1   **Goal congruency dominates reward value in accounting for behavioral and neural**

2   **correlates of value-based decision-making**

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Abstract

When choosing between options, whether menu items or career paths, we can evaluate

how rewarding each one will be, or how congruent each option is with our current choice goal

(e.g., identifying the best or worst ones). Past decision-making research has interpreted findings

through the former lens, but in these experiments the most rewarding option has always been

most congruent with one's goal (choosing the best option). It is therefore unclear to what extent

expected reward versus goal congruency can account for a wide body of behavioral and

neuroscientific research on choice value. Here, we deconfound these two variables. Contrary to

prevailing accounts, we find that goal congruency dominates choice behavior and neural activity.

We separately identify dissociable signals of expected reward. Our findings disentangle the roles

of rewards and goals in how people consider their options, and call for a reinterpretation of

previous research on value-based choice.

Keywords: reward, decision making, emotion, functional MRI

28   **Significance Statement**

Whether it is between restaurants or career paths, to make adaptive decisions we must evaluate

our options and identify those that are most conducive to our current goal. Dysfunctional

decision-making can therefore result from aberrant reward processing (e.g., impulse disorders) or

from aberrant goal processing (e.g., OCD, ADHD). By focusing only on how people choose their

favorite option in a choice set (when rewards and goals are aligned), past research has been

unable to distinguish the degree to which behavior and neural activity are determined by reward

versus goal processing. We disentangle these processes and show that behavior and fMRI

activity are differentially influenced by the promised rewards versus the degree to which those

rewards align with one's current goal.

Reward is central to the behavior of humans and animals alike (1). We repeat rewarded

behavior and strive to select actions that maximize reward (2, 3). Reward therefore serves dual

roles, as an end unto itself and as a guide towards a particular goal. The significance of this

distinction has often been overlooked.

A large body of research has examined how we evaluate the expected reward for an

object or a course of action, including how this reward value is shaped by feature integration (4,

5), context (6, 7), and motivational state (8). When people choose between multiple options with

varying reward, these studies have demonstrated consistent behavioral and neural correlates of

those reward values. People make faster decisions when the option they choose is much more

rewarding than the option(s) they forego (9, 10), and when the overall (i.e., average) reward

associated with those choice options is high (11-14). These behavioral findings are paralleled by

increasing relative and overall value related activity in regions of a well-characterized value

network (15, 16), that includes regions of ventral striatum and ventromedial prefrontal cortex.

These behavioral and neural indices have thus been taken to reflect variability in how *rewarding*

one's options are (3). However, this interpretation has overlooked a confound common to almost

all such experiments, between an option's promise of reward and its ability to support a person's

task goal.

Specifically, participants in studies of value-based decision-making are typically

instructed to choose the item they prefer most (9, 17-21) or dislike least (21-25). As a result, the

option that is most rewarding or least aversive also represents the option that is most congruent

with the participant's choice goal, and the remaining options represent those that are least

congruent with their goal. The relative *reward value* of a choice set (how much more rewarding

the chosen option is than the alternatives) is therefore identical to the relative *goal value* of those

options (how much more goal-congruent the chosen option is than the alternatives). Similarly,

having more rewarding options in one's choice set (higher overall reward value) also necessarily

means having more options that are congruent with one's goal (higher overall goal value). It is

thus unclear whether behavioral findings previously attributed to reward value, such as the

choice speeding effect of having options that are overall more rewarding, can instead be

accounted for by the *goal congruency* of these reward values. It is similarly unknown to what

extent putative neural correlates of relative and overall reward value instead reflect relative and

overall goal congruency. The present studies aim to address these open questions.

There are several a priori reasons to favor a strictly reward-based interpretation of past

research on choice value. First, research on Pavlovian conditioning shows that animals have an

innate drive to approach rewards and avoid punishments, such that they are faster to approach a

reward than to avoid it (26, 27). Second in an operant task (e.g., lever-pressing), the vigor (and

therefore speed) with which an animal responds scales with the average rate of reward in their

environment (28-30). Third, there is ample evidence that a network of brain regions – the *value*

network – processes both anticipated and experienced reward (15, 31). For instance, it has been

shown that activity in this network correlates with the reward value assigned to individual items

(32-35) and sets of items (10, 16, 21, 36), and that reward-related activity associated with those

items can be used to predict one's choice between multiple such options (31, 37, 38). As a result,

value network activity is generally assumed to encode a form of *reward value*, one that is

multifaceted (varying, e.g., with one's motivational state (8) and context (7, 39-41)) but which

should remain *directionally consistent* across choice goals. That is, an item with high reward

value should be encoded similarly by this network whether an individual is asked to indicate that

they like that item or that they do not dislike that item (42, 43). However, previous research on

choice-related behavior and neural activity have failed to dissociate reward-centered accounts of

these findings from accounts that are centered on the congruency of those rewards with one's

current goal.

Research on recognition memory provides an excellent example of how goal congruency

confounds can engender misinterpretations of a set of findings. Studies of frequency

discrimination required participants to indicate which of a pair of items had been presented more

frequently during an earlier study phase (44, 45). Researchers found that participants responded

faster when both options were high frequency items rather than when both were low frequency

items, suggesting that response speed was correlated with the overall frequency of the choice set.

However, this effect reversed when participants were instructed to choose the *less* frequent item

rather than the more frequent one; when given these instructions, participants were instead faster

to respond when the overall frequency was low. Thus, by manipulating choice instructions, this

study dissociated item frequency from goal congruency, and showed that behavior was driven by

the latter rather than the former.

Here, we employ a similar approach, asking participants to either choose the best (most

rewarding) or the worst (least rewarding) item in a value-based choice set, to test whether task

goals modulate the relationship reward value has with behavior and brain activity. Across four

studies, we find that a participant's decision speed is not accounted for by the reward value of

their options but rather the goal congruency of those options, including how congruent those

options are *overall* (a feature of the choice set that is irrelevant to goal achievement). We

demonstrate that these findings can be accounted for by modifying an established computational

model of choice dynamics to account for an individual's current choice goal. We further find that

the brain's value network tracks the relative value of a chosen item only in terms of its goal

congruency, while at the same time tracking both the overall goal congruency *and* overall reward

associated with the choice set. Within striatum, correlates of (overall) reward value and (relative)

goal congruency were spatially dissociated. Our findings suggest that, in common choice

paradigms, reward's relationship to behavior and neural activity is less direct than previously

thought, calling for a reinterpretation of previous findings in research on value-based decision-

making.

115   **Results**

116   Participants in two studies (Ns = 30) made a series of hypothetical choices between sets

117   of four consumer goods (Fig. 1A). Choice sets were tailored to each participant, based on how

118   rewarding they had rated each item individually earlier in the session, and these sets varied in

119   their overall reward value (how rewarding the items are on average) and in the similarity of

120   option values to one another (Fig. S4). Participants in Study 2 performed this choice task while

121   undergoing fMRI.

122   **The influence of overall value on choice speed is determined by goal congruency**

rather than reward value. Previous studies have consistently shown that participants are faster

to select their most preferred item out of a set as the overall reward value of their options

increases (11-14). These findings can be explained by accounts that posit a direct relationship

between potential rewards and choice speed (e.g., Pavlovian accounts) or they could be

explained by the fact that reward value was always congruent with one's task goal in these

studies (to choose the most rewarding item). To test these competing accounts, we explicitly

varied this goal while participants made choices between sets of options. For half of the trials, we

instructed participants to choose their *most* preferred from each choice set (i.e., the item with the

highest reward value; *Choose Best* task). In this task (the standard one used in research on value-

based choice), there is a positive relationship between reward value and *goal congruency* (which

we will also refer to as *goal value*) – the more rewarding an item, the more congruent it is with

the task goal. For the other half of trials, we reversed this instruction, asking participants to

instead choose their *least* preferred from a given set (i.e., the item with the lowest reward value;

Choose Worst task). Unlike the Choose Best task, for the Choose Worst task there was a *negative*

relationship between reward value and goal congruency (i.e., the least rewarding items were now

the most congruent with the task goal).

If choices are sped up by the overall reward value of a choice set (as inferred by previous

research), this should be true irrespective of whether participants were performing the Choose

Best or Choose Worst task (Fig. 1B, top). If these previous findings instead reflect variability in

choice speed with goal congruency, then RTs should decrease with overall reward value in the

Choose Best condition but *increase* with overall reward value in the Choose Worst condition

(Fig. 1B, bottom).

Figure 1. Overall value effects on RT are driven by goal congruency rather than
reward value. A) After evaluating each item in isolation (**left**), participants saw sets of
four options and (in separate blocks) were instructed to choose either the best or the worst
option (**right**). The same example is shown for both blocks but each choice sets was only
viewed once in a session. **B) Top:** A reward-based account predicts that RTs should
decrease with overall value of the set, irrespective of the choice goal. **Bottom:** A goal
congruency account predicts that RTs should decrease with overall value in Choose-Best
blocks but instead increase with overall value in Choose-Worst Blocks. **C)** Both Study 1
(behavioral) and Study 2 (fMRI) find the task-specific RT reversal predicted by a goal
congruency account (see also Study S1, Results 1). Shaded error bars show 95% confidence
intervals. **D)** Our empirical findings were captured by an LCA model that took goal values
(rather than reward values) as inputs.

158
Across both studies, we found strong evidence for a goal congruency account of choice

behavior. As an initial test, we ran linear mixed effects models separately on Choose-Best and

Choose-Worst trials. Consistent with previous studies (which instructed participants to choose

the best item in a set; 11, 12-14), we found that RTs decreased with overall reward value for our

Choose-Best task (Study 1: $b = -0.33$, $p < .001$, Study 2: $b = -0.39$, $p < .001$). However, for the

Choose-Worst task, we found that this relationship flipped – RTs instead *increased* rather than

decreased with overall value (Study 1: $b = 0.42$, $p < .001$, Study 2: $b = 0.38$, $p < .001$). Across all

trials, RTs therefore demonstrated an interaction between choice goal and overall reward value,

as predicted by a goal congruency account (Fig. 1C; Table 1, top; Study 1: $b = -0.75$, $t = -12.66$,

$p < .001$, Study 2: $b = -0.76$, $t = -14.39$, $p < .001$). Once accounting for this interaction, we did

not find residual main effects of overall value across the two tasks (Study 1: $t = 1.61$, $p = .108$;

Study 2: $t = -0.24$, $p = .809$), nor did we find a main effect of choice goal (Study 1: $t = -1.94$, p

$= .062$; Study 2: $t = -0.01$, $p = .736$). In other words, RTs varied with the overall reward value

inversely and *symmetrically* between the two tasks, but choices were not overall faster for one

task or another.

174   This pattern of findings is qualitatively consistent with the prediction that overall reward

value influences choice RT as a function of its goal congruency rather than its association with

potential rewards. In order to test this interpretation directly, we constructed an additional overall

value variable that encoded the overall goal congruency of one's options (overall goal value).

Overall goal value increases with overall reward value in the Choose-Best condition and

decreases with overall reward value in the Choose-Worst condition. We compared separate linear

mixed effects models predicting choice RT by overall reward value alone (Model 1), overall

reward value in interaction with task condition (Model 2), or overall goal value alone (Model 3).

We found that Models 2 and 3 substantially outperformed Model 1 (Table 2). Accordingly, when

including overall goal value and overall reward value in the same regression, we find a

significant effect only for overall goal value (Study 1: $b = -0.37$, $p < .001$; Study 2: $b = -0.38$, p

$< .001$) and not for overall reward value (Study 1: $b = 0.05$, $p = .108$; Study 2: $b = -0.01$, p

$= .809$; Table 1, bottom).

187   This effect of overall value on choice RT is independent of the effect of value *difference*,

188   i.e., the absolute difference between the goal value (highest item value for Choose-Best and

189   lowest item value for Choose-Worst) and the remaining items. As in previous studies, we find

190   that easier choices (those with higher value differences) are faster (Table 1), an effect that we

191   control for in all of our analyses.

192   Across both studies, analyzing choices as a function of value difference, overall value,

task and their interactions, we find that value difference also predicts the *consistency* (cf.

accuracy) of choices – as value difference increased, participants were more likely to choose the

item that achieved their current choice goal (the highest-value item in Choose-Best, the lowest-

value item in Choose-Worst; Study 1: $b = 3.76$, $z = 13.12$, $p < .001$, Study 2: $b = 3.65$, $z = 13.40$,

$p < .001$). The influence of value difference on choice accuracy did not differ between the two

choice goals (Study 1: $b = 0.58$, $z = 1.01$, $p = .311$, Study 2: $b = 0.53$, $z = 0.98$, $p = .326$). Unlike

value difference, the overall value of a set did not significantly predict choice consistency for

either task (Study 1: b $= .33$, $z = 1.08$, $p = .280$, Study 2: b $= 0.49$, $z = 1.70$, $p = .090$).

Table 1. *Comparison of overall reward vs overall goal value effects on log RT*

	Study 1					Study 2				
Predictors	*Estimates*	*CI*	*t*	*df*	*p*	*Estimates*	*CI*	*t*	*df*	*p*
OV_{reward} *by Choice Goal*										
(Intercept)	1.42	1.36 – 1.49	41.93	31.00	**<0.001**	1.52	1.44 – 1.59	38.93	31.00	**<0.001**
Value Difference	-0.49	-0.55 – -0.43	-16.48	3430.00	**<0.001**	-0.43	-0.52 – -0.34	-9.18	31.00	**<0.001**
Overall Value	0.05	-0.01 – 0.11	1.61	3448.00	0.108	-0.01	-0.06 – 0.05	-0.24	4222.00	0.809
Best - Worst Condition	-0.07	-0.14 – 0.00	-1.90	31.00	0.066	-0.01	-0.09 – 0.06	-0.33	31.00	0.740
Overall Value: Best - Worst	-0.75	-0.86 – -0.63	-12.66	3296.00	**<0.001**	-0.76	-0.86 – -0.65	-14.39	3914.00	**<0.001**
$OV_{goal} + OV_{reward}$										
(Intercept)	1.42	1.36 – 1.49	41.97	31.00	**<0.001**	1.52	1.44 – 1.59	38.93	31.00	**<0.001**
Value Difference	-0.49	-0.55 – -0.43	-16.48	3430.00	**<0.001**	-0.43	-0.52 – -0.34	-9.18	31.00	**<0.001**
Overall Goal Value	-0.37	-0.43 – -0.32	-12.66	3296.00	**<0.001**	-0.38	-0.43 – -0.33	-14.39	3914.00	**<0.001**
Overall Reward Value	0.05	-0.01 – 0.11	1.61	3448.00	0.108	-0.01	-0.06 – 0.05	-0.24	4222.00	0.809
Best - Worst Condition	-0.08	-0.15 – -0.01	-2.21	31.00	**0.035**	-0.03	-0.11 – 0.04	-0.85	31.00	0.402

203 **Table 2.** *Model comparison for OV_{reward} and OV_{goal} effects on RT across studies.*

	Study 1					Study 2				
Model	R^2	*AIC*	*dAIC*	X^2	*p*	R^2	*AIC*	*dAIC*	X^2	*p*
Model 0: VD + C_{b-w} (baseline)	0.22	4112				0.19	4396			
Model 1: baseline + OV_{reward}	0.22	4112	0	2.72	.099	0.19	4398	2	0.04	.847
Model 2: baseline + C_{b-w} : OV_{reward}	0.26	3956	-156	157.68	**< .001**	0.25	4197	-202	203.31	**< .001**
Model 3: baseline + OV_{goal}	0.26	3957	1	2.59	.108	0.25	4195	-2	0.06	.809

Note: For each study, models are compared sequentially, and dAIC is the difference in AIC of each model to the previous model.
VD = Value Difference; OV = overall value

The behavioral findings we observe across Studies 1 and 2 raise two potential concerns

that we addressed in follow-up studies. First, it could be the case that the novel findings we

observed in the Choose-Worst condition benefited from participants having the (traditional)

Choose-Best condition as a within-session reference point. While the counter-balancing of these

tasks mitigates this possibility, to rule out this possibility we performed a follow-up study in

which participants only performed the Choose-Worst task (Study S1), and replicated the overall

value findings we observed for this condition in Study 1 (Supplemental Results 1). Second, it is

possible that the dominance of the goal value over the reward value model observed in Studies 1-

2 reflected the hypotheticality of the choices in these studies, re-directing attention towards the

goal and away from potential rewards. While this would not straightforwardly account for

overall value effects observed in previous studies involving hypothetical choice (36), we sought

to rule this possibility out as well by performing a follow-up study in which choices were

incentivized rather than hypothetical, with participants receiving one of the products from the

study based on their indication that it was the most or least preferred item in a set (Study S2; see

Supplementary Methods). This study replicated the findings observed across Studies 1-2

(Supplementary Results 2); critically, RTs were still not influenced by overall reward value (p

= .838) and continued to be best accounted for by the goal value models (dAICs = -18 - -16).

Task-dependent effects of overall reward value can be collectively accounted for by

a goal value accumulator model. Previous research has shown that the dynamics of value-based

decision-making can be captured by models of accumulating evidence to bound (9, 10, 46, 47).

This work has demonstrated that a subset of such models can capture the speeding effects of both

overall reward value and value difference on choice RTs. For instance, these effects emerge

naturally from a model in which option values accumulate additively and compete with one

another (the leaky competing accumulator [LCA] model (48)). However, previous

implementations of this and related models of value-based decision-making have assumed that

activity in the model is driven by the *reward* value of one's options, and therefore predict that

overall value will only ever result in speeding of choice RTs. Accordingly, when we simulate

such a model we find that it replicates behavior on our Choose-Best task (consistent with past

work) but not our Choose-Worst task (Fig. S1).

To account for our findings, we developed a modified LCA model. Rather than

accumulating evidence of potential rewards, this model accumulates evidence in favor of one's

task goal (i.e., goal value). According to this model, higher reward associated with a given option

would constitute strong evidence in favor of one's goal during the Choose-Best task but weak

evidence in favor of one's goal in the Choose-Worst task. Choice behavior simulated by this

model qualitatively replicates all of our key findings across Studies 1 and 2: (1) the effect of

overall reward value on choice RTs reverses between Choose-Best trials (speeding) and Choose-

Worst trials (slowing) (Fig. 1D); (2) overall goal value has a consistent speeding effect, and is

sufficient to account for RTs across both tasks (Fig. S2); (3) value difference has a consistent

speeding effect across the two tasks (Fig. S3); and (4) overall choice RTs are similar across the

two tasks (Fig. S3).

The brain's value network differentially tracks reward and goal congruency.

Previous work has identified a network of brain regions – primarily comprising ventromedial

prefrontal cortex (vmPFC) and ventral striatum – that tracks both the overall reward value of a

choice set and the relative reward value of the chosen item (the *signed* difference between the

reward associated with the chosen versus unchosen items) (15, 31, 35, 36). However, because

these findings are based entirely on tasks that require participants to choose the best option, it is

unclear whether neural activity is being determined by the reward value of those items (as

implied by past studies) or by the goal congruency of those reward values. To disentangle these

two possibilities, we had participants in Study 2 perform our Choose-Best and Choose-Worst

tasks while undergoing fMRI. The fact that reward and goal congruency were orthogonalized by

design (Figure S4) allowed us to compare the degree to which activity in a previously defined

valuation network (15) was associated with overall and relative (chosen minus unchosen reward

value vs. chosen minus unchosen goal value) reward values and goal values.

To test the hypotheses that the value network tracks reward value (GLM-1) or goal

congruency (GLM-2), we performed separate regressions correlating single-trial BOLD activity

in this network with either relative and overall reward value (GLM-1) or relative and overall goal

value variables (GLM-2). We found that the goal value model (GLM-2) provided a significantly

better fit to neural activity ($X^2 = 5.67$, $p < $ 2e-16; Table 3), with significant effects of both overall

($b = 0.16$, $t = 2.45$, $p = .014$) and relative goal value ($b = 0.23$, $t = 3.50$, $p < .001$). Both goal

value variables continue to be significant when including reward-related variables in the same

model (GLM-3; Table 4; Fig. 2). Interestingly, in addition to tracking overall and relative goal

value, the value network *also* tracked overall reward value ($b = 0.21$, $t = 3.17$, $p = .001$; Table 4).

In other words, this network tracked overall and relative value of a choice set in terms of their

goal congruence, while also tracking the overall reward associated with those options

(irrespective of the choice goal). None of our GLMs revealed a significant correlation between

BOLD activity and relative *reward* value (i.e., chosen vs. unchosen reward).

This general pattern held across the two constituent regions of the valuation network

(ventral striatum and vmPFC, Fig. S5, Table S3), as well as posterior cingulate cortex (PCC,

Supplemental Results 4, Table S4), a region that often coactivates with the valuation network.

The only discernible difference between these regions was that vStr and PCC appeared to be

more weakly associated with overall goal value than vmPFC was.

277
278   **Figure 2. The valuation network tracks goal values and overall, but not relative**
279   **reward value. A)** Valuation Network Mask. **B)** Mixed-effects regression coefficients
280   show that the valuation network ROI (defined a priori based on Ref. 15) tracks both overall
281   and relative goal value, and also tracks overall (but not relative) reward value. Error bars
282   show standard error of the mean.
283

284   **Table 3.** *Model Comparison for Reward and Goal Value – related BOLD activity in the Valuation*
285   *Network ROI*

Model	R^2	AIC	BIC	dAIC	Chi^2	p
C_{b-w} + RT (baseline)	0.062	11976	12040			
GLM-1: baseline + RV_{reward} + OV_{reward}	0.066	11969	12045	-7	11	**0.004**
GLM-2: baseline+ RV_{goal} + OV_{goal}	0.065	11963	12040	-6	5.68	**<0.001**
baseline+ RV_{goal} + OV_{goal} + OV_{reward}	0.069	11955	12038	-8	10.2	**0.001**
GLM-3: baseline+ RV_{goal} + OV_{goal} +RV_{reward} + OV_{reward}	0.069	11957	12046	2	0.2	0.657

286   Note: RV = Relative Value, OV = Overall Value

287   **Table 4.** *Fixed Effects Summary GLM-3: Reward and Goal Values*

	BOLD Activity Value Network				
Predictors	*Estimates*	*CI*	*t*	*df*	*p*
(Intercept)	0.00	-0.08 – 0.08	0.01	31.00	0.992
Best - Worst Condition	-0.01	-0.10 – 0.09	-0.13	42.00	0.896
Overall Reward Value	0.21	0.08 – 0.34	3.15	3844.00	**0.002**
Relative Reward Value	-0.03	-0.16 – 0.10	-0.44	4225.00	0.657
Overall Goal Value	0.16	0.03 – 0.29	2.44	1632.00	**0.015**
Relative Goal Value	0.22	0.09 – 0.36	3.37	4173.00	**0.001**
RT	0.02	-0.08 – 0.12	0.37	32.00	0.715

Dissociable regions of striatum track reward versus goal congruency. Since we failed

to observe relative reward value signals in our a priori ROI, we performed a follow-up whole-

brain analysis (mirroring GLM-3) to test whether a subset of this network, or regions outside of

it, track this variable. We did not find such correlates, even at a liberal threshold (voxelwise

uncorrected $p < 0.01$, $k \geq 10$ voxels). However, our whole-brain analysis did reveal an

unexpected functional dissociation within the striatum, with more dorsal regions primarily

tracking overall reward value and more ventral regions primarily tracking relative goal value

(Fig. 3A). To test this dorsal-ventral dissociation directly, we defined ROIs for three sub-regions

along a dorsal-ventral axis previously shown to seed distinct resting-state networks (Figure 3B;

49). We found a significant interaction between value variables and striatal sub-regions (*F(4,*

25560) = 2.81, *p* = .024), such that sensitivity to overall reward value was greatest in the

dorsalmost sub-region (dorsal caudate; $b = -0.21$, $t = -2.10$, $p = .036$) whereas sensitivity to

relative goal value was greatest in the ventralmost sub-region (inferior ventral striatum; $b = 0.24$,

$t = 2.19$, $p = .029$, Fig. 3C, Supplemental Results 5, Table S5).

Unlike in striatum, an additional exploratory test of analogous differences across

subregions in vmPFC (rostral ACC versus medial orbitofrontal cortex (36)) did not reveal

significant interactions of any of our variables with subregion (Supplemental Results 6, Table

S6), though trends were observed differentiating rACC's tracking of overall reward value and

from mOFC's tracking of overall and relative goal value (Table S7).

Figure 3. Reward and goal value dissociate across the striatum's dorsal-ventral axis. A) Whole
brain results for relative goal value (blue) and overall reward value (green), thresholded at voxelwise
$p < 0.001$ and cluster-corrected $p < 0.05$. Despite being correlated with activity in our a priori value
network ROI, overall goal value did not survive the whole-brain threshold used for these follow-up
exploratory analyses. **B)** To interrogate our findings across regions of striatum, we selected
independent bilateral ROIs previously used as seeds for distinct resting-state networks (49): Dorsal
Caudate (dark orange; [x,y,z] = +/-12, 10, 8), Ventral Striatum, superior (orange; +/-8, 10, 1), and
Ventral Striatum, inferior (yellow; +/-10, 11, -9). **C)** Mixed-effects regression coefficients across
these ROIs demonstrate a dorsal-ventral dissociation, with dorsal regions more sensitive to overall
reward value and the ventralmost region more sensitive to relative goal value. Error bars show
standard error of the mean.

320
Correlates of overall reward value parallel those of choice appraisal. We have

previously shown that the overall reward value of an option set predicts how attractive a

participant will appraise that set to be (*set liking*), and that these liking ratings correlate with

value network activity (14, 36). Set liking correlated with activity in these regions irrespective of

whether participants were instructed to appraise the set on a given trial, suggesting that these

appraisals are triggered relatively automatically (32, 36, 50). Unlike previous work, our current

studies were able to distinguish whether set liking was driven by overall reward value or overall

goal value, and to compare the neural correlates of set liking with the neural correlates of both

types of overall and relative value. Across both studies, we found that set liking correlated with

overall reward value (Study 1: $b = 6.35$, $z = 25.44$, $p < .001$, Study 2: $b = 4.87$, $z = 20.74$, p

$< .001$) and not overall goal value (Study 1: $b = -0.07$, $z = -0.53$, $p = .599$, Study 2: $b = 0.09$, z

$= 0.68$, $p = .497$). Consistent with previous findings, in Study 2 we found that activity in the

value network was correlated with set liking ($b = 0.16$, $t = 2.22$, $p = .027$). While set liking

correlates did not differ significantly across regions of striatum ($F(2, 12295) = 0.72$, $p = .485$), it

is notable that the qualitative pattern of responses (numerically higher in the more dorsal regions)

most closely matched the pattern we observed for overall reward value (Fig. S6). Together these

findings provide tentative but convergent evidence that overall reward value signals may be

related to a process of automatic appraisal.

341   **Discussion**

342   In spite of the heterogeneity of approaches in research on value-based decision-making,

a consistent set of findings has emerged with respect to the behavioral and neural correlates of

choice values. However, up to this point it has been unknown whether these correlates reflect the

reward value of those options (i.e., how much reward one can expect for obtaining a given

option) or their goal congruency (i.e., to what degree does a given option support one's current

choice goal). To disentangle these accounts of behavioral and neural data, we had participants

make value-based choices under two different choice goals, either to choose the most rewarding

or least rewarding option. We found that choice RTs and neural activity in value-related regions

were accounted for by overall and relative *goal* value (increasing with the congruency of option

values with the current choice goal). We additionally identified neural (but not behavioral)

signatures of overall *reward* value. Together, these results provide a cognitive, rather than

reward-centered explanation of hallmark findings in value-based decision-making, and call for a

significant reinterpretation of past research.

Previous studies on value-based decision-making have consistently found that

participants make faster decisions as the overall reward value of their options increases (11-14,

51). It has been natural to assume that these speeding effects are related to the rewarding

properties of these choice options, particularly given that approach is known to be facilitated by

reward and impeded by punishment (26, 27) and response vigor is known to increase in high

reward rate environments (28-30). However, these studies confounded reward value and goal

congruency, concealing an alternative, cognitive explanation of these findings, that speeding

effects of overall value reflect increased activation of responses specific to one's goal (10, 45, 48,

52). Our findings support the latter account, demonstrating that a choice set's overall reward

value speeds choices only when the participant's goal is to choose the most rewarding item, and

that it has the opposite effect when the goal is to choose the least rewarding item.

Our neural findings serve to similarly qualify reward-based accounts of neural data.

Previous neuroimaging research has linked activity in a common set of brain regions to the

relative and overall reward value of a choice set. On their face, those findings could be

interpreted as reflecting a *directionally consistent* coding of potential rewards (31, 37, 42), such

that (a) one's preference for a given item could be read out from activity in these regions and (b)

the preferences coded in these regions could feed directly into a choice between the choice

options (reflected in relative value signals). On such an account, relative value signals should

always reflect whether the chosen option is generally more or less preferred than the unchosen

option(s), and overall value should always reflect how much those options are generally

preferred. This form of directionally consistent coding scheme is intuitive and can be seen as

evolutionarily adaptive because it enables an organism to rapidly determine which options are

most approach-worthy. Critically, this value code is also sufficient to perform our tasks –

providing participants just as much information about the least preferred items as it does the

most preferred – obviating the need for any additional, recoded value signals (53). We found

neural signals that are inconsistent with this coding scheme, but instead tracked options' goal

congruency. These overall and relative goal congruency signals would not be predicted by a

reward coding account but, together with our behavioral and modeling results, are consistent

with an alternate account whereby activity in these valuation networks reflects the accumulation

of goal-relevant information leading up to a choice (10, 13), including information about

potential rewards and their relationship to one's current task goal (54).

In addition to these neural signatures of goal congruency, we also observed neural signals

that tracked the overall value of a choice set in a goal-*independent* (i.e., directionally consistent)

manner. That is, the value network also tracks how much reward can be expected from the choice

set, across both Choose-Best and Choose-Worst tasks. Notably, unlike our goal-dependent

signals, these goal-independent overall value signals were not accompanied by goal-independent

relative value signals (i.e., chosen minus unchosen reward), as would be expected if this circuit

was comparing items according to their reward value. It may therefore be parsimonious to

interpret these overall reward signals as reflecting an appraisal process that occurs *in parallel*

with the choice process (14, 36, cf. 55, or preceding it; 56) rather than reflecting a component of

the decision-making process (36). With that said, we emphasize that these inferences pertain

specifically to BOLD signals within vmPFC and ventral striatum, and that it is entirely possible

that other coding schemes would be found at the single-unit level within these or other regions.

There is another potential explanation for the overall reward value and relative goal value

signals we found juxtaposed in the value network, which focuses on the *affective* qualities these

variables share (36). Specifically, we and others have shown that overall reward value predicts

how positively a person will feel about a set of items (14, 36) and that the value network tracks

such affective appraisals whether or not they are task-relevant (32, 36). Similarly, relative goal

value is a proxy for confidence in one's decision, as well as for the ease or fluency of a task.

Both of these engender positive affect (57-60) and both have been linked with activity in the

value network (36, 61-65). This shared association with affect offers a salient (but non mutually

exclusive) alternative to accounts of value network findings that focus on decision-making and

other cognitive processes. However, further research on this circuit is necessary to tease apart

affective and cognitive contributions to neural activity.

Within striatum, we found a surprising anatomical dissociation, whereby dorsomedial

versus ventral regions were differentially associated with overall reward value and relative goal

value. This dissociation was most strongly driven by an effect of overall reward value within

dorsal caudate that was absent for relative goal value (whereas both variables were associated, to

different degrees, with activity in ventral striatum). While not predicted, we speculate that this

pattern may reflect the differential relationship of the two variables with affective and

motivational states. Specifically, overall reward value is not only associated with affective

appraisal (as noted above), but can also serve to motivate approach behavior. Accordingly,

previous findings have linked dorsomedial striatum with action valuation (66-68) and motivation

(69-71). It is also noteworthy that in spite of their relatively close spatial proximity, these ROIs

fall within separate networks with distinct patterns of resting-state functional connectivity – the

ventralmost region is associated with the limbic network and the dorsalmost region is associated

with the frontoparietal control network (49, 72, 73). Therefore, an alternative account of this

dorsal caudate finding is that overall reward value, being a task-irrelevant variable (cf.

distractor), may elicit frontoparietal activity because of the increase in control demands. This

interpretation is consistent with the a similar dissociation recently observed by Fischer and

colleagues (74). They had participants perform a learning task and found that ventral striatum

tracked short-term rewards whereas dorsal striatum generally tracked long term rewards, which

had to inferred from the task environment (74). However, they found that dorsal striatum also

tracked short-term rewards and the extent to which it did so predicted *better* learning of long-

term rewards, which these authors interpreted as reflecting increased control that was triggered

by this region's encoding of salient information (short-term reward) that was irrelevant to the

task of inferring long-term reward. In spite of this converging evidence, we emphasize that the

interpretations of our striatal findings are post hoc and therefore should be taken with caution.

Nevertheless, they suggest avenues for further inquiry that could shed light on this intriguing

finding.

Our findings have important implications for future research on value-based decision-

making within healthy and clinical populations. We show that traditional approaches to studying

such decisions conflate two types of variables: those that are specific to rewards per se and those

that are related to how consistent rewards are with choice goals. We prescribe a method for

pulling these two types of signals apart, and show that doing so reveals distinct behavioral and

neural patterns. This method could, for instance, be employed to identify mechanisms that are

shared (e.g., goal-related) and distinct (e.g., outcome-related) when choosing among appetitive

options versus aversive options (25, 75).

In addition to informing basic research into distinct processes supporting value based

choice (reward processing vs. goal-driven behavior), our work carries important implications for

research on clinical populations that exhibit aberrant reward processing (70, 71, 76-78) or

impairments in translating rewards into goal-driven behavior (79). For instance, to the extent

these individuals exhibit impulsive behaviors linked to reward rather than goal values, we

provide an approach for uniquely identifying this variable of interest and honing in on the

relevant deficits. By decoupling these two processes, future research can identify the best

approaches for treating the relevant deficits, or at least rule out the worst ones.

452   **Method**

Participants

Participants were recruited from Brown University and the general community. For Study

1, 37 participants were recruited. Of these, seven were excluded, one due to previous

participation in a similar experiment, three due to incomplete sessions, and three due to

insufficient variance in product evaluation, precluding the generation of sufficient choice sets.

Thus, 30 participants (76.7 % female, M_{age} = 20.3, SD_{age} = 2.1) were included in our analyses.

For Study 2, 31 participants were recruited and one was excluded due to an incomplete session.

Thus, 30 participants (56.67% female, M_{age} = 21.87, SD_{age} = 4.40) were included in our analyses.

Participants in all studies gave informed consent in accordance with the policies of the Brown

University Institutional Review Board.

Procedure

Participants first evaluated a series of products according to how much they would like to

have them. Subsequently, participants viewed sets of four products and were instructed either to

choose the item they most preferred (Choose-Best condition; cf. 14) or to choose the item they

least preferred (Choose-Worst condition). The different choice goals (Choose-best vs. Choose

worst) were instructed in two separate blocks, and the order of the two blocks was

counterbalanced across participants (see Fig.1 and SI Methods for details). The general

procedure in Study 2 was identical to that in Study 1, except that the choice phase was performed

in the scanner.

fMRI Data Acquisition and Analysis

fMRI data were acquired on a Siemens 3T PRISMA scanner, and analyzed using a

combination of SPM12 and R (see SI Methods). After standard preprocessing, BOLD estimates

for each trial were extracted from our ROIs and analyzed using linear mixed effects models. Our

primary fMRI analyses focused on an a priori *value network* ROI, based on a meta-analysis of

the value-based decision-making literature by Bartra and colleagues (15). Secondary analyses

examined three additional ROIs within striatum, based on an analysis of resting-state networks

by Choi and colleagues (49).

Data Availability

The datasets generated and analyzed during the current study are available from the

corresponding author on reasonable request.

Code Availability

Scripts for all analyses are available under https://github.com/froemero/goal-congruency-

dominates-reward.

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Supplemental Information
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SI Methods

Procedure. The experiment was computerized using Matlab's Psychophysics Toolbox (1, 2). Participants first evaluated a series of products according to how much they would like to have each one. Items were presented individually, and participants used a mouse to indicate their evaluation of that item along a visual analog scale, anchored at 0 ("not at all") and 10 ("a great deal"). Participants were encouraged to use the entire range of the scale and reserve 0 and 10 for extreme values. Based on these subjective evaluations choice sets were created, such that half of the choices primarily varied in relative value (RV; e.g. best vs. average of the remaining items), while the other half primarily varied in overall set value (OV; average value of items in a set). Additional details regarding distribution of items and process of choice set construction can be found in (3). Participants then viewed sets of four products and were instructed either to choose the item they most preferred (Choose-Best condition; cf. 3, 4, 5) or to choose the item they least preferred (Choose-Worst condition). The different choice goals (Choose-Best vs. Choose-Worst) were performed in separate blocks, and the order of the two blocks was counterbalanced across participants. Stimuli remained on the screen until a choice was made. In Study 1, participants performed a total of 120 choices, 60 for each condition. They indicated their choice with a mouse click. There were no response deadline in the initial rating or choice. The general procedure in Study 2 was identical to that in Study 1 except that (1) choices were performed in the scanner; (2) they completed 144 choices (72 for each choice goal); (3) the interval between choices was varied across trials (2-7s, uniformly distributed); and (4) choices were made with a button press rather than with a mouse. Participants responded with the index and middle fingers of their left hand and right hand, using an MR-safe response keypad. Response keys were mapped to

locations on the screen (upper left, lower left, upper right and lower right). Participants practiced the response mappings prior to the choice phase by pressing the button corresponding to a given cued location until they gave the correct response on 15 consecutive trials.

Study S1. 19 participants were recruited from Brown University and the general community, out of which one had to be excluded due to technical issues. Thus, the final sample includes 18 participants (61.1% female, M_{age} = 20.1, SD_{age} = 3.6). The procedures in Study S1 were identical to Study 1, except that participants only performed the Choose Worst condition (120 trials).

Study S2. 15 participants were recruited from Brown University and the general community, out of which one had to be excluded due to technical issues. Thus, the final sample includes 14 participants (78.6 % female, M_{age} = 23.71, SD_{age} = 5.93). The procedures in Study S2 were identical to Study 1, except that choices were incentivized. Following the individual item rating and immediately prior to the choice phase, participants were informed that they will receive one of the items they rated following the experiment and were asked to provide delivery information. They were informed that these options went into a lottery which they could bias with their subsequent choices. Depending on the choice task, they were then informed that choosing an option will increase (choose best) or decrease (choose worst) the probability of that option to be drawn in the final lottery.

Imaging Parameters. Scans were acquired on a Siemens 3T PRISMA scanner with a 64-channel phase-arrayed head coil, using the following gradient-echo planar imaging (EPI) sequence parameters: repetition time (TR) = 2500 ms; echo time (TE) = 30 ms; flip angle (FA) = 90°; 3 mm voxels; no gap between slices; field of view (FOV): 192 x 192; interleaved acquisition; 39 slices. To reduce signal dropout in regions of interest, we used a rotated slice prescription (30° relative to AC/PC). The slice prescription encompassed all ventral cortical

structures but in a few participants omitted regions of dorsal posterior parietal cortex. Structural

data were collected with T1-weighted multi-echo magnetization prepared rapid acquisition

gradient echo image (MEMPRAGE) sequences using the following parameters: TR = 2530 ms;

TE = 1.69 ms; FA = 7°; 1.0 mm isotropic voxels; FOV = 256 X 256. Head motion was restricted

with a pillow and padding. Stimuli were generated using Matlab's Psychophysics Toolbox

(Matlab 2013a) presented on a 24'' BOLDscreen flat-panel display device (Cambridge Research

Systems) and were viewed through a mirror mounted on the head coil.

fMRI Analysis. fMRI data were processed using SPM12 (Wellcome Department of Imaging

Neuroscience, Institute of Neurology, London, UK). Raw volumes were realigned within

participants, resampled to 2mm isotropic voxels, non-linearly transformed to align with a

canonical T2 template and spatially smoothed with a 6 mm full-width at half-max (FWHM)

Gaussian kernel.

Trial-wise ROI analyses. Preprocessed data were submitted to linear mixed-effects analyses

using a two-step procedure. In the first step, we computed first-level general linear models

(GLM) in SPM to generate BOLD signal change estimates for each trial and participant. GLMs

modeled stick functions at the onset of each trial. Trials were concatenated across the two task

blocks and additional regressors were included to model within-block means and linear trends.

GLMs were estimated using a reweighted least squares approach (RobustWLS Toolbox; 6) to

minimize the influence of outlier time-points (e.g. due to motion). The obtained estimates were

transformed with the hyperbolic arcsine function (to achieve normality), and then analyzed using

LMMs using lme4 (7) in R (8).

Of primary interest in the present study is activation in the valuation network, which was

obtained using the conjunction mask from Bartra, McGuire and Kable (9). These results were

followed up with separate masks for the vmPFC and vStr clusters within this ROI. PCC activity was extracted from an ROI identified in a previous study (4).

To test for gradients within vStr, we constructed spheres (r = 4mm) around seeds in three regions along the caudate's dorsal-ventral axis most strongly associated with the frontoparietal control network (left hemi coordinates: -12, 10, 8), the default mode network (-8, 10, 1), and the limbic network, respectively (-10, 11, -9; 10). For the gradient analysis of these spheres, the data were best fit modeling linear as well as quadratic effects for region. ROIs for rACC and mOFC to test for differences within vmPFC were selected anatomically, analogous to a previous study (5).

Whole-brain analyses. We complemented the ROI analyses with whole-brain GLMs. For these analyses, we computed first-level GLMs, modeling stick function at stimulus onsets, and parametric regressors for 1) choice goal, 2) reward-related OV, 3) goal-related OV, 4) reward-related chosen versus unchosen value and 5) goal-related chosen versus unchosen value. Regressors were de-orthogonalized to let them compete for variance. As above, trials were concatenated across the two task blocks, additional regressors were included to model within-block means and linear trends, and GLMs were estimated using RobustWLS. Second level random effects analyses on first-level estimates were performed using SPM with voxel-wise thresholds of $p < .001$ and cluster-corrected thresholds of $p < .05$. Results were visualized using XJview (http://www.alivelearn.net/xjview).

Additional measures. The following measures were obtained in each experiment. With the exception of set appraisals, these additional evaluations are not of primary interest to the present study and were therefore not included in any analyses.

Affective Rating. Following the choices, participants sequentially viewed all choices again and for each set rated (1) how stressed/anxious that choice made them feel, (2) how confident they were in their choice, and (3) how much they liked the set as a whole (set appraisal).

Item reevaluation. At the end of the experiment, participants in Study S1and a subset of participants in Study 1 were asked to rate all items one more time with the same procedure as during subjective evaluation, but using a bipolar scale anchored at -10 ("dislike a great deal") and 10 ("like a great deal").

Individual difference measures. At the start of each session, participants completed the following personality inventories: the Behavioral Inhibition/Activation Scales (BIS/BAS), Neuroticism subscale of the NEO Five Factor Inventory, Intolerance for Uncertainty Scale, and Need for Cognition.

Analysis. Scripts for all analyses are available under https://github.com/froemero/goal-congruency-dominates-reward. Data were analyzed using lme4 package (7) for R (version 3.4.3, 8). RTs were analyzed with linear mixed effects models (LMMs) and choices were analyzed with generalized linear mixed effects models (GLMMs) with a binomial link function. We used the lmerTest package to obtain *p*-values based on Satterthwaite approximation of degrees of freedom. Choice consistency was determined based on whether participants chose the item with the highest value in Choose-Best, and conversely the lowest value in Choose-Worst. Analyses of choice consistency tested the probability of choosing the option that best satisfies the current choice goal (lowest-valued item for Choose-Best, highest-valued item for Choose-Worst); these

analyses exclude trials for which multiple options equally satisfy these instructions (i.e., when multiple options share the lowest or highest value). Response times were log-transformed to reduce skew. Value difference and overall value were mean-centered and rescaled to a range from zero to one prior to analysis. Choice condition (Choose-Best vs. Choose-Worst) was included as a factor to test for main effects of choice goal and interactions between goal and choice values. We used a sliding difference contrast that estimates the difference in means between subsequent factor levels with the intercept (baseline value) being estimated as the mean across both conditions. Random effects were specified as supported by the data, according to singular value decomposition (11, 12).

Computational model. To directly compare reward-based and goal-based accounts of our findings, we simulated data using two versions of a leaky competing accumulator model (LCA; 13). In an LCA model, evidence for each potential response (in our case, each of four choice options) accumulates noisily over time based on its respective input (I_{Option}). As the evidence for each response accumulates, it competes with the evidence accumulated for each of the other responses, according to a mutual inhibition term (w). Over time, accumulated evidence decays according to a leak (or forgetting) term (k). A decision is made once activation of one of the options reaches a decision boundary (z), and the time at which this threshold is crossed is recorded as the decision time. This decision time is combined with a fixed non-decision time (reflecting stimulus processing and response execution; t_0) to generate the response time for a given model simulation. Unlike the standard drift diffusion model, the LCA has additive properties that produce RTs that vary with the both the relative and overall value of its inputs (13).

We compared two versions of the LCA model to see which could better account for our findings. One version used reward values as inputs (as in previous studies; 14) while a second, modified form of this model used goal values as inputs. In the Choose Best condition, goal values are identical to reward values. For the Choose Worst condition goal values were computed by recoding reward values, such that lower reward values had higher goal value and vice versa. Both models otherwise used identical parameter, which we determined by performing gradient descent (using *fmincon*) to maximize the likelihood of Study 1 data (choices and RTs) given model predictions, collapsing across participants. We generated trial-wise predictions by simulating the LCA's accumulation-to-bound process 1000 times for each unique choice set in the experiment. The best-fit model parameters were as follows: $z = 2.243$, s (noise coefficient) = 0.587, $k = 0.153$, $w = 0.671$, $t_0 = 0.849$s (log likelihood = -12883.128). The goal value model simulations reported in the main text use these parameter values but all of the relevant qualitative predictions of these models are robust to variation in these parameters. We did not perform additional model fits or simulations for an analogous reward value-based LCA model because it is necessarily unable to incorporate information about the different choice goals and will hence inevitably always choose the best option regardless of the task. Model simulation and parameter estimation were performed using custom Matlab (2017b) scripts.

Visualization. Figures displaying projected values and confidence intervals of response times were generated using effects and ggplot2 packages based on the relevant LMMs. For viewing purposes, centered and scaled predictor variables are rescaled to their original values prior to plotting.

SI Results

1. The slowing effect of overall value in the Choose-Worst condition is not dependent on a Choose-Best reference point or hypotheticality of choices. In Study S1, participants only performed the Choose-Worst task. As in Studies 1-2, we found that RTs increased with increasing overall value (b = .52, t = 5.16, p < .001). As in these other studies, RTs were also faster for larger value differences (minimum value vs. average remaining, b = -.57, t = -11.10, p < 2e-16).

Study S2's participants performed the same tasks as in Study 1 but, rather than engaging in hypothetical choices, these participants had the opportunity to receive one of the items (products) from the study and their choices directly influenced which item they would receive. In spite of this difference, Study S2's results were qualitatively identical to Study 1. RTs decreased with increasing overall value in choose best (b = -0.22, p = .002) but increased with increasing overall value in choose worst (b= 0.20, p = .005). Therefore, we once again observed a significant interaction between choice goal and overall value (b = -0.41, p < .001, Table S1). Like in Study 1 and Study 2, goal values provided a parsimonious and sufficient account of the observed effects and there was no significant residual effect of overall reward value (b = -0.01, p = .838, Table S2).

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Table S1

Fixed effects of Overall Reward Values vary by Choice Goal

Predictors	log RT				
	Estimates	*CI*	*t*	*df*	*p*
(Intercept)	1.70	1.52 – 1.88	18.77	15.00	**<0.001**
Value Difference	-0.61	-0.71 – -0.51	-12.05	1630.00	**<0.001**
Overall Value	-0.01	-0.11 – 0.09	-0.20	1635.00	0.838
Best - Worst Condition	0.08	-0.07 – 0.22	1.00	15.00	0.335
Overall Value: Best - Worst	-0.41	-0.61 – -0.22	-4.20	1489.00	**<0.001**
Random Effects					
σ^2	0.21				
$\tau_{00 \; subj\_idx}$	0.10				
$\tau_{11 \; subj\_idx.isChooseBestTrial2-1}$	0.07				
$\rho_{01 \; subj\_idx}$	-0.30				
ICC	0.36				
$N_{\; subj\_idx}$	14				
Observations	1644				
Marginal R^2 / Conditional R^2	0.066 / 0.404				

Table S2

Model comparison for overall reward and overall goal value effects on RT

	R^2	*AIC*	*BIC*	*dAIC*	*Chi^2*	*p*
VD + C_{b-w} (baseline)	0.33	2255	2293			
baseline + OV_{reward}	0.33	2257	2300	2	0.17	0.676
baseline + OV_{goal}	0.35	2239	2282	-18	17.77	**<0.001**
baseline + C_{b-w}: OV_{reward}	0.35	2241	2289	2	0.04	0.837

Note: For each study, models are compared sequentially, and dAIC is the difference in AIC of each model to the previous model. VD = Value Difference; OV = overall value

Table S1
Fixed Effects Summary GLM-3: Reward and Goal Values within sub-regions of valuation network

Predictors	BOLD Activity vmPFC					BOLD Activity vStr				
	Estimates	*CI*	*t*	*df*	*p*	*Estimates*	*CI*	*t*	*df*	*p*
(Intercept)	-0.56	-0.70 – -0.42	-7.76	31.00	**<0.001**	-0.41	-0.55 – -0.27	-5.73	31.00	**<0.001**
Best - Worst Condition	-0.06	-0.21 – 0.09	-0.78	46.00	0.439	0.08	-0.10 – 0.27	0.90	37.00	0.374
Overall Reward Value	0.31	0.07 – 0.55	2.52	3747.00	**0.012**	0.28	0.08 – 0.48	2.80	4022.00	**0.005**
Relative Reward Value	0.05	-0.18 – 0.28	0.43	4202.00	0.669	-0.13	-0.32 – 0.07	-1.28	4237.00	0.200
Overall Goal Value	0.33	0.09 – 0.56	2.74	972.00	**0.006**	0.08	-0.12 – 0.28	0.76	2633.00	0.445
Relative Goal Value	0.33	0.09 – 0.57	2.70	4195.00	**0.007**	0.30	0.10 – 0.50	2.99	4133.00	**0.003**
RT	0.20	0.01 – 0.39	2.08	32.00	**0.045**	-0.17	-0.33 – -0.01	-2.12	32.00	**0.042**
Random Effects										
σ^2	3.20					2.15				
τ_{00}	0.13 $_{subj\_idx}$					0.13 $_{subj\_idx}$				
τ_{11}	0.05 $_{subj\_idx.isChooseBestTrial2-1}$					0.17 $_{subj\_idx.isChooseBestTrial2-1}$				
	0.11 $_{subj\_idx.cRT}$					0.08 $_{subj\_idx.cRT}$				
ρ_{01}	0.28					0.01				
	-0.09					0.09				
ICC	0.05					0.08				
N	30 $_{subj\_idx}$					30 $_{subj\_idx}$				
Observations	4270					4270				
Marginal R^2 / Conditional R^2	0.007 / 0.056					0.009 / 0.088				

Table S2
PCC Activity tracks reward and goal congruency similarly to the valuation network

Predictors	Estimates	CI	t	df	p
		BOLD Activity PCC			
(Intercept)	-0.76	-0.90 – -0.63	-11.02	31.00	**<0.001**
Best - Worst Condition	0.04	-0.13 – 0.22	0.51	41.00	0.616
Overall Reward Value	0.25	0.03 – 0.48	2.21	3766.00	**0.027**
Relative Reward Value	-0.09	-0.30 – 0.13	-0.77	4216.00	0.443
Overall Goal Value	0.12	-0.11 – 0.34	1.03	1829.00	0.301
Relative Goal Value	0.43	0.21 – 0.66	3.77	3999.00	**<0.001**
RT	-0.16	-0.32 – -0.01	-2.08	31.00	**0.046**
Random Effects					
σ^2	2.81				
$\tau_{00 \; subj\_idx}$	0.12				
$\tau_{11 \; subj\_idx.isChooseBestTrial2-1}$	0.12				
$\tau_{11 \; subj\_idx.cRT}$	0.04				
ρ_{01}	0.16				
	-0.01				
ICC	0.05				
N_{subj_idx}	30				
Observations	4270				
Marginal R^2 / Conditional R^2	0.008 / 0.060				

Table S3. *Changes in value effects across the dorsal-ventral axis in striatum*

	Estimates	SE	t	df	p
(Intercept)	-0.40	0.06	-6.26	29.61	**<0.001**
Region (linear)	-0.11	0.02	-6.28	25559.83	**<0.001**
Region (quadratic)	-0.08	0.02	-4.50	25559.83	**<0.001**
RT	-0.21	0.07	-3.00	30.15	**0.005**
Overall Reward Value	0.35	0.06	5.54	25479.49	**<0.001**
Relative Goal Value	0.20	0.07	2.99	25531.34	**0.003**
Region (linear): Overall Reward Value	-0.14	0.10	-1.41	25559.83	0.158
Region (quadratic): Overall Reward Value	-0.21	0.10	-2.10	25559.83	**0.036**
Region (linear): Relative Goal Value	0.24	0.11	2.19	25559.83	**0.029**
Region (quadratic): Relative Goal Value	0.02	0.11	0.20	25559.83	0.844

Table S6
Testing for Differences in Reward and Goal Congruency related Activity between rACC and mOFC

	Sum Sq	Mean Sq	NumDF	DenDF	F value	Pr(>F)
Best - Worst Condition	1.6	1.6	1	226.88	1.67	0.197
Overall Reward Value	5.38	5.38	1	6676.4	5.63	**0.018**
Relative Reward Value	0.06	0.06	1	8475.2	0.07	0.796
Overall Goal Value	2.58	2.58	1	5966.15	2.7	0.101
Relative Goal Value	6.61	6.61	1	8405.74	6.91	**0.009**
Region	0.02	0.02	1	29.39	0.02	0.890
RT	0.23	0.23	1	28.54	0.24	0.628
Region: Best - Worst Condition	0.87	0.87	1	8440.16	0.91	0.341
Region: Overall Reward Value	0.21	0.21	1	2585.16	0.22	0.641
Region: Relative Reward Value	0.91	0.91	1	8455.28	0.95	0.330
Region: Overall Goal Value	0.05	0.05	1	8445.82	0.06	0.811
Region: Relative Goal Value	1.29	1.29	1	8358.15	1.34	0.246

Table S7
Reward and Goal Congruency effects within rACC and mOFC

Predictors	BOLD Activity rACC					BOLD Activity mOFC				
	Estimates	*CI*	*t*	*df*	*p*	*Estimates*	*CI*	*t*	*df*	*p*
(Intercept)	-0.77	-0.87 – -0.67	-14.85	31.00	**<0.001**	-0.23	-0.34 – -0.12	-3.95	31.00	**<0.001**
Best - Worst Condition	-0.02	-0.13 – 0.09	-0.32	4087.00	0.751	-0.08	-0.20 – 0.03	-1.44	4154.00	0.150
Overall Reward Value	0.22	0.02 – 0.43	2.15	3253.00	**0.032**	0.12	-0.09 – 0.33	1.16	3562.00	0.247
Relative Reward Value	-0.06	-0.26 – 0.14	-0.61	4252.00	0.544	0.10	-0.11 – 0.31	0.95	4249.00	0.341
Overall Goal Value	0.02	-0.17 – 0.21	0.22	4216.00	0.830	0.22	0.03 – 0.42	2.24	4240.00	**0.025**
Relative Goal Value	0.18	-0.03 – 0.39	1.70	4203.00	0.089	0.24	0.03 – 0.45	2.19	4249.00	**0.028**
RT	-0.36	-0.53 – -0.18	-4.04	32.00	**<0.001**	0.30	0.10 – 0.49	2.98	33.00	**0.005**
Random Effects										
σ^2	2.37					2.50				
$\tau_{00\ subj\_idx}$	0.06					0.08				
$\tau_{11\ subj\_idx.cRT}$	0.11					0.17				
$\rho_{01\ subj\_idx}$	0.17					0.04				
ICC	0.03					0.04				
N_{subj_idx}	30					30				
Observations	4270					4270				
Marginal R^2 / Conditional R^2	0.014 / 0.047					0.009 / 0.052				

Figure S1. LCA-simulated RTs based on reward values of the items. Increasing overall value (left) and value difference (right) lead to reduced RTs irrespective of choice goal. Thus the model cannot replicate the observed interaction of overall value and choice goal.

Figure S2. LCA-simulated RTs based on goal values of the items. Increasing overall goal value captures RTs speeding across both choice goals.

Figure S3. LCA-simulated RTs based on goal values of the items replicates patterns in observed behavior. Increasing overall value (left) leads to reduced RTs for Choose Best, but increased RTs for Choose Worst. Increasing Value Difference leads to reduced RTs irrespective of choice goal.

Figure S4. Relationships between value-related predictors. **A.** Distributions of Overall and Relative Reward and Goal Values across all trials. **B.** Reward and Goal Values are orthogonal. Shown are Spearman correlation coefficients.

Figure S5. Effects of Relative and Overall Reward and Goal Values within constituting regions of the valuation network.

Figure S6. Set appraisal (liking) effects along the striatum's dorsal to ventral axis.