1   343
or 100. Both models predict that beliefs in the first latent state (corresponding primarily with acquisition)   344
remains relative high in the reconsolidation window, allowing the agent to update the acquisition memory to   345
be less fearful. Outside the reconsolidation window, the agent updates the second latent state (corresponding   346
primarily with extinction), leaving the acquisition memory relatively intact. While qualitatively similar   347
between models, the effect of retrieval is more pronounced in our model in terms of differences in magnitude   348
of cue associative strength during testing.   349

Model practicality   350

As a final test, we checked whether our model is a practical choice when looking to explain actual behavioral   351
data. We fit models to choices of participants (n=55) during a learning and decision-making task (see S1   352

Fig 8. Memory modification. Associative strength and beliefs are depicted for three different time delays (1,5, and 100) between a single retrieval and extinction. Both models correctly predict that certain time delays can weaken the associative strength upon testing. The bottom row depicts a 'reconsolidation window' of time delays after retrieval wherein the associative strength on testing is decreased.

Text). Model parameters are estimated using maximum likelihood estimation, assuming choice probability is related to model-predicted expectations of reward for each choice via a soft-max function. Our model provided better model fits in term of Akaike and Bayesian information criteria than other models, thus demonstrating that our model is indeed practical with respect to modeling real human behavior.

Discussion

We presented a computational model of latent-state learning, using the Rescorla-Wagner (RW) model as its foundation, and positing that latent states represent disparate associations between cues and rewards. In a series of simulation experiments, we tested the model's ability to reproduce learning experiments both famously explained and not explained by the RW model and to capture behavior in recent experiments testing latent-state inferences. We then used novel dataset (n=55) of a learning and decision-making task to demonstrate that our model is practical choice for explaining behavioral data. Lastly, we formally derived our latent-state model under the premise of computational rationality, i.e. a learning agent wants to predict

outcomes efficiently.

Our goal was to identify a model of latent-state learning that uses online updates and is sufficiently flexible for describing learning in many experiments. The resulting model makes five critical assumptions/predictions about how an agent learns cue-reward associations: an agent learns online, an agent uses latent states to predict rewards, associability depends on beliefs in latent states and cue novelty, latent states are relatively stable between trials, and beliefs are maintained over multiple latent states. Including these features helps to ensure that the proposed model could examine learning in numerous and diverse experiments, from traditional associative learning experiments to more recent experiments on latent-state inferences. We show that most features fall-out naturally when trying to develop an online approach to reward prediction which uses latent-state inferences. Other models of latent-state learning share some, but not all these features [11, 17, 19].

An agent who learns online, for example, requires only current observations rather than the entire history of observations to update parameters and quantities such as associative strength. Consequently, online approaches have computational and memory requirements that grow in the number of latent states rather than in trials, which could be a more realistic reflection of how humans learn. The infinite-mixture model also uses online updates for all its variables [17]. The Gershman (2017) model uses online updates for certain variables, but keeps track of the history of cues and MAP estimates to update the posterior probability of each latent state [19], whereas the Redish model (2007) updates the state-action value function online but keeps track of the history of cues to update the mean and covariance of cues per latent state [11]. For comparison, an optimal Bayesian approach to latent-state learning would require enumerating over all possible sequences of latent-states, a number that grows exponentially with each trial, imposing a computational and memory burden that is believed to be overly optimistic for human computation [30].

Critically, our latent-state model relies on an approximate Bayesian filter to maintain and update beliefs over latent-states online. Other models of latent-state learning use other approaches to reduce computational burden such serial representations of committed beliefs rather than distributing belief across multiple hypotheses [11], particle methods [15, 17, 19], or maximum a posteriori estimates [19]. Additional assumptions might also be used such as exchangeable trials and binary outcomes [17], one latent state [24], one latent-state transition [25], or known cue-reward associations [26]. Assumptions, however, limit the ability of any model to describe learning in varied experiments. For example, we can capture renewal, spontaneous recovery, and other learning phenomena that require a more general number of latent states and transitions because of our use of an approximate Bayesian filter [31]. We can also capture experiments such as [26] that require beliefs to be maintained over over multiple latent-states. Notably, the experiment [26]

suggested a link between log-beliefs and activity in the orbitofrontal cortex.

Interestingly, our model still updates associative strength in a similar manner to a RW model: change in associative strength is prediction error scaled by associability. In fact, the RW model is a specific case of our latent-state model when there is only one latent state and associability is fixed. For this reason, our latent-state model can reproduce similar learning effects as the RW model such as blocking, overexpectation, and conditioned inhibition. Some latent-state models cannot capture all of these classic learning features [11, 17]. Further, our latent-state model better explained participant behavior than the RW model and the Gershman (2017) model in [19].

However, our latent-state model goes beyond the RW model by allowing associability to vary across trials. This feature is thought to be important for describing learning effects not explained by the RW model, such as the Pearce-Hall effect, and is incorporated into newer models, such as the Hybrid RW model [9, 10, 47]. Associability in our latent-state model involves two components: effort matrices and beliefs in latent-states. Because associability depends on effort matrices, changes in associative strength are rotated into a direction which has not been previously learned, helping to explain backwards blocking. These matrices approximate Fisher information matrices, which are often incorporated into iterative approaches to maximum likelihood estimation [31, 48]. Meanwhile, because associability also depends on beliefs, associative strength can be updated quickly when a learning agent infers a sudden shift in experimental conditions. This feature allowed our latent-state model to capture experimental results from Experiment 1 in Wilson et al (1992) [25] and Experiments 1A–B in Rescorla (2000) [40].

Many of the assumptions/predictions of the presented model are testable. For example, associability for a latent-state is predicted to be directly dependent on the beliefs in that latent-state. Stronger beliefs lead to faster learning, whereas weaker beliefs lead to slower learning. An experiment could be designed to test, for example, if associability decreases by 50% when beliefs are split between two latent-states relative to when beliefs were concentrated on one latent-state. Further, associability is predicted to depend on whether cues had previously been presented together. New combination of cues can lead to quick learning even when individual cues making up the combination have already been presented. Last, latent-state beliefs are predicted to be updated in a non-optimal manner, suggesting one try to induce sub-optimal behavior in an experiment.

Several limitations of the proposed model should be considered. First, we anticipate there are learning effects that our model fails to describe or there are other models that could explain participant behavior. Second, we did not examine to what extent beliefs in latent-states and other variables from our model are encoded in various brain regions. Third, we assume beliefs are maintained over multiple latent states, but

some experiments suggest that an agent is committed to a state and alternates between committed ⁴²⁹ states [35]. One possibility is that beliefs in latent states influence which state is committed and over time, ⁴³⁰ the frequency of committed states mirrors these beliefs. Modalities such as fMRI might then suggest beliefs ⁴³¹ are maintained over multiple states, as in [26], because they lack the temporal resolution needed to detect ⁴³² the alternating committed states. Allowing beliefs to determine committed states might even lead to better ⁴³³ explanations of participant data, as it allows for more flexibility in how states are updated for a given ⁴³⁴ participant. Fourth, there may be better ways to integrate computational mechanisms of how memories are ⁴³⁵ formed into our model [49]. For example, we use online updates in order to avoid memory requirements that ⁴³⁶ grow in the number of trials, but it is possible that humans are either capable of such requirements or use ⁴³⁷ other ways than online updates to integrate past observations. Fifth, some effects (e.g., memory modification ⁴³⁸ effect [45,46]) reproduced by our model might be too subtle to ever be detected experimentally. Sixth, we ⁴³⁹ assume latent states are relatively stable between trials. This assumption may be accurate for describing ⁴⁴⁰ learning in experiments, since experiments often involve blocks of trials in which rewards/cues are generated ⁴⁴¹ by the same rules, but may be less accurate for real-world learning environments. Finally, we did not study ⁴⁴² model performance in terms of predicting rewards as a consequence of using an approximate Bayesian filter. ⁴⁴³ It may useful to know how far from optimal our approximate method is for maximum likelihood estimation ⁴⁴⁴ in various settings. ⁴⁴⁵

In sum, this work establishes the validity of an online model for latent-state inferences that generalizes ⁴⁴⁶ across experiments. Establishing such a framework is a necessary step towards quantifying normative and ⁴⁴⁷ pathological ranges of latent-state inferences (and mediating neurocircuitry) across varied contexts. Overall, ⁴⁴⁸ this work moves us closer to a precise and mechanistic understanding of how humans infer latent-states. ⁴⁴⁹

Methods ⁴⁵⁰

Simulation details ⁴⁵¹

Parameters were fixed throughout the simulation (Table 2). Parameters for alternative models were chosen ⁴⁵² from their respective papers, except for the maximum number of latent states which we fixed at 15 to reduce ⁴⁵³ memory requirements. Our model includes cues throughout simulation to represent pairwise interaction ⁴⁵⁴ terms (i.e. a binary indicator if a pair of cues are present or absent). Our model also centers rewards with ⁴⁵⁵ $R(t) = 1/2$ when a reward is presented and $R(t) = -1/2$ when a reward is not presented. That way, initial ⁴⁵⁶ associative strengths give rise to neutral (i.e. 50-50) expectations for reward. Accompanying code is ⁴⁵⁷

Model	Parameter	Description	Value
Our model	α_0	Associative strength learning rate	0.05
Our model	β_0	Variance learning rate	0.05
Our model	γ	Latent-state transitions	0.05
Our model	σ_0	Initial expected uncertainty	0.5
Our model	ν	Threshold for new state	0.2
Our model	δ	Unexpected uncertainty update	0.6
Our model	χ	Rumination steps	5
Rescorla-Wagner	α	Learning rate	0.15
Infinite-mixture	α	Concentration parameter	1
Infinite-mixture	M	Number of particles	100
Gershman (2017)	α	Concentration parameter	0.1
Gershman (2017)	η	Learning rate	0.3
Gershman (2017)	σ_r^2	Reward variance	0.4
Gershman (2017)	σ_x^2	Cue variance	1
Gershman (2017)	n/a	Rumination steps	3

Table 2. Fixed set of parameters used for all simulated experiments.

publicly-available at https://sites.google.com/site/amylouisecochran/. S1 Text provides further details on $_{458}$
the schedules for how cues and rewards were delivered during each simulation task and examines sensitivity $_{459}$
of our model's predictions to changes in parameters. $_{460}$

Mathematical justification $_{461}$

We justify our choice of model under the premise of *computational rationality* [50], i.e. a learning agent wants
to predict rewards relatively efficiently. An optimistic strategy for the learning agent would be to use
maximize likelihood estimation (MLE) to aid prediction. This would entail starting with a probabilistic
model of rewards defined up to an unknown parameter θ and finding θ to maximize the log-likelihood of
rewards (scaled by $1/t$):

$$\ell(\theta) = \frac{1}{t} \log f\left[R(1{:}t) \,|\, \theta\right]. \tag{12}$$

Treating rewards as a continuous variable, we use f to denote a probability density function. The estimate of $_{462}$
θ together with the reward model predict future rewards. With an eye towards efficiency, we focus on *online* $_{463}$
approaches, such as the Rescorla-Wagner model, and refer the reader to the work in [48] and [31] on $_{464}$
updating latent-variable models online. $_{465}$

Rescorla-Wagner model as an online approach to maximum likelihood estimation

Before we motivate our latent-state model, we explore how one could perform MLE online to predict rewards without latent-states. In the process, we show that the RW model is an online approach to MLE and illuminate a pathway for performing MLE online with latent-states. The most natural modeling choice for rewards is arguably a standard Gaussian regression model, ubiquitous in statistical inference:

$$R(t) = \vec{c}(t)'\vec{\mu} + \epsilon(t)$$

where $\vec{c}(t)$ is the cue vector on trial t (independent variables); $\vec{\mu}$ are unknown parameters (unknown
regression coefficients); and $\epsilon(t)$ is an independent Gaussian random variable (error) with mean zero and
unknown variance ν. Unknown parameters are collected in $\theta = \begin{bmatrix} \vec{\mu}' & \nu \end{bmatrix}'$.

Although we could maximize ℓ directly, we want an online approach to maximization and instead consider applying Newton's method for optimization (i.e., gradient descent), which involves repeatedly updating an estimate $\hat{\theta}$ of θ with a new estimate of the form:

$$\hat{\theta} - \lambda(t)H(t)^{-1}\nabla_\theta \ell(\theta)\Big|_{\theta=\hat{\theta}}$$

for some learning rate $\lambda(t)$ and an appropriate matrix $H(t)$. Matrix $H(t)$ can take many forms and is important for determining how quickly estimates converge to a maximum/minimum. An ideal choice is the Hessian resulting in the Newton-Raphson method, but the Hessian is often noninvertible in MLE. A common alternative is the negative of the Fisher information matrix $\mathcal{I}(\theta)$ scaled by $1/t$ and evaluated at the current estimate [48]. Even with Newton's method, rewards still need to be stored up to trial t, but Newton's method can be turned into an online algorithm using the well-known Robbins-Monro method [51]. We simply replace the gradient with an (stochastic) approximation that depends only on the current trial:

$$\nabla_\theta \ell(\theta) \approx \nabla_\theta \log f\left[R(t) \,|\, \theta\right]$$

Note that each side of the equation is equal in expectation when rewards are independent and
identically-distributed.

Upon replacing the gradient with a stochastic approximation, we arrive at the RW model if we use a

constant learning rate $\lambda(t) = \lambda_0$ and the matrix

$$H(t) = -\frac{1}{\hat{\nu}} \begin{bmatrix} I & 0 \\ 0 & \frac{1}{2\hat{\nu}} \end{bmatrix}.$$

For the Gaussian regression model, matrix $H(t)$ resembles

$$-\frac{1}{t}\mathcal{I}(\theta) = \frac{1}{t}\mathbb{E}\left[\nabla_\theta^2 \log f\left[R(1{:}t)\,|\,\theta\right]\Big|\,\theta\right] = -\frac{1}{\nu} \begin{bmatrix} \frac{1}{t}\sum_{s=1}^t \vec{c}(s)\vec{c}(s)' & 0 \\ 0 & \frac{1}{2\nu} \end{bmatrix}.$$

evaluated at $\hat{\theta} = \begin{bmatrix} \vec{\mu}' & \hat{\nu} \end{bmatrix}'$ We simply replaced $\frac{1}{t}\sum_t \vec{c}(t)\vec{c}(t)'$ with the identity matrix I, which ensures $H(t)$ is invertible. Putting the online approach together with the fact that

$$\nabla_\theta \log f\left[R(t)\,|\,\theta\right] = \frac{1}{\nu} \begin{bmatrix} \vec{c}(t)(R(t) - \vec{c}(t)'\vec{\mu}) \\ -\frac{1}{2} + \frac{1}{2\nu}(R(t) - \vec{c}(t)'\vec{\mu})^2 \end{bmatrix}$$

for the Gaussian regression model, we arrive at the following online update for the estimate $\hat{\theta}$:

$$\hat{\theta} + \lambda_0 \begin{bmatrix} \vec{c}(t)(R(t) - \vec{c}(t)'\vec{\hat{\mu}}) \\ (R(t) - \vec{c}(t)'\vec{\hat{\mu}})^2 - \hat{\nu} \end{bmatrix}.$$

Critically, estimates $\vec{\hat{\mu}}$ are updated exactly as in an RW model:

$$\vec{\hat{\mu}} + \lambda_0 \vec{c}(t)(R(t) - \vec{c}(t)\vec{\hat{\mu}}).$$

In other words, the RW model is an online approach to maximum likelihood estimation. 472

Latent-state learning as an online approach to expectation-maximization 473

Motivated by the previous argument, we now explore how one could perform MLE online to predict rewards with latent-states. A natural choice for modeling rewards is to use a Markov chain of Gaussian regression models. Latent random variable $X(t) \in \{1, \ldots, L\}$ represents the active hypothesis on trial t. Given the active hypothesis $X(t) = l$, rewards are described by a Gaussian linear regression model:

$$R(t) = \vec{c}(t)'\vec{\mu}_l + \epsilon(t)$$

where $\vec{\mu}_1, \ldots, \vec{\mu}_L$ are unknown parameters and $\epsilon(t)$ is an independent Gaussian random variable with mean
zero and unknown variance ν. The extension to latent-state learning thus leads to L different Gaussian
regression models for rewards rather than one. As discussed in the Model subsection, we assume $X(t)$ is a
Markov chain that transitions to itself from one trial to the next with probability $1 - \gamma(L-1)/L$ and
transitions to a new state (all new states being equally-likely) with probability $\gamma(L-1)/L$. Parameter γ and
initial probabilities for $X(0)$ are assumed to be known/given, leaving unknown parameters collected in
$\theta = \begin{bmatrix} \vec{\mu}_1' & \cdots & \vec{\mu}_L' & \nu \end{bmatrix}'$.

In a latent-variable setting, a popular algorithm to perform MLE is expectation-maximization (EM) [52]. An iterative algorithm, EM improves upon a current estimate $\hat{\theta}$ at each iteration in two steps. Fixing the current trial number t, an *expectation* step uses a current estimate $\hat{\theta}$ to calculate the expectation of the complete log-likelihood of rewards and latent states given rewards and scaled by $1/t$:

$$
\begin{aligned}
Q(\theta|\hat{\theta}) &= \frac{1}{t} \mathbb{E}\left[\log f\left[R(1{:}t), X(1{:}t) \,\middle|\, \theta\right] \middle| R(1{:}t), \hat{\theta}\right] \\
&= \frac{1}{t} \sum_{s=1}^{t} \mathbb{E}\left[\log f\left[R(s) \,\middle|\, X(s), \theta\right] + \log \mathbb{P}\left[X(s) \,\middle|\, X(s-1)\right] \middle| R(1{:}t), \hat{\theta}\right]
\end{aligned}
$$

A *maximization* step searches for a maximum of $Q(\theta|\hat{\theta})$ over θ. The maximum is then used as the estimate in
the next iteration. The crux of the EM method is that $Q(\theta|\hat{\theta})$ is easier to maximize than the log-likelihood
$\ell(\theta)$ of the observed data and that improvement in $Q(\theta|\hat{\theta})$ guarantees improvement in $\ell(\theta)$.

Mirroring our argument for the Rescorla-Wagner model, we can replace the maximization step with a Newton update:

$$
\hat{\theta} - \lambda(t) H(t)^{-1} \nabla_\theta Q(\theta|\hat{\theta}) \Big|_{\theta=\hat{\theta}}
$$

for appropriate learning rate $\lambda(t)$ and matrix $H(t)$. Even in a latent-variable setting, one choice for $H(t)$ is still the negative of the Fisher's information matrix scaled be $1/t$ and evaluated at $\hat{\theta}$, but Newton's method still requires all rewards up to trial t. To recover an online approach, we use a stochastic approximation to $\nabla_\theta Q(\theta|\hat{\theta}(t))$ in the maximization step:

$$
\begin{aligned}
\nabla_\theta Q(\theta|\hat{\theta}(t)) &\approx \nabla_\theta \mathbb{E}\left[\log f\left[R(t) \,\middle|\, X(t), \theta\right] + \log \mathbb{P}\left[X(t) \,\middle|\, X(t-1)\right] \middle| R(1{:}t), \hat{\theta}\right] \\
&= \sum_{l=1}^{L} \mathbb{P}\left[X(t) = l | R(1{:}t), \hat{\theta}\right] \nabla_\theta \log f\left[R(t) \,\middle|\, X(t) = l, \theta\right]
\end{aligned}
$$

We then replace $\mathbb{P}\left[X(t) = l \,|\, R(1{:}t), \hat{\theta}\right]$, which is computed using all the rewards, with latent-state beliefs $p_l(t)$ defined at Eq. 5 which is computed online. This replacement yields the stochastic approximation

$$\nabla_\theta Q(\theta|\hat{\theta}(t)) \approx \sum_{l=1}^{L} p_l(t) \nabla_\theta \log f\left[R(s) \,\big|\, X(s) = l, \theta\right]$$

$$= \frac{1}{\nu} \begin{bmatrix} p_1(t)\vec{c}_1(t)\left(R(t) - \vec{c}_1(t)'\vec{\mu}_l\right) \\ \vdots \\ p_L(t)\vec{c}_L(t)\left(R(t) - \vec{c}_L(t)'\vec{\mu}_L\right) \\ -\frac{1}{2} + \frac{1}{2\nu}\sum_{l=1}^{L} p_l(t)\left(R(t) - \vec{c}_l(t)'\vec{\mu}_l\right)^2 \end{bmatrix}$$

The last equality follows from our assumption that rewards are a Markov chain of Gaussian regression models. [484]

Upon replacing the gradient in Newton's method with this stochastic approximation, we can arrive at our latent-state learning model if we use $\lambda(t) = \alpha_0$ and matrix

$$H(t) = -\frac{1}{\hat{\nu}} \begin{bmatrix} B_1(t) & 0 & \cdots & 0 & 0 \\ 0 & B_2(t) & & 0 & 0 \\ \vdots & & \ddots & & \vdots \\ 0 & 0 & & B_L(t) & 0 \\ 0 & 0 & \cdots & 0 & \frac{1}{2\hat{\nu}} \end{bmatrix},$$

where we used the definition of $B_l(t)$ at Eq. 4. For a Markov chain of Gaussian regression models, matrix $H(t)$ resembles the negative of the Fisher information matrix scaled by $1/t$:

$$-\frac{1}{t}\mathcal{I}(\theta) = \frac{1}{t}\mathbb{E}\left[\nabla_\theta^2 \log f\left[R(1{:}t), X(1{:}t) \,\big|\, \theta\right]\Big|\theta\right]$$

$$= -\frac{1}{\nu} \begin{bmatrix} \frac{1}{t}\sum_{s=1}^{t} \rho_1(s,\theta)\vec{c}_1(s)\vec{c}_1(s)' & & \cdots & & 0 \\ & \ddots & & & \vdots \\ \vdots & & \frac{1}{t}\sum_{s=1}^{t} \rho_L(s,\theta)\vec{c}_L(s)\vec{c}_L(s)' & 0 \\ 0 & & \cdots & 0 & \frac{1}{2\nu} \end{bmatrix}$$

where $\rho_l(s,\theta) = \mathbb{P}\left[X(s) = l|\theta\right]$. We simply replaced diagonal blocks with $B_l(t)$ which ensures $H(t)$ is invertible and replaced $\rho_l(s,\theta)$ with latent state beliefs $p_l(s)$. With these choices for $\lambda(t)$ and $H(t)$, we

update $\hat{\theta}$ online according to:

$$\hat{\mu}_1 + \alpha_0 p_1(t) B_1(t)^{-1} \vec{c}_1(t) \left(R(t) - \vec{c}_1(t)' \vec{\hat{\mu}}_1 \right)$$

$$\vdots$$

$$\hat{\mu}_L + \alpha_0 p_L(t) B_L(t)^{-1} \vec{c}_L(t) \left(R(t) - \vec{c}_L(t)' \vec{\hat{\mu}}_L \right)$$

$$\hat{\nu} + \alpha_0 \left(\sum_{l=1}^{L} p_l(t) \left(R(t) - \vec{c}_l(t)' \vec{\hat{\mu}}_l \right)^2 - \hat{\nu} \right)$$

We thus arrive at our latent-state model for updating associative strengths at Eq.(1). We can also recover   485
our latent-state model for updating variance at Eq (7), if we replace the learning rate α_0 with a different   486
learning rate β_0. This completes our motivation for our latent-state model.   487

Supporting information   488

S1 Text. Additional support and details. This supplement is divided into five sections: Extensions,   489
Psychological experiment, Comparison to other learning models, Simulation details, and Sensitivity of   490
predictions to parameters. The first section discusses how to extend the latent-state model to allow for   491
multidimensional rewards and decision-making. The second section provides an example of model fits to   492
behavioral data from a psychological experiment. The third section examines simulated behavior from other   493
learning models. The fourth section provides additional details on simulated experiments. The last section   494
examines whether the ability of our model to reproduce target effects in simulation is sensitive to changes in   495
parameters.   496

References   497

1. Huys QJ, Guitart-Masip M, Dolan RJ, Dayan P. Decision-theoretic psychiatry. Clinical Psychological   498
Science. 2015;3(3):400–421.   499

2. Sutton RS, Barto AG, et al. Reinforcement learning: An introduction. MIT press; 1998.   500

3. Daw ND, O'doherty JP, Dayan P, Seymour B, Dolan RJ. Cortical substrates for exploratory decisions   501
in humans. Nature. 2006;441(7095):876.   502

4. Doll BB, Simon DA, Daw ND. The ubiquity of model-based reinforcement learning. Current opinion   503
in neurobiology. 2012;22(6):1075–1081.   504

5. Wilson R, Collins A. Ten simple rules for the computational modeling of behavioral data. 2019;.

6. Dayan P, Berridge KC. Model-based and model-free Pavlovian reward learning: revaluation, revision, and revelation. Cognitive, Affective, & Behavioral Neuroscience. 2014;14(2):473–492.

7. Gläscher J, Daw N, Dayan P, O'Doherty JP. States versus rewards: dissociable neural prediction error signals underlying model-based and model-free reinforcement learning. Neuron. 2010;66(4):585–595.

8. Rescorla RA, Wagner AR, et al. A theory of Pavlovian conditioning: Variations in the effectiveness of reinforcement and nonreinforcement. Classical conditioning II: Current research and theory. 1972;2:64–99.

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

10. Le Pelley ME. The role of associative history in models of associative learning: A selective review and a hybrid model. The Quarterly Journal of Experimental Psychology Section B. 2004;57(3b):193–243.

11. Redish AD, Jensen S, Johnson A, Kurth-Nelson Z. Reconciling reinforcement learning models with behavioral extinction and renewal: implications for addiction, relapse, and problem gambling. Psychological review. 2007;114(3):784.

12. Bouton ME. Context and behavioral processes in extinction. Learning & memory. 2004;11(5):485–494.

13. Vervliet B, Craske MG, Hermans D. Fear extinction and relapse: state of the art. Annual review of clinical psychology. 2013;9:215–248.

14. Redish AD. Addiction as a computational process gone awry. Science. 2004;306(5703):1944–1947.

15. Gershman SJ, Blei DM, Niv Y. Context, learning, and extinction. Psychological review. 2010;117(1):197.

16. Gershman SJ, Hartley CA. Individual differences in learning predict the return of fear. Learning & behavior. 2015;43(3):243–250.

17. Gershman SJ, Niv Y. Exploring a latent cause theory of classical conditioning. Learning & behavior. 2012;40(3):255–268.

18. Gershman SJ, Norman KA, Niv Y. Discovering latent causes in reinforcement learning. Current Opinion in Behavioral Sciences. 2015;5:43–50.

19. Gershman SJ, Monfils MH, Norman KA, Niv Y. The computational nature of memory modification. Elife. 2017;6:e23763.

20. Gershman SJ, Jones CE, Norman KA, Monfils MH, Niv Y. Gradual extinction prevents the return of fear: implications for the discovery of state. Frontiers in behavioral neuroscience. 2013;7:164.

21. Tolman EC. Cognitive maps in rats and men. Psychological review. 1948;55(4):189.

22. Stalnaker TA, Cooch NK, Schoenbaum G. What the orbitofrontal cortex does not do. Nature neuroscience. 2015;18(5):620.

23. Wikenheiser AM, Schoenbaum G. Over the river, through the woods: cognitive maps in the hippocampus and orbitofrontal cortex. Nature Reviews Neuroscience. 2016;17(8):513.

24. Niv Y, Daniel R, Geana A, Gershman SJ, Leong YC, Radulescu A, et al. Reinforcement learning in multidimensional environments relies on attention mechanisms. Journal of Neuroscience. 2015;35(21):8145–8157.

25. Wilson RC, Niv Y. Inferring relevance in a changing world. Frontiers in human neuroscience. 2012;5:189.

26. Chan SC, Niv Y, Norman KA. A probability distribution over latent causes, in the orbitofrontal cortex. Journal of Neuroscience. 2016;36(30):7817–7828.

27. Farrell S, Lewandowsky S. Computational models as aids to better reasoning in psychology. Current Directions in Psychological Science. 2010;19(5):329–335.

28. Hall G, Pearce JM. Latent inhibition of a CS during CS–US pairings. Journal of Experimental Psychology: Animal Behavior Processes. 1979;5(1):31.

29. Agrawal S, Goyal N. Thompson sampling for contextual bandits with linear payoffs. In: International Conference on Machine Learning; 2013. p. 127–135.

30. Angela JY, Dayan P. Uncertainty, neuromodulation, and attention. Neuron. 2005;46(4):681–692.

31. Cappé O. Online EM algorithm for hidden Markov models. Journal of Computational and Graphical Statistics. 2011;20(3):728–749.

32. Page ES. Continuous inspection schemes. Biometrika. 1954;41(1/2):100–115.

33. Granjon P. The CuSum algorithm-a small review. 2013;.

34. Mackintosh NJ. A theory of attention: variations in the associability of stimuli with reinforcement. Psychological review. 1975;82(4):276.

35. Rich EL, Wallis JD. Decoding subjective decisions from orbitofrontal cortex. Nature neuroscience. 2016;19(7):973.

36. Kamin LJ. Predictability, surprise, attention, and conditioning. 1967;.

37. Lattal KM, Nakajima S. Overexpectation in appetitive Pavlovian and instrumental conditioning. Animal Learning & Behavior. 1998;26(3):351–360.

38. Rescorla RA. Pavlovian conditioned inhibition. Psychological Bulletin. 1969;72(2):77.

39. Miller RR, Matute H. Biological significance in forward and backward blocking: Resolution of a discrepancy between animal conditioning and human causal judgment. Journal of Experimental Psychology: General. 1996;125(4):370.

40. Rescorla RA. Associative changes in excitors and inhibitors differ when they are conditioned in compound. Journal of Experimental Psychology: Animal Behavior Processes. 2000;26(4):428.

41. Wilson PN, Boumphrey P, Pearce JM. Restoration of the orienting response to a light by a change in its predictive accuracy. The Quarterly Journal of Experimental Psychology Section B. 1992;44(1b):17–36.

42. Capaldi E. The effect of different amounts of alternating partial reinforcement on resistance to extinction. The American Journal of Psychology. 1957;.

43. Jenkins HM. Resistance to extinction when partial reinforcement is followed by regular reinforcement. Journal of Experimental Psychology. 1962;64(5):441.

44. Theios J. The partial reinforcement effect sustained through blocks of continuous reinforcement. Journal of Experimental Psychology. 1962;64(1):1.

45. Monfils MH, Cowansage KK, Klann E, LeDoux JE. Extinction-reconsolidation boundaries: key to persistent attenuation of fear memories. science. 2009;324(5929):951–955.

46. Schiller D, Monfils MH, Raio CM, Johnson DC, LeDoux JE, Phelps EA. Preventing the return of fear in humans using reconsolidation update mechanisms. Nature. 2010;463(7277):49.

47. Li J, Schiller D, Schoenbaum G, Phelps EA, Daw ND. Differential roles of human striatum and amygdala in associative learning. Nature neuroscience. 2011;14(10):1250.

48. Cappé O, Moulines E. On-line expectation–maximization algorithm for latent data models. Journal of the Royal Statistical Society: Series B (Statistical Methodology). 2009;71(3):593–613.

49. Burgess N, Becker S, King JA, O'Keefe J. Memory for events and their spatial context: models and experiments. Philosophical Transactions of the Royal Society of London Series B: Biological Sciences. 2001;356(1413):1493–1503.

50. Gershman SJ, Horvitz EJ, Tenenbaum JB. Computational rationality: A converging paradigm for intelligence in brains, minds, and machines. Science. 2015;349(6245):273–278.

51. Robbins H, Monro S. A stochastic approximation method. In: Herbert Robbins Selected Papers. Springer; 1985. p. 102–109.

52. Dempster AP, Laird NM, Rubin DB. Maximum likelihood from incomplete data via the EM algorithm. Journal of the royal statistical society Series B (methodological). 1977; p. 1–38.