

Coding of behaviorally relevant and irrelevant cue features in the nucleus accumbens

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1 Abstract

2 to do

3 Significance Statement (120 words)

4 to do

5 Introduction

6 Theories of nucleus accumbens (NAc) function generally agree that this brain structure contributes to moti-
7 vated behavior, with some emphasizing a role in learning from reward prediction errors (Joel, Doya, Schultz;
8 see also the addiction literature on the effects of drug rewards; Nestler, Kalivas; Carelli) and others a role in
9 the modulation of ongoing behavior through stimuli associated with motivationally relevant outcomes (in-
10 vigorating, directing; Nicola, Floresco, Salamone). These proposals echo similar ideas on the functions of
11 the neuromodulator dopamine (Schultz, Berridge, Maia/Frank, Cools), with which the NAc is tightly linked
12 functionally as well as anatomically (Haber, Sesack, Takahashi).

13 Much of our understanding of NAc function comes from studies of how cues that predict motivationally
14 relevant outcomes (e.g. reward) influence behavior and neural activity in the NAc. Task designs that asso-
15 ciate such cues with rewarding outcomes provide a convenient access point eliciting conditioned responses
16 such as sign-tracking and goal-tracking (Robinson), pavlovian-instrumental transfer (Balleine) and enhanced
17 response vigor (Niv; McGinty), which tend to be affected by NAc manipulations (Flagel, Balleine, Chang;
18 although not always straightforwardly: Hauber, Chang). Similarly, analysis of reward prediction errors typ-
19 ically proceeds by establishing an association between a cue and subsequent reward, with NAc responses
20 transferring from outcome to the cue with learning (Schultz, Schoenbaum, Carelli). WHAT ABOUT HU-
21 MAN WORK

22 Surprisingly, although substantial work has been done on the coding of outcomes predicted by such cues
23 (e.g. reward value; Hollerman/Schultz, Roesch, Day; reward identity; Cooch), much less is known about
24 how reward-predictive cues themselves are encoded in the NAc (Hayden from primate realm). This is an
25 important issue for at least two reasons. First, in reinforcement learning, motivationally relevant outcomes
26 are typically temporally delayed relative to the cues that predict them. In order to solve the problem of as-
27 signing credit (or blame) across such temporal gaps, some trace of preceding activity needs to be maintained
28 (Maia?). Since NAc is a primary target of DA signals interpretable as RPEs, and NAc lesions impair RPEs

29 related to timing, its activity trace will help determine what can be learned when RPEs arrive (Takahashi).

30 Second, for ongoing behavior, the relevance of cues typically depends on context. In experimental settings,
31 context may include the identity of a preceding cue (occasion setter, Holland, Kesner), spatial or configural
32 arrangements (Good/Honey, Eichenbaum), and unsignaled rules as occurs in set shifting and other cognitive
33 control tasks (CITE). In such situations, the question arises how selective, context-dependent processing of
34 reward-predictive cues is implemented. For instance, is there a gate prior to NAc, or are all cues represented
35 in NAc but their current values dynamically updated (FitzGerald; WHAT ARE SOME THEORETICAL
36 POSITIONS ON THIS)?

37 Thus, both from a learning and a flexible performance perspective, it is of interest to determine how cue iden-
38 tity is represented in the brain, with NAc of particular interest given its anatomical and functional position
39 at the center of motivational systems. We sought to determine whether cue features signalling identity are
40 represented in the NAc, and if cue identity is represented alongside other motivationally relevant variables,
41 such as cue value (Figure 1). To address this question, we designed an experiment in which multiple, distinct
42 sets of cues predicted the same outcome. We recorded the activity of NAc neurons as rats performed this
43 task.

44 [Figure 1 about here.]

45 **Methods**

46 **Subjects:**

47 Adult male Long-Evans rats ($n = 4$, Charles River, Saint Constant, QC) were used as subjects. Rats were in-

48 individually housed with a 12/12-h light-dark cycle, and tested during the light cycle. Rats were food deprived
49 to 85-90% of their free feeding weight (weight at time of implantation was 440 - 470 g), and water restricted
50 4-6 hours before testing. All experimental procedures were approved by the the University of Waterloo An-
51 imal Care Committee (protocol# 11-06) and carried out in accordance with Canadian Council for Animal
52 Care (CCAC) guidelines.

53 **Overall timeline:**

54 Each subject was first handled for seven days where they were exposed to the running room, the sucrose
55 solution, and the click of the valves upon approach to the receptacles. They were then shaped to run on the
56 task for seven days where they were restricted to running in the clockwise direction by presenting a physical
57 barrier to running counterclockwise. Rats underwent hyperdrive implantation after showing discrimination
58 of approach behavior for rewarded and unrewarded cues for three consecutive days according to a chi square
59 test. Rats were allowed to recover for a minimum of five days before being retrained on the task, and
60 recording began once performance returned to pre-surgery levels. Upon completion of recording, animals
61 were sacrificed and recording sites were histologically confirmed.

62 **Behavioral task and training:**

63 Rats were trained to run clockwise on an elevated, square-shaped track (100x100 cm) containing four possi-
64 ble reward locations (Figure 2). Rats initiated a trial by running down the length of an arm, and triggering a
65 photobeam located 24 cm from the start of each arm. Upon trial initiation, a light or sound cue was presented
66 that signaled the presence of absence of a 12% sucrose water reward (0.1 mL) at the upcoming site. A trial
67 was classified as an approach trial if the rat turned left at the decision point and made a nosepoke at the
68 reward receptacle (40 cm from the decision point), while trials were classified as a skip trial if the rat instead
69 turned right at the decision point and triggered the photobeam to initiate the following trial. There was a 1
70 second delay between a rewarded nosepoke and subsequent reward delivery. Trial length was determined by

71 measuring the length of time from cue onset until nosepoke (~~for approach trials~~, or from cue onset until or
72 the start of the following trial(~~for skip trials~~). Trials could only be initiated through clockwise progression
73 through the series of arms, and each entry into the subsequent arm on the track counted as a trial. Each
74 day rats were trained in both a light and sound block for 100 trials each. Within a block, one cue signaled
75 reward was available on that trial, while the other signaled reward was not available. Light block cues were a
76 flashing white light, and a constant yellow light. Sound block cues were a 2 kHz sine wave and a 8 kHz sine
77 wave whose amplitude was modulated from 0 to maximum by a 2 Hz sine wave. Reward-cue associations
78 were counterbalanced across rats. Cue presentation was pseudorandomized so that the same cue could not
79 be presented more than twice in a row. Block order within each day was also pseudorandomized, such that
80 the rat could begin a session with the same block for more than two days in a row. Each training or testing
81 day consisted of a 5 minute pre-session period on a pedestal, followed by the first block, then the second
82 block, then a 5 minute post-session period on the pedestal. Accuracy was determined by the proportion of
83 trials a rat approached each cue. Perfect performance would be 100% approach on approach trials (reward
84 available), and 0% approach on skip trials (no reward available). Rats were trained daily until they could
85 distinguish between the rewarded and unrewarded cues for both light and sound blocks for three consecutive
86 days according to a chi-square test, at which point they underwent surgery. Furthermore, we generated linear
87 mixed effects models to investigate the relationships between cue type and our behavioral variables, with
88 cue type was used as a fixed effect, and the ~~addition~~addtion of an intercept for rat identity as a random
89 effect. Average proportion of trials approached and trial length for a session were used as response variables.
90 Contribution of cue type to behavior was determined by comparing the full model to a model with cue type
91 removed for each behavioral variable.

92 [Figure 2 about here.]

93 **Surgery:**

94 Surgical procedures were as described previously (Malhotra et al., 2015). Briefly, animals were anesthetized

95 with isoflurane, induced with 5% in medical grade oxygen and maintained at 2% throughout the surgery
96 (0.8 L/min). Rats were then chronically implanted with a hyperdrive consisting of 16 independently drivable
97 tetrodes, either all 16 targeted for the right NAc (AP +1.4 mm and ML +1.6 mm, relative to bregma; Paxinos
98 and Watson, 2005), or 12 in the right NAc and 4 targeted at the mPFC (AP +3.0 mm and ML +0.6 mm,
99 relative to bregma; only data from NAc tetrodes were analyzed). Following surgery, all animals were given
100 a least five days to recover and lower tetrodes to the target (DV -6.0 mm) before being reintroduced to the
101 behavioral task.

102 **Data acquisition and preprocessing:**

103 After recovery, rats were placed back on the task for recording. NAc signals were acquired at 20 kHz with
104 a RHA2132 v0810 preamplifier (Intan) and a KJE-1001/KJD-1000 data acquisition system (Amplipex).
105 Signals were referenced against a tetrode placed in the corpus callosum above the NAc.

106 Candidate spikes for sorting into putative single units were obtained by band-pass filtering the data between
107 600-9000 Hz, thresholding and aligning the peaks (UltraMegaSort2k, Hull et al., 2011). Spike waveforms
108 were then clustered with KlustaKwik using energy and the first derivative of energy as features, and manually
109 sorted into units (MClust 3.5, A.D. Redish et al.). Isolated units containing a minimum of 200 spikes within
110 a session were included for subsequent analysis. Units were classified as high firing neurons if they had high
111 tonic firing rates marked by an absence of interspike intervals (ISIs) > 2 s, while medium spiny neurons had
112 a combination of ISIs > 2 s and phasic activity with shorter ISIs (Barnes 2005, Atallah 2014).

113 **Data analysis:**

114 To investigate the contribution of various cue features on NAc firing rates we first determined whether firing
115 rates for a unit were modulated by the onset of a cue by collapsing across all cues and comparing the firing
116 rates for the 1 s preceding cue-onset with the 1 s following cue-onset. Single units were considered to be

cue-responsive if a Wilcoxon signed-rank test comparing pre- and post-cue firing had a $p < .01$. (Excluded:
the mean firing rate difference between pre- and post-cue onset was within the lower or upper 2.5% of a
shuffled distribution, as there is redundancy using both, and I did both because I was paranoid about just
using one.). Cue-modulated units were then classified as either increasing or decreasing in response to the
cue if the post-cue activity was higher or lower than the pre-cue activity, respectively.

To determine the relative contribution of various task parameters to firing rate variance for units whose firing
was modulated by cue-onset (as in Figures 5,6), a forward selection stepwise general linear model (GLM)
was fit to each cue-responsive unit. Cue modality, cue location, cue outcome, approach behavior, trial length,
trial number, and trial history were used as predictors, and the 1 s post-cue firing rate as the response variable.
Units were classified as being modulated by a given task parameter if addition of the parameter significantly
improved model fit using deviance as the criterion ($p < .01$). A comparison of the R-squared value between
the final model and the final model minus the predictor of interest was used to determine the amount of firing
rate variance explained by the addition of that predictor for a given unit.

To better visualize responses to cues and enable subsequent population level analyses (as in Figures 5,7,8),
spike trains were convolved with a Gaussian kernel, and peri-event time histograms (PETHs) were generated
by taking the average of the convolved spike trains across trials for a given task condition. For analysis
of population-level responses for cue features (Figure 7), convolved spike trains for all units where cue
modality, cue location, or cue outcome explained a significant portion of firing rate variance were z-scored.
Within a given cue feature, normalized spike trains were then separated according to the preferred and non-
preferred cue condition (e.g. light vs sound block), and averaged across units to generate population-level
averages.

To visualize NAc representation of task space within cue conditions, normalized spike trains for all units
were ordered by the location of their maximum or minimum firing rate for a specified cue condition (Figure
8). To compare representation of task space across cue conditions for a cue feature, the ordering of units

¹⁴¹ derived for one condition (e.g. light block) was then applied to the normalized spike trains for the other
¹⁴² condition (e.g. sound block).

¹⁴³ To identify the responsivity of units to different cue features at the time of a nosepoke into a reward receptacle, the same cue-responsive units from the cue-onset analyses were analyzed at the time of nosepoke using
¹⁴⁵ identical analysis techniques (Figures 9,10,11).

¹⁴⁶ Given that some of our analyses compare firing rates across time, particularly comparisons across blocks,
¹⁴⁷ we sought to exclude units with unstable firing rates that would generate spurious results reflecting a drift
¹⁴⁸ in firing rate over time unrelated to our task. To do this we ran a Mann-Whitney U test comparing the cue-
¹⁴⁹ evoked firing rates for the first and second half of trials within a block, and excluded units from analysis that
¹⁵⁰ showed a significant change for either block. All analyses were completed in MATLAB R2015a, and the
¹⁵¹ code is available on GitHub.

¹⁵² **Histology:**

¹⁵³ Upon completion of the experiment, rats were anesthetized with 5% isoflurane, then asphyxiated with carbon
¹⁵⁴ dioxide. Transcardial perfusions were performed, and brains were fixed and removed. Brains were sliced
¹⁵⁵ in 50 um coronal sections and stained with thionin. Slices were visualized under light microscopy, tetrode
¹⁵⁶ placement was determined, and electrodes with recording locations in the NAc were analyzed (Figure 3).

¹⁵⁷ [Figure 3 about here.]

158 **Results**

159 **Behavior**

160 Rats were trained to discriminate between cues signaling the availability and absence of reward on a square
161 track with four identical arms for two distinct sets of cues. ~~During each session, rats were presented with~~
162 ~~two behavioral blocks containing different sensory cues in sequential order, a light and a sound block, with~~
163 ~~each block containing a cue that predicted reward, and a cue that signalled the absence of reward.~~ An
164 example learning curve is seen in Figure 4A,B. All four rats learned to discriminate between the rewarded
165 and unrewarded cue for both the light and sound blocks as determined by reaching significance ($p < .05$)
166 on a daily chi-square test comparing approach behavior for rewarded and unrewarded cues for each block,
167 for at least three consecutive days(~~range for time to criteria: 22 – 57 days~~). ~~Maintenance of behavioral~~
168 ~~performance during recording sessions was assessed using linear~~ Linear mixed effects models ~~for both~~
169 ~~proportion of trials where the rat approached the receptacle, and trial length. Analyses~~ revealed that cue type
170 had an influence on the likelihood of a rat to make an approach ($p < .001$), but not for the length of time
171 taken to complete a trial ($p = .13$)(Figure 4C,D).

172 [Figure 4 about here.]

173 **NAc neurons encode behaviorally relevant and irrelevant cue features**

174 **General responses to cue:**

175 ~~We sought to address which parameters of our task were encoded by NAc activity, specifically whether the~~
176 ~~NAc encodes aspects of motivationally relevant cues not directly tied to reward, such as the identity and~~

177 location of the cue, and whether this coding is independent or integrated with coding of cue outcome. To
178 do this we recorded a total of 443 units were recorded in the NAc from 4 rats over 57 sessions while they
179 performed on the square track (Table 1). Units that had less than 200 spikes, or exhibited a drift in firing
180 rate over the course of either block were excluded from further analysis. The activity of 133 (30%) of these
181 was modulated by the cue, with more units showing a decrease in firing ($n = 124$) than an increase ($n =$
182 47) around the time of cue-onset (Table 2). Within this group, 32 were classified as HFNs, while 139 were
183 classified as SPNs. To determine whether these cue-evoked responses were modulated by various aspects
184 of the task, we fit Fitting a GLM to each cue-modulated unit. Fitting GLMs unit revealed that a variety of
185 task parameters accounted for a significant portion of firing rate variance in NAc cue-modulated units, with
186 all task parameters except trial history being predictors in 19-38% of cue-modulated units, and explaining
187 4-9% on average, although there was substantial variability across cells (Figures 5, 6, Table 2(Figures ??,
188 6). Notably, there were units that discriminated between whether the rat was performing in the light or
189 sound block($n = 37 / 133$), which arm the rat was currently on($n = 50 / 133$), and whether the rat was
190 engaged in a rewarded or unrewarded trial ($n = 34 / 133$), suggesting that the NAc independently encodes
191 features of reward-predictive cues separate from expected outcome (Figure 5A-F). Furthermore, interactions
192 Interactions between multiple cue features appeared as significant predictors of firing rate variance for 9%
193 cue-modulated units($n = 11 / 133$), although this effect was modest (1-2% variance explained), suggesting a
194 weak effect for integrated coding across various aspects of a cue (relatively modest) (Figure 5G,H). Fitting a
195 GLM to all recorded units...*Jimmie finish this* (data not shown).

196 [Table 1 about here.]

197 [Table 2 about here.]

198 [Figure 5 about here.]

199 [Figure 6 about here.]

200 **Population level responses:**

201 NAc units do not work in isolation, and some information about the cue may be hidden in single unit
202 responses. To get a sense of how NAc output encoded cue information cue information was encoded at
203 the population level, we normalized firing activity for each unit that was modulated by a given cue feature
204 , then generated the cue feature was z-scored, then the population average for a cue feature was aligned to
205 cue-onset aligned population average firing rate for each of the cue features was generated (Fig 7). This
206 analyses revealed overall that cells that showed an increase upon cue presentation had stronger responses for
207 the preferred cue condition (Figure 7A,C,E). Interestingly, units that were classified as decreasing in response
208 to the cue showed a biphasic response at the population level, with a small peak at a time in alignment with
209 entry into the arm, followed by a sustained dip after cue-onset (Figure 7B,D,F). Units that were visualization
210 of the data revealed units whose activity was modulated by cue modality showed by a stronger increase in
211 response to the preferred cue modality, as well as a higher tonic firing rate to the preferred cue modality, most
212 notably in units that decreased in firing rate a difference in firing rate across blocks that extended beyond
213 the transient response to the cue (Figure 7A,B). Units that were modulated by cue location showed a graded
214 response to locations of decreasing preference (Figure 7C,D). Additionally, units whom had exhibited a de-
215 crease in firing in response to the cue and whose activity was modulated by cue outcome, showed a sustained
216 discriminatory response to rewarded and unrewarded cues response that extended beyond cue-onset (Figure
217 7F). Together, these visualizations of the averaged population responses revealed nuanced differences in the
218 way NAc units are modulated by cue conditions across cue features.

219 [Figure 7 about here.]

220 **NAc units segment the task:**

221 Another method to get a sense of how the NAc is behaving at the population level is to analyze NAc unit
222 activity across time, and NAc NAc neurons have been shown to have correlates across an entire task space.

223 To look at the distribution of responses throughout our task space and see if this distribution is modulated by
224 cue features, we z-scored the firing rate of each unit and plotted the normalized firing rates of all units aligned
225 to cue-onset and according to peak firing rate([Figure 8](#)). We did this separately for both the light and sound
226 blocks, and found a nearly uniform distribution of firing fields in task space that was not limited to alignment
227 to the cue (Figure 8A). Furthermore, to see if this population level activity was similar across blocks, we
228 also organized firing during the sound blocks according to the ordering derived from the light blocks. This
229 revealed~~that while there was some preservation of order, that~~ the overall firing was qualitatively different
230 across the two blocks. Additionally, given that the majority of our units showed an inhibitory response to the
231 cue, we also plotted the firing rates according to the lowest time in firing,~~and again found some maintenance~~
232 ~~of order, but largely different ordering across the two blocks (Figure 8B)~~. This process was repeated for
233 cue location and cue outcomes,~~showing that NAc segmentation of the task is qualitatively different across~~
234 ~~various cue conditions (Figure 8C–F)~~.

235 [Figure 8 about here.]

236 **Encoding of cue features is not limited to cue-onset:**

237 In order to be useful for learning, a trace of the cue must be maintained until the outcome. Fitting a GLM
238 to the firing rates of cue-modulated units at the time of a nosepoke response showed that a variety of units
239 still discriminated firing according to various cue features, but not other task parameters,~~showing that NAc~~
240 ~~activity discriminates various cue conditions well into a trial~~ (Table 3, [Figure 9](#)[Figures 9, 10](#)). Fitting a GLM
241 to all recorded units...*Jimmie finish this* (data not shown). ~~Population level averages for units that increased~~
242 ~~to cue-onset showed a ramping up of activity that peaked upon nosepoke, whereas units that decreased to~~
243 ~~cue-onset showed a gradual reduction of firing activity that reached a minimum upon nosepoke (Figure~~
244 ~~10)~~. Additionally, a peak is seen for preferred cue outcome in decreasing units at 1 second post cue-onset
245 ~~when reward was received, demonstrating an integration of expected and received reward (Figure 10F)~~.
246 Furthermore, aligning normalized peak firing rates to nosepoke onset, revealed a clustering of responses

247 around reward receipt **for all cue conditions where the rat would have received reward** (Figure 11). Alongside
248 this, fitting a GLM to the firing rates of cue-modulated units at the time of reward receipt, revealed 10 units
249 (8%) where cue outcome accounted for an average of 32% of firing rate variance (data not shown).

250 [Table 3 about here.]

251 [Figure 9 about here.]

252 [Figure 10 about here.]

253 [Figure 11 about here.]

254 Discussion

255 The present study found evidence for coding of multiple identifying features of motivationally relevant stim-
256 uli; the sensory modality of the presented cue, as well as its physical location within the track. Furthermore,
257 this coding was both independent, and intermixed with coding for the associated outcome of the cue as well
258 as motivational vigor, measured by time to complete the trial. At the population level, a tiling of task struc-
259 ture was observed such that all points within our analyzed task space was accounted for by the ordered peak
260 firing rates of all cells, and this tiling differed between blocks where sound or light cues were presented.
261 Cells that discriminated across blocks were not simply due to drifting of the signal across trials, as cells that
262 showed a drift in firing between the first and second half within a block were excluded from the analysis.
263 Furthermore, even though actions were stereotyped during correct trials, such that the rat always turned left
264 at the decision point to approach for reward, and right to skip the receptacle and initiate the next trial, cells
265 that were modulated by the expected value of the cue maintained their specific firing patterns even during

266 error trials where the rat turned left after presentation of the unrewarded cue, suggesting that these signals
267 did not represent action values. Additionally, NAc signals have been shown to be modulated by response
268 vigor, to detangle this from our results we included the trial length as a predictor in our GLMs, and found
269 cells with correlates independent of trial length.

270 **Cue modality:**

271 Our finding that ventral striatal units can discriminate between cues from different sensory modalities ex-
272 pands upon an extensive literature examining neural correlates of conditioned stimuli. Perhaps the most
273 comparable work in rodents comes from a study that found distinct coding for an odor when it predicted
274 separate but equally valued rewards (Cooch). The present work is complementary to this as it shows that
275 ventral striatal cells have representations of identifiable aspects of the cue itself, in addition to the reward
276 it predicts. Another study paired separate cues with appetitive or aversive outcomes, and found separate
277 populations of cells that encode each cue, with many switching selectivity after reversal of the associations
278 between the cues and outcomes, providing evidence that the NAc encodes the biological significance of
279 stimuli. Once again, our study was different as we recorded neural responses to distinct cues encoding the
280 same anticipated outcome, suggesting that even when the biological relevance of these stimuli is similar, the
281 NAc dissociates their representations at the level of the single-unit (Setlow). Another possibility is that these
282 modality specific cells were encoding the context, rule, or sequence within a session as some cells responded
283 similar for both rewarded and unrewarded cues within a block. This interpretation is in alignment with a
284 recent paper from the primate realm that recorded ventral striatal responses during the Wisconsin Card Sort-
285 ing Task (WCST), a common set-shifting task used in both the laboratory and clinic, and found cells that
286 preferred firing to stimuli when a certain rule, or rule category was currently active (Sleazeer). Indeed, an
287 encoding of the current strategy could be an explanation as to why differentially tiling of task structure was
288 observed across blocks in the current study. Further support for a modulation of NAc responses by strategy
289 comes from an fMRI study that examined BOLD levels during a set-shifting task (FitzGerald et al., 2014). In
290 this task, participants learned two sets of stimuli-reward contingencies, a visual set and auditory set. During

291 testing they were presented with both simultaneously, and the stimulus dimension that was relevant was peri-
292 odically shifted between the two. Here, they found that bilateral NAc activity reflected value representations
293 of whatever the currently relevant stimulus dimension was, and not the irrelevant stimulus. The current find-
294 ing of separate, but overlapping, populations of cells encoding cue modality and expected value, suggests
295 that the fMRI finding is generated by the combined activity of several different functional cell types.

296 A caveat of the current study is that rats were never presented with both sets of cues simultaneously, and
297 thus never had to switch strategies, although extrapolating the data from the primate study, suggests that the
298 activity of the cue modality cells would be modulated by relevance. Keeping along this theme, the current
299 data set is unable to identify precisely what the modality-sensitive neurons were encoding, that is were they
300 tracking representations of stimulus identity, a preferred context, or even a macroscale representation of
301 progress through the session. Furthermore, their relevance for ongoing behavior is also uncertain. NAc core
302 lesions have been shown to impair shifting between different behavioral strategies, and it is possible that
303 selectively silencing the cells that prefer responding for a given modality or rule would impair performance
304 when the animal is required to use that information, or artificial enhancement of those cells would cause
305 them to use the rule when it is the inappropriate strategy.

306 **Encoding of position:**

307 Our finding that cue-evoked activity was modulated by cue location sides with some of the literature (Lavoie,
308 1994; Tabuchi, 2000; Strait, 2016). An alternative explanation for a pure spatial representation, is that
309 these are task segmentation correlates, keeping track of where in the task the rat is. A previous non-human
310 primate paper has shown that when reward is contingent upon completion of a series of trials, separate
311 populations of NAc neurons signal the start of a schedule, subsequent trials in the schedule, and the first
312 trial in extended schedules (Shidara et al., 1998). This signalling of position within a sequence has been
313 observed in subsequent studies, and it is possible that our rats were keeping track of which specific arm
314 they were in as part of a sequence of arms, and not just strictly a spatial representation (Mulder, 2004 and

315 2005; Khamassi et al., 2008; Berke, 2009). Also, given that our task is pseudo-random, it is possible that the
316 rats learned which cue to anticipate, and the neural activity could reflect this. However, this is unlikely as
317 including a previous trial variable in the analysis did not explain a significant amount of firing rate variance
318 in response to the cue for the vast majority of cells..

319 **Mixed selectivity:**

320 Several other papers have reported unit profiles that integrate different task-related variables. These papers
321 report integrated coding between expected value and subsequent motor responses, expected value and iden-
322 tity of a reward, and a combination of spatial-, movement-, and reward-related features (Roesch, Lavoie,
323 Cooch). However, our study is the first to show mixed selectivity among identifying features of a cue and
324 expected outcome or behavior. The presence of mixed selectivity responses confers a larger number of input-
325 output relationships that are available to a given neuron. A possible functional consequence of this attribute
326 of NAc units, is the combination and transformation of various motivationally relevant features into a sig-
327 nal informing downstream decoders such as the ventral pallidum about appropriate behaviors in obtaining
328 motivationally relevant goals and biasing action selection towards these behaviors. Mixed selectivity in the
329 NAc could be a consequence of synaptic integration from a variety of anatomically distinct inputs, as seen
330 in experiments examining the convergence of various NAc afferents at the level of synaptic transmission and
331 stimulation-induced firing (Goto and Grace 2008). In one such experiment it was shown that NAc cells that
332 responded to stimulation of either the fornix, amygdala, or PFC, typically responded to stimulation from all
333 inputs (ODonnell and Grace, 1995). Furthermore, an interaction between these inputs was observed such
334 that PFC stimulation failed to elicit spiking in the NAc neurons unless they were in a depolarized UP-state, a
335 state induced by hippocampal stimulation and was dependent on an intact fornix. Hippocampal-induced sup-
336 pression of other inputs has also been observed for the BLA (Mulder et al., 1998). Recently, it has also been
337 shown that train stimulation of PFC afferents reduces hippocampal-evoked NAc responses, suggesting that
338 there is competition between various inputs (Calhoon and ODonnell, 2013). These studies suggest that the
339 integration of the variables we saw could be the result of this gating observed in behaviorally-independent

340 preparations. However, given that we did not systematically manipulate these various limbic and cortical
341 afferents, comments on the anatomical origins of the observed mixed selectivity responses are speculative at
342 this point.

343 Integrating cue identity and value, as seen in the present study, could be one neural instantiation of how
344 value is associated with the appropriate predictive stimuli (credit assignment), keeping in mind that value
345 encoding is distributed, redundantly in some aspects, across various structures (Hayden Nat Neuro opinion).
346 Indeed, lesions of the NAc impair the ability to learn changes in reward value or identity in an unblocking
347 experiment, as well as disrupting dopamine RPEs generated by modification of timing of reward (McDannald
348 2011, Takahashi 2016). Would be interesting to see if uncoupling the integrated coding of stimulus features
349 and predictive properties of a cue has an effect on the ability of a rat to use reward-predictive cues to pursue
350 the associated reward.

351 **Tiling of task structure:**

352 Additionally, we found that the population of recorded units had a relatively uniform distribution of firing
353 fields within our task space, similar to what has been reported previously (Shidara, 1998; Berke, 2009;
354 Lansink, 2012). Uniquely, we found that this representation differed according to whether the rat was cur-
355 rently engaged in the light or sound block, suggesting that this could be a possible neural correlate for
356 encoding the currently relevant strategy in the NAc. During progress through a predictable trial series, neu-
357 rons represented state value of cue (Shidara 1998). Single-unit responses allowed the monkey to know how
358 it was progressing throughout the task. Likewise, the tiling we saw could be a consequence of upstream cor-
359 tical or limbic inputs informing the striatum of the current task rules. Another possibility is that the NAc not
360 only pays attention to progress throughout a task within a trial, but also higher-order task information, like
361 blocks. Cue location was a behaviorally irrelevant variable in the current experiment, but it is possible that
362 if this tiling is dependent on hippocampal input, or related to a state value representation, that making cue
363 location a relevant variable by adding positional contingencies such as only alternating arms are rewarded

364 in one block, would result in a further separation of the mapping within a block between the rewarded and
365 unrewarded arms. Furthermore, dopamine levels in the NAc fluctuate through a trial, and it is possible that
366 the observed tiling could be a NAc-representation of state value related to this temporally evolving dopamine
367 signal. Future experiments should monitor this mapping of task structure during the application of dopamine
368 antagonists. Finally, the presence of functional correlates not evident when looking at single-unit responses
369 time-locked to salient task events emphasizes the need to employ ensemble level analyses across all aspects
370 of a task.

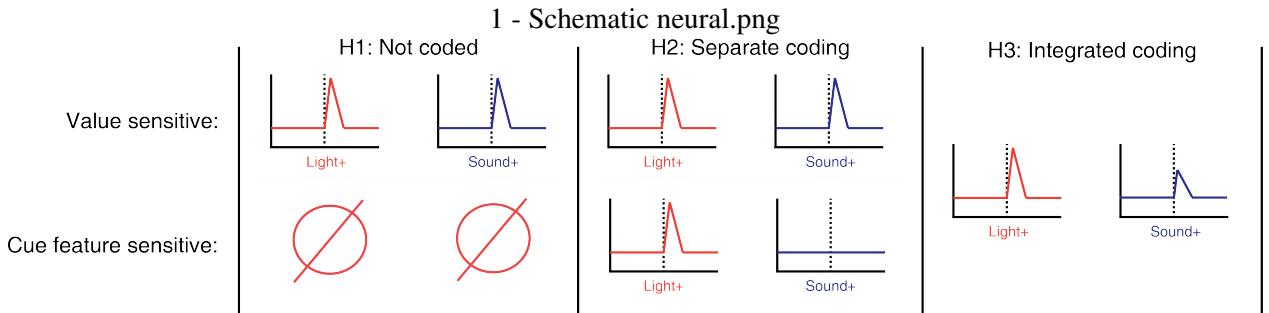


Figure 1: Schematic of potential coding strategies employed by single units in the NAc. **H1 (left panel):** Coding of identifying cue features ~~is could occur either be~~ absent in the NAc. ~~Top row: Unit A responds to a motivationally relevant variable, such as expected value, similarly across other cue features, such as stimulus modality or physical location. Hypothetical plot is firing rate across time. Light+ (red left) signifies a rewarded light cue, Sound+ (blue) a rewarded sound cue. Dashed line indicates onset of cue. Bottom row: No units within the NAc discriminate their firing across various conditions of these cue features.~~ **H2 (middle panel):** Coding of cue features occurs ~~occur~~ independently of encoding of motivationally relevant variables like expected value or subsequent vigor. ~~Top row: Same as H1. Bottom row: Unit B discriminates firing across conditions of a cue feature, depicted here as firing to Light+ but not Sound+. H3 (right panel middle): Coding of cue features is ~~and/or be~~ integrated with coding of other motivationally relevant variables (right). Hypothetical example demonstrating a unit that responds to reward-predictive cues, but firing rate is also modulated by which rewarded-predictive cue is presented, firing more for Light+ than Sound+.~~

2 - Schematic task.png

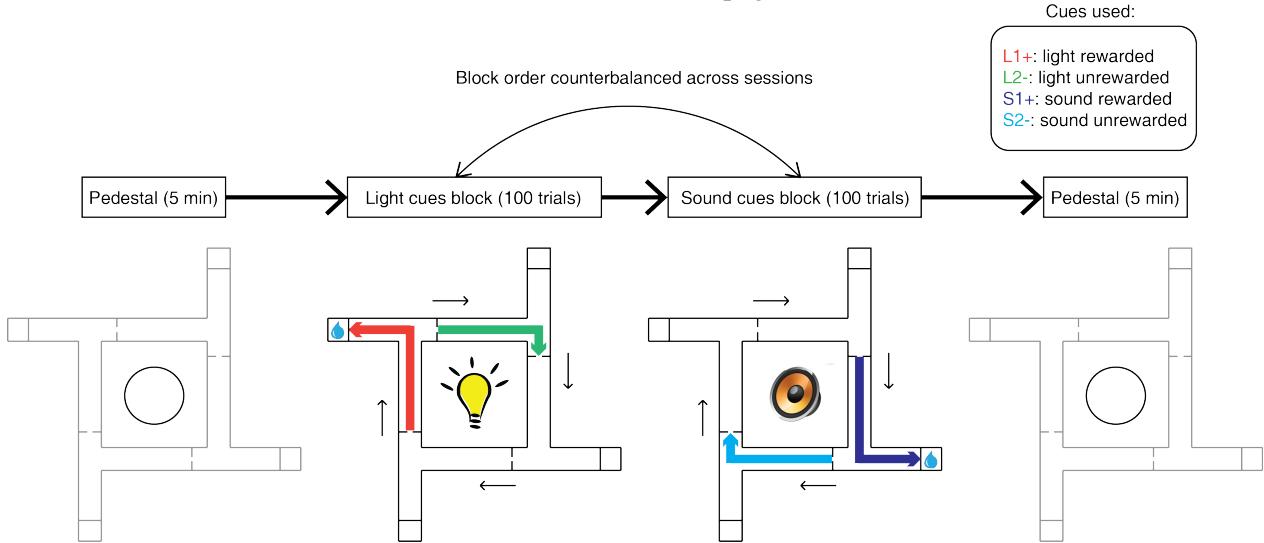


Figure 2: Schematic of behavioral task and procedure used in current study. **Left:** A session was started with a 5 minute recording period on a pedestal ~~placed~~ placed in the center of the apparatus (~~left~~). **Middle:** Rats then underwent two blocks of a cue discrimination task on a square track where they had to dissociate between a rewarded and unrewarded cue for a set of light cues and a set of sound cues for a target of 100 trials each (middle). Rats ran clockwise on the square track, and upon presentation of the cue had to turn left at the choice point to receive reward for a rewarded cue (approach trial; red and navy blue in figure), or turn right at the choice point to initiate the following trial for an unrewarded cue (skip trial; green and ~~light blue~~-cyan in figure). **Middle-left in figure depicts a light block, showing an example trajectory for a rewarded (red) and unrewarded (green) trial.** **Middle-right depicts a sound block, with a rewarded (navy blue) and unrewarded (light blue) trial.** Ordering of the light and sound blocks was counterbalanced across sessions. Dashed line within track indicates photobeam that triggered cue onset. Solid line within track indicates photobeam that registered a nosepoke. Receptacles containing sucrose solution are present within the squares at the end of each arm. Water droplet signifies reward receipt on correct trial. Arrows outside of track indicate correct running direction. **Right:** A session ended with another 5 minute recording period on the pedestal (right).

3 - Histology.png

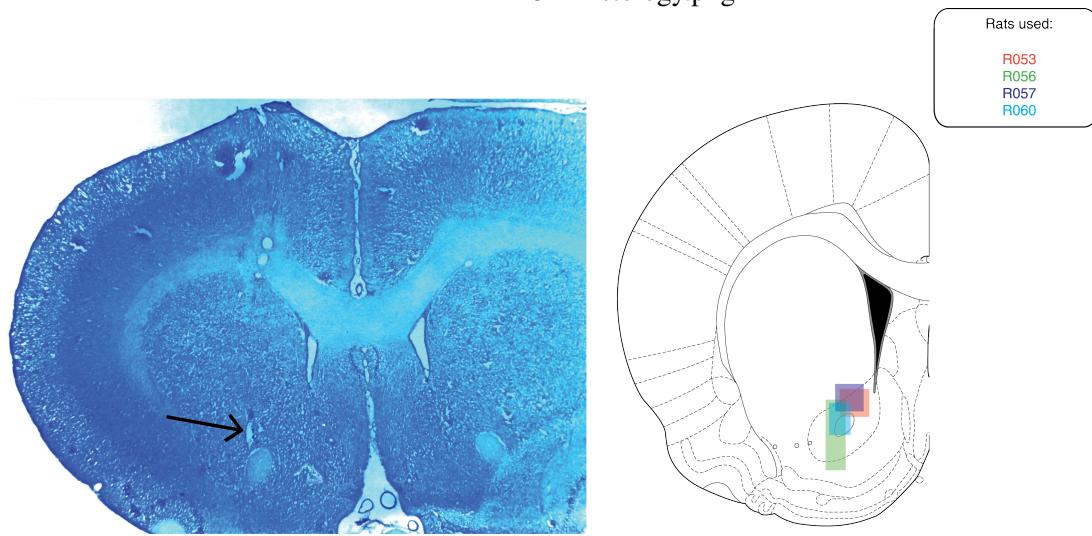


Figure 3: Histological verification of recording sites. Upon completion of experiments, brains were sectioned-sliced and tetrode placement was confirmed. A. Example section from R060 showing a recording site in the NAc core ~~just dorsal to the anterior commissure (white circle)~~. B. Schematic showing recording areas for the four rats used in the present study.

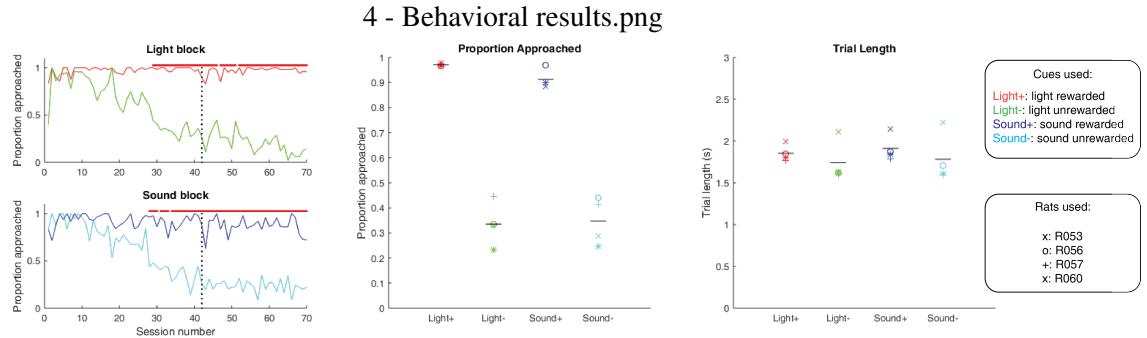


Figure 4: Performance on the behavioral task. A-B. Example learning curves ~~across sessions~~ from a single subject (R060) showing ~~acquisition and maintenance of performance in the proportion approached for rewarded (red line for light block, navy blue line for sound block (A) and unrewarded trials (green line for light block, light blue line for sound block (B) blocks. Fully correct performance corresponds to approach~~ Dependent measure is proportion of trials approached within a session for a given cue, with a value of 1 being perfect performance for rewarded trials/cues, and 0 being perfect performance for unrewarded trials/cues. Note that this subject starts out approaching on both rewarded and unrewarded trials, and learns with experience to skip non-rewarded trials. Red bars indicate days in which a rat statistically discriminated between rewarded and unrewarded cues, determined by a chi square test. Dashed line indicates time of ~~electrode-implant surgery~~ implantation. C-D. Summary of performance during recording sessions for each rat. C. Proportion approached for all rats, averaged across all recording sessions. Different columns indicate the different cues (rewarded (+) and unrewarded (-) light cues, rewarded and unrewarded sound cues). Different symbols correspond to individual subjects; black line shows the mean. All rats learned to discriminate between the rewarded and unrewarded cues, within a block as indicated measured by the clear difference a comparison of proportion approached between rewarded (0.9x approached) and unrewarded cues (0.3x approached) linear mixed effects models, interpreted as a decreased likelihood to make an approach for both blocks (see Results for statistics) an unrewarded cue. D. Average trial length of time to complete a trial for each cue. Note that the Rats on average showed a comparable length of time to complete a trial was comparable for the different cues regardless of cue.

5 - Neural examples.png

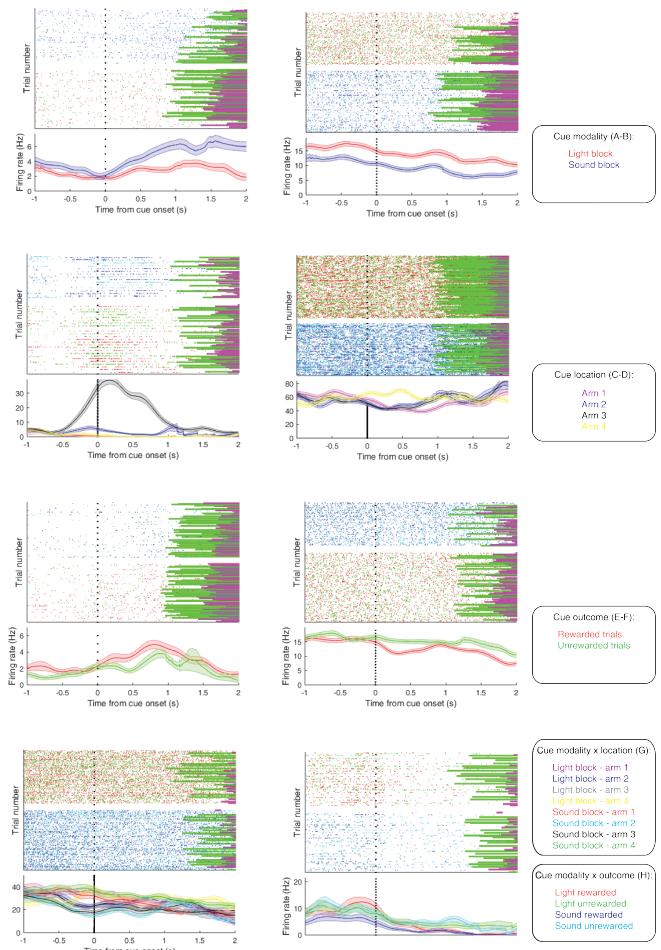


Figure 5: Individual examples of NAc units influenced by various task parameters. A. Example of a NAc unit that showed an increase in firing in response to the cue, and whose activity was influenced by which block the rat was in. Top: rasterplot showing the spiking activity across all trials aligned to cue-onset. Spikes across trials are color coded according to cue type (red: rewarded light; green: unrewarded light; navy blue: rewarded sound; light blue: unrewarded sound), using the same scheme as in previous figures. Green and magenta bars indicate trial termination when a rat initiated the next trial or made a nosepoke, respectively. White space halfway up the rasterplot indicates switching from one block to the next. Dashed line indicates cue-onset. Bottom: PETHs showing the average smoothed firing rate for the unit for trials during light (red) and sound (blue) blocks, aligned to cue-onset. Lightly shaded area indicates standard error of the mean. Note this unit showed a larger increase in firing to sound cues. B. Same as A, but for a unit that showed a decrease in firing to the cue. Note the sustained higher firing rate during the light block. C-D. Same as A-B for cue location, each color in the PETHs represents average firing response for a different cue location. C. This firing rate of this unit only changed on arm 3 of the task. D. Firing decreased for this unit on all arms but arm 4. E-F. Same as A-B for cue outcome, with the PETHs comparing rewarded (red) and unrewarded (green) trials. E. This unit showed a slightly higher

6 - GLM.png

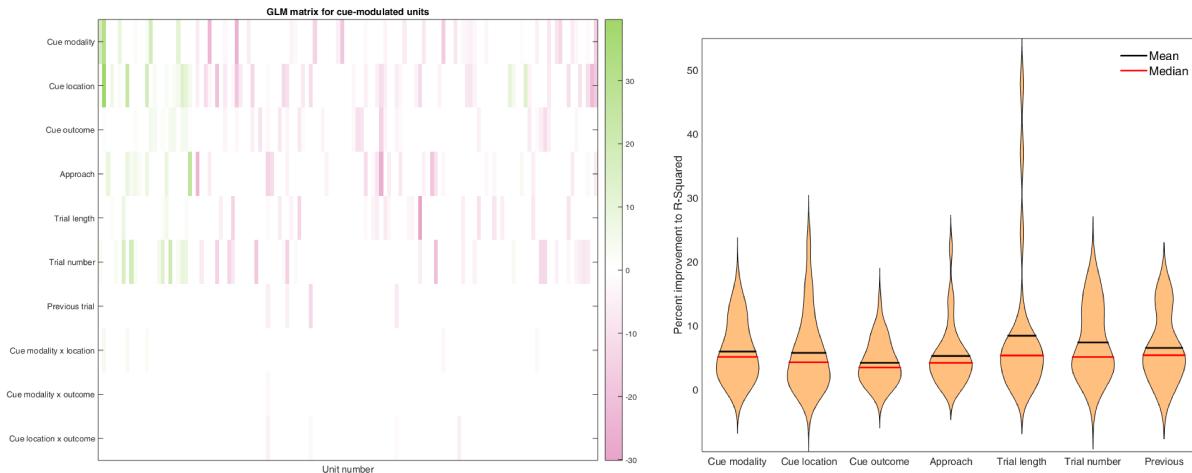


Figure 6: Summary of influence of various task parameters ~~on-of~~ cue-modulated NAc units after cue-onset. A. GLM matrix demonstrating impact of various task parameters on NAc firing rates. A ~~stepwise~~ GLM was fit to each unit that showed evidence of cue modulation by a Wilcoxon signed-rank test. Each row represents a given task parameter, and ~~each column is the x axis shows the influence of that the task parameter on a given parameters for each unit. Response variable is how much of the firing rate variance an individual predictor contributed to the model, as measured by differences in R-squared between the final model and the model minus the predictor of interest.~~ Ordering organized from left to right ~~for~~ MSNs that increased firing in response to the cue (green, left), MSNs with a decreasing response (red, left), FSIs with an increasing response (green, right), FSIs with a decreasing response (red, right). Darker shades indicate more ~~Response variable is how much of the firing rate variance explained an individual predictor contributed to the model, as measured by a given differences in R-squared between the final model and the model minus the predictor of interest.~~ B. ~~Violin plots Bar graph~~ demonstrating ~~changes average change~~ in R-squared ~~values value~~ with the addition of each of the individual predictors. ~~The mean, median, and distribution of changes in R-squared values is plotted for each of the seven task parameters used in the GLM.~~

7 - Population averages.png

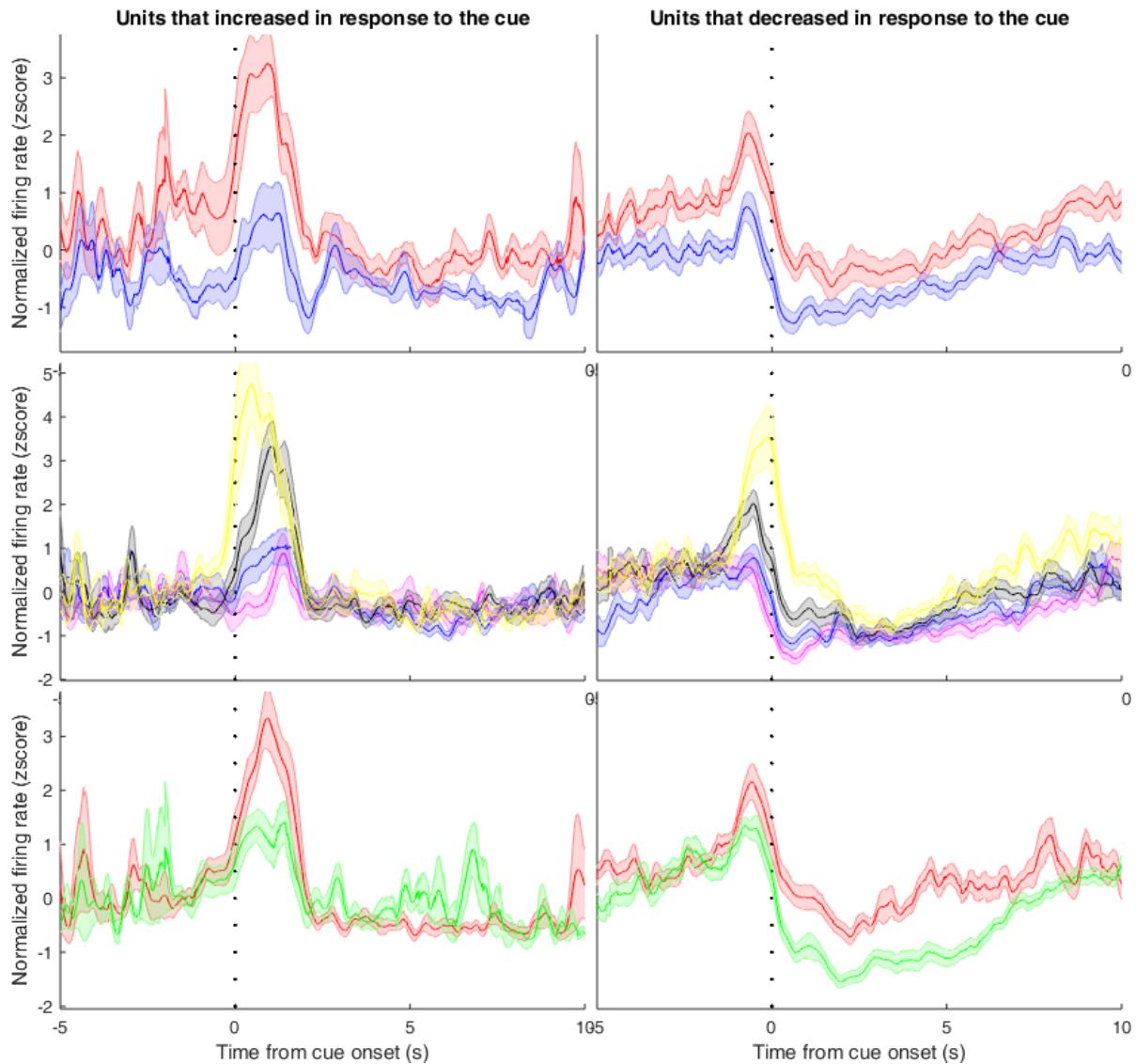


Figure 7: Population-level averages of cue feature sensitive NAc units. A. Average smoothed normalized activity for cue-modulated units where cue modality was a significant predictor in the GLM, aligned to cue-onset. Activity is plotted for preferred stimulus block (red) and nonpreferred stimulus block (blue). Dashed line indicates onset of cue. Lightly shaded area indicates standard error of the mean. Note larger increase to preferred stimulus block to nonpreferred stimulus block. B. Same as A-but for units that decreased in firing. Note population level activity reveals cells classified as decreasing in response to cue show a biphasic response at the population level, with a transient increase around the time the rat starts on the arm, followed by a minimum after cue onset. Also, note the sustained difference in firing between the two blocks. C-D. Same as A-B for cue location. Activity is plotted from most preferred arm (yellow), in decreasing order to least preferred arm (black, navy blue, magenta, respectively). Note the graded response to arms of decreasing preference. E-F. Same as A-B for cue outcome. Activity is plotted for preferred expected outcome (red), and nonpreferred outcome (green). Note the larger increase to the cue representing the cells preferred outcome (E), and the sustained decrease to the nonpreferred outcome (F).

8 - Task tiling.png

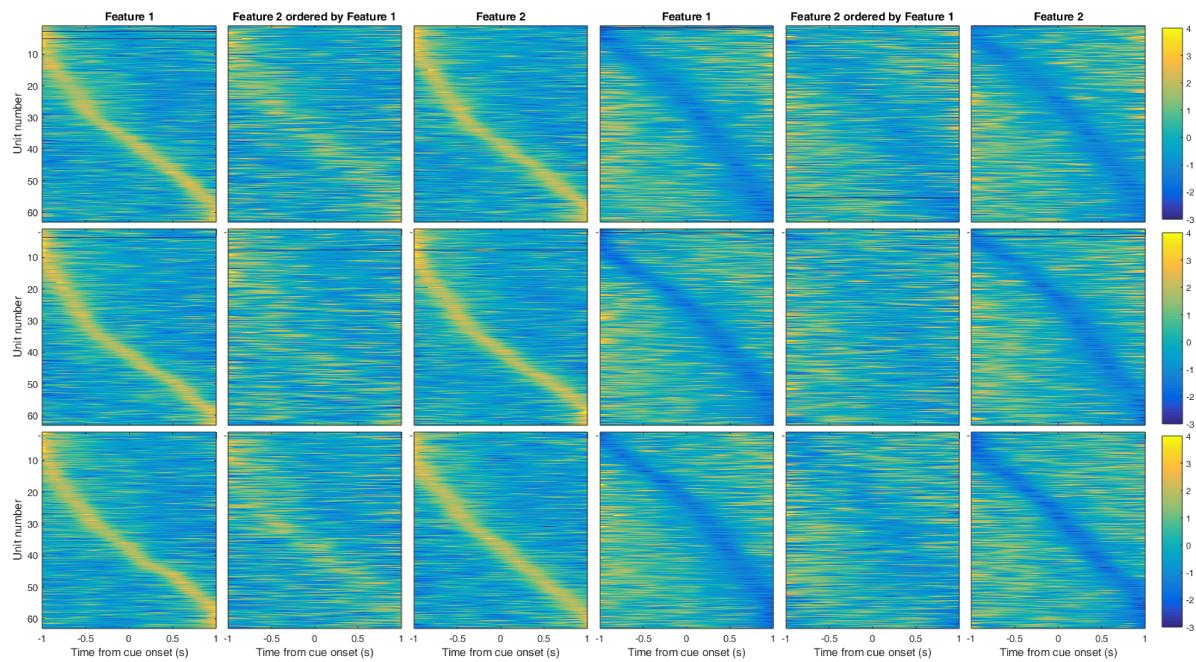


Figure 8: Distribution of NAc firing rates across time task space surrounding cue-onset. Each panel shows normalized firing A. Firing rates for all recorded NAc units (each row corresponds to one unit) as a function of time (time 0 indicates cue-onset), averaged across all trials for a specific cue type, indicated by text labels. A, left: Heat plot showing smoothed were normalized firing activity of all recorded NAc units and ordered according according to the time of their peak firing rate during the light block. Each row is a units average activity across time to the light (left) and sound block. Dashed line indicates cue onset. Notice the yellow band across time(right), indicating all aspects of visualized task space were captured by the peak firing rates of various units aligned to cue-onset. A, middle: Same units ordered according to the time Distribution of the peak firing rate during the sound block. Note that for both blocks, units tile time approximately uniformly with a clear diagonal of elevated firing rates. A, right: Unit firing rates taken from the sound block, ordered according to peak firing rate taken from the using ordering for light block. Note that a weaker but still discernible diagonal persists, indicating partial similarity between firing rates in the two blocks. B: Same layout as in A, but with units ordered according to the time of their minimum firing rate. C: C-D: Same layout as in A, except that the panels now compare two different locations on the track instead A-B for a comparison of two cue modalities. As for the different cue modalities, NAc units clearly discriminate between locations, but also maintain some similarity across locations, as evident from the visible diagonal in the right panel. Two example locations were used for display purposes; other location pairs showed a similar pattern. D: As for C, but with units ordered by their minimum firing rate. E: E-F: Same layout as in A, except that panels now compare A-B for comparison of rewarded and unrewarded trials cues. F: As for A, but with units ordered by their minimum firing rate. Overall, NAc units "tiled" experience on the task, as opposed to being confined to specific task events only. Units from all sessions and animals were pooled for this analysis.

9 - NP GLM.png

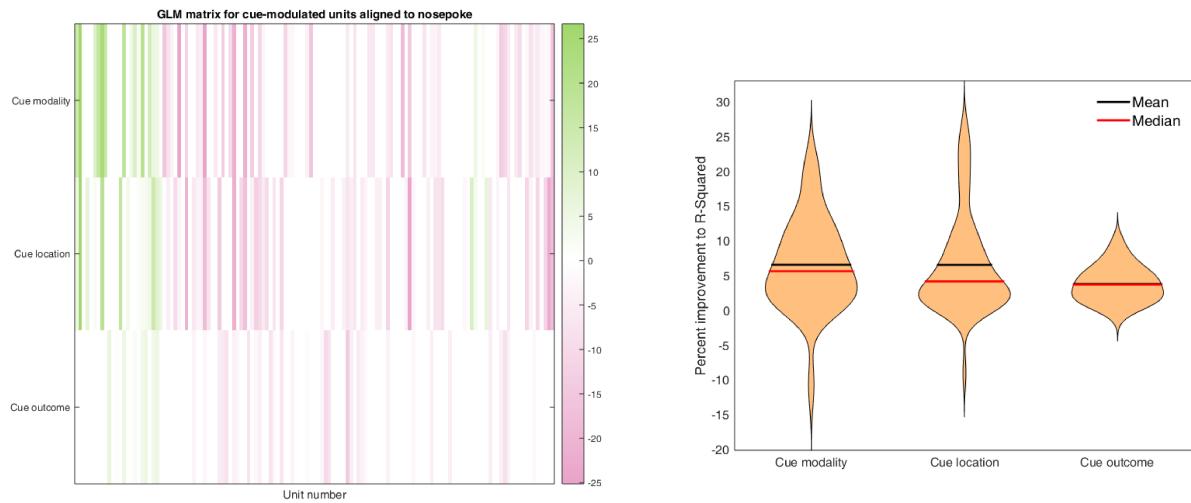


Figure 9: Summary of influence of various task parameters of cue-modulated NAc units during nosepoke. A. GLM matrix demonstrating impact of various task parameters on NAc firing rates. A stepwise GLM was fit to each unit that showed evidence of cue modulation by a Wilcoxon signed-rank test. Each row represents a given task parameter, and each column is the x axis shows the influence of that the task parameter on a given parameters for each unit. Response variable is how much of the firing rate variance an individual predictor contributed to the model, as measured by differences in R-squared between the final model and the model minus the predictor of interest. Ordering organized from left to right :for MSNs that increased firing in response to the cue (green , left), MSNs with a decreasing response (red , left), FSIs with an increasing response (green , right), FSIs with a decreasing response (red , right). Darker shades indicate more Response variable is how much of the firing rate variance explained an individual predictor contributed to the model, as measured by a given differences in R-squared between the final model and the model minus the predictor of interest. B. Violin plots Bar graph demonstrating changes average change in R-squared values value with the addition of each of the individual predictors. The mean, median, and distribution of changes in R-squared values is plotted for each of the three task parameters that were significant predictors in the GLM.

10 - NP population averages.png

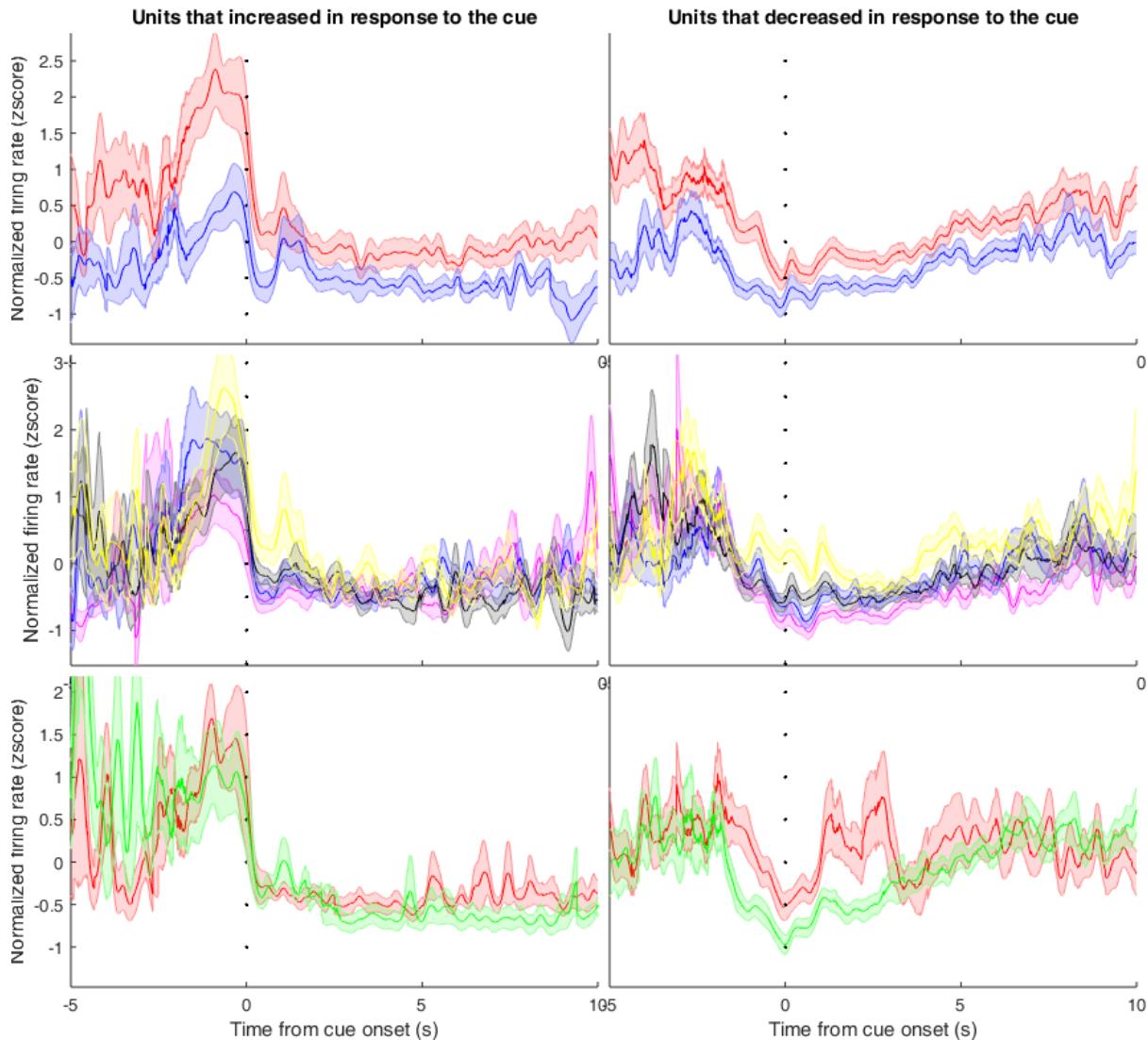


Figure 10: Population-level averages of cue feature sensitive NAc units during a nosepoke. A. Average ~~smoothed~~ normalized activity for cue-modulated units where cue modality was a significant predictor in the GLM, aligned to nosepoke with reward delivery occurring 1 s after nosepoke. Activity is plotted for preferred ~~stimulus~~ block (red) and nonpreferred ~~stimulus~~ block (blue). ~~Dashed line indicates nosepoke. Lightly shaded area indicates standard error of the mean. Note larger increase leading up to nosepoke to preferred stimulus block to nonpreferred stimulus block.~~ B. Same as A but for units that decreased in firing. ~~Note the sustained difference in firing between the two blocks.~~ C-D. Same as A-B for cue location. Activity is plotted from most preferred arm (~~yellowcyan~~), in decreasing order to least preferred arm (~~black, navy blue, magenta~~green, ~~red~~, respectively). ~~Note the subtle but graded response to arms of decreasing preference.~~ E-F. Same as A-B for cue outcome. Activity is plotted for preferred expected outcome (red), and nonpreferred outcome (green). ~~Note the peak after reward receipt for preferred outcome in decreasing cells (F).~~

11 - NP task tiling.png

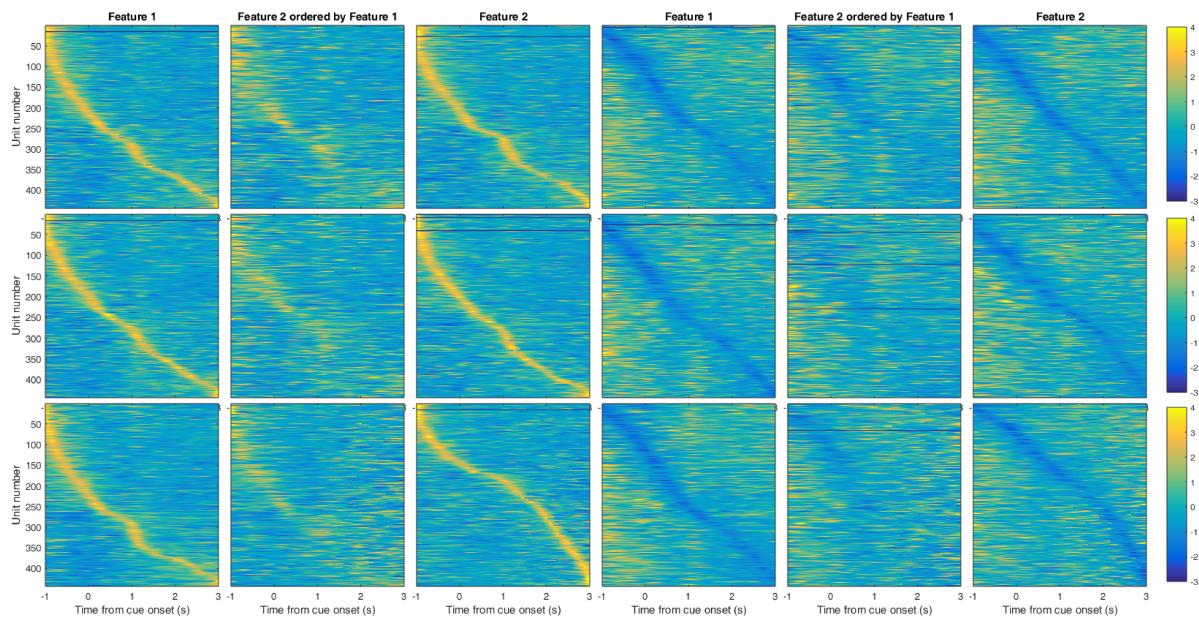


Figure 11: Distribution of NAc firing rates across task space during approach trials. **A.** Firing rates for all recorded NAc units (each row corresponds to one unit) as a function of time (time 0 indicates nosepoke), averaged across all approach trials for a specific cue type, indicated by text labels. **A, left:** Heat plot showing smoothed normalized firing activity of all recorded NAc units and ordered according to the time of their peak firing rate during the light block. **A, middle:** Same units ordered according to the time of their peak firing rate during the sound block. **A, right:** Unit firing rates taken from the sound block, ordered according to peak firing rate taken from the light block. Note that a weaker but still discernible diagonal persists, indicating partial similarity between firing rates in the two blocks. **B:** Same layout as in A, but with units ordered according to the time of their minimum firing rate. **C-D:** Same layout as in A, except that the panels now compare two different locations on the track instead of A-B for a comparison of two cue modalities. As for the different cue modalities, NAc units clearly discriminate between locations, but also maintain some similarity across locations, as evident from the visible diagonal in the right panel. Two example locations were used for display purposes; other location pairs showed a similar pattern. **D:** As for C, but with units ordered by their minimum firing rate. **E-F:** Same layout as in A, except that panels now compare A-B for comparison of rewarded and unrewarded trials. Notice the absence of a cluster of tiling around reward receipt for unrewarded trials. **F:** As for A, but with units ordered by their minimum firing rate. Overall, NAc units "tiled" experience on the task, as opposed to being confined to specific task events only. Units from all sessions and animals were pooled for this analysis.

Rat	Total	MSN (increasing)	MSN (decreasing)	FSI (increasing)	FSI (decreasing)
R053	145	51	79	4	11
R056	70	12	13	17	28
R057	136	55	75	3	3
R060	92	37	49	3	3

Table 1: Cells from each rat

Task parameter	Total	MSN (increasing)	MSN (decreasing)	FSI (increasing)	FSI (decreasing)
All cells	443	155	216	27	45
Cue modulated	133	24	85	6	18
Cue modality	37	7	21	1	8
Cue location	50	13	27	3	7
Cue outcome	34	10	18	0	6
Approach behavior	31	8	18	1	4
Trial length	25	5	18	0	2
Trial number	32	11	12	1	8
Recent trial history	5	0	5	0	0
Cue x cue interactions	11	3	7	0	1
Cue x behavior interactions	?	?	?	?	?

Table 2: Cells from GLM

Task parameter	Total	MSN (increasing)	MSN (decreasing)	FSI (increasing)	FSI (decreasing)
Cue modality	66	14	36	2	14
Cue location	66	14	40	3	9
Cue outcome	42	8	29	0	5
Trial length	0	0	0	0	0
Trial number	0	0	0	0	0
Recent trial history	0	0	0	0	0
Cue x cue interactions	0	0	0	0	0
Cue x behavior interactions	0	0	0	0	0

Table 3: Cells from nosepoke GLM