Distinct contributions of anterior and posterior orbitofrontal cortex to adaptive decision-making

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**Abstract**

The lateral orbitofrontal cortex (OFC) is critical for flexibly adjusting choices when outcome values change. This requires representations of stimulus-outcome associations and inferring the current value of outcomes, but whether and how different parts of OFC contribute to these functions has remained unclear. Here we used transcranial magnetic stimulation (TMS) to disrupt activity in functional networks centered on the anterior (aOFC) and posterior (pOFC) lateral OFC. Participants (n = 48) received aOFC or pOFC network-targeted TMS either before learning associations between visual stimuli and sweet or savory food odor rewards, or, on the next day, before a meal to selectively devalue one of these rewards. TMS targeting pOFC before the meal disrupted goal-directed behavior, as measured by choices of stimuli predicting non-sated rewards in a probe test, whereas disrupting aOFC before learning stimulus-outcome associations similarly impaired choices in the probe test. These findings demonstrate distinct contributions of different OFC subregions to goal-directed behavior.

# Introduction

Humans and animals effortlessly adapt to changing environments by flexibly adjusting their behavior. This adaptability relies on outcome-guided decision-making, where individuals can re-evaluate their choices in real time, simulating potential outcomes based on changes in outcome value (Daw, Niv, and Dayan 2005) rather than defaulting to habitual responses. For example, a restaurant chef might anticipate that a guest could experience an allergic reaction to certain ingredients and adjust the dish accordingly before an issue arises. To enable this flexibility, a detailed representation of the environment—commonly referred to as a cognitive map or model -is essential (Behrens et al. 2018). A chef with full knowledge of ingredients and associated allergies can efficiently modify recipes to accommodate allergies without compromising the dish. The orbitofrontal cortex (OFC) plays a central role in both processes, supporting adaptive behaviors through the formation of cognitive maps (Costa et al 2023; Wilson et al. 2014; M. Z. Wang and Hayden 2021) as well as their use to simulate potential outcomes (Howard et al. 2020; Rudebeck and Murray 2014).

The OFC is a heterogeneous region, comprising multiple subregions with varying anatomical and functional properties along both mediolateral and anterior-posterior axes (Price 2007; Wallis 2012; Kahnt et al. 2012; Izquierdo 2017; M. Z. Wang, Hayden, and Heilbronner 2022; Heilbronner et al. 2016; Walton et al. 2011; Mackey and Petrides 2010; Kringelbach and Rolls 2004; Neubert et al. 2015). In humans, studies on value-based decision-making have primarily focused on the functional distinctions between the medial and lateral OFC (Kringelbach & Rolls 2004; Wallis 2012; Kahnt et al. 2012; Walton et al. 2011; McNamee et al 2013; Howard et al. 2015; O’Doherty et al. 2001), whereas the anterior-posterior gradient has received comparatively less attention.

The current study aims to identify the distinct roles of anterior and posterior subregions within the lateral OFC in supporting adaptive behavior in an outcome devaluation task (Wilson et al. 2014; Howard et al. 2020; Colwill and Rescorla 1985; Balleine and Dickinson 1998; Baxter et al. 2000; Murray et al. 2015; Critchley and Rolls 1996; O’doherty et al. 2000; Gottfried, O’Doherty, and Dolan 2003; Howard and Kahnt 2017, 2021; Gallagher, McMahan, and Schoenbaum 1999; Pickens et al. 2003; Ostlund and Balleine 2007). Outcome devaluation assesses responses to predictive cues following the selective devaluation of their associated outcomes, thereby revealing the capacity to align choices with updated goals and contexts. In outcome-specific versions of this task, different stimuli are first associated with different but equally preferred rewards. Next, one of the outcomes is selectively devalued (for instance by feeding it to satiety), and then choices between stimuli are assessed in a probe test. While earlier theories emphasized the role of the OFC in signaling the current value of stimuli to guide response selection (Baxter et al. 2000), more recent accounts propose two complementary roles: one in using mental simulations to infer or update the value of outcome-predicting stimuli (Wilson et al. 2014; Murray et al. 2015; Howard et al. 2020), and another in constructing and modifying the relevant cognitive map that links stimuli to outcomes during initial learning (Costa et al. 2023). In the current work, we focus on these latter two mechanims, proposing a unified framework that integrates them within the lateral OFC and empirically test for functional specialization across subregions.

Based on previous work in non-human primates suggesting functional specializations for anterior and posterior OFC in goal-directed behavior (Murray et al. 2015), we hypothesize that anterior and posterior OFC subregions are differentially required during different phases of the outcome devaluation task. Specifically, we hypothesize that the anterior portion of the central/lateral OFC is required for the acquisition of specific stimulus-outcome associations during initial training, whereas the posterior portion is required for retrieving and using these associations to guide choices in the probe test. To test this, we applied network-targeted transcranial magnetic stimulation (TMS) with continuous theta burst stimulation (cTBS) before initial training or before the probe test in a multi-session within-participant study. This approach allowed us to test the specific roles of anterior and posterior portions of the lateral OFC network for learning the associative task structure and guiding choices based on current values.

Our findings reveal distinct roles for the anterior and posterior lateral OFC networks in goal-directed behavior. Disruption of the posterior but not anterior lateral OFC network before the probe test impaired adaptive choices, whereas disruption of the anterior but not posterior lateral OFC network before initial learning similarly impaired subsequent goal-directed choices in the probe test. Additionally, cTBS targeting either region disrupted value acquisition, but only during the first session. Together, these results suggest that anterior and posterior lateral OFC networks play complementary roles for goal-directed behavior, supporting the acquisition and use of outcome-specific stimulus-reward associations, respectively.

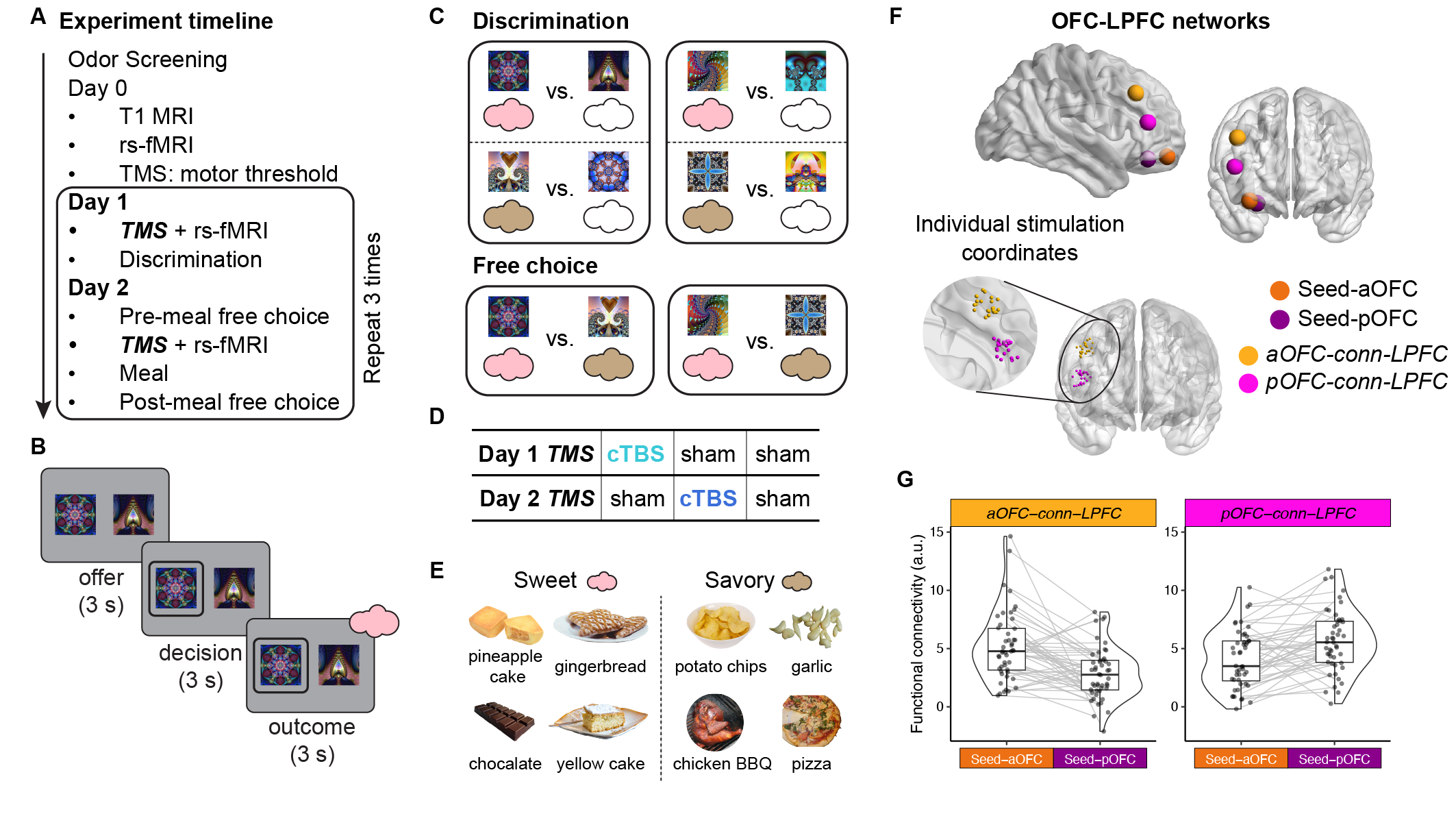
# Results

## Experimental design and outcome devaluation task.

This study follows a within-participant, multiple-session design, with 48 healthy human participants completing a two-day experiment, repeated across three separate sessions (spaced at least one week apart; [1](#fig-design)A). Each session involves the delivery of either cTBS on one day and sham TMS on the other, or sham TMS on both days, resulting in three conditions (Day 1-Day 2: cTBS-sham, sham-cTBS, sham-sham, order counterbalanced; [1](#fig-design)D).

On Day 1, participants learned to discriminate pairs of visual stimuli associated with desirable food odors (sweet or savory, equally valued based on pre-task ratings; [1](#fig-design)E) and clean air. They were asked to select the stimulus associated with any odor, meaning they were not required to encode the specific stimulus-outcome identity associations to perform the discrimination task ([1](#fig-design)B, C). On Day 2, participants chose between stimuli based on odor preferences, making choices between stimuli predicting sweet and savory odors, or between stimuli predicting odor and air. A pre-meal free choice task was followed by a meal, then by a post-meal free choice task. Participants received the odors during the Day 1 discrimination task and the Day 2 pre-meal free choice task. No odors were delivered during Day 2 post-meal free choice task. Participants also reported how much they liked each odor before and after the meal.

To explore the potentially distinct functional roles of OFC subregions in this task, TMS was administered at two different time points—either before the discrimination task on Day 1 or before the meal on Day 2 ([1](#fig-design)A)—and targeted either the anterior (aOFC) or posterior (pOFC) portions of the lateral OFC in different groups of subjects ([1](#fig-design)F). Stimulation targets were defined using MNI coordinates in the right hemisphere: aOFC at [34, 54, -14] and pOFC at [28, 38, -16]. Each target showed strong functional connectivity with isolated clusters in lateral prefrontal cortex (LPFC) ROIs (referred to as aOFC-conn-LPFC and pOFC-conn-LPFC, respectively). Based on resting-state fMRI data collected on Day 0, we individually selected LPFC stimulation sites with the highest connectivity to the respective aOFC or pOFC targets ([1](#fig-design)F). We confirmed the functional separation of these networks across all resting-state fMRI sessions: the aOFC-conn-LPFC showed stronger connectivity with the aOFC than the pOFC (, , Wilcoxon signed rank test, two-sided), and the pOFC-conn-LPFC showed stronger connectivity with the pOFC than the aOFC (, ) ([1](#fig-design)G).



**Experimental design and outcome devaluation task.** **A. Experiment timeline.** Following odor screening, participants completed T1 MRI, resting-state fMRI, and TMS motor threshold determination on Day 0. On Day 1, they received either continuous theta burst stimulation (cTBS) or sham TMS before a discrimination task. On Day 2, they performed a pre-meal free choice task, received TMS (cTBS or sham), consumed a meal, and then completed a post-meal free choice task. **B. Trial structure of discrimination and choice tasks.** Each trial started with an offer phase (3 s), presenting two visual stimuli paired with different outcomes, followed by a decision phase (maximum 3 s) where participants selected one stimulus. In the discrimination task, the trial concluded with an outcome phase (3 s) where participants received an odor or no odor, depending on their choice. **C. Task structure.** In the discrimination task, participants learned which stimuli predicted odors (colored clouds) versus non-odor (i.e., clean air, empty clouds) outcomes. In the free choice task, participants selected stimuli based on learned odor associations and their odor preference, but without immediate odor delivery. The free choice task also included trials comparing odor-predictive and non-odor-predictive stimuli, similar to the discrimination task. **D. TMS conditions.** Participants were assigned to one of three counterbalanced conditions: (1) cTBS on Day 1, sham on Day 2 (cTBS-sham), (2) sham on Day 1, cTBS on Day 2 (sham-cTBS), and (3) sham on both days (sham-sham). **E. Odor stimuli.** Eight food-related odors (savory and sweet). One savory and one sweet odor was selected per participant to match pleasantness ratings. **F. OFC-LPFC networks.** Stimulation coordinates within LPFC for each participant, selected to maximize functional connectivity with either the aOFC (tangerine) or pOFC (magenta) seed region. **G. Functional connectivity estimates.** Half-violin plots depict distribution of connectivity estimates between stimulated LPFC regions and OFC seed regions. Dots represent individual connectivity estimates, and lines indicate within-subject comparison across different ROI combinations.

## Selective satiation affects free choices.

We conducted a proof-of-concept analysis to determine whether choices on Day 2 were influenced by selective satiation, specifically by feeding participants an odor-matched meal. We examined participants’ choices between stimuli predicting sated (SA) and non-sated (NS) odor options in savory-sweet pairs that had not been previously trained.

Participants’ odor pleasantness ratings decreased after the meal across all sessions and participants (, [2](#fig-choices)A). This reduction was unaffected by TMS condition (sham vs. cTBS), TMS target site (aOFC vs. pOFC), session number (1st, 2nd, 3rd), or sated odor type (savory/sweet) (all ; [6](#EDFig_odor)). Importantly, these results suggest that TMS did not impair participants’ ability to update the value of reward outcomes (Izquierdo, Suda, and Murray 2004; Rhodes and Murray 2013; Howard et al. 2020)

When collapsing across sessions, post-meal choices of SA stimuli were significantly reduced relative to pre-meal in both the aOFC (Wilcoxon signed rank test, one-sided, ) and pOFC () groups ([2](#fig-choices)B), confirming an effect of selective satiation on free choices. SA choices were significantly correlated with the pleasantness difference between sated and non-sated odors, both before and after the meal ([7](#EDfig_choices)A, B), indicating that participants made their choices based on relative odor preference, as anticipated. We further calculated the change in pleasantness for both sated and non-sated odors (post-meal minus pre-meal), and then subtracted the change in non-sated odor ratings from that of sated odors. This “selective satiation index” was significantly correlated with the corresponding change in SA choices (Pearson’s , ; [7](#EDfig_choices)C), again supporting the behavioral impact of the change of subjective odor value.

In addition to savory-sweet odor choices, we examined participants’ choices between odors and clean air, which had been associated with outcomes during the Day 1 discrimination task. Participants generally preferred odors over clean air ([2](#fig-choices)C), consistent with successful learning of odor-outcome associations.

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**Selective satiation affects free choices.** **A**. Change of rated odor pleasantness before and after the meal, for sated and non-sated odors. **B.** Choice of sated odors in sweet-savory choices for sham-sham and sham-cTBS conditions, under aOFC-targeted and pOFC-targeted cTBS. **C.** Choice of odors vs. clean air, for sated odors and non-sated odors, pre-meal and post-meal. **D.** Choice of sated odors options with value difference. Dot size represents the number of trials with such value difference.

Additionally, although not part of our original hypothesis—and not typically examined in outcome devaluation studies—we found that individual choices were also influenced by the learned value of each stimulus. We estimated these values based on participants’ behavior during the Day 1 discrimination task (see Section [2.5](#subsec:disc) for details). The probability of choosing the SA over the NS option significantly increased with the value difference between the two stimuli () (Pearson’s , ; [2](#fig-choices)D, [7](#EDfig_choices)D).

Therefore, when evaluating the effects of cTBS (applied on Day 1 or Day 2) on SA choices during Day 2, we included both the learned value difference () and the selective satiation index as regressors to account for factors influencing behavior beyond the effects of TMS.

## Posterior, but not anterior, OFC-targeted cTBS before the free choice impairs outcome devaluation

To examine the role of the aOFC and pOFC in outcome devaluation during the test phase, we focused on the “sham-sham” and “sham-cTBS” TMS conditions. We found a significant interaction between stimulation target (aOFC vs. pOFC) and TMS condition (sham vs. cTBS on Day 2, Day 1 fixed at sham) in predicting SA choices (), according to logistic mixed-effects models on post-meal SA choices, with the session odor preference baseline, satiation status, and the value difference () accounted for. We further separately analyzed the aOFC and pOFC group ([3](#fig_day2)A) and found that cTBS significantly increased SA choices — indicating poorer adaptation to the current goal — only in the pOFC group (), but not in the aOFC group (). Additionally, we confirmed that the effect of pOFC-targeted cTBS on SA choices remained robust regardless of session order ([10](#EDFig_day2)B).

We conducted additional analyses to assess whether the effect of TMS on SA choices was driven by other factors, such as satiation status or perceived TMS discomfort or intensity. The across-participant correlations between pleasantness ratings and SA choices were unchanged by Day 2 cTBS (all ; [7](#EDfig_choices)C), suggesting that the effect of Day 2 cTBS on SA choices was not modulated by satiation status. Moreover, the changes in SA choices induced by cTBS could not be explained by perceived TMS discomfort or intensity, as incorporating TMS ratings into the regression models did not alter any of the findings ([9](#EDFig_corr)).

We also examined choices made between an odor and clean air to see if TMS had any effect on those choices. Following the meal, preference for stimuli predicting sated odors (vs. clean air) decreased, while choices for stimuli predicting non-sated odors (vs. clean air) remained unchanged. This decrease in sated odor selection was significantly stronger after sham stimulation (Wilcoxon signed-rank test, , two-sided) but not after cTBS on Day 2 (; [10](#EDFig_day2)B). SA choices were marginally lower after sham compared to cTBS but only with TMS targeting pOFC () and not aOFC (; [3](#fig_day2)B). These findings align with results from savory-sweet choices, indicating that pOFC-targeted cTBS on Day 2 also impaired choice updating for non-sated odors. This further highlights the critical role of pOFC for adaptive decision-making, even for previously well learned stimuli.

Together, this suggests that pOFC-targeted cTBS before the free choice phase impaired outcome devaluation, as indicated by the continued selection of sated odor-predicting stimuli. In contrast, aOFC-targeted cTBS had no such effect, highlighting the specificity of the pOFC involvement.

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**Posterior, but not anterior, OFC-targeted cTBS before the free choice impaired outcome devaluation**. **A.** Change of choice of sated odors in sweet-savory choices from pre-meal to post-meal test. **B.** Change of preference of sated odors relative to non-sated odors, by comparing odor choices between odor vs. clean air.

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**Anterior, but not posterior, OFC-targeted cTBS on Day 1 impaired subsequent outcome devaluation**. **A**. Probability of sated odor selection after the meal, after adjusting modeled contributions of value difference, selective satiation effects, pre-meal odor preference, compared between sham-sham and cTBS-sham sessions. Overlayed points are the model fitted values of the sated odor selection for each trial collapsing across participants.

## Anterior, but not posterior, OFC targeted cTBS before discrimination learning impaired subsequent outcome devaluation

We explored whether cTBS targeting aOFC and pOFC before learning could affect outcome devaluation measured on Day 2, as would be expected if cTBS disrupted the latent learning of stimulus-reward identity associations during discrimination training.

To assess Day 1 cTBS effect on post-meal choices of sated odors on sweet-savory choices, we focused on “sham-sham” and “cTBS-sham” TMS conditions. For the aOFC group, both TMS condition and session number significantly influenced post-meal sated odor choices, with a significant interaction between the two. Specifically, the cTBS-sham condition significantly increased the selection of sated odors relative to sham-sham ([4](#fig-day1)A; , SE = 0.625, ), and this effect diminished over sessions (, SE = 0.290, ). Choices also increased with session number (, SE = 0.165, ). Additional covariates, including selective satiation index, value difference, and pre-meal odor preference, were significant predictors. Overall, this shows that aOFC-targeted cTBS on Day 1 increased post-meal choices of stimuli predicting sated odors, with the effect moderated by session number.

In contrast, similar analyses in the pOFC group revealed no significant difference between the sham-sham and cTBS-sham stimulation conditions (p=), regardless of whether session numbers were considered as a covariate ([4](#fig-day1)A; all ). However, pre-meal odor preference and value difference were significant predictors of post-meal choices, while the selective satiation index was not (). Additionally, no interaction between stimulation location and TMS condition was identified ().

These findings support our hypothesis that the aOFC plays a critical role in learning the specific stimulus-outcome associations on Day 1, even when the task does not explicitly require it (i.e., latent learning). Notably, this result is independent of the Day 2 TMS, emphasizing the aOFC’s importance in constructing cognitive maps that are later used to guide behavior.

## Posterior and anterior OFC-targeted cTBS disrupted value acquisition during the first session

The discrimination task on Day 1 required participants to select the stimulus associated with desirable food odors (vs. clean air) from a pair of stimuli, reflecting a process of value acquisition. Over five runs, participants significantly improved in selecting odor-predictive stimuli (). This improvement was influenced by the TMS condition applied before the task (cTBS vs. sham; ), session number (1st, 2nd, 3rd session; ), and their interaction (), according to logistic mixed-effects models with participants as a random factor (Line plot and error bar, [5](#fig_disc)A). Response times decreased significantly across runs (), and this decrease was affected by session number (, [11](#EDFig_disc)A) but was not by TMS condition (), according to linear mixed-effects models with participants as a random factor.

To further examine the effects of TMS condition and session number on discrimination behavior, we grouped participants by the order in which they received cTBS or sham on Day 1 ([5](#fig_disc)B). This analysis revealed that the impairment in discrimination due to cTBS was only observed when cTBS was applied during the first session (). We additionally investigated whether the effect of cTBS varied depending on whether cTBS targeted the anterior or posterior OFC but found no evidence for a differential effect (all ).

To quantify and compare the learning process, we fitted a Rescorla-Wagner model to the discrimination behavior using a hierarchical Bayesian approach (Myung, Karabatsos, and Iverson 2005) (see **Supplementary Note** for details). We compared three models: one with condition-specific learning rates, one with session-specific learning rates, and one with fixed learning rates across sessions/conditions. Model comparison showed that the session-specific learning rate model provided the best fit (deviance information criterion (Spiegelhalter et al. 2002); DIC; session-specific learning rates = 13161.95, condition-specific learning rates = 13544.84, fixed learning rates = 14045.46). The winning model well captured the data, as illustrated by the shaded fit overlaid on the experimental data ([5](#fig_disc)A). We examined the estimated learning rates from the winning model and compared them across TMS conditions for each participant group. Wilcoxon signed-rank tests revealed that learning rates were significantly lower after cTBS compared to sham, but only for participants who received cTBS during their first session (; [11](#EDFig_disc)B). We explored if the low learning rates in this group were correlated with perceived TMS discomfort and intensity reported by the participants ([11](#EDFig_disc)C) but found no significant correlation (, ).

Overall, cTBS targeting both posterior and anterior OFC impaired value acquisition in the discrimination task, but only when applied during the first session. This may reflect a general effect of cTBS on participants’ ability to perform the discrimination. As noted in the earlier sections of the results, we included the estimated difference in learned values as regressors when assessing the effects of cTBS (Day 1 or Day 2) on Day 2 choices when estimating cTBS effects on SA choices.

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**Posterior or anterior OFC-targeted cTBS disrupted value acquisition, when applied during the first session.** **A. Discrimination accuracy across runs**. This is plotted by day 1 TMS conditions (cTBS, sham), and session numbers (1st, 2nd, 3rd), separated by different OFC targeted locations (aOFC, pOFC). Line plots and error bars display the experimental data while the shade displays the 95% confidence interval of simulated accuracy using the posterior estimates of learning rates. **B.** Discrimination accuracy across runs, separated by session numbers and the session order of Day 1 TMS.

# Discussion

In this study, we used a three-session times two-day design with network-targeted TMS to selectively modulate activity in anterior and posterior subregions of the human lateral OFC. Using an outcome devaluation task requiring adaptive decision-making based on learned stimulus-outcome identity associations, we found that TMS targeting the pOFC (but not the aOFC) prior to the meal disrupted adaptive behavior, as evidenced by continued choices of stimuli predicting sated rewards in the probe test. Conversely, disrupting the aOFC (but not the pOFC) before learning stimulus-outcome associations on Day 1 also impaired behavior in the probe test on the following day. These findings demonstrate that the aOFC facilitates adaptive decision-making by supporting the acquisition of stimulus-outcome associations, while the pOFC supports their use.

Our findings suggest that the anterior OFC plays a key role in enabling individuals to learn specific stimulus-outcome structures (e.g., associating visual stimuli with specific odors) even when the current task does not explicitly require it. This aligns with prior work indicating that the OFC represents the current task state (Wilson et al. 2014; Vaidya and Badre 2022). However, stimulus-outcome associations in our study were directly observable, contrasting with only partially observable problems where states have to be inferred or retained in memory (e.g. Zhou et al. 2019; Schuck et al. 2016). The function of the anterior OFC in cognitive map construction identified here bears more resemblance with previous research indicating that both humans and animals are driven by curiosity to explore and learn about the environment, known as latent learning (M. Z. Wang and Hayden 2021; Tolman 1948), constructing a representation of the world even in the absence of direct rewards (M. Z. Wang and Hayden 2021; O’keefe and Nadel 1978; Kidd and Hayden 2015). Such cognitive maps, once formed, provide a foundation for guiding goal-directed behaviors (Behrens et al. 2018; Tolman 1948). Importantly, although discrimination training in our task involved reward, learning the specific identity of the reward was not required or reinforced. In that sense, our work parallels a previous study in rats showing that chemogenetic inhibition of lateral OFC caused a deficit in credit assignment during map construction (Costa et al. 2023). Notably, our findings highlight the specific role of the anterior part of the lateral OFC in forming cognitive maps. This work is also in line with recent studies in both rodents and humans suggesting that the lateral OFC supports learning the identity of rewards associated with stimuli (Costa et al. 2023; Howard et al. 2015; Howard and Kahnt 2018; Liu et al. 2024; McDannald et al. 2014; Namboodiri et al. 2019). However, the current study offers a novel and unique contribution by showing that aOFC remains essential even when individuals are not explicitly tasked with encoding such identity information. Moreover, the effect of disrupting learning of identity can be revealed in later stages, when the encoded information becomes crucial for adaptive decision making.

Consistent with previous work (Howard et al. 2020), we found that the posterior part of the lateral OFC is critical for goal-directed behavior during the probe test. Without an intact posterior OFC, individuals fail to change their choices after selective satiation, continuing to choose stimuli predicting devalued outcomes. This suggests that the posterior OFC may support retrieving and applying the cognitive map to guide current behavior. Importantly, our findings show that this effect is specific for the posterior OFC network and does not occur when stimulating a network centered on the adjacent anterior OFC. This provides important information on the specific roles of different OFC subregions and highlights the regional specificity of network-targeted TMS more generally.

Our findings align with a range of studies demonstrating distinct roles of OFC subregions across various tasks and across species, including goal-directed choices with outcome devaluation (Murray et al. 2015), two-choice probabilistic tasks (Stoll and Rudebeck 2024), differential information encoding in the OFC (Rich and Wallis 2017), and the specific contributions of central OFC subregions to economic decision-making (M. Z. Wang, Hayden, and Heilbronner 2022). Particularly relevant is work in non-human primates examining the differential roles of OFC subregions in flexible behavior (Murray et al. 2015), demonstrating that the anterior OFC (area 11) is more involved in goal selection during choice, while the posterior OFC (area 13) primarily supports outcome value updating. In contrast to (Murray et al. 2015), our study focused on the differential involvement of lateral OFC subregions in learning and using stimulus-outcome identity associations to guide adaptive behavior. While a precise cross-species mapping of our defined anterior and posterior OFC regions to animal models remains challenging, our study is, to our knowledge, the first causal investigation to differentiate the functional roles of the human OFC along the anterior-posterior gradient. Recognizing these functional differences represents a substantial advance in our understanding of this brain area and will help guide future studies assessing the role of OFC in learning and decision-making. In human subjects research, this distinction is particularly important for neuroimaging studies and neuromodulation approaches targeting the OFC (Howard and Kahnt 2021; Howard et al. 2020; Liu et al. 2024; F. Wang et al. 2020; Tegelbeckers et al. 2023; Ouellet et al. 2015).

Although not part of our initial hypothesis, we found that cTBS targeting both the anterior and posterior OFC disrupted performance in the discrimination task, but only during the first session, with no impact in later sessions. This challenges the view that OFC is not important for simple Pavlovian acquisition (Murray, O’Doherty, and Schoenbaum 2007; Delamater 2007; Stalnaker et al. 2015), line with recent rodent studies suggesting that OFC’s role in Pavlovian acquisition may be more nuanced than previously thought (Panayi and Killcross 2021). Interpreting this result is further complicated by our within-participant design, as the deficit emerged only in the first session. This initial impairment likely reflects difficulty in grasping the basic task structure. Once this fundamental task structure was learned, it could be reused in subsequent sessions with different stimulus sets (Behrens et al. 2018; Harlow 1949), potentially explaining why TMS had no effect on task performance in later sessions. To account for these effects, we included the stimulus-level learned values of each option in the analysis of SA choices, instead of simply assuming “perfect" learning during the discrimination task (Murray et al. 2015; Howard et al. 2020).

One limitation of this study is the within-participant design, which enhances statistical power but may introduce interpretive challenges. For instance, participants completing the first session could learn that odor identity would be relevant for the Day 2 task, potentially altering their approach to processing odor identity in later sessions. To test this possibility, we compared groups of participants based on the order of cTBS and sham stimulation. Importantly, our findings were not driven by stimulation order, speaking to the robustness of our results. However, the small sample size within each session-order group may limit the ability to detect subtle order effects. Another limitation is the difference in perceived TMS discomfort and intensity between cTBS and sham conditions as reported in the current and our previous work (Liu et al. 2024). However, we found no differences in these ratings between anterior and posterior sites, and individual differences did not account for the observed behavioral effects.

# Conclusion

In conclusion, our study reveals distinct roles of the anterior and posterior OFC network in cognitive map formation and its use for goal-directed behavior in humans. These findings contribute to a better understanding of the functional role of OFC subregions in adaptive decision-making. Additionally, this work offers valuable insights for research in rodents and non-human primates, advancing our understanding of the neural mechanisms underlying adaptive decision-making across species.

# Methods

## Participants

Eighty-eight healthy, right-handed participants (ages 18-40) with no history of psychiatric or neurological disease provided written informed consent to participate in this study. Of these, 48 participants (16 males; ages 18-40, mean = 25.17, SD = 4.14) completed all sessions. Due to a technical error, behavioral data from the cTBS-sham session were unavailable for one participant, but data from the other two sessions were included in the analysis where applicable. MRI data for five resting-state scans were not acquired and excluded from analysis. All participants fasted for at least four hours before each study visit.

## Study design

The study consisted of eight visits ([1](#fig-design)A, D), with Day 1 and Day 2 occurring on consecutive days. The two-day experiment was repeated across three sessions. Sessions were spaced at least one week apart, with a median interval of 13.5 days, a mean of 18.02 days (SD = 9.09), and a range of 7 to 63 days. On each Day 1 and Day 2, participants received either continuous theta-burst stimulation (cTBS, labeled C) or sham stimulation (S). Over the three sessions, they experienced three TMS conditions: cTBS-sham (CS), sham-cTBS (SC), and sham-sham (SS). The order of these conditions was counterbalanced, with 9 participants receiving CS-SC-SS, 7 receiving CS-SS-SC, and the remaining 32 equally assigned to one of the other four possible sequences (SC-CS-SS, SC-SS-CS, SS-CS-SC, and SS-SC-CS).

To prevent differences in stimulation location from affecting participants’ experience across sessions, each participant received TMS targeting either the anterior or posterior portion of the lateral OFC throughout all three sessions. Among the participants, 16 of 32 females and 9 of 16 males received TMS targeted to the posterior portion. Additionally, the order of satiation conditions was counterbalanced: half of the participants received a sweet meal in their first session, while the other half received a savory meal. The sated odor type alternated for each participant across the three sessions (e.g., savory-sweet-savory or sweet-savory-sweet).

## Screening session

After providing informed consent and completing eligibility screening, participants rated the pleasantness of eight food odors. These odors, supplied by International Flavors and Fragrances (New York, NY), included four savory (garlic, potato chip, pizza, barbecue) and four sweet (chocolate, yellow cake, pineapple cake, gingerbread) odors. In each trial, participants smelled a food odor for 2 seconds and rated their liking on a visual analog scale ranging from “Most Disliked Sensation Imaginable” to “Most Liked Sensation Imaginable.” Ratings were made using a scroll wheel and keyboard press. Each odor was presented three times in a pseudo-randomized order, and ratings were averaged per odor. Based on these ratings, two odors (one savory, one sweet) that were pleasant (above neutral) and closely matched were selected for the discrimination and choice tasks. These odors were used across all three sessions. Participants were excluded if no suitable odors were identified.

A custom-built, computer-controlled olfactometer was used to deliver the odors with precise timing to nasal masks worn by participants. The olfactometer directed medical-grade air through the headspace of amber bottles containing the odor solutions at a constant flow rate of 3.2L/min. Using two independent mass flow controllers (Alicat, Tucson, AZ), the device enabled precise dilution of the odorized air with odorless air. Throughout the experiment, a constant stream of odorless air was delivered, and odorized air was mixed in at specific time points without altering the overall flow rate or causing somatosensory stimulation.

## Day 0: Scan & Motor threshold

We acquired a T1-weighted structural MRI scan to assist with TMS neuronavigation and an 8 min multi-echo resting-state fMRI scan (310 volumes, TR = 1.5s) to individually define the OFC-targeted cTBS coordinates (see section [5.8](#coordination-selection)). The same scanning parameters were used for other resting-state scans. We then measured resting motor threshold (rMT) by administering single TMS pulses to the hand area of the left motor cortex. rMT was defined as the lowest stimulator output required to evoke 5 visible thumb movements from 10 pulses.

## Day 1: Discrimination task

Participants first underwent a TMS session (cTBS or sham, see section [5.9](#tms)) followed by a resting-state scan. Then they completed five runs of a discrimination task. In each trial, participants chose between two fractal stimuli: one associated with a savory or sweet odor, and the other with clean air. Stimuli were displayed for 3 seconds, followed by a choice phase (maximum 3 seconds). If participants selected a stimulus leading to an odor, the odor was delivered for 2 seconds. The inter-trial interval ranged from 4 to 8 seconds. Each run consisted of 24 trials, using four groups of stimulus pairs: two sets (A and B) crossed with sweet/savory odors. Each combination had three non-overlapping stimulus pairs, resulting in 24 distinct fractals. Each pair was presented twice to counterbalance left and right positions on the screen. Choice and response times were recorded for each trial, and different fractals were used across the three sessions.

## Day 2: Meal consumption and free choice task

Day 2 started with an odor pleasantness rating followed by a choice task (pre-meal) where participants selected between pairs of stimuli. Afterwards, participants underwent a TMS session and then had a meal carefully matched in flavor to either the sweet or savory food odor used in their task. Following the meal, participants completed another set of odor pleasantness ratings and the post-meal free choice task. Both pre-meal and post-meal choice tasks instructed participants to choose based on their current odor preferences.

The pre-meal free choice task included 30 trials, all from set A, consisting of 3 sweet vs. clean air pairs, 3 savory vs. clean air pairs, and 9 savory vs. sweet pairs. Each pair was presented twice to counterbalance left and right positions on the screen. The post-meal choice task included 60 trials from both sets A and B. In both pre- and post-meal choice tasks, similar to the discrimination task, every trial began with a pair of stimuli presented for 3 seconds, followed by a decision phase of up to 3 seconds. In the pre-meal free choice task, if participants selected a stimulus linked to an odor, the odor was delivered for 2 seconds after their choices. No odors were delivered during the post-meal free choice task participants received the odors chosen in five randomly selected trials at the end of the task. The inter-trial interval ranged from 4 to 8 seconds, and choice and response times were recorded from all trials. Pre- and post-meal free choices for both set A and set B stimuli were highly correlated ([8](#EDFig_sets)), indicating consistent choices across sets based on odor preferences. Thus, to assess the satiation effect on choices, we used the pre-meal average choice from set A as a session-wise odor preference baseline and compared it with the post-meal choices.

## MRI data acquisition

MRI data were acquired on a Siemens 3T PRISMA system equipped with a 64-channel head-neck coil. Each TMS session on Day 1 and Day 2 was immediately followed by a resting-state MRI scan. Resting-state fMRI data were collected across all seven sessions with the same multi-echo sequence (310 volumes; TR = 1.5s; TE1-TE3 = 14.60ms, 39.04ms, 63.48ms). The short TE of the first echo is beneficial to mitigate signal dropout near the OFC, as demonstrated in previous studies using both resting-state and task-based fMRI (Fernandez et al. 2017; Poser et al. 2006; Kirilina et al. 2016; Zhao et al. 2024). Other scanning parameters included: flip angle, 72°, slice thickness, 2mm (no gap), multi-band acceleration factor 4, 60 slices with interleaved acquisition, matrix size 104 x 104 voxels, and field of view 208mm x 208mm. A 1mm isotropic T1-weighted structural scan was acquired on Day 0 session for neuronavigation during TMS and to aid spatial normalization.

## Coordination selection for network-targeted TMS

The stimulation coordinates were computed based on the multi-echo resting-state MRI data collected from the Day 0 session. We defined our stimulation targets in the right hemisphere’s aOFC and pOFC using MNI coordinates: aOFC [34, 54, -14] and pOFC [28, 38, -16]. The pOFC coordinates were identical to those used in our previous network-targeted TMS studies (Howard et al. 2020; Liu et al. 2024; F. Wang et al. 2020; Tegelbeckers et al. 2023). Each targeted coordinate in the aOFC and pOFC exhibited strong functional connectivity with isolated clusters in the LPFC with peak coordinates of [44, 28, 38] and [46, 38, 14], respectively, as determined in data from Neurosynth.org involving a sample of 1,000 subjects.

We first generated spherical masks of 8-mm radius around these four coordinates in MNI space, each inclusively masked by the gray matter tissue probability map provided by SPM12 (thresholded at > 0.1). We then transformed these four masks to each subject’s native space using the inverse deformation field generated during the normalization of the T1 anatomical image. We then specified two resting-state fMRI functional connectivity analyses (one per region) for each subject, using individual aOFC and pOFC masks as the seed regions and motion parameters from the realignment of the first echo as regressors of no interest. Finally, stimulation coordinates were defined as the voxels within the right LPFC masks with the strongest functional connectivity to the right aOFC and pOFC seed regions, respectively. We used infrared MRI-guided stereotactic neuronavigation (LOCALITE) to apply stimulation to these two individual LPFC coordinates.

## Transcranial magnetic stimulation

Similar to our previous work, the target coordinates were defined as the locations in the right LPFC with the strongest functional connectivity with the corresponding right OFC seed regions (see details above). The Figure-eight coil was tilted so that its long axis was approximately perpendicular to the long axis of the middle frontal gyrus. TMS was administered at 80% of the rMT using a cTBS protocol. This protocol involved delivering bursts of three pulses at 50 Hz every 200 ms (5 Hz) for a total of 600 pulses over approximately 40 seconds. Stimulation was applied using a MagPro X100 stimulator equipped with a MagPro Cool-B65 A/P butterfly coil (MagVenture). Previous work has demonstrated that this cTBS protocol at 80% MT has inhibitory aftereffects which persist for 50–60 min over primary motor cortex (Huang et al. 2005). Whereas cTBS was delivered by positioning the active side of the A/P coil to modulate neural tissue, sham cTBS was applied with the placebo side of the A/P coil, producing similar somatosensory and auditory experiences for the participant without modulating neural tissue. Electrodes were placed on participants’ forehead and direct current stimulation was applied in synchrony with the TMS pulses to mask TMS effects and enhance the similarity between cTBS and sham sessions.

Participants were informed about potential muscle twitches in the face, eyes, and jaw during simulation. To assess tolerability, two single pulses were applied over the stimulation coordinates before administering cTBS. Discomfort and perceived stimulation intensity were evaluated after each TMS session. The cTBS sessions was generally rated as more uncomfortable and intense compared to the sham sessions. On a scale from 0 (not uncomfortable at all) to 10 (extremely uncomfortable), mean discomfort ratings were 3.38 for sham and 5.8 for cTBS sessions (, linear mixed effects model). Similarly, on a scale from 0 (not strong at all) to 10 (extremely strong), mean intensity ratings were 3.79 for sham and 6.23 for cTBS sessions (, linear mixed effects model). Discomfort and intensity ratings did not differ between aOFC- or pOFC-targeted cTBS (all 0.6). For analyses involving cTBS effects (Day 1 or Day 2 TMS), standardized discomfort and intensity ratings were used to examine correlations or regressions against other variables, assessing if the observed cTBS effects were driven by subjective discomfort or perceived TMS intensity, but none of the effects can be explained by those ratings (see [9](#EDFig_corr)).

## Meal consumption

On Day 2, participants consumed a meal following the TMS session to selectively satiate one of the two food odors. The meal items were carefully chosen to closely match the corresponding food odors, and water was provided. Participants were instructed to eat until they felt very full and were then left alone for 15 minutes. Immediately afterward, they rated the pleasantness of the odors and proceeded to the post-meal choice task. On average, participants consumed 669.89 ± 44.16 calories (SEM). Before analyzing the relationship between odor ratings and task behavior, we standardized the ratings within each participant across sessions.

## Modeling value learning

We used a standard Rescorla-Wagner model (Rizley and Rescorla 1972) to describe learning in the discrimination task, where participants chose between two stimuli—one predicting an odor and the other leading to clean air. Since stimulus pairs had no overlap, we assumed that learning was primarily driven by the odor-predictive stimulus rather than the stimulus associated with clean air. Accordingly, we modeled the learned value of the odor-predictive stimulus across trials.

The model updated of the odor-predictive stimulus based on prediction error, defined as the difference between the actual outcome () and the expected value on each trial. The learning rate determined how quickly adjusted across trials. Initially, was set to 0.5, with indicating complete learning of the odor-predictive stimulus. We estimated a separate learning rate for each odor-predictive stimulus, with priors constrained by session-wise or condition-wise hyper parameters in a hierarchical Bayesian framework (Myung, Karabatsos, and Iverson 2005). This approach allowed us to obtain learned value estimates for each odor-predictive stimulus, which were then used to aid the analysis of the free-choice task data. The session-wise hyper learning rate parameters are correlated with TMS ratings in [11](#EDFig_disc). Details of the model specification and estimation are provided in **Supplementary Text 1.**

## Multi-echo MRI data processing

Preprocessing of the multi-echo resting-state fMRI data involved several steps. First, all functional images from the smallest echo across all rs-fMRI runs were realigned to the first volume of the first echo, and the resulting voxel-to-world mapping matrix was applied to the other two echoes, volume by volume. All functional images were then resliced for each echo. Next, the images across the three echoes were combined using temporal signal-to-noise ratio (tSNR) weighting, following parallel-acquired inhomogeneity desensitized (PAID) approach (Poser et al. 2006). Specifically, voxel-wise tSNR maps were computed for each echo, multiplied by the echo time (TE), and normalized across the three echoes to generate weight maps. These weight maps were then used to combine the resliced images by multiplying each volume by its respective weight map. Lastly, the combined data underwent coregistration, normalization, and smoothing using a 6 mm FWHM Gaussian kernel.

We analyzed participants’ motion during the resting-state scan after different types of TMS (sham vs. cTBS) and stimulation targeted locations (anterior vs. posterior OFC). Framewise displacement (FD) was calculated per volume and summed across volumes (Power et al. 2012). No significant differences were observed between TMS types or stimulation locations (all 0.8). FD for cTBS was 38.3mm (±10.8mm) at the anterior OFC and 41.3mm (±17.8mm) at the posterior OFC, while for sham, FD was 41.0mm (±16.7mm) at the anterior OFC and 39.6mm (±15.8mm) at the posterior OFC.

Analysis of Behavioral Data

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# Extended Data

![](data:application/pdf;base64,)

**Extended Data Fig.** : **Supplementary results on odor pleasantness ratings.** **A.** Odor pleasantness ratings separated by TMS conditions and stimulation locations. **B.** Odor pleasantness ratings separated by session numbers and stimulation locations.

![](data:application/pdf;base64,)

**Extended Data Fig.** : **Free choices are influenced by learned stimulus values and selective satiation effects.** **A.** Scatter plots showing correlations between the choice of stimuli predicting sated odors and odor pleasantness ratings of sated minus non-sated odors before (left) and after the meal (right), separated by the three TMS conditions. **B.** Scatter plot showing the change in SA choices against the change of odor pleasantness difference (sated minus non-sated) after eating the meal. **C.** Choice of sated odors options associated with each of the learned weight of the combination of sated and non-sated options. Dot size represents the number of trials per value combination (log scale), with missing dots indicating unobserved combinations.

![](data:application/pdf;base64,)

**Extended Data Fig.** : **Scatter plots showing correlations of the choice for selecting sated odors across post-meal, pre-meal, set A and set B.** **A.** Relationship between pre-meal and post-meal of set A. **B.** Relationship between pre-meal and post-meal of set B. **C.** Relationship between pre-meal set A and post-meal of set B.

![](data:application/pdf;base64,)

**Extended Data Fig.** : **Relationship between perceived TMS discomfort and intensity and sated odor (SA) choices.** **A.** Correlation between SA choices and TMS ratings, separated by Day 2 TMS conditions (sham-cTBS vs. sham-sham) and TMS targeted regions (aOFC, pOFC). A positive correlation was observed between TMS ratings and SA choices in the aOFC group, but including ratings of TMS perception into the regression models did not alter the observed TMS effects on SA choices. **B.** Same as **A**, but focus on Day 1 TMS effect (sham-sham vs. cTBS-sham). **C.** Scatter plot showing the relationship between the condition-wise difference (sham-cTBS vs. sham-sham) of SA choices and condition-wise difference of TMS ratings from Day 2 TMS. There was a significant positive correlation in the aOFC group (Pearson’s , ) **D.** Same as **B**, but focus on Day 1 TMS effect (sham-sham vs. cTBS-sham). Shaded areas represent 95% confidence intervals estimated using robust linear regression. Marginal distributions are shown on the top and right axes. Pearson correlation coefficients (R) and p-values are reported for each TMS condition.

![](data:application/pdf;base64,)

**Extended Data Fig.** : **Supplementary results of Day 2 cTBS effect.** **A.** Choice of sated odors for participants experiencing different Day 2 TMS orders within each stimulation location group (aOFC and pOFC). **B.** Change in the choice of odors during odor-air choices, separated by sham-cTBS and sham-sham TMS conditions and sated/non-sated odors. **C.** Correlation of the baseline odor preference between derived from savory-sweet choices and from odor-air choices, for each of TMS condition.

![](data:application/pdf;base64,)

**Extended Data Fig.** : **Supplementary results on cTBS effect on discrimination learning.** **A.** Change of response times across runs as a function of TMS (top) and session number (bottom). **B.** Effect of cTBS on estimated learning rates, separated by Day 1 TMS order. **C.** Relationship between estimated learning rates and perceived TMS discomfort/intensity, separated by Day 1 TMS order.

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