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Figs. S1 to S4
Tables S1 and S2
References

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Dissociable Roles of Ventral and Dorsal Striatum in Instrumental Conditioning

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Instrumental conditioning studies how animals and humans choose actions appropriate to the affective structure of an environment. According to recent reinforcement learning models, two distinct components are involved: a "critic," which learns to predict future reward, and an "actor," which maintains information about the rewarding outcomes of actions to enable better ones to be chosen more frequently. We scanned human participants with functional magnetic resonance imaging while they engaged in instrumental conditioning. Our results suggest partly dissociable contributions of the ventral and dorsal striatum, with the former corresponding to the critic and the latter corresponding to the actor.

The ability to orient toward specific goals in the environment and control actions flexibly in pursuit of those goals is a hallmark of adaptive behavior. Instrumental conditioning, the most basic form of such behavior, allows an organism to learn contingencies between its own responses and rewarding or punishing outcomes (1–5). Models of reinforcement learning, such as the actor-critic (6) or advantage learning model (7), provide a two-process account of instrumental conditioning. One component, the critic, uses a temporal difference prediction error signal to update successive predictions of future reward associated with being at a state of the external and internal environment (determined by the arrangement of stimuli). The other component, the actor, uses a similar signal to modify stimulus-response or stimulus-response-reward associations in the form of a policy, so that actions associated with greater long-term reward are chosen more frequently on subsequent trials (8–11).

A putative neuronal correlate of these temporal difference prediction error signals is the phasic activity of dopamine neurons (12–

14), which send prominent projections to the ventral and dorsal striatum. Lesion and human imaging studies suggest that the ventral and dorsal striatum may have distinct functions. The former is implicated in reward and motivation (15). The latter is implicated in motor and cognitive control (16–19), specifically the learning of stimulus-response associations. On the basis of these findings, a putative neural substrate for reinforcement learning has been proposed (20), according to which dopaminergic projections to ventral striatum might be involved in reward prediction, corresponding primarily to the critic component of instrumental learning, whereas dopaminergic projections to dorsal striatum might be involved in the modulation of stimulus-response or stimulus-response-reward associations, corresponding to the instrumental actor.

We analyzed functional magnetic resonance imaging (fMRI) data from human participants performing an instrumental conditioning task. We used a reinforcement learning model called advantage learning (21) to calculate a reward prediction error signal and tested for correlations between that signal and evoked neural activity in the striatum. To dissociate stimulus-response learning from value prediction learning itself, we used a yoked Pavlovian conditioning task as a control condition. This task involves the same value predictions (critic), without ac-

tion selections (actor). If the ventral striatum corresponds to the critic, then this region should show prediction error activity during both the instrumental and Pavlovian conditioning tasks. If the dorsal striatum corresponds to the actor, then we would expect it to manifest stronger prediction error-related activity during instrumental than during Pavlovian conditioning.

The instrumental task was composed of two trial types: reward and neutral. In the reward trials, participants had to choose between one of two stimuli: one associated with a high probability of obtaining a juice reward (on 60% of occasions) and the other with a low probability of obtaining a juice reward (on 30% of occasions). In neutral trials, participants had to choose between two other stimuli associated with either a high (60%) or low (30%) probability of obtaining an affectively neutral solution. The Pavlovian task was identical to the instrumental task (with both reward and neutral trials), except that the computer made the selection and the participant's task was to indicate which stimulus had been chosen by the computer (Fig. 1A).

Participants rated the fruit juice as significantly more pleasant than the control tasteless solution in both the instrumental and Pavlovian conditioning tasks ($P < 0.001$; Fig. 1B). In the reward trials of the instrumental task, participants chose the high-probability action significantly more frequently than the low-probability action, but they showed no preference for the high-probability action in the neutral trials (Fig. 1C). There was evidence of "response matching" in the instrumental task (22) in that the ratio of responses made to the high-probability and low-probability stimuli was 1.92:1 during reward trials, a value very close to the actual 2:1 ratio of reward probabilities associated with the two stimuli.

To obtain a behavioral measure of learning in the Pavlovian conditioning task, we tested for differences in reaction times between responses in the reward and neutral trials (pooling over responses to high- and low-probability stimuli) between early and late phases of the session. Participants were faster to respond during the reward trials than during the neutral trials by the second block of trials (Fig. 1D).

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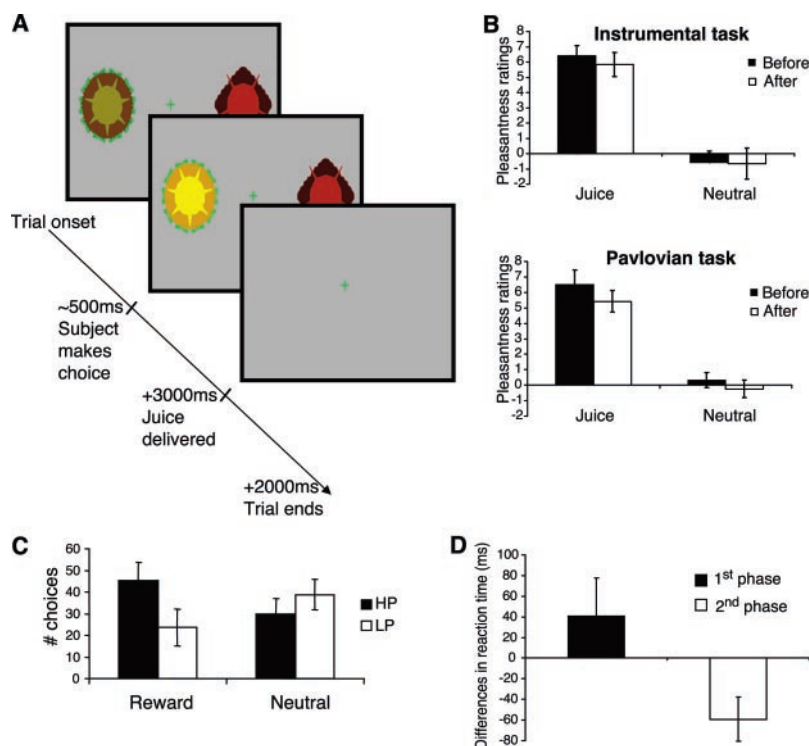


Fig. 1. (A) Illustration of instrumental task. Participant chose one of two fractals, which on each trial were randomly assigned to the left or right of the fixation cross. After the choice, the chosen fractal was illuminated, and 2000 ms later the outcome occurred. After another 3000 ms, the next trial was triggered. (B) Pleasantness ratings for the fruit juice and control tasteless solutions. Ratings were taken before and after the instrumental and Pavlovian conditioning sessions (+10, very pleasant; 0, neutral; -10, very unpleasant). Participants found the fruit juice to be significantly more pleasant than the control tasteless solution ($P < 0.001$). (C) Choices of high- versus low-probability actions in the instrumental task. Plot shows total number of choices of the high-probability (HP) and low-probability (LP) actions averaged across participants in both the reward and neutral trials of the instrumental conditioning task. Participants chose the high-probability action significantly more often than the low-probability action in reward trials ($P < 0.05$). (D) Reaction times during the Pavlovian conditioning task. Differences in reaction times are shown plotted between the reward and neutral trials during the Pavlovian conditioning task. In the second phase of the experiment, participants were faster to respond during reward trials than neutral trials (approaching significance at $P = 0.054$). This provides a behavioral measure of learning, providing some evidence that participants did acquire the Pavlovian associations. Error bars show mean \pm SEM.

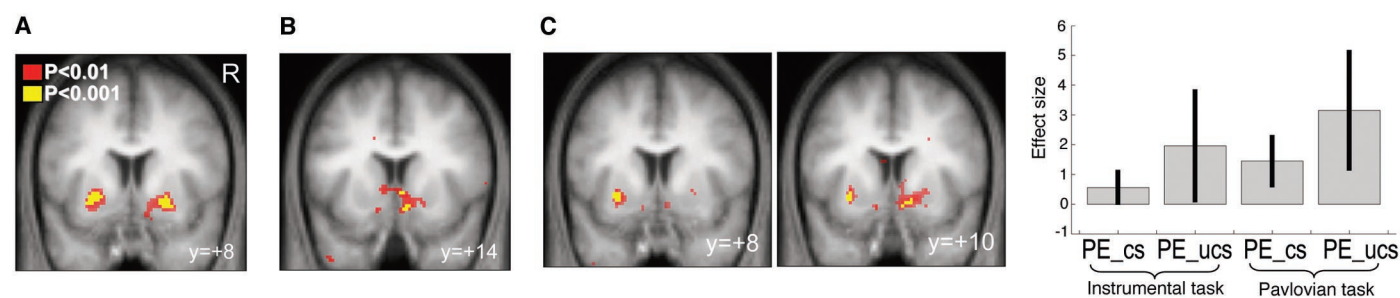
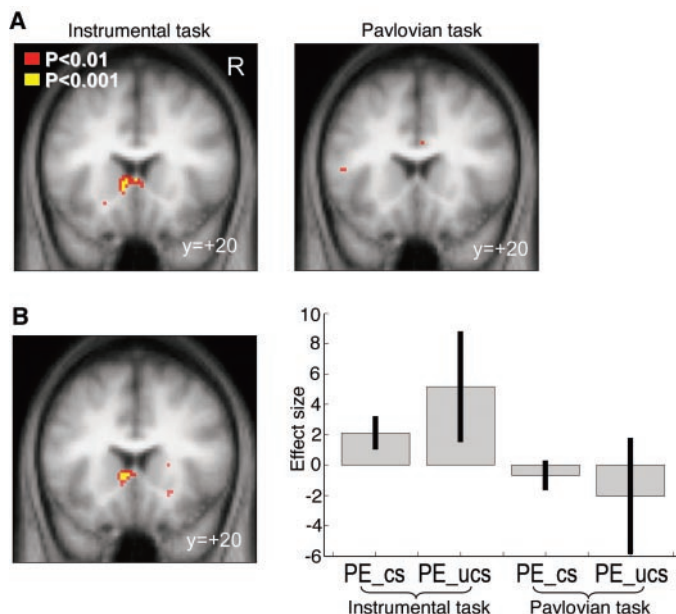


Fig. 2. Ventral striatum correlating with prediction error signal during Pavlovian and instrumental conditioning. (A) Reward prediction error responses in bilateral ventral striatum (ventral putamen) during Pavlovian conditioning in reward compared to neutral trials (left hemisphere coordinates: -26, 8, -4 mm; peak z -score = 3.98; right hemisphere coordinates: 26, 6, -8 mm; $z = 4.167$). Effects significant at $P < 0.001$ are shown in yellow, and effects significant at $P < 0.01$ are shown in red to illustrate the full extent of the activation. R, right. (B) Reward prediction error responses in ventral striatum (nucleus accumbens) during instrumental conditioning (right hemisphere coordinates: 6, 14, -2 mm; $z = 3.43$). (C) Results are shown for the conjunction of the prediction error signal for both types of conditioning.

Significant effects were found in bilateral ventral striatum [in the bilateral ventral putamen (left hemisphere coordinates: -28, 8, -6 mm; $z = 3.73$; right hemisphere coordinates: 20, 12, -8 mm; $z = 3.54$) and in the right nucleus accumbens (14, 10, -10 mm; $z = 3.21$)] at $P < 0.001$. Images in (A), (B), and the left and middle panels of (C) show coronal slices through different sections of ventral striatum (at $y = 8$ mm, $y = 14$ mm, $y = 8$ mm, $y = 10$ mm, respectively). A plot of the contrast estimates is also shown (bar chart, right) for the peak voxel in the conjunction analysis with prediction error (PE) effects at the time of presentation of the cue or conditioned stimulus (cs) and at the time of presentation of the reward or unconditioned stimulus (ucs), plotted separately for each type of conditioning.

Fig. 3. Dorsal striatum correlating with prediction error signal during instrumental conditioning. (A) Results depict the correlation of the prediction error signal with neural activity in the dorsal striatum for the instrumental task (left) and the Pavlovian task (right). Significant activations were found in the left anterior caudate nucleus ($-8, 22, 0$; $z = 3.84$). No significant effects were observed in the Pavlovian task at $P < 0.001$ or even $P < 0.01$. R, right. (B) Area of the dorsal striatum (anterior caudate nucleus) showing significantly greater prediction error responses in instrumental conditioning than in Pavlovian conditioning ($P < 0.001$; $-6, 22, 2$; $z = 3.78$) (left). A plot of the contrast estimates is also shown (right) for the peak voxel.



tion error responses in the left caudate nucleus were significantly enhanced in instrumental conditioning at $P < 0.001$ (Fig. 3B). The finding of enhanced temporal-difference prediction error-related responses in dorsal striatum during instrumental conditioning compared with Pavlovian conditioning is consistent with our hypothesis that this region plays a central role in implementing the instrumental actor.

Activity in both the ventral and dorsal striatum during instrumental-styled tasks has been shown previously in which a response is required to obtain an outcome (25, 26) but in which, notably, there is no explicit choice between different allowed responses. Our task embodies the essence of instrumental conditioning in that the participants had to choose between two options (overall favoring the high-probability action), which required their active engagement. An fMRI study of gambling in which participants were engaged in decision making has also reported activity in the dorsal striatum (27). Our results differentiate ventral and dorsal striatum according to their relative contributions to stimulus-reward and stimulus-response (or stimulus-response-reward) learning. Anatomical distinctions, such as those made between matrisomes and striasomes within the dorsal striatum (28, 29), have also been implicated in the implementation of critic and actor learning (9, 30) on the basis of their

differential control over and innervation by dopamine. However, these lie at a finer spatial scale than is accessible with the resolution of our neuroimaging technique.

Advantage learning, which underlies the prediction error signals used in instrumental conditioning, has been suggested as a bridge between goal-directed action selection, in which actions are chosen with reference to an explicit representation of the incentive value of the outcome or goal state, and habitual action selection, in which actions are elicited by the presentation of a specific stimulus, without incorporating a representation of the outcome itself (5, 31). Although we did not directly test it here, goal-directed forms of instrumental learning may rely on structures in the prefrontal cortex (4, 32).

Reinforcement learning links psychological ideas of stimulus-reward and stimulus-response-reward learning to computational and engineering ideas about adaptive optimal control (6) and a putative dopaminergic substrate (20, 33). The present study on instrumental conditioning, together with previous studies on Pavlovian conditioning (23, 24), suggests how different aspects of learning are parsed among parts of the basal ganglia.

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Materials and Methods

Figs. S1 to S4

References and Notes

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