RESEARCH ARTICLE

Ambiguity drives higher-order Pavlovian learning

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Abstract

In the natural world, stimulus-outcome associations are often ambiguous, and most associations are highly complex and situation-dependent. Learning to disambiguate these complex associations to identify which specific outcomes will occur in which situations is critical for survival. Pavlovian occasion setters are stimuli that determine whether other stimuli will result in a specific outcome. Occasion setting is a well-established phenomenon, but very little investigation has been conducted on how occasion setters are disambiguated when they themselves are ambiguous (i.e., when they do not consistently signal whether another stimulus will be reinforced). In two preregistered studies, we investigated the role of higher-order Pavlovian occasion setting in humans. We developed and tested the first computational model predicting direct associative learning, traditional occasion setting (i.e., 1st-order occasion setting), and 2nd-order occasion setting. This model operationalizes stimulus ambiguity as a mechanism to engage in higher-order Pavlovian learning. Both behavioral and computational modeling results suggest that 2nd-order occasion setting was learned, as evidenced by lack and presence of transfer of occasion setting properties when expected and the superior fit of our 2nd-order occasion setting model compared to the 1st-order occasion setting or direct associations models. These results provide a controlled investigation into highly complex associative learning and may ultimately lead to improvements in the treatment of Pavlovian-based mental health disorders (e.g., anxiety disorders, substance use).

Author summary

In everyday life, we learn to associate various situations with various outcomes. For example, perhaps a specific person usually receives praise (outcome) when giving a public speech (situation), but if they give a speech after a particularly charismatic speaker...
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Introduction
Real-life learning is filled with associations that range from being very simple to very complex. For example, most people have learned that when a streetlight turns green, cars will start driving. This association is simple and consistent with very few exceptions. As a more complicated example, imagine a person who is about to give a public speech. Will the speech go well? It depends. What is the speech topic? How competent is the speaker on that topic? Who is the audience? How much did the speaker prepare? How well did the speaker sleep the night before? There are many situational factors that influence whether a speech will go well, and there are many life examples like this in which it is unclear or ambiguous as to whether a given stimulus/situation will result in a specific outcome. To date, most research has focused on simple or moderately complex associative learning, but very little work has been conducted on highly complex associative learning. By investigating highly complex and ambiguous associations, we take an important step towards understanding how humans learn about real-life associations, which are often highly complex.

The most prominent experimental paradigm for learning stimulus-outcome associations is Pavlovian conditioning, in which the individual learns associations between conditional stimuli (CSs) and their outcomes (i.e., unconditional stimuli; USs). The CS+ is a CS that is paired with the US, whereas the CS− is paired with the absence of the US. In many experiments, the CSs tend to be unambiguous stimuli—meaning, they always (or almost always) predict the presence or absence of the US. However, in the real world, it is rare for CSs to be truly unambiguous. For example, whether a speech (CS) will result in a specific outcome—such as praise, rejection, or something in between—is usually ambiguous and depends on many situational factors.

One of the primary Pavlovian experimental designs investigating ambiguous CS/US associations is occasion setting [1–8], in which one stimulus (i.e., the occasion setter) indicates whether an ambiguous CS will result in the US. For example, perhaps a specific child is usually talkative during dinner, but the parent is interested in figuring out which situations lead the child to be quiet during dinner. From a Pavlovian perspective, the parent is interested in learning which stimuli, situational factors, or contexts determine whether the child (CS) will talk
(US) or be silent (no US) during dinner. In this case, perhaps when the child spends time with a particular friend in the afternoon, the child is quiet during dinner. Theoretically, spending time with the friend (occasion setter) would signal that the child (CS) will be quiet during dinner (no US), but not spending time with the friend would signal that the child (CS) will be talkative during dinner (US).

There are two types of occasion setters: positive and negative. Positive occasion setters signal that the CS will predict the US (i.e., the CS predicts the US only if the positive occasion setter was presented). The public speech example would be positive occasion setting if the person believes their speech (CS) will result in rejection (US) only if they give a speech after a particularly charismatic and engaging speaker (positive occasion setter). Negative occasion setters signal that the CS will not predict the US (i.e., the CS predicts the US unless the negative occasion setter was presented). The friend/child example above is an example of negative occasion setting, where the friend (negative occasion setter) signals that the child (CS) will be quiet during dinner (no US).

Occasion setting is thought to operate via modulation [1,7], where the occasion setter affects the CS/US association. The modulation account posits that stimuli are arranged hierarchically, where higher-order learning (i.e., occasion setting) affects lower-order learning (i.e., direct associations: learning that a CS directly predicts the presence or absence of the US). Many studies have been conducted on traditional occasion setting, described above ([1,2,7–18]; hereafter referred to as 1st-order occasion setting), but there is very limited research on 2nd-order occasion setting, including studies investigating it directly [19] or indirectly [9,16,20,21]. We define 2nd-order occasion setters as stimuli that determine how ambiguous 1st-order occasion setters will affect the CS/US association. Using our example above, the child (CS) is ordinarily talkative during dinner (US) unless they see their friend that day (1st-order negative occasion setter). Converting this to a 2nd-order occasion setting example, perhaps the friend only sometimes causes the child to be quiet during dinner, making the friend an ambiguous 1st-order occasion setter. A 2nd-order occasion setter would determine whether the friend (1st-order occasion setter) will cause the child (CS) to be quiet during dinner (no US). Perhaps the child’s grandparent (2nd-order positive occasion setter) gives the child good advice regarding the importance of family time that causes the child (CS) to talk during dinner (US)–even after the child sees their friend in the afternoon (1st-order occasion setter). This type of example more closely resembles the complexities of real-life associative learning, where stimulus-outcome associations are usually ambiguous and dependent on situational factors. See Fig 1 for our model of 2nd-order occasion setting, in which higher-order learning can only occur when lower-order stimuli are ambiguous (i.e., when lower-order stimuli are sometimes followed by the US).

The Present report

While 2nd-order occasion setting is a theoretically plausible learning process, there are no clear demonstrations of it. Additionally, there are no formal models that predict 2nd-order occasion setting—perhaps because 2nd-order occasion setting has not been explicitly demonstrated. Thus, there were two goals in the present experiments: 1) determine whether 2nd-order occasion setting can be learned (as a model of highly complex associative learning), and 2) evaluate our computational model of 2nd-order occasion setting to see if it is a more accurate predictor of learning than simpler models (i.e., 1st-order occasion setting or direct learning). To this end, we conducted two mirror-image experiments: a 2nd-order negative occasion setting experiment (Experiment 1) and a 2nd-order positive occasion setting experiment (Experiment 2). To address the first goal, we trained multiple stimuli across discriminations intended to
Fig 1. Hierarchical Model of 2nd-Order Occasion Setting. Using example from main text of child, friend, and grandparent: a) direct associative learning, b) 1st-order occasion setting, and c) 2nd-order occasion setting. Panels d-j display: d) direct excitation, e) direct inhibition, f) 1st-order positive occasion setting, g) 1st-order negative occasion setting, h) 2nd-order positive occasion setting, i) 2nd-order negative occasion setting, and j) the total model with all associations. Our mathematical model is presented at the bottoms of panels d-j, where black/bold variables are active (i.e., values greater than 0), and gray variables are inactive (i.e., values are 0). See Table 3 for details on formulas. In the figure, circles are stimuli: unconditional stimulus (US), conditional stimulus (CS), 1st-order occasion setter (OS1), and 2nd-order occasion setter (OS2). Blue arrows indicate direct excitation; blue line segments indicate positive occasion setting; red line segments indicate direct inhibition or negative occasion setting; yellow glow indicates CS ambiguity; purple glow indicates OS1 ambiguity; blue USs indicate US delivery; and red USs indicate US omission. While we suggest that stimulus ambiguity is a dimensional, learned property, we present it as present/absent in the figure for simplicity. Thick arrows/lines indicate activated pathways; thin arrows/lines indicate deactivated pathways. Stimulus ambiguity is required for higher-order associative learning: 1st-order occasion setting is learned only if the CS is ambiguous and has been trained with an OS1; 2nd-order occasion setting is only learned if the CS and OS1 are ambiguous and if the CS has been trained with an OS2. CSs have a direct predictive relationship with the US. If the CS is ambiguous (i.e., sometimes predicts the US, sometimes predicts absence of the US), then attention is broadened to other stimuli or contextual factors (i.e., to the OS1); if a stimulus that disambiguates CS reinforcement is identified and is less salient than the CS, it becomes an OS1. The OS1 modulates the CS/US association. If OS1 consistently excites the CS/US association, then OS1 is a positive OS1; if OS1 consistently inhibits the CS/US association, then OS1 is a negative OS1. If OS1 sometimes excites and sometimes inhibits the CS/US association (i.e., OS1 is ambiguous), then attention is broadened to other stimuli or contextual factors (i.e., OS2) that disambiguate how the OS1 affects the CS/US association. If a stimulus disambiguates the effect of OS1 on the CS/US association and is presumably less salient than the OS1, 2nd-order occasion setting is learned. If the OS2 consistently disables the OS1’s 1st-order positive occasion setting ability, the OS2 is a 2nd-order negative occasion setter. If OS2 consistently disables OS1’s 2nd-order negative occasion setting ability, then OS2 is a 2nd-order positive occasion setter. Additionally, each hierarchical level
produce learning across three hierarchical levels: direct learning, 1st-order occasion setting, and 2nd-order occasion setting. We conducted specific tests (i.e., transfer tests) to test whether 1st-order and 2nd-order occasion setting were indeed learned, in which we would expect the occasion setters to transfer their effects to lower-order stimuli who underwent similar occasion setting training but not to stimuli that did not undergo occasion setting training. As an additional assessment, we predicted that each hierarchical level would be orthogonal—meaning, a stimulus could signal outcomes in each of the three levels (e.g., Stimulus A would be a CS+ when presented alone, a 1st-order positive occasion setter when preceding a CS, and a 2nd-order negative occasion setter when preceding a 1st-order occasion setter and CS; [1,2,7,15,22]). To address the second goal, we conducted Bayesian hierarchical modeling to evaluate model fit using our novel 2nd-order occasion setting model and contrasted its predictions with our simplified 1st-order occasion setting model and our even more simplified direct associations model. If the data fit our 2nd-order occasion setting model better than the 1st-order occasion setting model and direct associations model, this would provide computational support that 2nd-order occasion setting was indeed learned. For a list of all specific hypotheses and analyses details, please see our pre-registrations (Experiment 1: https://osf.io/n2c6v, Experiment 2: https://osf.io/hxcfs).

In both experiments, we trained three families of stimuli to produce direct associative learning, 1st-order occasion setting, and/or 2nd-order occasion setting (Fig 2). In Experiment 1, the stimulus families were the “ABC” stimuli, “TJK” stimuli, and “Direct Learning” stimuli (i.e., G+, H-, and R+). When multiple stimuli were presented within a trial, they were presented serially with inter-stimulus intervals to facilitate occasion setting [23]. Specifically, within the “ABC” stimulus family, C was trained as an ambiguous CS that predicted no US on its own but predicted the US when preceded by putative 1st-order positive occasion setter B (i.e., C-, BC+). Stimulus A acted as a putative 2nd-order negative occasion setter, so when A preceded BC, no US was delivered (i.e., AB+, A+). Additionally, B was non-reinforced on its own, and A acted as B’s putative 1st-order positive occasion setter (i.e., A+, AB+). Lastly, A was reinforced when presented on its own (i.e., A+). Thus, the stimulus contingencies were C-, BC+, ABC-, B-, AB+, A+. We hypothesized that stimulus A would have three hierarchical values (2nd-order negative occasion setter, 1st-order positive occasion setter, unambiguous CS+), B would have two hierarchical values (1st-order positive occasion setter, ambiguous CS-), and C would have one hierarchical value (i.e., ambiguous CS-). This training occurred prior to Transfer Test 1 and Transfer Test 2.

Also in Experiment 1, the “TJK” family was trained with identical meanings as the “ABC” family, substituting T for A, J for B, and K for C. The difference was that JK was trained without T before Transfer Test 1 (i.e., JK was trained only in 1st-order positive occasion setting), but JK was trained with T before Transfer Test 2 (i.e., in 2nd-order negative occasion setting). Thus, whereas the ABC family was trained in 2nd-order occasion setting prior to both transfer tests, the TJK family was trained just in 1st-order positive occasion setting before Transfer Test 1 (i.e., JK+, K-, J-) and was trained in 2nd-order negative occasion setting prior to Transfer Test 2 (i.e., K-, JK+, TJK-, J-, TJ+, T+). When testing A’s ability to transfer its 2nd-order negative occasion setting ability to JK+ (i.e., AJK), we predicted A would have less of an effect on JK+ at Transfer Test 1 than Transfer Test 2 because at Transfer Test 1, JK+ had not undergone 2nd-order occasion setting training yet. Lastly, G+ and H- were trained as an unambiguous CS+ and CS-, respectively, prior to Transfer Test 1. R was trained as an unambiguous
CS+ prior to Transfer Test 1 as part of ABR+ and TJR+ (where “AB” and “TJ” had no influence on R’s reinforcement). This was done so that participants would not simply assume that A or T followed by two other stimuli would result in no US.

Due to the large number of trial types trained and tested, we conducted several sub-phases in uniform order across participants (see Table A in S7 Text for details). Within each sub-

Fig 2. Experiment 1 (2nd-Order Negative Occasion Setting) Trial Design. Each colored box represents a trial type. Gray boxes represent what was shown visually on screen. Inter-trial intervals (ITIs) and inter-stimulus intervals (ISIs) included a gray screen with a fixation cross (“+”). Duration of each trial component is shown at top of each trial type. Rating slide is shown in abbreviated form, and visual analog scale was used to rate US Expectancy. Images of nature scenes, shapes, and auditory stimuli indicate experimental stimuli (i.e., CSs, occasion setters). Auditory stimuli are indicated below slides in horizontal auditory band. Violin symbol indicates violin sound, static screen indicates white noise, and dollar sign indicates cash register sound. None of the auditory symbols were shown on screen during the experiment. Gold coin indicates monetary reward (US). Black arrow pointing to the right for each trial type indicates chronological component sequence during trials. All stimuli are counterbalanced across participants within stimulus category: G/H (Unambiguous CSs), C/K (Ambiguous CSs), B/J (1st-order occasion setters), and A/T (2nd-order occasion setters). All trial types for Experiment 1 are shown except TJR+, which is identical to ABR+, except T and J stimuli are substituted for A and B. Experiment 2 (2nd-order POS) design is mirror image of Experiment 1 in which all trial types reinforced in Experiment 1 were not reinforced in Experiment 2, and all trial types not reinforced in Experiment 1 were reinforced in Experiment 2. The only exceptions are G+ and H-, which remained a CS+ and CS-, respectively, in each study. Images of violin, gold coin, confetti, fractals, and dollar sign were obtained from https://openclipart.org.

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phase, each trial type received usually 5–10 training trials, and trial order was pseudo-randomized. During the first training phase (Training 1), participants were first trained in 1\(^{st}\)-order positive occasion setting with the “JK” stimuli (J-, K-, JK+) and then the “BC” stimuli (B-, C-, BC+); they were then trained in 2\(^{nd}\)-order negative occasion setting with the “ABC” stimuli (C-, BC+, ABC-, B-, AB+, A+); afterwards, they were trained in direct learning with ABR+ and then G+ and H-, where each of these stimuli/combinations always predicted the US (ABR+, G+) or its absence (H-). After this, participants engaged in the Reminder 1 training phase in which all previously trained stimuli were trained within one sub-phase.

Participants then conducted Transfer Test 1 in which we tested responding to the trained stimuli and novel combinations of stimuli. These novel combinations were designed to test whether 1\(^{st}\)-order occasion setting and 2\(^{nd}\)-order occasion setting were indeed learned, since an occasion setter will only affect stimuli that were previously trained in occasion setting (e.g., an occasion setter would not affect an unambiguous CS+ or CS-). We hypothesized that occasion setters would not transfer to stimuli that were not trained in that form of occasion setting (i.e., B and J would not transfer to H-; A would not transfer to JK+, G+, or H-; AB would not transfer to G+). This would be shown by more similar responding between the novel transfer combinations to the trained target than to the trained occasion setting combination (e.g., ABG would be more similar to G+ than ABC++; AJK would be more similar to JK+ than ABC-).

In the next training phase (Training 2), participants were trained with a new 2\(^{nd}\)-order negative occasion setter (T) within the “TJK” family of stimuli. Note that “JK” was already trained in 1\(^{st}\)-order positive occasion setting but not 2\(^{nd}\)-order negative occasion setting; it would now be trained with T in 2\(^{nd}\)-order negative occasion setting (TJK-, JK+, K-, J-, TJ+, T+). Participants were then trained in direct learning with TJR+ and engaged in the Reminder 2 training phase in which multiple trial types from the “ABC” and “TJK” families were trained within the same sub-phase (TJK-, JK+, K-, ABC-, BC+, C-, A+).

Finally, the experiment concluded with Transfer Test 2, in which three trained trial types were tested (ABC-, TJK-, JK+) and one untrained combination testing 2\(^{nd}\)-order negative occasion setting (AJK). Of critical importance: AJK was tested during both Transfer Test 1 and Transfer Test 2. At the time of Transfer Test 1, JK had only been trained in 1\(^{st}\)-order positive occasion setting (i.e., it had not been trained with 2\(^{nd}\)-order negative occasion setter T yet). During Transfer Test 1, we expected A would have little effect on JK. Conversely, after Transfer Test 1, JK was trained with 2\(^{nd}\)-order negative occasion setter T, so we hypothesized A would have a relatively greater effect on JK during Transfer Test 2. If this occurred, it would provide strong evidence that A was indeed a 2\(^{nd}\)-order negative occasion setter, as it would only affect JK after JK had been trained with a different 2\(^{nd}\)-order negative occasion setter.

Experiment 2 followed the same structure as Experiment 1, but the US reinforcement contingencies were largely reversed. The “ABC” family became the “DEF” family, and reinforcement was reversed (i.e., C-, BC+, ABC- vs F+, EF-, DEF+; B- and AB+ vs E+ and DE-; A+ vs D-); the “TJK” family became the “UMN” family, and reinforcement was reversed; and the “Direct Learning” family largely remained as such, where G+ and H- remained the same, but R became S with reinforcement reversed (i.e., ABR+, TJR+ vs DES-, UMS-). For concision, we will not write the same details for Experiment 2 as mentioned above for Experiment 1. We instead refer the reader to Table A in S7 Text for those details, which are congruent across both experiments.

Additionally, we created a novel computational model to i) explain 2\(^{nd}\)-order occasion setting for the first time (as well as 1\(^{st}\)-order occasion setting and direct learning); ii) to posit, operationalize, and test stimulus ambiguity as a mechanism of higher-order learning; and iii) to predict the specific transfer effects that are observed in these forms of learning. Our computational model is related to our theoretical model in Fig 1. The details of our computational
model are found in the Materials and Methods section, but we provide a brief overview here. Our computational model has separate learning variables for excitation and inhibition at each hierarchical level (six variables total: V, V̅, P, N, P2, N2). Additionally, because occasion setters only affect CSs that have been trained in occasion setting, we have four learning variables representing the CS’s ability to be modulated in excitatory or inhibitory directions by 1st- or 2nd-order occasion setters (P, N, P2, N2). A special aspect of our model is that 1st-order occasion setting can only be learned if the CS is ambiguous (i.e., sometimes predicts the US; sometimes does not). This is reflected in one learning variable (γ1). Similarly, 2nd-order occasion setting can only be learned if the 1st-order occasion setter is also ambiguous (sometimes excites the CS/US association; sometimes inhibits it). This is reflected in one learning variable (γ2). Our model also has two free parameters: α (learning rate) and ι (leaky memory). The α parameter estimates the learning rate, and the ι parameter(s) set the degree of retention and functionally can move the individual’s asymptote of learning. If empirically supported, the goal of our model would be to provide insight into learning mechanisms and stimulate research on highly complex Pavlovian learning (see Materials and Methods for details on our computational model).

Results
Training
Results from training phases are shown in Fig 3 (see Tables A and B in S4 Text for full statistical details). The critical test of reinforcement learning was the Reminder phases, as this was the end of each training section. This test was conducted by comparing the self-reported US expectancy ratings from each trial across stimuli. Overall, in both experiments, participants correctly learned which stimuli were reinforced and which were not for all trial types–direct associations, 1st-order occasion setting, and 2nd-order occasion setting. The most important and novel of these results was 2nd-order occasion setting: as hypothesized, 2nd-order occasion setting trial types had significantly lower (Experiment 1) and greater (Experiment 2) responding than their respective 1st-order occasion setting trial types during Reminder (e.g., Experiment 1: ABC- vs BC+; Experiment 2: UMN+ vs MN-; ps < .001).

Transfer test
Transfer tests are used to determine whether occasion setting was indeed learned, in which a putative occasion setter is trained with one stimulus and is tested with a separately trained stimulus [1,2,7]. If the putative occasion setter only developed direct associative properties, it would summate with other CSs, producing responding equal to the sum of the presented stimuli. Conversely, true occasion setters transfer their occasion setting properties to stimuli that were separately trained in occasion setting, although this transfer is usually strong but incomplete. Thus, if a stimulus were a true occasion setter, we would expect it to have a greater effect on other stimuli trained in the same type of occasion setting than stimuli not trained in that form of occasion setting. We examine this in our transfer tests below.

See Tables A and B in S5 Text for details on statistical analyses. In total, all of our hypotheses were supported in Experiment 1, and most hypotheses were supported in Experiment 2. Each of the significant results below supporting our hypotheses survived Holm-Bonferroni correction for multiple comparisons [24].

Replicating previous research, we hypothesized that a 1st-order occasion setter would not affect an unambiguous CS [10,25,26]. To this end, we tested whether the novel stimulus combination was closer to one trained stimulus than another (e.g., whether the novel BH
Fig 3. Experiment 1 and 2 Training Results. a, b, c) Experiment 1 Training results generally reflect direct CS/US associations, 1\textsuperscript{st}-order positive occasion setting, and 2\textsuperscript{nd}-order negative occasion setting. d, e, f) Experiment 2 Training results generally reflect direct CS/US associations, 1\textsuperscript{st}-order negative occasion setting, and 2\textsuperscript{nd}-order positive occasion setting. Congruent conditions/panels are displayed horizontally between experiments. Results in both experiments showed that participants correctly learned which stimuli were (non)reinforced. Error bands reflect standard error. Generally, “cool” colors (blues, greens, purples) indicate hypothesized higher values, whereas “warm” colors (reds, oranges, yellows) indicate hypothesized lower values. Additionally, because not all stimuli were trained in every phase of the experiment (e.g., the “ABC” and “DEF” stimuli were not trained during 2\textsuperscript{nd} Training), there are some empty spaces in the graphs of stimuli being shown either earlier or later in the experiment (e.g., G+ and H- were trained in the first half of the experiment). The reason that trial numbers vary between stimuli is to balance thoroughness of training and concision. During reminder phases, we wanted to remind participants of the most critical trial types relevant for the upcoming transfer test. For example, J- was not in the 2\textsuperscript{nd} Reminder
combination was closer to the trained H- or the trained BC+) because novel stimulus combinations would reasonably produce more uncertain responding than trained stimuli simply because they are novel (i.e., novel stimuli would have values closer to “3” on our 1–5 US expectancy scale, where 1 = “Certain No Bonus,” 3 = “Completely Uncertain,” and 5 = “Certain Yes Bonus”). Thus, we examined whether the novel stimulus combination was closer to one trained stimulus than another (see main text discussion and S5 Text for further discussion). This was tested three times in each experiment. In Experiment 1, all three 1st-order positive occasion setting transfer tests supported the hypotheses; in Experiment 2, one of three 1st-order negative occasion setting tests supported the hypotheses (constituting our only two null transfer test results across all hypotheses in both experiments). First, in Experiment 1, all three 1st-order occasion setting tests supported the hypotheses, as evidenced by BH (the transfer stimulus combination) having more similar responding to H- than BC+ (Fig 4A; p < .001), JH having more similar responding to H- than JK+ (Fig 4B; p < .001), and AH having more similar responding to H- than G+ (Fig 4G; p < .001). Experiment 2 showed that one of three 1st-order negative occasion setting transfer tests supported the hypotheses. Specifically, responding to EG was equidistant between EF and G+ (Fig 4F; p = .588), and responding to MG was equidistant between MN- and G+ (Fig 4G; p = .197). Conversely, our third test showed DG had more similar responding to G+ than H- (Fig 4H; p < .001), supporting the hypothesis.

Second, as part of our novel hypotheses, we hypothesized that 2nd-order occasion setters would only affect a CS if a 1st-order occasion setter were present; thus, we tested 2nd-order occasion setters on CSs in absence of 1st-order occasion setters. For example, during ABC trials, A was trained as a 2nd-order negative occasion setter with B as the 1st-order positive occasion setter and C as the CS. We would expect A to only affect a CS if a 1st-order occasion setter were present (e.g., if “B” were present from ABC). We thus tested the 2nd-order occasion setter in absence of a 1st-order occasion setter with a separately trained CS that had a learning value opposite to the 2nd-order occasion setter (i.e., 2nd-order negative occasion setter A (inhibitory) tested with G+ (excitatory); 2nd-order positive occasion setter D (excitatory) tested with a H- (inhibitory)). Results supported this hypothesis in both experiments. In Experiment 1, AG showed more similar responding to G+ than H- (Fig 4C; p < .001), and in Experiment 2, DH showed more similar responding to H- than G+ (Fig 4H; p < .001). This suggests that the 2nd-order occasion setters had minimal effects on the CSs in absence of 1st-order occasion setters.

Third, one of our critical novel tests was whether a 2nd-order occasion setter would affect unambiguous lower-order stimuli not trained in 2nd-order occasion setting; we hypothesized it would not. This was assessed using i) the trained 2nd-order occasion setter/1st-order occasion setter combination with an unambiguous CS (e.g., testing AB with G+ in ABG), as well as testing ii) the 2nd-order occasion setter with a trained unambiguous 1st-order occasion setter/CS combination (e.g., testing A with JK+ in AJK). In each case and in both experiments, all hypotheses were supported. Specifically, in Experiment 1, AJK had more similar responding to JK+ than ABC- (Fig 4D; p = .005), and ABG had more similar responding to G+ than ABC- (p = .005). Congruently, in Experiment 2, DMN had more similar responding to MN- than DEF+ (Fig 4I; p = .002), and DEH had more similar responding to H- than DEF+ (p < .001).
Fourth, our other critical novel test was to evaluate whether the ability of 2nd-order occasion setters to affect lower-order stimuli depended on whether the lower-order stimuli were ambiguous and trained in 2nd-order occasion setting. In the previous paragraph, we demonstrated that 2nd-order occasion setting setters had little effect on lower-order 1st-order occasion setter/CS combinations that were unambiguous and not trained with a 2nd-order occasion setter (e.g., A tested with JK+ in AJK; D tested with MN- in DMN). In each experiment, we later trained those same 1st-order occasion setter/CS combinations with a 2nd-order occasion setter (i.e., JK+ was later trained with T in TJK--; MN- was later trained with U in UMN+). We then tested whether a different 2nd-order occasion setter could affect the 1st-order occasion setter/CS combination more than it did before—now that the 1st-order occasion setter/CS combination had been trained with a 2nd-order occasion setter. Thus, the exact same stimulus combinations (i.e., Experiment 1: AJK; Experiment 2: DMN) were each tested twice—before and after the 1st-order occasion setter/CS combinations (JK+, MN-) were trained with a 2nd-order occasion setter (T, U). We hypothesized there would be a greater effect of 2nd-order occasion setters (i.e., A, D) on the 1st-order occasion setter/CS combinations (JK+, MN-) after the latter was trained with a different 2nd-order occasion setter than before—now that the 1st-order occasion setter/CS combination had been trained with a 2nd-order occasion setter. This hypothesis was supported in both experiments. In Experiment 1, the 2nd-order negative occasion setter (A) had a greater effect on JK after JK was trained with T (i.e., TJK--; Fig 4E; see AJK1 vs AJK2 comparison; p < .001). Congruently, in Experiment 2, the 2nd-order positive occasion setter (D) had a greater effect on MN- after MN was trained with U (i.e., UMN+; Fig 4J; see DMN1 vs DMN2 comparison; p < .001).

**Computational modeling**

The behavioral results above demonstrate that 2nd-order occasion setting was learned, and these behavioral results can be bolstered by further evaluation of the underlying learning processes using computational modeling. We tested a computational model that allowed occasion setters to impact US expectancy only if lower-order stimuli were ambiguous and trained with occasion setters. That is, the influence of 1st-order occasion setters on CSs was dependent on CS ambiguity and CS training with a 1st-order occasion setter, and the influence of 2nd-order occasion setters on 1st-order occasion setters and CSs was dependent on 1st-order occasion setter ambiguity, CS ambiguity, and the CS's training with a 1st-order and 2nd-order occasion setter. We compared models limited to each hierarchical level: our full 2nd-order occasion setting model (which also included 1st-order occasion setting and direct associations), our 1st-order occasion setting model (which also included direct associations), and our direct associations model (i.e., direct excitation and direct inhibition).

**Parameter recovery**

For each model, we simulated random learning rate (α) and leaky memory (ı) parameters and evaluated the models’ ability to estimate those parameters accurately. In short, α sets the
learning rate, \( i \) sets the retention rate, and \( i \) was estimated after \( \alpha \). All models in both experiments showed high correlations between simulated and recovered parameters (\( rs > .939; \) see Fig A in S6 Text), indicating that we were able to accurately estimate individual subjects’ parameter values.

**Model fit**

We used Watanabe-Akaike Information Criterion (WAIC; [27]) to measure model fit. Our results showed that the 2\(^{nd}\)-order occasion setting models outperformed the 1\(^{st}\)-order occasion setting models, and both occasion setting models outperformed the direct associations models (Fig 5A). This suggests that our 2\(^{nd}\)-order occasion setting model more closely resembles the underlying computations and learning process engaged in by the participants compared to the 1\(^{st}\)-order occasion setting model or direct associative learning model. Secondary to the WAIC results, we estimated model \( R^2 \) scores as a measure of our models’ explanatory power (Fig 5B). These results were fully congruent with the WAIC scores, where our 2\(^{nd}\)-order occasion setting models had the highest \( R^2 \) values at \( \approx .6 \) (Experiment 1: \( R^2 = .632 \); Experiment 2: \( R^2 = .560 \)). To illustrate our 2\(^{nd}\)-order occasion setting model’s predictions, we plotted example participants’ real data and model-predicted data (Fig 5C and 5D). These examples show a high degree of overlap between our model’s predictions and the participants’ behavior.

**Discussion**

This report investigates highly complex Pavlovian learning in order to experimentally model the complex associative learning that might occur in real-life circumstances. Pavlovian 1\(^{st}\)-order occasion setting is a form of learning in which a 1\(^{st}\)-order occasion setter signals whether a conditional stimulus (CS) will predict an outcome (the unconditional stimulus; US). Occasion setting occurs when the CS has a mixed (i.e., ambiguous) association in predicting the US and when the CS has been trained with a second stimulus that reduces the ambiguity (1\(^{st}\)-order occasion setter). In our experiments, we examined the form of learning that occurs when both the CS and 1\(^{st}\)-order occasion setter are ambiguous (i.e., both provide mixed signals of US (non)occurrence), positing that a third stimulus (the 2\(^{nd}\)-order occasion setter) will signal how the 1\(^{st}\)-order occasion setter modulates the CS/US association. We hypothesize three hierarchies of learning would be learned, in which the higher-order factors modulate the lower-order factors. In ascending order, these hierarchical levels are direct associative learning, 1\(^{st}\)-order occasion setting, and 2\(^{nd}\)-order occasion setting. Additionally, we created the first computational model predicting all three hierarchical levels of learning and evaluated its performance in two experiments.

A primary test of whether occasion setting was indeed learned is a transfer test, in which a putative occasion setter is tested with a separately trained stimulus. We would expect an occasion setter to only affect ambiguous lower-order stimuli trained in the same fashion with a different occasion setter. We conducted transfer tests in our experiments in which occasion setters were tested with i) the stimuli they were trained with and ii) separately trained stimuli. Our behavioral results can be summarized as the following: i) in most cases, a 1\(^{st}\)-order occasion setter minimally affected responding to an unambiguous CS; ii) in all cases, a 2\(^{nd}\)-order occasion setter minimally affected responding to an unambiguous CS or an unambiguous 1\(^{st}\)-order occasion setter / CS combination; iii) in all cases, a 2\(^{nd}\)-order occasion setter affected responding to an ambiguous 1\(^{st}\)-order occasion setter / CS combination in the hypothesized direction; iv) the successful transfer of the 2\(^{nd}\)-order occasion setter to the 1\(^{st}\)-order occasion setter / CS combination was present but incomplete, which is the expected effect since occasion setters maximally affect the CS/US association they were trained with and have less effect
Fig 5. Computational Modeling Results. a) Model fit was determined with Watanabe-Akaike Information Criterion (WAIC), where lower scores indicate more accurate models. In both experiments, we tested three models of hierarchical learning: our direct associative learning model, our 1st-order occasion setting model (which also included direct associative learning), and our 2nd-order occasion setting model (which also included 1st-order occasion setting and direct associative learning). Results show that, in both experiments, the 2nd-order occasion setting model (bold colors) outperformed the 1st-order occasion setting model and direct associations model. b) As secondary/complementary results to our WAIC analyses, we also estimated median R² for each model, finding that our 2nd-order occasion setting models had the greatest R² (0.632, 0.560). c) Experiment 1 exemplar participant responding, model-predicted responding, and “perfect learning” prediction for the 2nd-order occasion setting model. d) Experiment 2 exemplar participant responding, model-predicted responding, and “perfect learning” prediction for 2nd-order occasion setting model. Transfer Test 1 = trials 205–249 and Transfer Test 2 = trials 331–342, shown between the vertical green hashed lines; remaining trials were Training/Reminder.

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on separately trained stimuli [1,7,14,15]; and v) participants learned that a given stimulus could have both excitatory and inhibitory meanings across all three hierarchies (direct learning, 1st-order occasion setting, and 2nd-order occasion setting). Our behavioral results were bolstered by the computational modeling results, which showed that the 2nd-order occasion setting model provided better fit (i.e., WAIC score) than the 1st-order occasion setting or direct associations model and, as a secondary/complementary analysis, had greater explanatory power ($R^2 \approx .6$). Integrating the behavioral and computational results, the evidence supports that 2nd-order occasion setting was learned and that our model is an accurate depiction of 2nd-order occasion setting.

In our theoretical and computational model, lower-order stimulus ambiguity (i.e., an inconsistent association between a stimulus and the US) is the gateway to higher-order learning; without ambiguity, occasion setting will not be learned (see Materials and Methods for computational model details). It is important to note that lower-order stimulus ambiguity is necessary but not sufficient for 1st-order or 2nd-order occasion setting to occur. In our model, the ambiguous CS must also be trained with a 1st-order occasion setter or 2nd-order occasion setter in order for each form of learning to occur. This assumes that a discernable occasion setter is present and is salient enough to acquire a learning value (which was the case in our experiments). This is exemplified by conducting occasion setter transfer tests. In our case, we evaluated transfer to an unambiguous, consistently (non)reinforced CS (i.e., CS+, CS-), in which we predicted and observed low transfer in most cases [25,26] (see next paragraph for discussion). Additionally, we compared 2nd-order occasion setter transfer to a 1st-order occasion setter/CS combination prior to and after the latter’s training with a different 2nd-order occasion setter. We predicted greater transfer after vs before 2nd-order occasion setting training [1,7,13–15,28], which we observed in both experiments. Conversely, a partially reinforced CS with no discernable occasion setter will be much less affected by occasion setters [13,29].

Our computational model predicts these transfer effects by allowing the CS to obtain the ability to be affected by 1st- or 2nd-order occasion setters (i.e., $P$, $N$, $P^2$, $N^2$), which requires the CS to be ambiguous ($\gamma_1$ and $\gamma_2$). Thus, only stimuli that are ambiguous and trained with an occasion setter can be affected by an occasion setter of the same type. Furthermore, while we are not the first to posit stimulus ambiguity as being a requirement of occasion setting [1,3–5,30] or to experimentally assess it [31], we are the first to extend it to 2nd-order occasion setting, investigate its robustness across learning hierarchies, and to create a theoretical and computational model operationalizing CS ambiguity and 1st-order occasion setter ambiguity. The data from the present experiments supports our computational model and its unique approach to operationalizing CS ambiguity (i.e., CS ambiguity = direct excitation \ direct inhibition) and 1st-order occasion setter ambiguity (i.e., 1st-order occasion setter ambiguity = 1st-order positive occasion setting \ 1st-order negative occasion setting) as a requirement of higher-order learning.

Furthermore, our statistical approach during transfer tests was to assess whether responding to the novel transfer stimulus combination was closer to one trained stimulus/combination or the other trained stimulus/combination. As described two paragraphs above, the results largely supported our hypotheses. However, we observed differences in responding comparing the novel combinations of stimuli to the unambiguous trained stimuli that were theoretically hypothesized to have similar responding to each other (see S5 Text for statistical details). For example, H- was a trained unambiguous CS-, BC+ was trained as a 1st-order positive occasion setting combination (from B-, C-, BC+), and novel combination BH showed responding that was significantly closer to H- than BC+ but was greater than H- (Fig 4A). This effect was generally observed across most novel vs trained transfer test stimuli. A likely reason that the novel
stimulus combinations (e.g., BH) were significantly different from the trained unambiguous stimulus (e.g., H-) is that the novelty of the transfer test stimulus combinations could produce US expectancy ratings closer to the midpoint of “Completely Uncertain” [32], which is what we observed. This interpretation seems fairly intuitive: uncertainty will be greater with novel stimuli compared to well-trained stimuli. However, we did not include a separate manipulation in our experimental design to test whether the novelty of the stimulus combinations indeed is what produced this change in responding, so we cannot conclude this with certainty.

A second possibility is that the occasion setters developed a small degree of direct associative properties with the US when presented in combination with their trained CSs. For example, perhaps “B” from BC+ acquired largely 1st-order positive occasion setting abilities in the BC+ combination and a small degree of direct excitation with the US in this combination, as well. This direct excitation could then summate/transfer when tested with novel CSs (e.g., when H- was tested with B as BH). A third possibility is that the occasion setters had a small degree of transfer of their occasion setting properties to the unambiguous stimuli. This could happen if our “unambiguous” stimuli developed some degree of ambiguity, such as through generalization of learning from stimuli that were experimentally designed to be ambiguous stimuli trained with occasion setters (e.g., generalization from ambiguous stimuli C or K to unambiguous stimuli H- or G+). This interpretation would be consistent the broader occasion setting model since it would mean that the properties of ambiguous stimuli trained with occasion setters might generalize to “unambiguous” stimuli and allow the latter to be relatively minorly influenced by an occasion setter. A fourth explanation generally relies on imperfect knowledge and awareness from the participants (i.e., human error). Even if the environment and stimuli are designed to be deterministically ambiguous or unambiguous (as in our experiments), participants’ experiences might be different, for example, if they forget certain associations or miscredit which stimulus/combination led to the US or not. These inconsistencies could produce ambiguity in unambiguous stimuli, resulting in partial transfer of occasion setting properties (e.g., mistakenly thinking that H- led to the US or G+ led to no US could make them somewhat ambiguous and possibly allow occasion setters to minorly influence responding to them). In total, each of the explanations above is plausible, and they are not mutually exclusive. Perhaps one or more of them contributed to the partial transfer effects we observed, and future experiments could be designed to distinguish between these explanations.

Moreover, we believe that the CS is the primary target stimulus when presented with occasion setters, so we posit that the real-time moment of effect of an occasion setter occurs during the CS. For example, when a 1st-order positive occasion setter was presented (which also functioned as a CS- in our experiment), responding to this stimulus by itself was inhibitory, but when it was presented with the CS it was trained with in 1st-order positive occasion setting, responding was excitatory. Conversely, when the 1st-order positive occasion setter was followed by a separately trained unambiguous CS-, responding was inhibitory, thus suggesting that the CS is the target stimulus that activates the form of responding. Indeed, this is consistent with previous literature showing that the CS (rather than the occasion setter) controls response form [33]. It is also unlikely that participants were responding to blocks of cues, as we would expect very little transfer of the 2nd-order occasion setter to the separately trained 1st-order occasion setter / CS combination. In both experiments, we found strong transfer when expected.

Additionally, our model theoretically claims that mixed CS/US reinforcement will lead to broadening of attention to other situational factors, contexts, or stimuli to disambiguate whether the CS will predict the US on a given trial (i.e., γ1; [3,5,34,35]); we theorize the same for 1st-order occasion setters when they send mixed modulatory signals of CS/US reinforcement (i.e., γ2). Thus, our model posits when a search for 1st- or 2nd-order occasion setters will
begin, but it is less clear when the search will end. In our experiments, the occasion setters deterministically signaled whether lower-order stimuli would predict the US, and there was only one occasion setter that was trained with a given lower-order stimulus or stimulus combination, so the search could theoretically end with perfect predictive accuracy once the occasion setter was identified and fully trained. It is less clear what would happen in non-deterministic occasion setting or in which there are multiple occasion setters present across trials. We presume the search for additional occasion setters would not occur while a trained, deterministic occasion setter is present (i.e., the occasion setter would block learning to other potential occasion setters) [36,37]. However, for example, if a 1st-order positive occasion setter’s presence perfectly signals that the CS will predict the US, but the CS still receives some mixed reinforcement when presented alone, we presume that the individual will continue searching for other occasion setters that will disambiguate the CS alone (perhaps there is an additional 1st-order positive or negative occasion setter the individual has not found yet). Thus, we posit that while mixed reinforcement is ongoing, the search for occasion setters is possible or likely. However, real-world constraints (e.g., time demands, energy) or other factors (e.g., relative importance of fully disambiguating this CS/US association vs focusing on other CS/US associations or engaging in other behaviors/tasks; whether the individual believes it is even possible to deterministically or near-deterministically predict the US) will likely affect whether an individual chooses to search for and learn about occasion setters that disambiguate US (non)occurrence. Ultimately, if an individual determines there are no occasion setters that can disambiguate the CS any further, they may infer that the unexplained mixed reinforcement of the CS is simply due to partial reinforcement. Furthermore, our free parameters of learning rate ($\alpha$) and memory retention ($\tau$) estimate and apply constraints that may affect how quickly an individual ends their search for occasion setters. For example, an individual who learns quickly and remembers strongly that an occasion setter affects the CS/US association may end their search more quickly than someone who learns slowly and remembers poorly. Our experiments were not designed to test when/how the search for occasion setters concludes, but future research could test this, and doing so would be very applicable to real-world scenarios in which there are presumably many occasion setters and CSs and in which learning rate and memory retention will likely affect the search for occasion setters.

The present results provide numerous clinical implications across many disorders that have strong Pavlovian components (e.g., anxiety, substance use). For example, anxious individuals seem to have deficits in discriminating safety from danger with direct associations [38–40] and increased fear of 1st-order occasion setting compounds [8]. The latter could be due to working memory deficits in anxious individuals [41–43] since 1st- and 2nd-order occasion setting are presumably more demanding of working memory than direct associative learning (i.e., needing to remember if an occasion setter was presented or not). Anxious individuals also have elevated intolerance of uncertainty [44,45]. We define “uncertainty” as relative difficulty predicting US (non)occurrence from the presented stimulus(i), whereas “ambiguity” is a different construct in which the individual learns that the stimulus(i) has/have a mixed association with the US. Due to the relative complexity of learning occasion setting vs direct associations, perhaps individuals experience greater uncertainty during occasion setting training or experience the uncertainty for more trials due to the complexity and ambiguity of occasion setting. This may lead anxious individuals to have greater fear of occasion setting combinations or depressed individuals to expect less reward from occasion setting compounds [8]. Second, our model claims that 1st-order occasion setters can be ambiguous (as has been shown elsewhere; e.g., [9,21]), and it argues that CS responding will be minimal if the CS has direct inhibition, is presented with a 1st-order negative occasion setter, and is absent of 1st-order positive occasion setters. This is relevant for anxiety disorders, as conventional exposure
therapy focuses primarily on direct CS+ extinction with some effort to increase 1st-order negative occasion setters via context variability [46], but there is no explicit emphasis on 1st-order positive occasion setter extinction. Extinction of 1st-order positive occasion setting has been investigated elsewhere [16,20,36], demonstrating that extinction of the 1st-order positive occasion setter with the CS together (as opposed to extinction with the CS alone or the 1st-order positive occasion setter alone) is required in order to reduce responding to the 1st-order positive occasion setter/CS combination. For example, conducting exposures to public speeches (CS) could reduce fear to the CS in absence of any positive occasion setters, but if the client gives a speech (CS) after a charismatic speaker (unextinguished 1st-order positive occasion setter), fear will increase. From our model’s perspective, when a 1st-order positive occasion setter is extinguished, a stimulus/context could become a 2nd-order negative occasion setter (if it is salient enough and valid). This is relevant for exposure therapy for anxiety disorders, as it suggests that exposure therapists should not only encourage CS extinction, but 1st-order positive occasion setter extinction, as well. Perhaps one approach using this example would be to conduct speeches (CS) after a charismatic speaker (1st-order positive occasion setter) in a variety of contexts/situations to extinguish both, though this awaits testing in clinical studies.

Furthermore, our computational model uses an elemental approach both conceptually and mathematically [47]. Each sound, image, or perceptual occurrence is a separate stimulus that can contain multiple learning values (e.g., V, V, P, N, P2, N2; see model in Materials and Methods) that get expressed differently depending on the presence or absence of other stimuli. Other models exist, such as latent cause models [48,49], in which individuals do not learn the associations between stimuli (e.g., CS, US, occasion setters); rather, they learn that latent causes predict the presence or absence of the stimuli, including the US. While we are unaware of a latent cause model that can predict 2nd-order occasion setting, and while evaluating its utility is beyond the scope of this report, it would be interesting to explore in future work. On the one hand, perhaps latent cause models of 2nd-order occasion setting would be parsimonious since they tend to cluster stimuli together as part of the latent causes, obviating the need for multiple associative weights. On the other hand, latent cause models might be complex because 2nd-order occasion setting is inherently complex; the latent cause model could require individuals to infer multiple latent causes. For example, in Experiment 1, stimulus A predicted the US as a CS+ and 1st-order positive occasion setter but predicted the US’s absence as a 2nd-order negative occasion setter. The latent cause model would need to account for the multiple predictive abilities of each stimulus, presumably requiring multiple latent causes to be inferred. It is also unclear how a latent cause model would predict the pattern of occasion setting transfer effects that are known to occur [1,7].

Additionally, our model is capable of predicting partial reinforcement of CSs or occasion setters using the leaky memory [50] parameters. These parameters allow learning values to leak and to provide an estimate of the participants’ inferred reinforcement rate. While Experiments 1 and 2 did not have partial reinforcement, we used a separate database of 75%/25% reinforcement in 1st-order positive occasion setting (as opposed to the traditional 100%/0% reinforcement rate) to test the utility of these leaky memory parameters (see S8 Text). Our results showed that including the parameters provided better model fit than excluding them. We additionally included the parameters in Experiments 1 and 2 for completeness, which are the models presented in the main text.

Furthermore, our computational model has some assumptions and limitations worth discussing. First, it engages in trial-by-trial learning rather than real-time learning [51,52]. However, the goal of our model is not to evaluate learning in real-time but rather to explain 2nd-order occasion setting for the first time (and the lower-order learning hierarchies), to posit
and operationalize stimulus ambiguity as a mechanism of higher-order learning, and to predict the specific transfer effects that are observed in these forms of learning. Including real-time learning would require even more parameters and further complicate the model. Second, our model assumes that a discernable occasion setter is present in order for occasion setting to be learned. In the present experiments, these stimuli were clearly identifiable since they were tangible and salient (i.e., sounds, images). However, in situations of less salient (but valid) occasion setters, there would likely be individual differences in whether participants would detect and learn about the occasion setters. Third, our model assumes that less salient stimuli that provide information about CS (non)reinforcement are occasion setters. In our experiments, stimuli were designated as CSs or occasion setters based on previous work demonstrating which situations lead to occasion setting vs direct associative learning—all converging on the signal validity of the putative occasion setter and its relative salience to the CS [1,2,23,53]. Thus, we specified that temporal distance from the US (non)occurrence was positively associated with learning hierarchy [53].

Lastly, the neurocircuitry of 1st-order occasion setting remains largely unknown [1], and research on the neurocircuitry of 2nd-order occasion setting is non-existent. Because the learning process of 2nd-order occasion setting is theoretically similar to 1st-order occasion setting, it seems more likely and parsimonious that both forms of learning involve the same brain regions. However, this awaits empirical testing. Based on the limited research investigating the neurocircuitry of 1st-order occasion setting, candidate brain regions include the entorhinal cortex (EC), hippocampus, subiculum, orbitofrontal cortex (OFC), basolateral amygdala (BLA), and ventromedial prefrontal cortex (vmPFC). In short, the EC is involved in learning about temporally separated cues (like in occasion setting) and communicates this with the hippocampus [54–61]. Lesioning both the hippocampus and subiculum prevents occasion setting learning [25,62–66]. The OFC and BLA work together to flexibly learn changing associations between stimuli [1,67–82], and the OFC in particular has been shown to be active in occasion setting [83]. Furthermore, because the vmPFC is associated with extinction learning with simple CSs (which is arguably a form of 1st-order negative occasion setting) [84–88], it may also be associated with extinction of occasion setting (i.e., 2nd-order negative occasion setting). Neuroanatomically, the EC, hippocampus, subiculum, BLA, OFC, and vmPFC are connected [55,89–98], providing the structural ability to learn about changing cue meaning (BLA, OFC) for temporally separates cues (EC, hippocampus, subiculum) to produce occasion setting.

In conclusion, our experiments investigated highly complex associative learning and were the first to explicitly demonstrate the existence of 2nd-order occasion setting. We experimentally showed that lower-order stimulus ambiguity (i.e., conditional stimuli, 1st-order occasion setters) was necessary for higher-order learning (i.e., 1st- and 2nd-order occasion setting) to occur and that a 2nd-order occasion setter only transferred to a 1st-order occasion setter/CS combination when the latter was ambiguous and trained with a different 2nd-order occasion setter. Our stimuli were also trained in both excitatory and inhibitory directions across hierarchies, which would not be possible if occasion setting was not learned. These results in total are strong indications that 2nd-order occasion setting was indeed learned. Additionally, our computational model showed validity in predicting direct associations, 1st-order occasion setting, and 2nd-order occasion setting, and our 2nd-order occasion setting model outperformed simpler models based on 1st-order occasion setting and/or direct associative learning. These experiments are important to draw research attention to an additional layer of Pavlovian learning (2nd-order occasion setting), to more accurately model complex real-life associative learning, and to inform the treatment of disorders with strong Pavlovian components (e.g., anxiety, substance use).
Materials and methods

Ethics statement

This study was deemed exempt by the California Institute of Technology Institutional Review Board, and all participants provided written informed consent prior to commencing the study.

Participants

Prolific [99] was used to recruit and collect human participant data online (Experiment 1: final N = 58; Experiment 2: final N = 67). Because 2nd-order occasion setting has not been investigated in a design like this, we did not have a strong basis for a power analysis. However, we pre-registered collecting 50–75 participants per experiment based on the strong effect size of 1st-order occasion setting as measured by US expectancy from transfer tests in our previous study (N = 80, d = .580 to 1.132; [8]). In the present experiments, we conducted post-hoc analyses of power for our most critical transfer tests (e.g., AJK2 vs AJK1, stimuli from t-tests comparing difference scores; see Tables A and B in S5 Text). In Experiment 1, the average power for our significant results was .9409 (range: .8257–1.0000); in Experiment 2, the average power for our significant results was .9768 (range: .8899–1.0000). Thus, our experiments were well-powered with our final sample sizes.

Participant eligibility criteria included being age 18–65, healthy or corrected vision, United States residents, English fluent, no hearing difficulties, and a Prolific approval rating of ≥95%; participants were only allowed to participate in one of the experiments. Across both experiments, demographics information included gender (53.60% female, 45.60% male, 0.80% agender), age (mean = 30.18, SD = 10.88, min = 18, max = 63), and ethnicity (10.40% Black or African-American, 8% Central/East Asian (e.g., Chinese, Japanese, Korean), 4.80% Hispanic or Latin(x), 6.40% South Asian (e.g., Indian, Pakistani, Sri Lankan), 62.40% White, and 8% Multiracial). Participants were paid $19.42 in Experiment 1 and $19.06 in Experiment 2 for completing the study. This amount was achieved by US presentations at the end of reinforced trials, where each US was a $0.12 USD increase in payment (as well as $2.50 for completing questionnaires).

Additionally, prior to data collection, we pre-registered each study (Experiment 1: https://osf.io/n2c6v, Experiment 2: https://osf.io/hxcfs). Based on our pre-registered exclusion criteria (i.e., automatic/invariant responding), we excluded one participant each from Experiment 1 and 2; also, an additional participant from Experiment 2 was excluded because of technical difficulties.

Design

There were no between-subjects conditions; within-subjects conditions included trial number and stimuli with direct associations with the US (i.e., CSs), 1st-order occasion setters, and 2nd-order occasion setters. Experiment 1 was a 2nd-order negative occasion setting design, which included 1st-order positive occasion setting and 2nd-order negative occasion setting. Experiment 2 was a 2nd-order positive occasion setting design, which included 1st-order negative occasion setting and 2nd-order positive occasion setting. Thus, Experiments 1 and 2 are mirror opposites of each other. Both experiments included CSs with direct associations with the US. Our dependent variable was US expectancy, measured at the end of every trial.

Materials and apparatus

The Pavlovian conditioning procedure was programmed using PsychoPy 2020.1.3. All learning stimuli (CSs, 1st-order occasion setters, 2nd-order occasion setters) were 4sec audio or visual
stimuli. When multiple stimuli were presented within the same trial, they were presented serially with a 4sec inter-stimulus interval (ISI) between them. Serial presentation (rather than simultaneous presentation) is conducive to learning occasion setting rather than direct associations [23,100]. The US was a 1.25sec audio/visual stimulus showing an image of a gold coin with “12¢” written on it, confetti surrounding it, and an auditory cash register sound (i.e., “cha-ching!”). Inter-trial intervals (ITIs) were 1.25sec. ITIs and ISIs included a fixation cross, which was also displayed uninterrupted during audio stimuli. All trials ended with a US expectancy rating, which had no time constraint.

Unambiguous CSs (i.e., G+, H-; not trained with occasion setters) were images of fractals, ambiguous CSs (Experiment 1: C, K; Experiment 2: F, N; trained with occasion setters) were images of a blue triangle and orange circle, 1st-order occasion setting setters (Experiment 1: B, J; Experiment 2: E, M) were a violin sound and white noise sound, and 2nd-order occasion setting setters (Experiment 1: A, T; Experiment 2: D, U) were images of a desert and forest. Within each category, stimuli were counterbalanced across participants. Using different stimulus modalities (e.g., auditory, visual) between hierarchical levels facilitates distinction between direct and occasion setting learning [101]. Given that we only had two modalities to use, we made the 2nd-order occasion setting level visual (to distinguish from 1st-order occasion setting) but qualitatively different from the CS images (i.e., context images vs shapes/fractals). An additional unambiguous CS (Experiment 1: R; Experiment 2: S) was an image of a three-dimensional white gem; this stimulus was used to facilitate 2nd-order occasion setting (see S3 Text). Notably, the above list is the most hierarchically advanced function of each stimulus, but a given stimulus may have had more than one hierarchical meaning. For instance, each “1st-order occasion setter” listed above was also a CS with a direct association with the US, and this stimulus was modulated by a “2nd-order occasion setter” listed above, which acted as a 1st-order occasion setter in that case (e.g., Experiment 1: “B” was a 1st-order occasion setter (C-, BC+) but also a CS (B-, AB+); “A” was a 2nd-order negative occasion setter (ABC-, BC+, C-) but also a 1st-order positive occasion setter (B-, AB+) and a CS+ (A+)). This allowed us to test the specific hierarchical functions of each stimulus and determine whether independence between hierarchical levels was learned. Lastly, the following is a list of congruent stimuli between each Experiment (listed as Experiment 1/Experiment 2): A/D, B/E, C/F, J/M, K/N, R/S, T/U. G+ and H- were identical across both studies.

US expectancy. Participants used a visual analog scale to rate, “How certain are you that you are about to receive a bonus payment?” The values ranged from 1 = “Certain No Bonus”, 3 = “Completely Uncertain,” and 5 = “Certain Yes Bonus.” The visual analog scale did not display numerical values, but it displayed the anchors mentioned above. US expectancy was measured at the end of every trial using the mouse to click on the scale with unlimited time to respond.

Procedure

Participants attended one experimental session online lasting approximately 1 hour 45 minutes, where they provided informed consent, completed questionnaires, and completed the Pavlovian learning experiment. In the experiments (Tables 1 and 2), participants were informed, “Your goal in this experiment is to learn which sounds and images predict receiving bonus payments.” During Training and Reminder phases, participants experienced US (non) reinforcement, which resulted in increases in their payment. Importantly, to maintain the Pavlovian nature of the experiment (rather than instrumental), participant responses did not affect their payment. During Transfer Test, participants were informed that they would not know whether they receive the US on these trials, which was accomplished by using an image of a
Table 1. Summary of Experiment 1 Design. When three stimuli are listed (e.g., ABC-), 2nd-order occasion setting is hypothesized to be learned (except for ABR+ and TJR+). When two stimuli are listed (e.g., BC+), 1st-order occasion setting is hypothesized to be learned. When one stimulus is presented (e.g., C-), direct associative learning is hypothesized to be learned. All stimuli within a trial were presented serially. “+” indicates reinforcement; “-” indicates no reinforcement; “OS2” indicates 2nd-order occasion setting; “OS1” indicates 1st-order occasion setting; “Direct” indicates direct associative learning. Notably, the “ABC” family was trained in 2nd-order occasion setting before Transfer Test 1 and 2; the “TJK” family was only trained in 1st-order occasion setting before Transfer Test 1 but was trained in 2nd-order occasion setting before Transfer Test 2. This allowed the examination of whether “A” was indeed a 2nd-order occasion setter by testing “A” with “JK” before and after “JK” was trained with 2nd-order occasion setter “T.” We predicted “A” would affect “JK” more strongly after “JK” was trained with “T” than before that training.

| Stimulus Type       | Training 1/Reminder 1 | Transfer Test 1 | Training 2/Reminder 2 | Transfer Test 2 |
|---------------------|-----------------------|-----------------|-----------------------|-----------------|
|                     | OS2 | OS1 | Direct | OS2 | OS1 | Direct | OS2 | OS1 | Direct | OS2 | OS1 |
| “ABC” Stimulus Family | ABC- | BC+ | C-     | ABC- | BC+ | C-     | ABC- | BC+ | C-     | ABC- | BC+ |
|                     | AB+ | B-  |       | A+  |     |       | A+  |     |       |
| “TJK” Stimulus Family | JK+ | K-  | JK    | TJK- | JK+ | K-     | TJK- | JK+ | K-     | TJK- | JK+ |
|                     | J-  |     | T+    |     |     |       |     |     |       |
| Unambiguous CSs     | G+  | G   | TJR+  | H-  | H   |       | ABR+ |     | ABR    |
| Transfer Test Stimuli | ABG | BH  | A+    | AJK | JH  | AIK   | AG  |     | AH     |
|                     |     |     |       |     |     |       |     |     |       |

Table 2. Summary of Experiment 2 Design. When three stimuli are listed (e.g., DEF+), 2nd-order occasion setting is hypothesized to be learned (except for DES- and UMS-). When two stimuli are listed (e.g., EF-), 1st-order occasion setting is hypothesized to be learned. When one stimulus is presented (e.g., F+), direct associative learning is hypothesized to be learned. All stimuli within a trial were presented serially. “+” indicates reinforcement; “-” indicates no reinforcement; “OS2” indicates 2nd-order occasion setting; “OS1” indicates 1st-order occasion setting; “Direct” indicates direct associative learning. Notably, the “DEF” family was trained in 2nd-order occasion setting before Transfer Test 1 and 2; the “UMN” family was only trained in 1st-order occasion setting before Transfer Test 1 but was trained in 2nd-order occasion setting before Transfer Test 2. This allowed the examination of whether “D” was indeed a 2nd-order occasion setter by testing “D” with “MN” before and after “MN” was trained with 2nd-order occasion setter “U.” We predicted “D” would affect “MN” more strongly after “MN” was trained with “U” than before that training.

| Stimulus Type       | Training 1/Reminder 1 | Transfer Test 1 | Training 2/Reminder 2 | Transfer Test 2 |
|---------------------|-----------------------|-----------------|-----------------------|-----------------|
|                     | OS2 | OS1 | Direct | OS2 | OS1 | Direct | OS2 | OS1 | Direct | OS2 | OS1 |
| “DEF” Stimulus Family | DEF+ | EF- | F+ | DEF | EF | DEF+ | EF- | F+ | DEF+ |
|                     | DE- | E+ | D- |     |     |     |     |     |     |
| “UMN” Stimulus Family | MN- | N+ | MN | UMN+ | MN- | N+ | UMN | MN |
|                     | M+ |     |     | UMN- | M+ |     |     |
| Unambiguous CSs     | G+  | G   |     | H-  | H   |     |     |     |
| Transfer Test Stimuli | DES- |     |     | DEH | EG  |     |     |     |
|                     | DMN | MG | DMN | DGH | DGH |     |     |

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curtain to cover the location on screen where the US image would otherwise occur and by muting the US sound. This curtain/muted modification was done so that no learning and no reinforcement/non-reinforcement would occur during Transfer Test, allowing us to test the underlying learning processes that occurred during training with many Transfer Test trial types (see S1 Text). We used the unambiguous CS+ and CS- as transfer test targets in many of our transfer tests based on previous experiments that exclusively used these stimuli as transfer test targets [8,25,26]. We additionally used the (un)ambiguous 1st-order occasion setter/CS combination as transfer test targets in our remaining transfer tests. Experimental phases followed a particular sequence, and within each sequence, stimulus presentation order was pseudo-randomized (see Table A in S7 Text for details on sequence and number of trials). In sum, Training/Reminder were conducted for participants to learn which stimuli predict (non)reinforcement, and Transfer Test was conducted to investigate how participants learned which stimuli predict (non)reinforcement.

Data analysis

We used Stata 15.1 multilevel modeling for inferential statistics. For US expectancy during the Training phase, Level 1 predictors were Stimulus, Linear Slope, and Quadratic Slope. If the Quadratic Slope was not significant, it was removed from the model and re-run as a linear model. If the Linear Slope was not significant, it was removed and collapsed across Stimulus. For Transfer Test, the Level 1 predictor was Stimulus using the average of all three trials from a given block. We conducted multilevel modeling to examine whether one stimulus/compound was significantly different from another stimulus/compound. For analyses of the 2nd-order occasion setting compounds that were evaluated before and after 2nd-order occasion setting (i.e., AJK1 vs AJK2; DMN1 vs DMN2), we conducted multilevel modeling to determine if responding to AJK2 was lower than AJK1 and whether responding to DMN2 was greater than DMN1 (as hypothesized). For the remaining Transfer Test analyses, we used two sets of difference scores between the relevant stimuli and conducted paired samples t-tests on those difference scores. For example, in Experiment 1, when assessing whether the transfer test stimulus combination (BH) was significantly closer to the unambiguous CS (H-) than the putative 1st-order positive occasion setting compound (BC+) as hypothesized, we took a difference score of the higher value minus the lower value (e.g., BH minus H-; BC+ minus BH) and analyzed whether the difference scores were significantly different from each other. In this case, we examined whether BH minus H- was significantly smaller than BC+ minus BH; if so, this would indicate that BH was significantly closer to H- than BC+ as hypothesized. Within all analyses, we conducted Holm-Bonferroni corrections [24] to correct for multiple comparisons.

Furthermore, computational modeling was conducted using Python 3.7.6 to evaluate our theoretical model’s fit with the data using all trials. All models were fit using a hierarchical Bayesian approach, assuming subject-level parameters were drawn from group-level distributions, with parameters estimated using variational inference implemented in PyMC3 with 25,000 iterations. In our models, our free parameters were α (learning rate) and multiple ι parameters (leaky memory; this allows learning values to “leak” between trials and allows for the prediction of partial reinforcement). All other variables in the model were automatically calculated via the formulas, the stimuli presented, and the participants’ responses.

To evaluate parameter recovery, we separately simulated random α values ranging 0–1 and recovered them using our model with our sample size. We did the same with ι values depending on which model it was (see S6 Text for details). For all models, the correlation of simulated vs recovered α (rs > .999) and ι (rs > .939) parameters were very high (see Fig A in S6 Text).
Model comparison was performed using Watanabe-Akaike Information Criterion (WAIC) scores [27], which provide a goodness of fit measure for Bayesian models, penalizing for increasing numbers of free parameters in the model (lower scores indicate better model fit). We chose our best-fitting model in each experiment based on the WAIC scores. To provide secondary/complementary model results, we calculated median $R^2$ for each model using r2_score from the sklearn package.

Our 2nd-Order occasion setting model

Ambiguity of the CS is one of the major purported mechanisms through which occasion setting as a learning process is theorized to occur [5,31,102,103]. Importantly, we view stimulus ambiguity as a dimensional and learned phenomenon—meaning, an individual needs to learn that a stimulus is ambiguous, and stimuli can vary in the degree to which they are ambiguous. According to the ambiguity hypothesis, a CS that always predicts the US has no ambiguity and therefore needs no other stimuli (i.e., occasion setters) to resolve which outcome it will predict on a given trial. In support, the evidence suggests that occasion setters can affect (i.e., transfer to) CSs that have undergone occasion setting training but have little-to-no effect on unambiguous CSs (e.g., CS+, CS-) (i.e., occasion setters generally do not affect responding to unambiguous CSs) [8,9,11,13,22,104–109]. CSs with partial US reinforcement but not trained with an obvious occasion setter are also ambiguous, but transfer of occasion setting to these CSs tends to not occur [13,29], presumably because the CS was not trained with an identifiable occasion setter. This suggests that CS ambiguity is necessary but not sufficient for occasion setting to occur. Another ambiguous stimulus is an extinguished CS+, where after having been trained with a particular outcome, the outcome is no longer delivered. Transfer of occasion setters to extinguished CSs is small but mixed, ranging from none [10,13] to partial [12,13,17]. This transfer is presumably greater than transfer to a partially reinforce CS because the extinguished CS+ was trained with an identifiable negative occasion setter (e.g., the physical or temporal extinction context), though generalization decrements between occasion setters can mitigate this transfer (e.g., if the other occasion setters are cues, such as a light or sound, rather than more diffuse physical or temporal extinction contexts). Lastly, transfer is highest between similar occasion setters trained with similar CSs and USs [7,10,13–15,17,18,28], ranging from partial [10,28] to complete [13,18,28]. Thus, a major way to determine whether occasion setting was indeed learned is to conduct transfer tests, where we would expect little-to-no transfer to unambiguous CSs and strong but not necessarily complete transfer to CSs that were trained with similar occasion setters and similar USs. Indeed, occasion setters are thought to have strong but incomplete transfer to CSs trained with other occasion setters, suggesting that occasion setters operate on the specific CS/US association they were trained with [1,2,7]. In contrast, if the putative occasion setter does not actually acquire occasion setting properties, we would expect simple summation of its direct associative value with other stimuli.

One foundational process through which occasion setting is learned is likely attentional. According to attentional theories of ambiguity [3,5,34,35], once a CS predicts more than one outcome, attention broadens to other stimuli or the context to find what will determine which outcome the CS predicts on a given trial. Because ambiguity is likely a learned and dimensional phenomenon, the degree to which attention is broadened to search for a disambiguating stimulus (i.e., an occasion setter) is also dimensional (i.e., for stimuli that are only slightly ambiguous, little effort is likely used to search for disambiguating stimuli; for stimuli that are highly ambiguous, more effort is likely used to search for disambiguating stimuli). We operationalize CS ambiguity as having a mixed direct association with the US (i.e., direct excitation * direct inhibition), and we operationalize 1st-order occasion setter ambiguity as having mixed
modulation of the CS/US association in excitatory and inhibitory directions (i.e., 1st-order positive occasion setting * 1st-order negative occasion setting). This means that if a CS is ambiguous, a 1st-order occasion setter may modulate whether it predicts the US, and if a 1st-order occasion setter is ambiguous, a 2nd-order occasion setter may determine how the 1st-order occasion setter modulates the CS/US association.

Our model follows a prediction error format [47] with several learning variables ranging 0 to 1 (see Table 3). The variables are ultimately combined into the “final” formula that predicts responding (i.e., R, ranging -1 to 1), in which a) direct excitation, 1st-order positive occasion setting, and 2nd-order positive occasion setting are added, and b) direct inhibition, 1st-order negative occasion setting, and 2nd-order negative occasion setting are subtracted. Many of these variables are all automatically calculated by the learning formulas and the inputted data. The only free parameters are α (learning rate) and i (leaky memory). Additionally, see S1 Fig to download an interactive html file of our computational model, where the user can modify formula values to see the expected behavioral output. We suggest the reader uses this tool to more easily learn how our formulas work. For the reader’s convenience, we have provided example output of our computational model with varying levels of direct learning, 1st-order occasion setting, and 2nd-order occasion setting in the main text (see Fig 6). Note that 1st-order occasion setters will not affect responding unless both direct excitation and direct inhibition are > 0, and 2nd-order occasion setters will not affect responding unless direct excitation, direct inhibition, 1st-order positive occasion setting, and 1st-order negative occasion setting are all > 0.

The following is a more detailed description of the mechanics and assumptions of our model, with points 8 and 9 providing a verbal explanation of how responding (R) is conceptually and mathematically predicted.

1. Given that a stimulus adds unique predictive power with regards to whether the US will occur, we assume that stimulus salience is a determinant of whether the stimulus acquires occasion setting or direct associative properties [1,2,53]. Salience can be determined/manipulated in a number of ways [1,23,53], though the most common is to present occasion setters serially with their CSs, in which the CS’s onset is more proximal to the US, and the occasion setter’s onset is more distal [23]. In the present report, we assume that temporal distance from the US is positively associated with hierarchical learning (from direct learning to 1st-order occasion setting to 2nd-order occasion setting) if the more distant stimuli provide information regarding US occurrence beyond the information provided by the stimuli in the lower hierarchical levels. Thus, on a three-stimulus trial, the stimulus most distant to the US (non)occurrence is the 2nd-order occasion setter, the middle stimulus is the 1st-order occasion setter, and the most proximal stimulus is the CS. A similar pattern follows for two-stimulus trials (1st-order occasion setter is more distant from US than the CS). Although the present model assumes that temporal factors are critical in determining the learning accrued to each stimulus, this is not a real-time model. All that we assume here is that temporal ordering of stimulus’ presentation affects learning, but we do not manipulate the presentation time of each stimulus nor the inter-stimulus interval.

2. In our model, we use a delta rule of trial-by-trial learning with prediction error as a learning mechanism [47]. Prediction error is identical for our variable V as with the Rescorla-Wagner model’s V (i.e., λ–V_{sum}) [47], where λ is the occurrence of the US (1 if US occurs, 0 if US does not occur; the US can also have values between 0 and 1 if using the same US with different intensities). Our model views excitation and inhibition as separate learning processes, and prediction error for inhibition is based on $\lambda$ (i.e., the absence of an expected
Table 3. Formulas for Learning Direct Learning, 1st-Order Occasion Setting, and 2nd-Order Occasion Setting. In the table, subscripts are stimulus names (e.g., A, B, C; "sum" for all stimuli present on a trial); superscripts are trial numbers. For "R" formula, superscript 'n' for all variables, and subscript 'sum' for all variables except "R." Formulas are arranged in column format for readability. Responding (R) formula ultimately predicts behavioral responding and learning. R operates by adding excitation and subtracting inhibition (formula in dark gray). Light gray columns highlight the similar variables used in the "recipe" across our learning formulas. Hierarchical control of a) 2nd-order occasion setting (2nd OS) on 1st-order occasion setting (1st OS) and b) 1st OS on direct associative learning (i.e., CSs) is accomplished with modulation, in which the higher-order stimulus affects the lower-order stimuli’s signal of US (non)occurrence. The gateway to higher-order learning (from direct learning to 1st OS, and from 1st OS to 2nd OS) is lower-order stimulus ambiguity. Mechanism through which 1st OS is learned is γ1, which is the degree to which the CS is ambiguous (i.e., that the CS has both direct excitation and direct inhibition). If the present CS is unambiguous (e.g., only excitatory or only inhibitory), γ1 remains at 0, and no 1st OS is learned. Once a given CS is ambiguous (i.e., the CS is both excitatory and inhibitory), γ1 becomes positive, allowing P and N to increase from zero and for 1st OS to be learned. CS ambiguity is necessary but not sufficient for 1st-order occasion setting to be learned. In order for 1st OS to be learned, the individual must also learn that the CS can be modulated by a 1st-order positive or negative occasion setter, respectively. Thus, a CS must be both ambiguous and trained with a 1st-order occasion setter in order to be modulated by other 1st-order occasion setters (e.g., a simple partially reinforced CS will not be affected by a 1st-order occasion setting because its P and N will equal 0, causing the 1st OS terms in the R formula to equal 0).

| Description | Formula | Range of Normal Values |
|-------------|---------|------------------------|
| Responding (R) | | -1 to 1 |
| 1st-Order Occasion Setting | | |
| Direct Excitation (V) | \( \overline{\Delta V}^n_a = \overline{A}^n_a \bar{\alpha} (\bar{\alpha} \overline{\Delta V}^n + [V_{\text{sum}}]_{\text{(n)}} - [V_{\text{max}}]_{\text{(n)}}) \) | 0 to 1 |
| Direct Inhibition (V; "V Bar") | \( \overline{\Delta V}^n_s = \overline{A}^n_s \bar{\alpha} (\bar{\alpha} \overline{\Delta V}^n + [V_{\text{sum}}]_{\text{(n)}} - [V_{\text{max}}]_{\text{(n)}}) \) | 0 to 1 |
| CS Ambiguity (γ1; "Gamma 1") | \( \overline{\gamma}1^n_a = V_{\text{A}}^{n} \overline{\Delta V}^n \) | 0 to 1 |
| 2nd-Order Occasion Setting | | |
| 2nd-Order Positive Occasion Setting (P) | \( \overline{\Delta P}^n_a = \overline{A}^n_a \bar{\alpha} \overline{\Delta P}^n + [P_{\text{sum}}]_{\text{(n)}} - [P_{\text{max}}]_{\text{(n)}} \) | 0 to 1 |
| 2nd-Order Negative Occasion Setting (N) | \( \overline{\Delta N}^n_a = \overline{A}^n_a \bar{\alpha} \overline{\Delta N}^n + [N_{\text{sum}}]_{\text{(n)}} - [N_{\text{max}}]_{\text{(n)}} \) | 0 to 1 |
| Ability for CS to be Modulated by 1st-Order Positive Occasion Setter (P) | \( \overline{\Delta P}^n_a = \overline{A}^n_a \bar{\alpha} \overline{\Delta P}^n + [P_{\text{sum}}]_{\text{(n)}} \) | 0 to 1 |
| Ability for CS to be Modulated by 1st-Order Negative Occasion Setter (N) | \( \overline{\Delta N}^n_a = \overline{A}^n_a \bar{\alpha} \overline{\Delta N}^n + [N_{\text{sum}}]_{\text{(n)}} \) | 0 to 1 |
| 1st-Order Occasion Setting Ambiguity (γ2; "Gamma 2") | \( \overline{\gamma}2^n_a = P_{B}^{n} + N_{B}^{n} \) | 0 to 1 |
| US-Related Variables | | |
| US Presentation (λ; "Lambda") | λ | Usually: if US occurs, λ = 1; if not, λ = 0 | 0 to 1 |
| Absence of Expected US (λ; "Lambda Bar") | \( \overline{\lambda} = \overline{V_{\text{sum}}}^{(n-1)} \) | 0 to 1 |
| Excitatory Learning Gating (A; "Big Lambda") | A | If US occurs, A = 1; if not, A = 0 | binary 0 or 1 |
| Inhibitory Learning Gating (A; "Big Lambda Bar") | \( \overline{A} = \overline{A} = 1 \) if A = 0 and \( \overline{V_{\text{sum}}}^{(n-1)} > 0 \); otherwise, \( \overline{A} = 0 \) | binary 0 or 1 |
| Free Parameters | | |
| Learning Rate (\( \alpha \); "Alpha") | Higher = Faster Learning Rate | 0 to 1 |
| Leaky Memory (β; "Iota") | Higher = Greater Retention of Learning | 0 to 1 |

The above recipe of CS ambiguity and the CS’s ability to be modulated by a 1st-order occasion setter is extended to 2nd OS. The mechanism through which 2nd OS is learned is γ2, which is the degree to which a 1st-order occasion setter is ambiguous (i.e., that the 1st-order occasion setter is both a 1st-order positive and negative occasion setter). If the 1st-order occasion setter is unambiguous (e.g., either a positive or negative 1st-order occasion setter), γ2 remains at 0, and no 2nd OS is learned. Once a given 1st-order occasion setter is ambiguous (i.e., is both positive and negative), γ2 becomes positive, allowing P2 and N2 to increase from 0 and for 2nd OS to be learned. 1st-order occasion setter ambiguity is necessary but not sufficient for 2nd OS to be learned. In order for 2nd OS to be learned, the individual must learn that the CS can be modulated by a 2nd-order positive or negative occasion setter (N2). This will occur if a stimulus/context (i.e., the 1st-order occasion setter) provides information about the CS's (non)reinforcement and if the stimulus/context is less salient than the CS itself. Usually: if US occurs, the CS is positive or negative (e.g., if \( P_{B} = 0.75 \); this will reduce P's value to a maximum of .75), thereby enabling estimation of partial reinforcement. Employing multiple i values across hierarchies, excitation, and inhibition allows for accurate prediction of participant responding.

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Fig 6. Examples of Formula Inputs and Outputs. The formula variables (e.g., V, V, P) represent presence/absence of stimuli on a hypothetical trial and their training history. Bar graphs indicate predicted responding (i.e., R) to CS based on its training history and presence/absence of occasion setters. Left column provides names of formula variables. Across each row from these variables are values of 0 or 1, corresponding to the values of the variables’ names on the left. The values used by each figure are located in a column directly to the left of each figure. For example, the top-left figure shows a CS with 1 for direct excitation (V) and 0 for all other values (i.e., this is a CS+). POS = positive occasion setting; NOS = negative occasion setting. Excitation is color-coded as teal; inhibition is color-coded as purple. a) Examples of Direct Associative Learning and Successful Occasion Setting. We arranged inputs and outputs in a 2x3 grid, where the first column shows direct learning, the second column shows successful 1st-order occasion setting, and the third column shows successful 2nd-order occasion setting. First row shows excitatory responses, and second row shows inhibitory responses. b) Examples of Unsuccessful Occasion Setting. In bottom-left figure, we provide an example to demonstrate that 1st-order occasion setters do not affect responding if either direct excitation or direct inhibition are 0 (in our example, a 1st-order negative occasion setter does not affect a CS+, whose direct inhibition = 0). Congruently, 2nd-order occasion setters do not affect responding if any of the following are 0: direct excitation, direct inhibition, 1st-order positive occasion setting, or 1st-order negative occasion setting. As examples, in the bottom-middle plot, we show a 2nd-order negative occasion setter will not affect a CS unless an ambiguous 1st-order occasion setter is present (i.e., no 1st-order occasion setter is present, so P and N = 0). In our bottom-right plot, a 2nd-order positive occasion setter will not affect a CS- for multiple reasons, such as direct excitation = 0 and having no 1st-order occasion setter present (i.e., P and N = 0).

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3. Our model includes gating variables which only allow excitation to be learned on reinforced trials (A; “big lambda”) and inhibition to be learned on non-reinforced trials (A; “big lambda bar”). These variables are binary, where $A = 1$ if the US occurs, and $A = 1$ if no US occurs and if $V_{\text{sum}} > 0$ on that trial (i.e., there was at least some expectation the US would occur).

4. Our model has $\alpha$ (“alpha”) as a free learning rate parameter. It also has $\iota$ (“iota”) as a free leaky memory parameter [50]. The $\alpha$ parameter is multiplied within the $\Delta$ formulas to estimate the speed of prediction error learning (e.g., $\Delta P_{B_n} = \Lambda_n \cdot \gamma_1 \Lambda_n \cdot (\Lambda_{n-1} P_{\text{sum}}(n-1))$). The $\iota$ parameters are multiplied by the $\Delta$ formulas to regulate how much excitation or inhibition is retained across trials at each hierarchical level (e.g., $P_{B_n} = P_{B_{n-1}} + \Delta P_{B_n} + \iota$). Chronologically, $\alpha$ is calculated before $\iota$, allowing $\alpha$ to set the learning rate and $\iota$ to maintain or reduce what was learned. Functionally, $\iota$ can maintain or lower the asymptote of predicted responding, allowing the asymptote to occur at any reinforcement rate (e.g., for 75% reinforcement, asymptote would be at .75 instead of a traditional asymptote of 1 for 100% reinforcement). Separate $\iota$s can be used with each learning variable (e.g., $V$, $V$, $P$, $N$, $P_2$, $N_2$) so they decrease independently, which enables different rates of partial reinforcement across excitation and inhibition at each hierarchical level. Thus, rather than each learning value reaching an asymptote of 1, $\iota$ allows them to asymptote at values between 0 and 1, which would reflect the individual’s perceived reinforcement rate of the stimulus(i) presented (see S8 Text for an example). In the main text experiments, we implemented $\iota$s parameters at the 1st- and 2nd-order occasion setting levels (i.e., $P$, $N$, $P_2$, $N_2$).

5. $\gamma_1$ and $\gamma_2$ are stimulus-specific ambiguity variables, where $\gamma_1$ measures CS ambiguity and $\gamma_2$ measures 1st-order occasion setter ambiguity. $\gamma_1$ equals direct excitation multiplied by direct inhibition (i.e., $V \cdot V$), and $\gamma_2$ equals 1st-order positive occasion setting multiplied by 1st-order negative occasion setting (i.e., $P \cdot N$). We conceptualize ambiguity as a learned phenomenon (i.e., an individual must learn that a stimulus is ambiguous). In accordance with attentional models of ambiguity [5,31,34,102,103], we believe $\gamma_1$ and $\gamma_2$ result in increased attention to other stimuli, the background, context, or situational factors that will disambiguate which outcome the CS and/or 1st-order occasion setter will signal on a given trial. This allows attention to be directed towards concrete objects (e.g., other stimuli, contexts) or abstract concepts (e.g., time as a context, the absence of specific stimuli) to disambiguate the meaning of the CS or 1st-order occasion setting. $\gamma_1$ is only included in the trial-by-trial $\Delta$ calculation for 1st-order occasion setting (i.e., $\Delta P$, $\Delta P$, $\Delta N$, and $\Delta N$), and $\gamma_2$ is only included in the trial-by-trial $\Delta$ calculations to learn 2nd-order occasion setting (i.e., $\Delta P_2$, $\Delta P_2$, $\Delta N_2$, $\Delta N_2$). In other words, if a CS is ambiguous, 1st-order occasion setting can be learned. If 1st-order occasion setter is ambiguous, 2nd-order occasion setting can be learned. As long as $\gamma_1 = 0$, no 1st-order occasion setting can be learned (and 2nd-order occasion setting cannot be learned); as long as $\gamma_2 = 0$, no 2nd-order occasion setting can be learned. Importantly, whether stimulus B can become a 1st-order occasion setter depends on the $\gamma_1$ for a different stimulus (e.g., stimulus A). For example, if stimulus A is ambiguous as a CS, then stimulus B can become its 1st-order occasion setter. The $\gamma_1$ and $\gamma_2$ formulas are calculated as the mean ambiguity of CSs or 1st-order occasion setters, respectively, present on a given trial, maintaining their 0–1 range. Similarly, whether stimulus C can become a 2nd-order occasion setter depends on the $\gamma_1$ and $\gamma_2$ for different stimulus (e.g., stimuli A
and B); if A is ambiguous as a CS and B is ambiguous as a 1\textsuperscript{st}-order occasion setter, then C can become a 2\textsuperscript{nd}-order occasion setter. Additionally, while “ambiguity” is a learned phenomenon that a stimulus has a mixed prediction of the US, “uncertainty” is a different construct in which the individual has relative difficulty predicting the US from a stimulus(i).

An ambiguous stimulus can be highly uncertain or have little-to-no uncertainty after training. For example, a 50% partially reinforced CS would be ambiguous (i.e., mixed association with the US) and relatively high in uncertainty (because we would not know which trial would lead to the US or not); conversely, an occasion-set CS (which is 100% reinforced on OS/CS trials and 0% reinforced on CS alone trials in our experiments) would be similarly ambiguous (i.e., mixed association with the US) but relatively low in uncertainty (because we can determine whether the CS will be reinforced based on the presence or absence of the occasion setter(s)).

6. Only CSs trained with a 1\textsuperscript{st}-order occasion setter can be affected by a 1\textsuperscript{st}-order occasion setter (i.e., \(P\) and \(N\)), and only CSs trained with a 2\textsuperscript{nd}-order occasion setter can be affected by a 2\textsuperscript{nd}-order occasion setter (i.e., \(P_2\) and \(N_2\)). This effect is most notably observed in transfer tests, where occasion setters will only affect a CS trained with a different occasion setter. Thus, our model predicts transfer only to CSs that are ambiguous and were trained with the specific type of occasion setter being tested.

7. Independence of direct associations and occasion setting has been repeatedly demonstrated across occasion setting studies [1,2,8]. Thus, in our model, direct excitation, direct inhibition, 1\textsuperscript{st}-order positive occasion setting, 1\textsuperscript{st}-order negative occasion setting, 2\textsuperscript{nd}-order positive occasion setting, and 2\textsuperscript{nd}-order negative occasion setting are (largely) independent from each other—meaning, a given stimulus can simultaneously have any value of \(V\), \(\bar{V}\), \(P\), \(N\), \(P_2\), and \(N_2\). For example, stimulus A can be a direct excitor (\(V = 1\)), a 1\textsuperscript{st}-order negative occasion setter (\(N = 1\)), and a 2\textsuperscript{nd}-order positive occasion setter (\(P_2 = 1\)). The only level of dependence between these variables in our model is that the rate at which inhibition is learned is dependent on the level of excitation (e.g., \(\bar{\lambda} = V_{\text{sum}}\)); otherwise, they are orthogonal.

8. The learning variables \(V\), \(\bar{V}\), \(P\), \(N\), \(P_2\), \(N_2\), \(\bar{P}\), \(\bar{N}\), \(\bar{P}_2\), and \(\bar{N}_2\) are computed together in a general “excitation minus inhibition” structure to produce a single output: R (i.e., responding). In the R formula, we first calculate direct excitation minus direct inhibition (i.e., \(V - \bar{V}\)). Functionally, this means direct excitation and direct inhibition will summate. To produce 1\textsuperscript{st}-order positive occasion setting, we multiply 1\textsuperscript{st}-order positive occasion setting (\(P\)) by direct inhibition (\(\bar{V}\)) and the CS’s ability to be modulated by a 1\textsuperscript{st}-order positive occasion setter (\(\bar{P}\)) (i.e., \(P * \bar{V} * \bar{P}\)). Functionally, this means that 1\textsuperscript{st}-order positive occasion setting will nullify direct inhibition if the CS has been trained with a 1\textsuperscript{st}-order positive occasion setter, leading to an excitatory response. To produce 1\textsuperscript{st}-order negative occasion setting, we multiply 1\textsuperscript{st}-order negative occasion setting (\(N\)) by direct excitation (\(V\)) and the CS’s ability to be modulated by a 1\textsuperscript{st}-order negative occasion setter (\(\bar{N}\)) (i.e., \(N * V * \bar{N}\)). Functionally, 1\textsuperscript{st}-order negative occasion setting will nullify direct excitation if the CS has been trained with a 1\textsuperscript{st}-order negative occasion setter, leading to an inhibitory response.

9. To produce 2\textsuperscript{nd}-order positive occasion setting, we multiply 2\textsuperscript{nd}-order positive occasion setting (\(P_2\)) by the ability of the CS to be modulated by a 2\textsuperscript{nd}-order positive occasion setter (\(P_2\)) and the entire 1\textsuperscript{st}-order negative occasion setting term (i.e., \(N * V * \bar{N}\)), resulting in \(P_2 * \bar{P}_2 * N * V * \bar{N}\). Functionally, this means that the 2\textsuperscript{nd}-order positive occasion setter will nullify 1\textsuperscript{st}-order negative occasion setting if the CS has been trained with a 2\textsuperscript{nd}-order...
positive occasion setter, leading to an excitatory response. To produce 2nd-order negative occasion setting, we multiply 2nd-order negative occasion setting (N2) by the ability of the CS to be modulated by a 2nd-order negative occasion setter (N2) and the entire 1st-order positive occasion setting term (i.e., P * V * P), resulting in N2 * N2 * P * V * P. Functionally, this means that the 2nd-order negative occasion setter will nullify 1st-order positive occasion setting if the CS has been trained with a 2nd-order negative occasion setter, leading to an inhibitory response.

10. Within a typical 1st-order positive occasion setting paradigm (e.g., B➔A+, A-), 1st-order positive occasion setting as a learning process is purported to occur, but our model also predicts that 1st-order negative occasion setting will occur on trials with the CS alone. While B is the 1st-order positive occasion setter, the absence of B becomes a 1st-order negative occasion setter. The opposite occurs with 1st-order negative occasion setting (B➔A-, A+), where B is a 1st-order negative occasion setter, and the absence of B is a 1st-order positive occasion setter. The same occurs for 2nd-order occasion setting. The ability to learn that the absence of a stimulus is an occasion setter corresponds with attentional models of ambiguity [5,31,34,102,103], where attention can be directed to abstract contextual or situational factors to determine whether a stimulus will predict an outcome on a given trial. It also follows an intuitive verbal explanation for predicting the US (e.g., “when A is preceded by B, the US occurs, but when A is presented alone, the US does not occur”).

11. In the context of an experiment in which participants receive reinforcement depending on the trial type, we assume that inhibitory stimuli (i.e., CS-) acquire inhibition. This is based on studies in which the CS- acts as a control stimulus for the CS+ and develops inhibitory properties [8,110,111]. Thus, for CS-s that were never reinforced (or CSs that were initially not reinforced), we yoked their excitation to relevant CSs with excitation in order for the CS-s to acquire inhibition. Specifically, the yoked pairs in both experiments were H- to G+ (both experiments); the yoked pairs in Experiment 1 were B- to C and J- to K; and the yoked pairs in Experiment 2 were D- to F, U- to N, and S- to F. This produced the expected form of responding.

Supporting information

S1 Fig. Zbozinek et al 2nd-Order Occasion Setting Formulas. Interactive figure of our computational model, where the user can enter inputs for the learning variables and observe the model’s predicted outputs [112].

S1 Text. Curtain during Transfer Test.

S2 Text. Data Quality Assurance.

S3 Text. Stimuli R and S.

S4 Text. Detailed Training Results. Contains two tables: i) Table A. Experiment 1 Training and Reminder Phase Statistical Analyses, and ii) Table B. Experiment 2 Training and Reminder Phase Statistical Analyses.
S5 Text. Detailed Transfer Test Results. Contains two tables: i) Table A. Experiment 1 Transfer Test Statistical Analyses, and ii) Table B. Experiment 2 Transfer Test Statistical Analyses. (DOCX)

S6 Text. Simulated vs Recovered Parameters and Parameter Designation. Contains one figure: Fig A. Simulated and Recovered Learning Rate (α) and Leaky Memory (β) Parameters. (DOCX)

S7 Text. Detailed Trial Sequence. Contains one table: Table A. Experiment Training Sequence. (DOCX)

S8 Text. Supplemental Experiment: 1st-Order Positive Occasion Setting with 75%/25% Reinforcement Rate. Contains two figures: i) Fig A. Supplementary Experiment Training Results, and ii) Fig B. Supplementary Experiment Real vs Model-Predicted Responding. (DOCX)

S9 Text. Additional Factors That Could Affect Whether Occasion Setting Is Learned. (DOCX)

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References
1. Fraser KM, Holland PC. Occasion setting. Behav Neurosci. 2019; 133: 145–175. https://doi.org/10.1037/bne0000306 PMID: 30907616
2. Trask S, Thrailkill EA, Bouton ME. Occasion setting, inhibition, and the contextual control of extinction in Pavlovian and instrumental (operant) learning. Behav Processes. 2017; 137: 64–72. https://doi.org/10.1016/j.beproc.2016.10.003 PMID: 27720958
3. Bouton ME. Context, time, and memory retrieval in the interference paradigms of Pavlovian learning. Psychol Bull. 1993; 114: 80–99. https://doi.org/10.1037/0033-2909.114.1.80 PMID: 8346330
4. Bouton ME. Context, ambiguity, and unlearning: sources of relapse after behavioral extinction. Biol Psychiatry. 2002; 52: 976–986. https://doi.org/10.1016/s0006-3223(02)01546-9 PMID: 12437938
5. Rosas JM, Aguilera JEC, Álvarez MMR, Abad MJF. Revision of Retrieval Theory of Forgetting: What does Make Information Context-Specific? Int J Psychol Psychoth. 2006; 6: 147–166.

6. Swartzentruber D. Perspectives on modulation: Modulator- and target-focused views. Occasion setting: Associative learning and cognition in animals. Washington, DC, US: American Psychological Association; 1998. pp. 167–197. https://doi.org/10.1037/10298-006

7. Bonardi C, Robinson J, Jennings D. Can existing associative principles explain occasion setting? Some old ideas and some new data. Behav Processes. 2017; 135: 5–18. https://doi.org/10.1016/j.behproc.2016.07.007 PMID: 27425659

8. Zbozinek TD, Wise T, Perez O, Qi S, Fanselow MS, Mobs D. Pavlovian occasion setting in human fear and appetitive conditioning: Effects of trait anxiety and trait depression. Behav Res Ther. 2021; 103986. https://doi.org/10.1016/j.brat.2021.103986 PMID: 34740100

9. Holland PC. Transfer of control in ambiguous discriminations. J Exp Psychol Anim Behav Process. 1991; 17: 231–248. https://doi.org/10.1037/0097-7403.17.3.231 PMID: 1890383

10. Holland PC. Transfer after serial feature positive discrimination training. Learn Motiv. 1986; 17: 243–268. https://doi.org/10.1016/0023-9690(86)90013-5

11. Holland PC. Transfer of negative occasion setting and conditioned inhibition across conditioned and unconditioned stimuli. J Exp Psychol Anim Behav Process. 1989; 15: 311–328. https://doi.org/10.1016/0023-9690(89)90014-6 PMID: 2794868

12. Morell JR, Davidson TL. Transfer across unconditioned stimuli in serial feature discrimination training. J Exp Psychol Anim Behav Process. 1986; 14: 380–386. https://doi.org/10.1016/0023-9690(86)90015-8 PMID: 3770157

13. Lamarre J, Holland PC. Transfer of inhibition after serial feature negative discrimination training. Learn Motiv. 1987; 18: 319–342. https://doi.org/10.1016/0023-9690(87)90014-4

14. Bonardi C, Bartle C, Jennings D. US specificity of occasion setting: Hierarchical or configural learning? Behav Processes. 2012; 90: 349–366. https://doi.org/10.1016/j.beproc.2012.03.005 PMID: 22459561

15. Bonardi C, Ward-Robinson J. Occasion Setters: Specificity to the US and the CS–US Association. Learn Motiv. 2001; 32: 349–366. https://doi.org/10.1006/lmot.2001.1089

16. Franssen M, Claes N, Vervliet B, Beckers T, Hermans D, Vervliet B. Reinstatement after human feature-positive discrimination learning. Behav Processes. 2017; 137: 73–83. https://doi.org/10.1016/j.beproc.2017.01.007 PMID: 28108345

17. Baeyens F, Vansteenwegen D, Hermans D, Vervliet B, Eelen P. Sequential and simultaneous feature positive discriminations: Occasion setting and configural learning in human Pavlovian conditioning. J Exp Psychol Anim Behav Process. 2001; 27: 279–295. https://doi.org/10.1016/S0023-9690(00)72179-7 PMID: 11497326

18. Baeyens F, Vervliet B, Vansteenwegen D, Beckers T, Hermans D, Eelen P. Simultaneous and sequential Feature Negative discriminations: Elemental learning and occasion setting in human Pavlovian conditioning. Learn Motiv. 2004; 35: 136–166. https://doi.org/10.1016/S0023-9690(03)00058-4

19. Arnold HM, Grahame NJ, Miller RR. Higher order occasion setting. Anim Learn Behav. 1991; 19: 58–64. https://doi.org/10.3758/BF03197860

20. Rescorla RA. Extinction of facilitation. J Exp Psychol Anim Behav Process. 1986; 12: 16.

21. Holland PC, Reeve CE. Acquisition and transfer of control by an ambiguous cue. Anim Learn Behav. 1991; 19: 113–124. https://doi.org/10.3758/BF03197867

22. Rescorla RA. Facilitation and excitation. J Exp Psychol Anim Behav Process. 1986; 12: 325–332. https://doi.org/10.1016/0023-9690(86)90018-6

23. Holland PC. Temporal determinants of occasion setting in feature-positive discriminations. Anim Learn Behav. 1986; 14: 111–120. https://doi.org/10.3758/BF03200045

24. Holm S. A Simple Sequentially Rejective Multiple Test Procedure. Scand J Stat. 1979; 6: 65–70.

25. Holland PC, Lamoureux JA, Han J-S, Gallagher M. Hippocampal lesions interfere with Pavlovian negative occasion setting. Hippocampus. 1999; 9: 143–157. https://doi.org/10.1002/(SICI)1098-1063(1999)2:4<143::AID-HIPO6>3.0.CO;2-Z PMID: 10226775

26. Lamarre J, Holland PC. Acquisition and transfer of feature-negative discriminations. Bull Psychon Soc. 1985; 23: 71–74. https://doi.org/10.3758/BF03329783

27. Watanabe S. Asymptotic Equivalence of Bayes Cross Validation and Widely Applicable Information Criterion in Singular Learning Theory. J Mach Learn Res. 2010; 11: 3571–3594.

28. Davidson TL, Rescorla RA. Transfer of facilitation in the rat. Anim Learn Behav. 1986; 14: 380–386. https://doi.org/10.3758/BF03320082

29. Wilson PN, Pearce JM. Selective transfer of responding in conditional discriminations. Q J Exp Psychol Sect B. 1990; 42: 41–68. https://doi.org/10.1080/14640749008401870 PMID: 2326493
30. Swartzentruber D. Modulatory mechanisms in Pavlovian conditioning. Anim Learn Behav. 1995; 23: 123–143. https://doi.org/10.3758/BF03199928
31. de Brugada I, Garcia-Hoz V, Bonardi C, Hall G. Role of stimulus ambiguity in conditional learning. J Exp Psychol Anim Behav Process. 1995; 21: 275–284. https://doi.org/10.1037/0097-7403.21.4.275 PMID: 7595235
32. Kagan J. Categories of Novelty and States of Uncertainty. Rev Gen Psychol. 2009; 13: 290–301. https://doi.org/10.1037/a0017142
33. Ross RT, Holland PC. Conditioning of simultaneous and serial feature-positive discriminations. Anim Learn Behav. 1981; 9: 293–303. https://doi.org/10.3758/BF03197835
34. Bouton ME. Context, Ambiguity, and Classical Conditioning. Curr Dir Psychol Sci. 1994; 3: 49–53. https://doi.org/10.1111/1467-8721.ep10769943
35. Miller RR, Oberling P. Analogies between occasion setting and Pavlovian conditioning. Occasion setting: Associative learning and cognition in animals. Washington, DC, US: American Psychological Association; 1998. pp. 3–35. https://doi.org/10.1037/10298-001
36. Alfaro F, Malleja J, Laborda MA, Cañete A, Miguez G. Assessing the blocking of occasion setting. Behav Processes. 2018; 154: 52–59. https://doi.org/10.1016/j.beproc.2018.02.008 PMID: 29444454
37. Duits P, Cath DC, Lissek S, Hox JJ, Hamm AO, Engelhard IM, et al. Updated meta-analysis of classical fear conditioning in the anxiety disorders. Depress Anxiety. 2015; 32: 239–253. https://doi.org/10.1002/da.22353 PMID: 25703487
38. Lissek S, Powers AS, McClure EB, Phelps EA, Woldehawariat G, Grillon C, et al. Classical fear conditioning in the anxiety disorders: a meta-analysis. Behav Res Ther. 2005; 43: 1391–1424. https://doi.org/10.1016/j.brat.2004.10.007 PMID: 15895564
39. Craske MG, Woltzky-Taylor KB, Mineka S, Zinbarg R, Waters AM, Yehsek-Schallhorn S, et al. Elevated responding to safe conditions as a specific risk factor for anxiety versus depressive disorders: Evidence from a longitudinal investigation. J Abnorm Psychol. 2012; 121: 315–324. https://doi.org/10.1037/a0025738 PMID: 21988452
40. Darke S. Anxiety and working memory capacity. Cogn Emot. 1988; 2: 145–154. https://doi.org/10.1080/02699938808408071
41. Shackman AJ, Sarinopoulos I, Maxwell JS, Pizzagalli DA, Lavric A, Davidson RJ. Anxiety selectively disrupts visuospatial working memory. Emotion. 2006; 6: 40–61. https://doi.org/10.1037/1528-3542.6.1.40 PMID: 16637749
42. Eysenck M, Payne S, Derakshan N. Trait anxiety, visuospatial processing, and working memory. Cogn Emot. 2005; 19: 1214–1228. https://doi.org/10.1080/02699930500260245
43. Carleton RN, Mulvogue MK, Thibodeau MA, McCabe RE, Antony MM, Asmundson GJG. Increasingly certain about uncertainty: Intolerance of uncertainty across anxiety and depression. J Anxiety Disord. 2012; 26: 468–479. https://doi.org/10.1016/j.janxdis.2012.01.011 PMID: 22366534
44. Jensen D, Cohen JN, Mennin DS, Fresco DM, Heimberg RG. Clarifying the unique associations among intolerance of uncertainty, anxiety, and depression. Cogn Behav Ther. 2016; 45: 431–444. https://doi.org/10.1016/j.cbt.2016.1197308 PMID: 227314213
45. Craske MG, Treanor M, Conway CC, Zbozinek T, Vervliet B. Maximizing exposure therapy: An inhibitory learning approach. Behav Res Ther. 2014; 58: 10–23. https://doi.org/10.1016/j.brat.2014.04.006 PMID: 24864005
46. Rescorla RA, Wagner AR. A theory of Pavlovian conditioning: Variations in the effectiveness of reinforcement and nonreinforcement. Class Cond II Curr Res Theory. 1972; 2: 64–99.
47. Gershman SJ, Niv Y. Exploring a latent cause theory of classical conditioning. Learn Behav. 2012; 40: 255–268. https://doi.org/10.3758/s13420-012-0080-0 PMID: 22927000
48. Gershman SJ. Context-dependent learning and causal structure. Psychon Bull Rev. 2017; 24: 557–565. https://doi.org/10.3758/s13423-016-1110-x PMID: 27418259
49. Wise T, Michely J, Dayan P, Dolan RJ. A computational account of threat-related attentional bias. PLOS Comput Biol. 2019; 15: e1007341. https://doi.org/10.1371/journal.pcbi.1007341 PMID: 31600187
50. Vogel EH, Ponce FP, Wagner AR. A theoretical analysis of transfer of occasion setting: SOP with replaced elements. Behav Processes. 2017; 137: 19–32. https://doi.org/10.1016/j.beproc.2016.06.013 PMID: 27346424
52. Zackheim J, Myers C, Gluck M. A temporally sensitive recurrent network model of occasion setting. Occasion setting: Associative learning and cognition in animals. Washington, DC, US: American Psychological Association; 1998. pp. 319–342. https://doi.org/10.1037/10298-011

53. Holland PC. Occasion setting with simultaneous compounds in rats. J Exp Psychol Anim Behav Proc. 1989; 15: 183–193. https://doi.org/10.1037/0097-7403.15.3.183

54. Esclassan F, Coutureau E, Scala GD, Marchand AR. A Cholinergic-Dependent Role for the Entorhinal Cortex in Trace Fear Conditioning. J Neurosci. 2009; 29: 8087–8093. https://doi.org/10.1523/JNEUROSCI.0543-09.2009 PMID: 19553448

55. Suh J, Rivest AJ, Nakashiba T, Tominaga T, Tonegawa S. Entorhinal Cortex Layer III Input to the Hippocampus Is Crucial for Temporal Association Memory. Science. 2011; 334: 896–901. https://doi.org/10.1126/science.1244634 PMID: 24457215

56. Kitamura T, Pignatelli M, Suh J, Kohara K, Yoshiki A, Abe K, et al. Island Cells Control Temporal Association Memory. Science. 2014; 343: 8087–8093. https://doi.org/10.1037/JNEUROSCI.0543-09.2009 PMID: 19553448

57. Kitamura T, Macdonald CJ, Tonegawa S. Entorhinal–hippocampal neuronal circuits bridge temporally discontinuous events. Learn Mem. 2015; 22: 438–443. https://doi.org/10.1101/lm.038687.115 PMID: 26286654

58. Ryou J-W, Cho S-Y, Kim H-T. Lesions of the Entorhinal Cortex Impair Acquisition of Hippocampal-Dependent Trace Conditioning. Neurobiol Learn Mem. 2001; 75: 121–127. https://doi.org/10.1006/nlme.2000.3966 PMID: 11222054

59. Tanninen SE, Morrissey MD, Takehara-Nishiuchi K. Unilateral Lateral Entorhinal Inactivation Impairs Memory Expression in Trace Eyeblink Conditioning. PLOS ONE. 2013; 8: e84543. https://doi.org/10.1371/journal.pone.0084543 PMID: 24367674

60. Morrissey MD, Maal-Bared G, Brady S, Takehara-Nishiuchi K. Functional Dissociation within the Entorhinal Cortex for Memory Retrieval of an Association between Temporally Discontiguous Stimuli. J Neurosci. 2012; 32: 5356–5361. https://doi.org/10.1523/JNEUROSCI.5227-11.2012 PMID: 22514300

61. Suter EE, Weiss C, Disterhoft JF. Perirhinal and postrhinal, but not lateral entorhinal, cortices are essential for acquisition of trace eyeblink conditioning. Learn Mem. 2013; 20: 80–84. https://doi.org/10.1101/lm.028894.112 PMID: 23322556

62. Davidson TL, Jarrard LE. Retention of concurrent conditional discriminations in rats with ibotenate lesions of hippocampus. Psychobiology. 1989; 17: 49–60. https://doi.org/10.3758/BF03327217

63. Jarrard LE, Davidson TL. Acquisition of concurrent conditional discriminations in rats with ibotenate lesions of hippocampus and of subiculum. Psychobiology. 1990; 18: 68–73. https://doi.org/10.3758/BF03327217

64. Jarrard LE, Davidson TL. On the hippocampus and learned conditional responding: Effects of aspiration versus ibotenate lesions. Hippocampus. 1991; 1: 107–117. https://doi.org/10.1002/hipo.4500110 PMID: 1669340

65. Ross RT, Orr WB, Holland PC, Berger TW. Hippocampectomy disrupts acquisition and retention of learned conditional responding. Behav Neurosci. 1984; 98: 211–225. https://doi.org/10.1037/0735-7044.98.2.211 PMID: 6721923

66. Yoon T, Graham LK, Kim JJ. Hippocampal lesion effects on occasion setting by contextual and discrete stimuli. Neurobiol Learn Mem. 2011; 95: 176–184. https://doi.org/10.1016/j.nlm.2010.07.001 PMID: 20643219

67. Baxter MG, Murray EA. The amygdala and reward. Nat Rev Neurosci. 2002; 3: 563–573. https://doi.org/10.1038/nnr875 PMID: 12094212

68. Gallagher M, Holland PC. The amygdala complex: multiple roles in associative learning and attention. Proc Natl Acad Sci. 1994; 91: 11771–11776. https://doi.org/10.1073/pnas.91.25.11771 PMID: 7991534

69. Janak PH, Tye KM. From circuits to behaviour in the amygdala. Nature. 2015; 517: 284–292. https://doi.org/10.1038/nature14188 PMID: 25592533

70. Wassum KM, Izquierdo A. The basolateral amygdala in reward learning and addiction. Neurosci Biobehav Rev. 2015; 57: 271–283. https://doi.org/10.1016/j.neubiorev.2015.08.017 PMID: 26341938

71. Chaudhri N, Sahuque LL, Janak PH. Context-Induced Relapse of Conditioned Behavioral Responding to Ethanol Cues in Rats. Biol Psychiatry. 2008; 64: 203–210. https://doi.org/10.1016/j.biopsych.2008.03.007 PMID: 18423574

72. Chaudhri N, Woods CA, Sahuque LL, Gill TM, Janak PH. Unilateral inactivation of the basolateral amygdala attenuates context-induced renewal of Pavlovian-conditioned alcohol-seeking. Eur J Neurosci. 2013; 38: 2751–2761. https://doi.org/10.1111/ejn.12278 PMID: 23758059
73. Fuchs RA, See RE. Basolateral amygdala inactivation abolishes conditioned stimulus- and heroin-induced reinstatement of extinguished heroin-seeking behavior in rats. Psychopharmacology (Berl). 2002; 160: 425–433. https://doi.org/10.1007/s00213-001-0997-7 PMID: 11919670

74. Fuchs RA, Evans KA, Ledford CC, Parker MP, Case JM, Mehta RH, et al. The Role of the Dorsomedial Prefrontal Cortex, Basolateral Amygdala, and Dorsal Hippocampus in Contextual Reinstatement of Cocaine Seeking in Rats. Neuropsychopharmacology. 2005; 30: 296–309. https://doi.org/10.1038/sj.npp.1300559 PMID: 15483559

75. Hobin JA, Goosens KA, Maren S. Context-Dependent Neuronal Activity in the Lateral Amygdala Represents Fear Memories after Extinction. J Neurosci. 2003; 23: 8410–8416. https://doi.org/10.1523/JNEUROSCI.23-23-08410.2003 PMID: 12968003

76. Lasseter HC, Wells AM, Xie X, Fuchs RA. Interaction of the Basolateral Amygdala and Orbitofrontal Cortex is Critical for Drug Context-Induced Reinstatement of Cocaine-Seking Behavior in Rats. Neuropsychopharmacology. 2011; 36: 711–720. https://doi.org/10.1038/npp.2010.209 PMID: 21124303

77. Hara K, Adams A, Milland K, Savage S, Callison-Burch C, Bigham JP. A Data-Driven Analysis of Workers’ Earnings on Amazon Mechanical Turk. Proceedings of the 2018 CHI Conference on Human Factors in Computing Systems. New York, NY, USA: Association for Computing Machinery; 2018. pp. 1–14. Available: https://doi.org/10.1145/3173574.3174023

78. Balleine BW, Leung BK, Ostlund SB. The orbitofrontal cortex, predicted value, and choice. Ann N Y Acad Sci. 2011; 1239: 43–50. https://doi.org/10.1111/j.1749-6632.2011.06270.x PMID: 22145874

79. Keiflin R, Reese RM, Woods CA, Janak PH. The Orbitofrontal Cortex as Part of a Hierarchical Neural System Mediating Choice between Two Good Options. J Neurosci. 2013; 33: 15989–15998. https://doi.org/10.1523/JNEUROSCI.0026-13.2013 PMID: 24083559

80. Rudebeck PH, Murray EA. The Orbitofrontal Oracle: Cortical Mechanisms for the Prediction and Evaluation of Specific Behavioral Outcomes. Neuron. 2014; 84: 1143–1156. https://doi.org/10.1016/j.neuron.2014.10.049 PMID: 25521376

81. Sharpe MJ, Schoenbaum G. Back to basics: Making predictions in the orbitofrontal–amygdala circuit. Neurobiol Learn Mem. 2016; 131: 201–206. https://doi.org/10.1016/j.nlm.2016.04.009 PMID: 27112314

82. Wilson RC, Takahashi YK, Schoenbaum G, Niv Y. Orbitofrontal Cortex as a Cognitive Map of Task Space. Neuron. 2014; 81: 267–279. https://doi.org/10.1016/j.neuron.2013.11.005 PMID: 24462094

83. Shobe JL, Bakhurin K, Claar LD, Masmanidis SC. Selective Modulation of Orbitofrontal Network Activity during Negative Occasion Setting. J Neurosci. 2017; 37: 9415–9423. https://doi.org/10.1523/JNEUROSCI.0572-17.2017 PMID: 28847808

84. Phelps EA, Delgado MR, Nearing KI, LeDoux JE. Extinction Learning in Humans: Role of the Amygdala and vmPFC. Neuron. 2004; 43: 897–905. https://doi.org/10.1016/j.neuron.2004.08.042 PMID: 15363399

85. Dunsmoor JE, Kroes MCW, Li J, Daw ND, Simpson HB, Phelps EA. Role of Human Ventromedial Prefrontal Cortex in Learning and Recall of Enhanced Extinction. J Neurosci. 2019; 39: 3264–3276. https://doi.org/10.1523/JNEUROSCI.2713-18.2019 PMID: 30782974

86. Milad MR, Quirk GJ. Neurons in medial prefrontal cortex signal memory for fear extinction. Nature. 2002; 420: 70–74. https://doi.org/10.1038/nature01138 PMID: 12422216

87. Milad MR, Wright CI, Orr SP, Pitman RK, Quirk GJ, Rauch SL. Recall of fear extinction in humans activates the ventromedial prefrontal cortex and hippocampus in concert. Biol Psychiatry. 2007; 62: 446–454. https://doi.org/10.1016/j.biopsych.2006.10.011 PMID: 17217927

88. Lange I, Goossens L, Michelesi S, Bakker J, Vervliet B, Marcelis M, et al. Neural responses during extinction learning predict exposure therapy outcome in phobia: results from a randomized-controlled trial. Neuropsychopharmacology. 2020; 45: 534–541. https://doi.org/10.1038/s41386-019-0467-8 PMID: 31352467

89. Sparta DR, Smithius J, Stamatakis AM, Jennings JH, Kantak PA, Ung RL, et al. Inhibition of projections from the basolateral amygdala to the entorhinal cortex disrupts the acquisition of contextual fear. Front Behav Neurosci. 2014; 8: https://doi.org/10.3389/fnbeh.2014.00129 PMID: 24834031

90. Pitkänen A, Pikkarainen M, Nurminen N, Ylinen A. Reciprocal Connections between the Amygdala and the Hippocampal Formation, Perirhinal Cortex, and Postrhinal Cortex in Rat: A Review. Ann N Y Acad Sci. 2000; 911: 369–391. https://doi.org/10.1111/j.1749-6632.2000.tb06738.x PMID: 10911886

91. Amunts K, Kedo O, Kindler M, Pieperhoff F, Mohlberg H, Shah NJ, et al. Cytoarchitectonic mapping of the human amygdala, hippocampal region and entorhinal cortex: intersubject variability and probability maps. Brain Struct Funct. 2005; 5–6: 343–352. https://doi.org/10.1007/s00429-005-0025-5 PMID: 16208455
92. Yang Y, Wang J-Z. From Structure to Behavior in Basolateral Amygdala-Hippocampus Circuits. Front Neural Circuits. 2017;11. https://doi.org/10.3389/fncir.2017.00086
93. Witter MP, Doan TP, Jacobsen B, Nilssen ES, Ohara S. Architecture of the Entorhinal Cortex A Review of Entorhinal Anatomy in Rodents with Some Comparative Notes. Front Syst Neurosci. 2017;11. https://doi.org/10.3389/fnsys.2017.00046 PMID: 28701931
94. Hoover WB, Vertes RP. Projections of the medial orbital and ventral orbital cortex in the rat. J Comp Neurol. 2011; 519: 3766–3801. https://doi.org/10.1002/cne.22733 PMID: 21800317
95. Kondo H, Witter MP. Topographic organization of orbitofrontal projections to the parahippocampal region in rats. J Comp Neurol. 2014; 522: 772–793. https://doi.org/10.1002/cne.23442 PMID: 23897637
96. Catenoix H, Magnin M, Guénot M, Isnard J, Mauguière F, Ryvlin P. Hippocampal-orbitofrontal connectivity in human: An electrical stimulation study. Clin Neurophysiol. 2005; 116: 1779–1784. https://doi.org/10.1016/j.clinph.2005.03.016 PMID: 16002335
97. Carmichael ST, Price JL. Limbic connections of the medial orbital and prefrontal cortex: Comparative anatomy in macaque monkeys. J Comp Neurol. 1995; 363: 615–641. https://doi.org/10.1002/cne.903630408 PMID: 8847421
98. Öngür D, Ferry AT, Price JL. Architectonic subdivision of the human orbital and prefrontal cortex. J Comp Neurol. 2003; 460: 425–449. https://doi.org/10.1002/cne.10609 PMID: 12692859
99. Peer E, Brandimarte L, Samat S, Acquisti A. Beyond the Turk: Alternative platforms for crowdsourcing behavioral research. J Exp Soc Psychol. 2017; 70: 153–163. https://doi.org/10.1016/j.jesp.2017.01.006
100. Holland PC. Differentia effects of reinforcement of an inhibitory feature after serial and simultaneous feature negative discrimination training. J Exp Psychol Anim Behav Process. 1984; 10: 461–475. https://doi.org/10.1037/0097-7403.10.4.461 PMID: 6941607
101. Holland PC. Acquisition and transfer of conditional discrimination performance. J Exp Psychol Anim Behav Process. 1989; 15: 154–165. https://doi.org/10.1037/0097-7403.15.2.154
102. Molet M, Urcelay GP, Miguez G, Miller RR. Using context to resolve temporal ambiguity. J Exp Psychol Anim Behav Process. 2010; 36: 126–136. https://doi.org/10.1037/a0016055 PMID: 20141323
103. Bouton M, Nelson J. Context-Specificity of Target Versus Feature Inhibition in a Feature-Negative Discrimination. J Exp Psychol Anim Behav Process. 1994; 20: 51–65. PMID: 8308493
104. Rescorla RA. Facilitation based on inhibition. Anim Learn Behav. 1988; 16: 169–176. https://doi.org/10.3758/BF03290961
105. Rescorla RA. Separate reinforcement can enhance the effectiveness of modulators. J Exp Psychol Anim Behav Process. 1991; 17: 259–269. https://doi.org/10.1037/0097-7403.17.3.259
106. Rescorla RA. Transfer of inhibition and facilitation mediated by the original target stimulus. Anim Learn Behav. 1991; 19: 65–70. https://doi.org/10.3758/BF03197861
107. Swartzentruber D, Rescorla RA. Modulation of trained and extinguished stimuli by facilitators and inhibitors. Anim Learn Behav. 1994; 22: 309–316. https://doi.org/10.3758/BF03209839
108. Holland PC, Coldwell SE. Transfer of inhibitory stimulus control in operant feature-negative discriminations. Learn Motiv. 1993; 24: 345–375. https://doi.org/10.1006/ldmot.1993.1020
109. Morell JR, Holland PC. Summation and transfer of negative occasion setting. Anim Learn Behav. 1993; 21: 145–153. https://doi.org/10.3758/BF03213394
110. Spence KW. The nature of discrimination learning in animals. Psychol Rev. 1936; 43: 427–449. https://doi.org/10.1037/h0056975
111. Rescorla RA. Stimulus generalization of excitation and inhibition. Q J Exp Psychol. 2006; 59: 53–67. https://doi.org/10.1080/17470210500162094 PMID: 16556558
112. Bokeh Development Team. Bokeh: Python library for interactive visualization. 2018. Available: http://www.bokeh.pydata.org