

Curiosity Satisfaction Increases Event-related Potentials Sensitive to Reward

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Abstract

■ Successful learning depends on various factors such as depth of processing, motivation, or curiosity about information. A strong drive to learn something or the expectation of receiving a reward can be crucial to enhance learning. However, the influence of curiosity on the processing of new information and its similarity with reward processing is not well understood. This study examined whether states of curiosity influence specific ERPs associated with reward processing and whether these ERPs are related with later memory benefits. In an initial screening phase, participants indicated their curiosity and confidence in prior knowledge about answers to various trivia questions. In a subsequent study phase, we targeted different time windows related to reward processing during the

presentation of trivia answers containing the reward positivity (RewP; 250–350 msec), the P3 (250–500 msec), and the late-positive-potential (LPP; 600–1000 msec). In a following surprise memory test, we found that participants recalled more high- than low-curiosity answers. The RewP, P3, and LPP showed greater positive mean amplitudes for high compared with low curiosity, reflecting increased reward processing. In addition, we found that the RewP and the P3 showed more positive mean amplitudes for later recalled compared with later forgotten answers, but curiosity did not modulate this encoding-related results. These findings support the view that the satisfaction of curiosity resembles reward processing, indicated by ERPs.

INTRODUCTION

Why did you start reading this article? Maybe because you stumbled across it by accident, maybe because you have to, or because you are curious about its content. Curiosity is an intrinsic motivation that drives us to collect new information and has been shown to enhance memory formation (Gruber & Ranganath, 2019; Galli et al., 2018; Kidd & Hayden, 2015; Loewenstein, 1994). For example, if someone asks you "What is the longest river in the European Union?" you might be curious and try to come up with the correct answer. In the last decade, initial research on states of curiosity have targeted the neural underpinnings of the elicitation of curiosity (e.g., when you read a question without knowing the answer; Poh et al., 2022; Lau, Ozon, Kuratomi, Komiya, & Murayama, 2020; Oosterwijk, Snoek, Tekoppele, Engelbert, & Scholte, 2020; Ligneul, Mermillod, & Morisseau, 2018; Gruber, Gelman, & Ranganath, 2014; Kang et al., 2009), but the literature still lacks findings as to what specific processes take place during the satisfaction of curiosity (e.g., when you receive the answer "Danube").

Most concepts about states of curiosity have in common that the awareness of a knowledge gap leads to increased exploration for new information (Metcalfe, Schwartz, & Eich, 2020; Gruber & Ranganath, 2019; Loewenstein,

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1994; Berlyne, 1954). New information satisfies curiosity, reduces uncertainty, can improve predictions in the future, and can be described as an unconditioned reward stimulus (Kang et al., 2009). Therefore, curiosity reflects an intrinsically anticipated expectation of reward that intensifies the processing of upcoming information (Gruber & Ranganath, 2019; Marvin & Shohamy, 2016; Kidd & Hayden, 2015; Loewenstein, 1994). Investigating ERPs during the satisfaction of curiosity may possibly explain how curiosity influences the processing of information, what similarities exist regarding to reward processing, and how curiosity satisfaction modulates memory formation.

There is evidence showing that the neural correlates of curiosity resemble those of reward anticipation (Lau et al., 2020; Oosterwijk et al., 2020; Gruber et al., 2014) and reward processing (Ligneul et al., 2018; Jepma, Verdonschot, van Steenbergen, Rombouts, & Nieuwenhuis, 2012). For example, examining curiosity satisfaction with ambiguous visual images, Jepma and colleagues (2012) showed that the resolution of the images was associated with increased activity of the striatum, which has been related to reward processing. Furthermore, investigating effects of intrinsic (curious) and extrinsic (rewarding) motivation on memory, Duan, Fernández, van Dongen, and Kohn (2020) found that curiosity-driven memory formation was associated with the ventral striatal reward network and the frontoparietal attention system, whereas extrinsic-driven

memory effects were correlated with deactivation in parietal midline regions. Therefore, it is not clear whether reward and curiosity are really the same processes. To investigate this question, we therefore assumed that curiosity satisfaction should be similar with reward processing and predicted that reward-sensitive ERPs should also be modulated by states of curiosity.

As the answer to a question can be seen as an unconditioned reward stimulus, we chose to investigate the reward positivity (RewP; 250–350 msec; Proudfit, 2015), also known as feedback-related-negativity (FRN; Höltje & Mecklinger, 2020; Peterburs, Kobza, & Bellebaum, 2016; Cohen & Ranganath, 2007). According to Proudfit (2015, p. 449), the FRN "reflects a reward-related positivity that is absent or suppressed following non-reward." Because in this study we assume that the answer represents a reward, we favor the interpretation as RewP. This component is assumed to reflect a better outcome than expected, an increase in dopaminergic signals, and reward processing at fronto-central sites (Glazer, Kelley, Pornpattananangkul, Mittal, & Nusslock, 2018; Heydari & Holroyd, 2016; Proudfit, 2015; Sambrook & Goslin, 2015). In addition, the RewP seems to be correlated with neural activity in reward- mediating regions such as the ventral striatum, anterior cingulate cortex, and medial prefrontal cortex (mPFC; Becker, Nitsch, Miltner, & Straube, 2014). We hypothesized that when curiosity is high, a new information should be perceived as more relevant, reflecting a greater reward, and should therefore result in a larger mean amplitude of the RewP.

After early reward processing reflected by the RewP, the P3 (P300/FB-P3; further: P3; 250-500 msec; Polich, 2007) could be also manipulated because of high- and lowcuriosity levels. The P3 is usually found at central and parietal electrodes and assumed to reflect the allocation of neural resources based on reward effects, stimulus relevance, as well as context updating in working memory (San Martín, 2012; Van Petten & Luka, 2012). In addition, it displays attention-driven categorization and increased processing of new information because of motivational salience. The P3 amplitude is larger if new information is infrequent or a reward is greater than expected (Hajcak, Moser, Holroyd, & Simons, 2006). For high curiosity, it is assumed that there should be a greater mean amplitude of the P3 during curiosity satisfaction because of higher motivational relevance for the updating of the stimulus context, increased attention, and a subjective greater reward than for low curiosity (Donaldson, Oumeziane, & Foti, 2016; Polich, 2007).

The late-positive-potential (LPP; 600–1000 msec; Glazer et al., 2018) is a centro-parietal positive-going ERP component that has been associated with the processing of emotional stimuli as well as extended cognitive and attentional processing based on reward expectancy and magnitude (Hajcak & Foti, 2020; Glazer et al., 2018; Meadows, Gable, Lohse, & Miller, 2016; Gable, Adams, & Proudfit, 2015; Weinberg, Ferri, & Hajcak, 2013; Schupp et al., 2000).

Furthermore, the LPP appears to be manipulated by the significance of a stimulus, which is determined by the activation of motivational appetitive systems (Bradley, 2009). On the basis of these findings, we expect the mean amplitude of the LPP to be more positive for high-than for low-curiosity answers.

To generate states of high and low curiosity, we used the so-called trivia paradigm and adopted an identical experimental design as in a previous fMRI study from our laboratory (Gruber et al., 2014). In an initial screening phase, participants were presented with general knowledge questions and were asked to indicate their curiosity and confidence in prior knowledge about the answer. In the following study phase, participants were presented with a selected set of trivia questions from the screening phase and a few seconds later with the associated answers (i.e., the satisfaction of curiosity). ERPs were computed timelocked to the onset of the answers. After a 5-min break, participants were asked to recall the answers in a surprise memory test.

On the basis of previous findings (e.g., Poh et al., 2022; Wade & Kidd, 2019; Marvin & Shohamy, 2016; Gruber et al., 2014; Kang et al., 2009), we predicted memory performance to be better for answers associated with high compared with low curiosity. Furthermore, we expected that ERPs recorded during the presentation of the answer should differentiate between remembered and forgotten ones. To address this question, we used the memory test results to investigate subsequent memory effects (SMEs), for which the ERPs are sorted according to whether the answers are later remembered or forgotten. In general, subsequently remembered items should show a more positive ERP amplitude during encoding than those that have been forgotten (for a review, see Mecklinger & Kamp, 2023; Cohen et al., 2015). Accordingly, we predicted more positive amplitudes for remembered answers. Furthermore, we also explored the interaction between curiosity and memory in relation to the ERP amplitudes.

METHODS

Participants

Thirty healthy young adults participated in the experiment. We required a sample size of n=24, which was determined with a power analysis for a repeated-measures ANOVA (G*Power, Version 3.1.9.7; Faul, Erdfelder, Buchner, & Lang, 2009), that is, the ERP amplitude difference between high and low curiosity, based on the assumption of a medium effect size according to Cohen's f (1988), f=0.25, $\alpha=.05$, $1-\beta=0.8$. Eight participants had to be excluded because of either poor memory performance (in at least one condition fewer than eight memorized answers) or incomplete EEG or memory data. Because our final data set was slightly below the targeted sample size, we additionally calculated the Bayes factor for all nonsignificant results. This

tests whether potential null results are because of inconclusive data or whether there is actual support for a null hypothesis (Dienes, 2014). The final sample included n=22 participants (11 women, 11 men, age range = 18–30 years, M=21.23 years). One participant was left-handed, and twenty-one were right-handed. All participants had normal or corrected-to-normal vision and were English native speakers. All participants were students at University California (UC) Davis and received money or course credits for their attendance. The UC Davis institutional review ethics committee approved the experiment.

Material

The questions and their matching answers for the screening and study phase were randomly drawn from a pool of 381 items (https://osf.io/he6t9/), which in turn were acquired from online resources about general knowledge questions. In addition, care was taken to select from as many different subject areas as possible (i.e., sports, food, science, nature, TV/movies, music, history). The stimuli were always presented centered in black on a gray background.

Procedure

The experiment was programmed with the Cogent 2000 Toolbox (Wellcome Laboratory of Neurobiology) and carried out in the Center for Neuroscience at UC Davis. The experiment is a version of the trivia paradigm (cf. Kang et al., 2009) and was divided into a screening, study, and

test phase (for an almost identical task procedure, see Gruber et al., 2014; for an overview of the procedure, see Figure 1).

In the screening phase, randomly chosen questions were presented for 6 sec and evaluated by participants according to their knowledge confidence and perceived level of curiosity. This intra-individual rating of the questions was necessary because curiosity levels for each trivia question and its respective answer differ between individuals. First, participants were asked to indicate how confident they were that they knew the answer to the trivia question (1-6; 1 ="I am confident that I do not know the answer" and 6 = "I am confident that I know the answer"). As a second judgment, participants indicated their level of curiosity about the answer (1-6; 1 = "I am not interested at all in the]answer" and 6 ="I am very much interested in the answer"). Questions were presented until 56 items could be attributed to high (curiosity rating 4-6) and 56 to low curiosity (curiosity rating 1–3). However, if the participants had given a "6" for the confidence rating, the question was not included, because it was assumed that participants already knew the answer. While participants were completing the screening phase, experimenters were attaching the EEG cap to the participant and were preparing all electrodes to record the EEG. The screening phase was followed by a short rest period (approx. 5 min) during which participants looked at a fixation cross.

In the study phase, 112 previously selected questions were presented again (3 sec) and after an anticipation period (11 sec; the time between question presentation

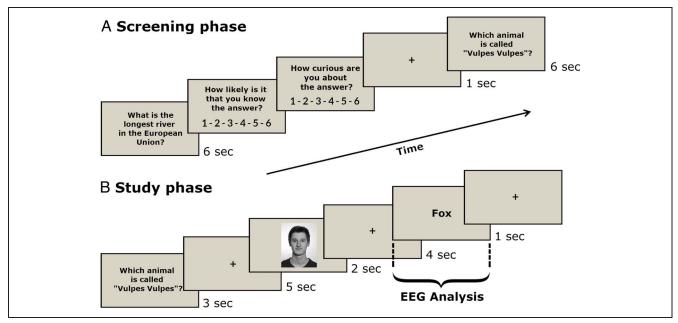


Figure 1. Experimental procedure. (A) In the screening phase, trivia questions were presented and evaluated by participants according to their knowledge confidence and perceived level of curiosity without reading the corresponding answers. (B) In the following study phase, participants are presented with a subset of these questions again and 11 sec later with their corresponding answers. Between questions and answers, emotionally neutral faces were presented. The instruction of the participants was to "always try to anticipate the possible answer and give a response ("YES" or 'NO") whether the person can help you." The EEG was recorded during the study phase, and ERPs were computed time-locked to the onset of the answer. (C) At last, participants were asked to recall all answers in a surprise memory test (not shown).

offset until answer presentation onset), the matching answers were displayed (1 sec). Six trials in each condition (~10%) were catch trials, during which a sequence of letters "xxxxx" were shown as answer, to keep participants' attention high. To create consistency to our prior work using the identical paradigm in a fMRI study (see Gruber et al., 2014), emotionally neutral faces (2 sec) were presented during the anticipation period. The instruction of the participants was to "always try to anticipate the possible answer and give a response ('YES' or 'NO') whether the person can help you." Because our article focuses on the satisfaction of curiosity (i.e., during answer presentation) and thus on specific ERPs that previously have been shown to be sensitive to reward processing, the analysis of the faces during the anticipation of curiosity were not part of the present study.

After a further 5-min rest period, during which participants looked at a fixation cross, a surprise memory test was carried out in the test phase. Participants were given a randomized list with all trivia questions from the study phase. They were encouraged to type in as many answers as possible within 20 min without guessing the answers. Participants also took part in a recognition memory test for all previously encoded face images, which are not part of this investigation (see Gruber et al., 2014, for face analysis). Each participant took part in each phase once and received money or course credits after the test phase.

EEG Recording and Processing

During the study phase, the EEG was recorded with the ActiveTwo EEG recording system (Biosemi) at 1024 Hz from 64 Ag/AgCl scalp electrodes, which were arranged according to an extended version of the International 10–20 electrode system (Jasper, 1958). The electrodes were offline rereferenced to the averaged mastoid electrodes. The vertical and horizontal EOG was recorded from four electrodes placed above and below the left eye and at the canthi of the left and right eye.

The analysis of the EEG data was performed using MATLAB (Version 7.10.0; Mathworks, Inc.) and the toolboxes EEGLAB (Delorme & Makeig, 2004) and ERPLAB (Lopez-Calderon & Luck, 2014). Electrodes were referenced to the average of the left and right mastoid electrodes. EEG data were downsampled to 500 Hz and bandpass filtered at 0.1-40 Hz using a second order Butterworth filter. The epochs started at -200 msec before stimulus presentation and ended at 1000 msec thereafter. In addition, data segments outside of the area of interest were discarded and an independent component analysis was applied to correct for artifacts. Segments that were still associated with ocular and noise artifacts were rejected manually for each participant. In addition, segments containing artifacts were removed based on the following criteria: The maximum voltage threshold was set between -150 and $+150 \mu V$, the maximal permissible difference of values at 150 µV during intervals of 200 msec, and the

maximum allowed voltage difference between two time points was 30 μ V. To exclude data with no signal, trials that did not exceed the limits -0.7 and +0.7 μ V within a period of 200 msec were discarded. On average, 83.93% of trials were retained (M=94, trial range = 40–110, high curiosity = 46.89, low curiosity = 47.11) and 16.07% were rejected (M=18, range = 2–72, high curiosity = 9.11, low curiosity = 8.89). These cleaned and prefiltered data were pruned and merged based on curiosity level and memory performance. We calculated the mean amplitudes for the components of interest over all trials for each participant.

The RewP is typically measured at fronto-central sites and thus activity at electrodes FC1, FCz, and FC2 in the time range of 250–350 msec was investigated (Proudfit, 2015). The P3 is usually found at central and parietal electrodes, which is why activity at electrodes Cz, CPz, and Pz between 250–500 msec was analyzed (Polich, 2007). The LPP is most pronounced at centro-parietal electrodes, and hence, activity at electrodes CP1, CPz, and CP2 in the time range of 600–1000 msec after stimulus onset was explored (Glazer et al., 2018; Gable et al., 2015). The time window of the LPP can also be broader, but our time window was limited to 1000 msec.

Statistical Analyses

The statistical analyses were performed using R Studio software (R Core Team, 2019). The significance level was set to $\alpha = .05$, and t tests were one-tailed based on specific hypotheses. For the behavioral analyses, we used dependent t tests and regression models with curiosity and confidence as predictors of memory. The ERP data were analyzed using 2 (Curiosity: high, low) \times 2 (Memory: remembered, forgotten) repeated-measures ANOVAs, with the values averaged across the electrodes. To address the question of sufficient evidence for nonsignificant results, we additionally computed the Bayes factor (Dienes, 2014). For interpretation, we used the widespread labels by Jeffreys (1961): $BF_{10} < 0.3$ as substantial evidence for H_0 and $BF_{10} > 3$ as substantial evidence for H_1 . As measures of effect sizes, partial eta squared (η_p^2) is reported for ANOVA and Cohen's d (Rasch, Friese, Hofmann, & Naumann, 2021; Cohen, 1988) was calculated for dependent t tests.

RESULTS

Behavioral Results

The analysis of memory performance showed a significant mean recall difference between high- (M=53.32%, SD=15.57%) and low-curiosity answers (M=37.15%, SD=16.19%), t(21)=7.37, p<.001, d=1.57, which replicates the previously found high-curiosity related memory benefit (Gruber et al., 2014; Kang et al., 2009). The overall memory performance was M=42.00% (lowest = 16.96%; highest = 67.86%), and the curiosity benefit was M=16.17% (high-curiosity answers – low-curiosity

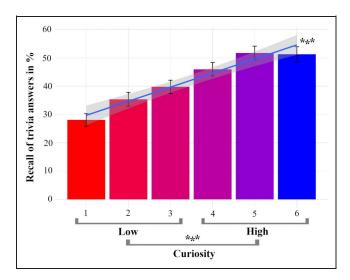


Figure 2. Curiosity-related memory benefit. Better memory performance for trivia answers in the high-compared with the low-curiosity condition. Error bars indicate the standard error of the mean. The line represents a linear model between curiosity and memory. *p < .05, **p < .01, ***p < .001.

answers). In addition, a follow-up behavioral analysis using the 6-scale curiosity ratings to predict memory was conducted to check whether a linear relationship exists between both variables and thus the classification to high and low curiosity can be supported. The regression model shows that as curiosity increases, memory performance also increases continuously ($\beta=0.21, SE=0.03, z=8.26, p<.001$). Accordingly, analyzing curiosity with a two-level factor (high and low) is an adequate approach for the ERP analyses (see Figure 2).

Moreover, calculating the confidence ratings for high (M=2.26,SD=.45) and low curiosity (M=1.50,SD=.34) revealed a significant difference, t(21)=7.64,p<.001,d=1.63, indicating that participants were more confident they knew the answers in relation to high curiosity. A correlation between curiosity and confidence was significant, with r=.35,p<.001, showing that 12.25% of variance caused by curiosity can be explained by confidence. These analyses show that curiosity and confidence in prior knowledge are partly related, but most of their variance exists independently.

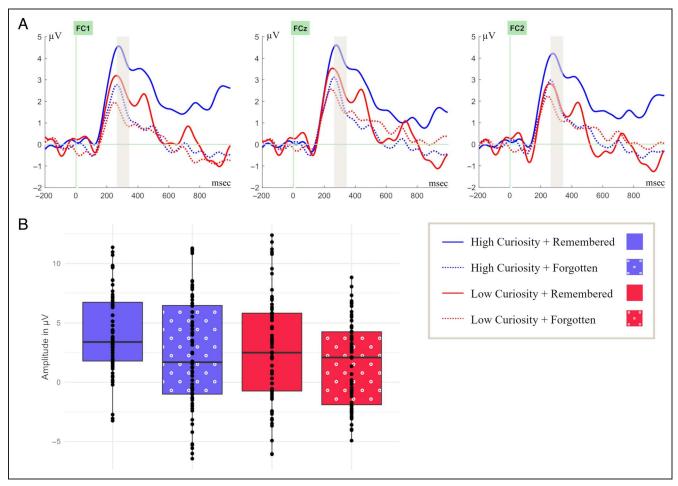


Figure 3. Curiosity and memory independently modulate the RewP component during answer presentation at fronto-central electrode sites. (A) ERPs were time-locked to the onset of the answers in the study phase at electrode sites FC1, FCz, and FC2 in the colored time window of 250–350 msec. (B) Box plots are bounded by the first and third quartiles, and the black line represents the median. The points correspond to the measures for individual participants.

Most importantly, to make sure that our curiosity effects on learning remain regardless of confidence in prior knowledge, we ran a follow-up logistic regression analysis. Controlling for each other, both *z*-scored curiosity ($\beta = 0.26$, SE = 0.05, z = 5.69, p < .001) and *z*-scored confidence ($\beta = 0.26$, SE = 0.05, z = 5.45, p < .001) were significant predictors of memory. Critically, there was no interaction between curiosity and confidence in predicting memory ($\beta = -0.08$, SE = 0.05, z = -1.66, p = .096; BF₁₀ = 0.05). Our behavioral results, along with previous findings in the literature (Wade & Kidd, 2019; Stare, Gruber, Nadel, Ranganath, & Gómez, 2018), suggest that although confidence has an impact on memory, the effect of curiosity remains independent of confidence in prior knowledge.

Electrophysiological Results

RewP (250-350 msec)

To investigate reward-related processes during the presentation of trivia answers, the RewP was analyzed in a

2 (Curiosity: high, low) \times 2 (Memory: remembered, forgotten) repeated-measures ANOVA. This analysis vielded a significant main effect for Curiosity, F(1, 21) = 6.29 p =.020, $\eta_p^2 = .23$, and for Memory, F(1, 21) = 6.43, p = .019, $\eta_p^2 = .23$ (Figure 3). Notably, a two-way interaction between Curiosity and Memory was not significant, F(1,21) = 0.24, p = .63, $\eta_p^2 = .01$; BF₁₀ = 0.15. The RewP amplitude was more positive for high-curiosity (M =3.26, SD = 3.54) compared with low-curiosity answers (M = 2.07, SD = 3.56; t(21) = 2.51, p = .010, d =0.54) as well as for remembered answers (M = 3.41SD = 3.51) compared with forgotten answers (M =1.91, SD = 3.78; t(21) = 2.54, p = .009, d = 0.54). It can be assumed that the greater RewP amplitude for high-curiosity answers indicates increased rewardrelated processing of the answers in comparison to low curiosity. Furthermore, a greater amplitude of the RewP was predictive for better memory, but we did not find that increased reward processing benefitted later memory, that is, there was no interaction between curiosity and memory for the RewP.

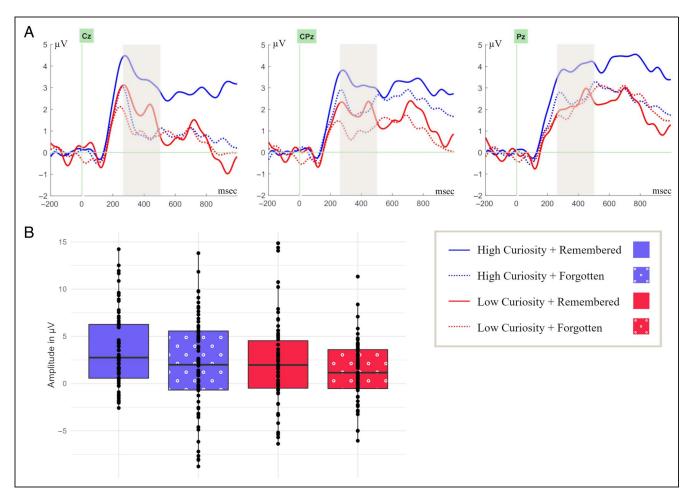


Figure 4. Curiosity and memory independently modulate the P3 component during answer presentation at central and parietal electrode sites. (A) ERPs were time-locked to the onset of the answers in the study phase at electrode sites Cz, CPz, and Pz in the colored time window of 250–500 msec. (B) Box plots are bounded by the first and third quartiles, and the black line represents the median. The points correspond to the measures for individual participants.

P3 (250-500 msec)

For examining the allocation of neural resources based on reward effects, stimulus relevance, and context updating in working memory, the P3 was analyzed in a 2 (Curiosity: high, low) \times 2 (Memory: remembered, forgotten) repeated-measures ANOVA. Similarly, to the findings of the RewP, this analysis yielded a significant main effect for Curiosity, F(1, 21) = 4.70, p = .042, $\eta_p^2 = .18$, as well as for Memory, F(1, 21) = 6.70, p = .017, $\eta_p^2 = .24$ (Figure 4). The two-way interaction between Curiosity and Memory was not significant, $F(1, 21) = 0.26, p = .61, \eta_D^2 =$.01; $BF_{10} = 0.16$. The amplitude was more positive for high-curiosity (M = 2.83, SD = 3.99) compared with low-curiosity answers (M = 1.76, SD = 3.31; t(21) =2.17, p = .021, d = 0.46) as well as for remembered answers (M = 2.92 SD = 3.66) compared with forgotten answers (M = 1.67, SD = 3.65; t(21) = 2.59, p = .008, d =0.55). Most relevantly, the greater P3 amplitude for high curiosity suggests increased stimulus relevance as well as more pronounced context updating in working memory.

In addition, as with the RewP, a greater amplitude of the P3 was predictive for better memory.

LPP (600-1000 msec)

The analysis of extended cognitive and attentional processing based on reward expectancy and magnitude was implemented in a 2 (Curiosity: high, low) \times 2 (Memory: remembered, forgotten) repeated-measures ANOVA on the LPP. This analysis yielded a significant main effect for Curiosity, F(1, 21) = 5.16, p = .034, $\eta_p^2 = .20$, but not for Memory, F(1, 21) = 0.83, p = .37, $\eta_p^2 = .04$, BF₁₀ = 0.27 (Figure 5). There was no two-way interaction between Curiosity and Memory, F(1, 21) = 1.26, p = .27, $\eta_p^2 = .06$; BF₁₀ = 0.26. The amplitude was more positive for high-curiosity (M = 2.66, SD = 5.40) compared with low-curiosity answers (M = 1.32, SD = 4.61; t (21) = 2.27, p = .017, d = 0.48). The greater LPP amplitude for high- compared with low-curiosity answers is assumed to indicate extended cognitive and attentional processing based on reward magnitude.

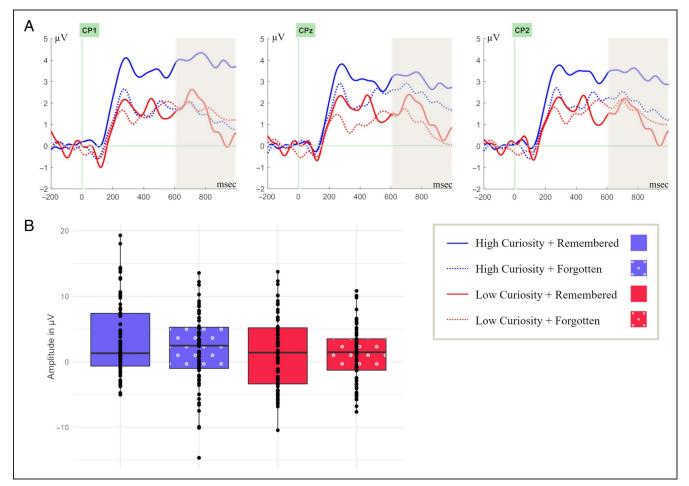


Figure 5. Only curiosity (but not memory) modulates the LPP during answer presentation at centro-parietal electrode sites. (A) ERPs were time-locked to the onset of the answers in the study phase at electrode sites CP1, CPz, and CP2 in the colored time window of 600–1000 msec. (B) Box plots are bounded by the first and third quartiles, and the black line represents the median. The points correspond to the measures of individual participants.

DISCUSSION

The present article addressed the question of whether curiosity satisfaction modulates specific ERPs that are sensitive to reward. In the initial screening phase, participants were presented with general knowledge questions and were asked to indicate their level of curiosity and confidence of one's own prior knowledge about the respective answer. In a subsequent study phase, the influence of curiosity levels on reward-related ERPs was examined via the RewP, P3, and LPP. At the end of the experiment, there was an unannounced memory test in which the participants had to repeat the previously presented answers to the trivia questions.

Behaviorally, we found better memory performance for high-curiosity in contrast to low-curiosity answers. In the same study setup, Gruber and colleagues (2014) found similarly large differences between high- and low-curiosity memory performance. In general, the reported memory benefit for high curiosity is in line with the findings of several other studies (Poh et al., 2022; Fandakova & Gruber, 2021; Lau et al., 2020; Galli et al., 2018; Stare et al., 2018; Marvin & Shohamy, 2016; Gruber et al., 2014; Kang et al., 2009). Our analyses also showed that curiosity and confidence in one's own prior knowledge influence memory independently and tend to share only a small proportion of variance. It can therefore be assumed that the results of the ERP analyses can be attributed to the different levels of curiosity independent of confidence.

ERP analysis aimed at assessing differences in the online processing of high- compared with low-curiosity answers and processes predictive of successful memory formation. We found that the amplitudes of the RewP, P3, and LPP were more positive in the high- than in the low-curiosity condition. Furthermore, the RewP and P3 (but not the LPP) showed greater positive amplitudes for later remembered compared with later forgotten answers. No interaction between curiosity levels and subsequent memory were found for the examined ERP components. However, this interaction results should be viewed with caution because of the exploratory approach. In general, the results suggest that the satisfaction of curiosity via new information resembles reward processing.

Reward Processing

First, we assumed that information—here the answer to a trivia question—represents a rewarding stimulus that reduces uncertainty and closes a knowledge gap (Loewenstein, 1994). Several behavioral and neuroimaging studies support the view that states of high curiosity and extrinsically motivated reward depend on similar mechanisms and activate overlapping brain areas (FitzGibbon et al., 2020; Marvin & Shohamy, 2016; Gruber et al., 2014; Bromberg-Martin & Hikosaka, 2009; Kang et al., 2009). Moreover, the examined RewP is seen as an indicator of reward processing (Anderson, 2017; Heydari & Holroyd, 2016; Sambrook & Goslin, 2015).

Accordingly, the discovered differences of the RewP amplitude based on curiosity levels can be seen as a neurophysiological correlate of reward processing, where answers with high curiosity triggered greater reward responses.

In addition, the Prediction, Appraisal, Curiosity, and Exploration framework (Gruber & Ranganath, 2019) suggests that higher expectation of reward and curiosity are accompanied by increased dopaminergic activity, which leads to facilitated hippocampus-based memory formation (Gruber & Ranganath, 2019; Smith, Starck, Roberts, & Schuman, 2005; Wittmann et al., 2005; Holroyd & Coles, 2002; Schultz, 2002). Although the present work does not allow to draw any direct conclusions about dopaminergic circuit activity, the identical experimental paradigm as in our previous fMRI study on curiosity (Gruber et al., 2014) permits to assume an increase in dopaminergic activity. Furthermore, the RewP can be an indication of dopaminergic signals itself (Baker & Holroyd, 2011). The origin of the dopaminergic activity echoed by the RewP has been localized in the basal ganglia and other rewardrelated brain areas, and positive reward seems to be associated with increased mPFC activity (Becker et al., 2014). Moreover, van Lieshout, Vandenbroucke, Müller, Cools, and de Lange (2018) found increased mPFC activity during curiosity satisfaction and Murphy, Ranganath, and Gruber (2021) investigated functional connectivity between the hippocampus and mPFC during answer presentation, which predicted the curiosity-related memory benefit. In summary, the increased positivity of the RewP in the high curiosity condition indicates that the answers were perceived as more rewarding, which in turn seems to have strengthened reward-related processing.

Second, larger P3 and LPP amplitudes in the high curiosity condition may reflect increased allocation of neural resources and attention based on reward processing. A more positive amplitude of the P3 for high-compared with low-curiosity answers suggests that there was an enhanced allocation of neural resources and thereby increased attention (San Martín, 2012; Debener, Makeig, Delorme, & Engel, 2005; Friedman, Cycowicz, & Gaeta, 2001). Suitable for this, the increased activity of the dopaminergic system (indicated by the RewP) can lead to more attention to upcoming rewards (Anderson, 2016; Hickey, Chelazzi, & Theeuwes, 2010). First evidence for this was provided by fMRI studies that showed increased activity at high curiosity in frontal and parietal brain areas associated with attentional and cognitive control (van Lieshout et al., 2018; Jepma et al., 2012). The modulation of the LPP also suggests that the perceived relevance of the information (reward magnitude) might have been greater during states of high curiosity because of the activation of appetitive motivational systems through curiosity (Hajcak & Foti, 2020). Synoptical, the greater positivity in the high curiosity condition of the P3 (San Martín, 2012; Van Petten & Luka, 2012) and the LPP (Weinberg et al., 2013; Schupp et al., 2000) can also be regarded as a marker of an increased reward processing.

Third, the generally more positive amplitudes with high curiosity are in line with other studies examining answer processing after the presentation of trivia questions. Investigating the tip-of-the-tongue phenomenon (see Metcalfe, Schwartz, & Bloom, 2017; Schwartz, 2006)—a subjective feeling associated with the state of high curiosity—Bloom, Friedman, Xu, Vuorre, and Metcalfe (2018) found an enhanced positivity during answer presentation at centro-parietal electrode sites, when participants reported to be in a tip-of-the-tongue state. This study supports our conclusions that identifying a gap in knowledge can enhance processing of new information, as seen in the increased positive amplitudes of the ERPs.

To ensure consistency to our prior work using the identical paradigm in an fMRI study (see Gruber et al., 2014), we also adopted the time intervals of stimulus presentation. However, from the literature on feedback processing, it is known that a long delay in presentation of a response (11 sec in this study), can lead to a reduction of the RewP/FRN (Höltje & Mecklinger, 2018; Yin, Wang, Zhang, & Li, 2018; Peterburs et al., 2016). For example, Yin and colleagues (2018) showed that the RewP decreases when feedback is delayed. At the same time, Höltje and Mecklinger (2018) were able to show that the FRN amplitude is reduced, but still reliable even with long delays (6.5 sec). Given that we found significant differences between curiosity conditions for RewP despite this longer response delay, this is evidence that curiosity might affect the RewP in a robust way.

Furthermore, the instruction of the participants was to "always try to anticipate the possible answer." Therefore, one cannot completely exclude the influence of expectation in this reward processing study. For example, the increased amplitude of the LPP (comparable with the P600 between 600 and 1000 msec; Kuperberg, Brothers, & Wlotko, 2020; DeLong, Quante, & Kutas, 2014) by high curiosity can potentially also be interpreted as increased integration effort of the answers based on expectancy. Late positive potentials have proven to be relevant EEG markers for the processing and integration of new information into context (Aurnhammer, Delogu, Schulz, Brouwer, & Crocker, 2021; Brouwer, Crocker, Noortje, Venhuizen, & Hoeks, 2017; Kos, van den Brink, & Hagoort, 2012; Kuperberg, 2007). Because neither the contextual prior knowledge nor expectancy errors were collected in this study, this idea could be addressed in future research. Moreover, this also applies vice versa to studies that examine expectations, where feedback can also be rewarding. According to Wu and Zhou (2009), the FRN/RewP is more affected by expectancy related processing than the P300. As we found the curiosity effect on both components (RewP and P300), we take this as evidence that these effects reflect the influence of curiosity rather than feedback expectation.

Memory Effects

A SME was found for the amplitudes of the RewP and P3 without an interaction with curiosity. On the one hand, the

difference in the amplitudes between remembered and forgotten words implies that the processes reflected by the RewP/P3 contribute to successful memory formation. On the other hand, the lack of an interaction does not allow for a direct connection between the curiosity modulation of the RewP/P3 and the behavioral curiosity-related memory benefits.

The SMEs could have been influenced by previous occurring processes, such as increased hippocampal activity or dopaminergic activity. It is important to note that Gruber and colleagues (2014) also did not find any significant interaction between curiosity-related brain activity and memory during answer presentation in the dopaminergic regions and the hippocampus. Critically, activation of dopaminergic areas and the hippocampus during curiosity elicitation but not satisfaction seems to support the curiosity-related memory benefit. During curiosity satisfaction, it seems that integration processes between hippocampus and large-scale cortical networks (e.g., with the default mode network, Murphy et al., 2021) might contribute to the curiosity-related memory benefit. It is also conceivable that high curiosity levels might have enhanced the distinctiveness of events in a given processing context and the more distinctive items were more efficiently encoded and gave rise to an SME in the P3 time window (Otten & Donchin, 2000).

The absence of the SME for the LPP can have several reasons. On the one hand, it is possible that SMEs were too small to modulate the investigated ERP components significantly because of the sample size. On the other hand, SME effects for the LPP with onset latencies beyond 600 msec have typically been reported when intentional and elaborative encoding strategies emphasizing associative processing of multiple study features are required (Kamp & Zimmer, 2015; Cheng & Rugg, 2010). These processing characteristics were presumably not initiated by the presentation of the answers to the trivia questions. Taken together, the SMEs in this work turned out to be greater at the beginning of the answer presentation (RewP, P3) and to vanish when it comes to a later time window (LPP).

Conclusion

In the present work, we specifically targeted the RewP, P3, and LPP because of their relationship with reward processing and found meaningful connections between these ERPs and curiosity. The findings show increased positive amplitudes for all examined ERPs for high compared with low curiosity, suggesting increased reward processing. The SMEs for the RewP and P3 indicate that these reward-related processes contribute to successful memory encoding. According to this first investigation of ERPs during curiosity satisfaction, our findings support the view that curiosity satisfaction resembles reward processing, opening up the investigation of further curiosity-related ERPs studies in future research. Moreover, the present study has shown that not only the state of curiosity but also

its satisfaction is an important object of research to better understand the processes underlying curiosity and its similarity to reward. Furthermore, understanding which processes are normally influenced by the satisfaction of curiosity could help to detect clinical conditions that are accompanied by a lack of motivation and dopaminergic circuit activity. In addition, ERPs are valuable tools to explore effects of curiosity because of their high temporal resolution. They allow to track the time course of curiosity-related processing while it unfolds. In summary, our findings suggest that the satisfaction of curiosity with new information is rewarding.

Acknowledgments

The authors thank Gerrit Höltje (Experimental Neuropsychology Department, Saarland University) for helping during the statistical analysis and writing.

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Data Availability Statement

The behavioral and ERP data are available at https://osf.io/9k3tv/. The stimuli are available at https://osf.io/he6t9/. The experiment was not preregistered.

Author Contributions

C. R. and M. J. G. designed research, J. C.-D. and M. J. G. performed research, T. R. analyzed the data, K. C. J. E. and J. C.-D. assisted and contributed to data analysis, A. M. and M. J. G. supervised the project. T. R. drafted the paper, T. R., A. M., K. C. J. E., and M. J. G. interpreted and wrote the paper, and all other authors gave comments on the paper.

Funding Information

This work was supported by a Wellcome Trust (https://dx.doi.org/10.13039/100010269) and Royal Society Sir Henry Dale Fellowship, grant number: 211201/Z/18/Z awarded to M. J. G. and a German Research Foundation Research Fellowship (https://dx.doi.org/10.13039/501100001659), grant number: 442588275 awarded to K. C. J. E. For the purpose of Open Access, the authors have applied a CC-BY public copyright license to any author accepted manuscript version arising from this submission.

Diversity in Citation Practices

Retrospective analysis of the citations in every article published in this journal from 2010 to 2021 reveals a persistent pattern of gender imbalance: Although the proportions of

authorship teams (categorized by estimated gender identification of first author/last author) publishing in the *Journal of Cognitive Neuroscience* (*JoCN*) during this period were M(an)/M = .407, W(oman)/M = .32, M/W = .115, and W/W = .159, the comparable proportions for the articles that these authorship teams cited were M/M = .549, W/M = .257, M/W = .109, and W/W = .085 (Postle and Fulvio, *JoCN*, 34:1, pp. 1 -3). Consequently, *JoCN* encourages all authors to consider gender balance explicitly when selecting which articles to cite and gives them the opportunity to report their article's gender citation balance. The authors of this article report its proportions of citations by gender category to be: M/M = .508; W/M = .317; M/W = .079; W/W = .095.

REFERENCES

- Anderson, B. A. (2016). The attention habit: How reward learning shapes attentional selection. *Annals of the New York Academy of Sciences*, *1369*, 24–39. https://doi.org/10.1111/nyas.12957, PubMed: 26595376
- Anderson, B. A. (2017). Reward processing in the value-driven attention network: Reward signals tracking cue identity and location. Social Cognitive and Affective Neuroscience, 12, 461–467. https://doi.org/10.1093/scan/nsw141, PubMed: 27677944
- Aurnhammer, C., Delogu, F., Schulz, M., Brouwer, H., & Crocker, M. W. (2021). Retrieval (N400) and integration (P600) in expectation-based comprehension. *PLoS One*, *16*, e0257430. https://doi.org/10.1371/journal.pone.0257430, PubMed: 34582472
- Baker, T. E., & Holroyd, C. B. (2011). Dissociated roles of the anterior cingulate cortex in reward and conflict processing as revealed by the feedback error-related negativity and N200. *Biological Psychology*, 87, 25–34. https://doi.org/10.1016/j .biopsycho.2011.01.010, PubMed: 21295109
- Becker, M. P. I., Nitsch, A. M., Miltner, W. H. R., & Straube, T. (2014). A single-trial estimation of the feedback-related negativity and its relation to bold responses in a time-estimation task. *Journal of Neuroscience*, *34*, 3005–3012. https://doi.org/10.1523/JNEUROSCI.3684-13.2014, PubMed: 24553940
- Berlyne, D. E. (1954). An experimental study of human curiosity. *British Journal of Psychology*, *45*, 256–265. https://doi.org/10.1111/j.2044-8295.1954.tb01253.x, PubMed: 13219284
- Biosemi, Neurospec. (n.d.). ActiveTwo | NEUROSPEC AG Research neurosciences. https://www.neurospec.com /Products/Details/1000/activetwo
- Bloom, P. A., Friedman, D., Xu, J., Vuorre, M., & Metcalfe, J. (2018). Tip-of-the-tongue states predict enhanced feedback processing and subsequent memory. *Consciousness and Cognition*, 63, 206–217. https://doi.org/10.1016/j.concog .2018.05.010, PubMed: 29887295
- Bradley, M. M. (2009). Natural selective attention: Orienting and emotion. *Psychophysiology*, *46*, 1–11. https://doi.org/10.1111/j.1469-8986.2008.00702.x, PubMed: 18778317
- Bromberg-Martin, E. S., & Hikosaka, O. (2009). Midbrain dopamine neurons signal preference for advance information about upcoming rewards. *Neuron*, *63*, 119–126. https://doi.org/10.1016/j.neuron.2009.06.009, PubMed: 19607797
- Brouwer, H., Crocker, M. W., Venhuizen, N. J., & Hoeks, J. C. J. (2017). A neurocomputational model of the N400 and the P600 in language processing. *Cognitive Science*, 41(Suppl. 6),

- 1318–1352. https://doi.org/10.1111/cogs.12461, PubMed: 28000963
- Cheng, S.-K., & Rugg, M. D. (2010). Event-related potential correlates of gist and verbatim encoding. *International Journal of Psychophysiology*, 77, 95–105. https://doi.org/10.1016/j.ijpsycho.2010.04.010, PubMed: 20460139
- Cogent. (n.d.). Wellcome Laboratory of Neurobiology, University College London. https://www.vislab.ucl.ac.uk/cogent.php
- Cohen, J. (1988). Statistical power analysis for the behavioral sciences (2nd ed.). Hillsdale, NJ: L. Erlbaum Associates.
- Cohen, M. X., & Ranganath, C. (2007). Reinforcement learning signals predict future decisions. *Journal of Neuroscience*, 27, 371–378. https://doi.org/10.1523/JNEUROSCI.4421-06.2007, PubMed: 17215398
- Cohen, N., Pell, L., Edelson, M. G., Ben-Yakov, A., Pine, A., & Dudai, Y. (2015). Peri-encoding predictors of memory encoding and consolidation. *Neuroscience & Biobehavioral Reviews*, 50, 128–142. https://doi.org/10.1016/j.neubiorev .2014.11.002, PubMed: 25446944
- Debener, S., Makeig, S., Delorme, A., & Engel, A. K. (2005). What is novel in the novelty oddball paradigm? Functional significance of the novelty P3 event-related potential as revealed by independent component analysis. *Cognitive Brain Research*, 22, 309–321. https://doi.org/10.1016/j.cogbrainres.2004.09.006, PubMed: 15722203
- DeLong, K. A., Quante, L., & Kutas, M. (2014). Predictability, plausibility, and two late ERP positivities during written sentence comprehension. *Neuropsychologia*, 61, 150–162. https://doi.org/10.1016/j.neuropsychologia.2014.06.016, PubMed: 24953958
- Delorme, A., & Makeig, S. (2004). EEGLAB: An open source toolbox for analysis of single-trial EEG dynamics including independent component analysis. *Journal of Neuroscience Methods*, *134*, 9–21. https://doi.org/10.1016/j.jneumeth.2003.10.009. PubMed: 15102499
- Dienes, Z. (2014). Using Bayes to get the most out of non-significant results. Frontiers in Psychology, 5, 781. https://doi.org/10.3389/fpsyg.2014.00781, PubMed: 25120503
- Donaldson, K. R., Oumeziane, B. A., Hélie, S., & Foti, S. H. D. (2016). The temporal dynamics of reversal learning: P3 amplitude predicts valence-specific behavioral adjustment. *Physiology & Behavior*, 161, 24–32. https://doi.org/10.1016/j.physbeh.2016.03.034, PubMed: 27059320
- Duan, H., Fernández, G., van Dongen, E., & Kohn, N. (2020). The effect of intrinsic and extrinsic motivation on memory formation: insight from behavioral and imaging study. *Brain Structure & Function*, 225, 1561–1574. https://doi.org/10.1007/s00429-020-02074-x, PubMed: 32350643
- Fandakova, Y., & Gruber, M. J. (2021). States of curiosity and interest enhance memory differently in adolescents and in children. *Developmental Science*, 24, e13005. https://doi.org/10.1111/desc.13005, PubMed: 32524703
- Faul, F., Erdfelder, E., Buchner, A., & Lang, A.-G. (2009). Statistical power analyses using G*Power 3.1: Tests for correlation and regression analyses. *Behavior Research Methods*, 41, 1149–1160. https://doi.org/10.3758/BRM.41.4 .1149, PubMed: 19897823
- FitzGibbon, L., Lau, J. K. L., & Murayama, K. (2020). The seductive lure of curiosity: Information as a motivationally salient reward. *Current Opinion in Behavioral Sciences*, 35, 21–27. https://doi.org/10.1016/j.cobeha.2020.05.014
- Friedman, D., Cycowicz, Y. M., & Gaeta, H. (2001). The novelty P3: An event-related brain potential (ERP) sign of the brain's evaluation of novelty. *Neuroscience & Biobehavioral Reviews*, 25, 355–373. https://doi.org/10.1016/s0149-7634(01)00019-7, PubMed: 11445140

- Gable, P. A., Adams, D. L., & Proudfit, G. H. (2015). Transient tasks and enduring emotions: The impacts of affective content, task relevance, and picture duration on the sustained late positive potential. *Cognitive, Affective, & Behavioral Neuroscience*, *15*, 45–54. https://doi.org/10.3758/s13415-014-0313-8, PubMed: 25139377
- Galli, G., Sirota, M., Gruber, M. J., Ivanof, B. E., Ganesh, J., Materassi, M., et al. (2018). Learning facts during aging: The benefits of curiosity. *Experimental Aging Research*, 44, 311–328. https://doi.org/10.1080/0361073X.2018.1477355, PubMed: 29787342
- Glazer, J. E., Kelley, N. J., Pornpattananangkul, N., Mittal, V. A., & Nusslock, R. (2018). Beyond the FRN: Broadening the time-course of EEG and ERP components implicated in reward processing. *International Journal of Psychophysiology*, 132, 184–202. https://doi.org/10.1016/j .ijpsycho.2018.02.002, PubMed: 29454641
- Gruber, M. J., Gelman, B. D., & Ranganath, C. (2014). States of curiosity modulate hippocampus-dependent learning via the dopaminergic circuit. *Neuron*, 84, 486–496. https://doi.org/10 .1016/j.neuron.2014.08.060, PubMed: 25284006
- Gruber, M. J., & Ranganath, C. (2019). How curiosity enhances hippocampus-dependent memory: The Prediction-Appraisal-Curiosity-Exploration (PACE) framework. *Trends in Cognitive Sciences*, 23, 1014–1025. https://doi.org/10.1016/j.tics.2019.10 .003, PubMed: 31706791
- Hajcak, G., & Foti, D. (2020). Significance? ... Significance! Empirical, methodological, and theoretical connections between the late positive potential and P300 as neural responses to stimulus significance: An integrative review. *Psychophysiology*, 57, e13570. https://doi.org/10.1111/psyp .13570, PubMed: 32243623
- Hajcak, G., Moser, J. S., Holroyd, C. B., & Simons, R. F. (2006). The feedback-related negativity reflects the binary evaluation of good versus bad outcomes. *Biological Psychology*, 71, 148–154. https://doi.org/10.1016/j.biopsycho.2005.04.001, PubMed: 16005561
- Heydari, S., & Holroyd, C. B. (2016). Reward positivity: Reward prediction error or salience prediction error? *Psychophysiology*, *53*, 1185–1192. https://doi.org/10.1111/psyp.12673, PubMed: 27184070
- Hickey, C., Chelazzi, L., & Theeuwes, J. (2010). Reward changes salience in human vision via the anterior cingulate. *Journal of Neuroscience*, 30, 11096–11103. https://doi.org/10.1523 /JNEUROSCI.1026-10.2010, PubMed: 20720117
- Holroyd, C. B., & Coles, M. G. H. (2002). The neural basis of human error processing: Reinforcement learning, dopamine, and the error-related negativity. *Psychological Review*, 109, 679–709. https://doi.org/10.1037/0033-295X.109.4.679, PubMed: 12374324
- Höltje, G., & Mecklinger, A. (2018). Electrophysiological reward signals predict episodic memory for immediate and delayed positive feedback events. *Brain Research*, 1701, 64–74. https://doi.org/10.1016/j.brainres.2018.07.011, PubMed: 30017716
- Höltje, G., & Mecklinger, A. (2020). Feedback timing modulates interactions between feedback processing and memory encoding: Evidence from event-related potentials. *Cognitive*, *Affective*, & *Behavioral Neuroscience*, 20, 250–264. https:// doi.org/10.3758/s13415-019-00765-5, PubMed: 31900874
- Jasper, H. (1958). The 10–20 electrode system of the international federation. *Electroencephalography and Clinical Neurophysiology*, 10, 371–375. https://ci.nii.ac.jp/naid/10010880516/
- Jeffreys, H. (1961). The theory of probability (3rd ed.). Oxford, England: Oxford University Press.
- Jepma, M., Verdonschot, R. G., van Steenbergen, H., Rombouts, S. A. R. B., & Nieuwenhuis, S. (2012). Neural mechanisms

- underlying the induction and relief of perceptual curiosity. *Frontiers in Behavioral Neuroscience*, 6, 5. https://doi.org/10.3389/fnbeh.2012.00005, PubMed: 22347853
- Kamp, S.-M., & Zimmer, H. D. (2015). Contributions of attention and elaboration to associative encoding in young and older adults. *Neuropsychologia*, 75, 252–264. https://doi .org/10.1016/j.neuropsychologia.2015.06.026, PubMed: 26115601
- Kang, M. J., Hsu, M., Krajbich, I. M., Loewenstein, G., McClure, S. M., Wang, J. T. Y., et al. (2009). The wick in the candle of learning: Epistemic curiosity activates reward circuitry and enhances memory. *Psychological Science*, 20, 963–973. https://doi.org/10.1111/j.1467-9280.2009.02402.x, PubMed: 19619181
- Kidd, C., & Hayden, B. Y. (2015). The psychology and neuroscience of curiosity. *Neuron*, 88, 449–460. https://doi. org/10.1016/j.neuron.2015.09.010, PubMed: 26539887
- Kos, M., van den Brink, D., & Hagoort, P. (2012). Individual variation in the late positive complex to semantic anomalies. *Frontiers in Psychology*, 3, 318. https://doi.org/10.3389/fpsyg .2012.00318, PubMed: 22973249
- Kuperberg, G. R. (2007). Neural mechanisms of language comprehension: Challenges to syntax. *Brain Research*, 1146, 23–49. https://doi.org/10.1016/j.brainres.2006.12.063, PubMed: 17400197
- Kuperberg, G. R., Brothers, T., & Wlotko, E. W. (2020). A tale of two positivities and the N400: Distinct neural signatures are evoked by confirmed and violated predictions at different levels of representation. *Journal of Cognitive Neuroscience*, 32, 12–35. https://doi.org/10.1162/jocn_a_01465, PubMed: 31479347
- Lau, J. K. L., Ozon, H., Kuratomi, K., Komiya, A., & Murayama, A. (2020). Shared striatal activity in decisions to satisfy curiosity and hunger at the risk of electric shocks. *Nature Human Behaviour*, 4, 531–543. https://doi.org/10.1038/s41562-020-0848-3, PubMed: 32231281
- Ligneul, R., Mermillod, M., & Morisseau, T. (2018). From relief to surprise: Dual control of epistemic curiosity in the human brain. *Neuroimage*, 181, 490–500. https://doi.org/10.1016/j.neuroimage.2018.07.038, PubMed: 30025853
- Loewenstein, G. (1994). The psychology of curiosity: A review and reinterpretation. *Psychological Bulletin*, *116*, 75–98. https://doi.org/10.1037/0033-2909.116.1.75
- Lopez-Calderon, J., & Luck, S. J. (2014). ERPLAB: An open-source toolbox for the analysis of event-related potentials. *Frontiers in Human Neuroscience*, *8*, 213. https://doi.org/10.3389/fnhum.2014.00213, PubMed: 24782741
- Marvin, C. B., & Shohamy, D. (2016). Curiosity and reward: Valence predicts choice and information prediction errors enhance learning. *Journal of Experimental Psychology: General*, *145*, 266–272. https://doi.org/10.1037/xge0000140, PubMed: 26783880
- Meadows, C. C., Gable, P. A., Lohse, K. R., & Miller, M. W. (2016). The effects of reward magnitude on reward processing: An averaged and single trial event-related potential study. *Biological Psychology*, *118*, 154–160. https://doi.org/10.1016/j.biopsycho.2016.06.002, PubMed: 27288743
- Mecklinger, A., & Kamp, S.-M. (2023). Observing memory encoding while it unfolds: Functional interpretation and current debates regarding ERP subsequent memory effects. *Neuroscience and Biobehavioral Reviews*, *153*, 105347. https://doi.org/10.1016/j.neubiorev.2023.105347, PubMed: 37543177
- Metcalfe, J., Schwartz, B. L., & Bloom, P. A. (2017). The tip-of-the-tongue state and curiosity. *Cognitive Research*:

- *Principles and Implications*, 2, 31. https://doi.org/10.1186/s41235-017-0065-4, PubMed: 28776003
- Metcalfe, J., Schwartz, B. L., & Eich, T. S. (2020). Epistemic curiosity and the region of proximal learning. *Current Opinion in Behavioral Sciences*, 35, 40–47. https://doi.org /10.1016/j.cobeha.2020.06.007, PubMed: 33709011
- Murphy, C., Ranganath, C., & Gruber, M. J. (2021). Connectivity between the hippocampus and default mode network during the relief—but not elicitation—of curiosity supports curiosity-enhanced memory enhancements. *bioRxiv*. https://doi.org/10.1101/2021.07.26.453739
- Oosterwijk, S., Snoek, L., Tekoppele, J., Engelbert, L. H., & Scholte, H. S. (2020). Choosing to view morbid information involves reward circuitry. *Scientific Reports*, *10*, 15291. https://doi.org/10.1038/s41598-020-71662-y, PubMed: 32943668
- Otten, L., & Donchin, E. (2000). Relationship between P300 amplitude and subsequent recall for distinctive events: Dependence on type of distinctiveness attribute. *Psychophysiology*, *37*, 644–661. https://doi.org/10.1111/1469-8986.3750644, PubMed: 11037041
- Peterburs, J., Kobza, S., & Bellebaum, C. (2016). Feedback delay gradually affects amplitude and valence specificity of the feedback-related negativity (FRN). *Psychophysiology*, *53*, 209–215. https://doi.org/10.1111/psyp.12560, PubMed: 26459164
- Poh, J.-H., Vu, M.-A. T., Stanek, J. K., Hsiung, A., Egner, T., & Adcock, R. A. (2022). Hippocampal convergence during anticipatory midbrain activation promotes subsequent memory formation. *Nature Communications*, 13, 6729. https://doi.org/10.1038/s41467-022-34459-3, PubMed: 36344524
- Polich, J. (2007). Updating P300: An integrative theory of P3a and P3b. *Clinical Neurophysiology*, 118, 2128–2148. https://doi.org/10.1016/j.clinph.2007.04.019, PubMed: 17573239
- Proudfit, G. H. (2015). The reward positivity: From basic research on reward to a biomarker for depression. *Psychophysiology*, 52, 449–459. https://doi.org/10.1111/psyp.12370, PubMed: 25327938
- R Core Team. (2019). R: A language and environment for statistical computing. Vienna, Austria: R Foundation for Statistical Computing. https://www.R-project.org/
- Rasch, B., Friese, M., Hofmann, W., & Naumann, E. (2021). Quantitative methoden 2. Einführung in die Statistik für Psychologie, Sozial- & Erziehungswissenschaften. Heidelberg: Springer. https://doi.org/10.1007/978-3-662 -63284-0
- Sambrook, T. D., & Goslin, J. (2015). A neural reward prediction error revealed by a meta-analysis of ERPs using great grand averages. *Psychological Bulletin*, 141, 213–235. https://doi. org/10.1037/bul0000006, PubMed: 25495239
- San Martín, R. (2012). Event-related potential studies of outcome processing and feedback-guided learning. *Frontiers* in Human Neuroscience, 6, 304. https://doi.org/10.3389 /fnhum.2012.00304, PubMed: 23162451
- Schultz, W. (2002). Getting formal with dopamine and reward. *Neuron*, 36, 241–263. https://doi.org/10.1016/s0896-6273(02)00967-4, PubMed: 12383780
- Schupp, H. T., Cuthbert, B. N., Bradley, M. M., Cacioppo, J. T., Ito, T., & Lang, P. J. (2000). Affective picture processing: The late positive potential is modulated by motivational relevance. *Psychophysiology*, *37*, 257–261. https://doi.org/10.1111/1469-8986.3720257, PubMed: 10731776
- Schwartz, B. L. (2006). Tip-of-the-tongue states as metacognition. *Metacognition and Learning*, *1*, 149–158. https://doi.org/10.1007/s11409-006-9583-z
- Smith, W. B., Starck, S. R., Roberts, R. W., & Schuman, E. M. (2005).Dopaminergic stimulation of local protein synthesis enhances surface expression of GluR1 and synaptic transmission in

- hippocampal neurons. *Neuron*, 45, 765–779. https://doi.org/10.1016/j.neuron.2005.01.015, PubMed: 15748851
- Stare, C. J., Gruber, M. J., Nadel, L., Ranganath, C., & Gómez, R. L. (2018). Curiosity-driven memory enhancement persists over time but does not benefit from post-learning sleep. *Cognitive Neuroscience*, *9*, 100–115. https://doi.org/10.1080/17588928.2018.1513399, PubMed: 30124373
- van Lieshout, L. L. F., Vandenbroucke, A. R. E., Müller, N. C. J., Cools, R., & de Lange, F. P. (2018). Induction and relief of curiosity elicit parietal and frontal activity. *Journal of Neuroscience*, *38*, 2579–2588. https://doi.org/10.1523/JNEUROSCI.2816-17.2018, PubMed: 29439166
- Van Petten, C., & Luka, B. J. (2012). Prediction during language comprehension: Benefits, costs, and ERP components. *International Journal of Psychophysiology*, 83, 176–190. https://doi.org/10.1016/j.ijpsycho.2011.09.015, PubMed: 22019481
- Wade, S., & Kidd, C. (2019). The role of prior knowledge and curiosity in learning. *Psychonomic Bulletin & Review*, 26, 1377–1387. https://doi.org/10.3758/s13423-019-01598-6, PubMed: 31079309

- Weinberg, A., Ferri, J., & Hajcak, G. (2013). Interactions between attention and emotion: Insights from the late positive potential. In M. D. Robinson, E. Watkins, & E. Harmon-Jones (Eds.), *Handbook of cognition and emotion* (pp. 35–54). The Guilford Press.
- Wittmann, B. C., Schott, B. H., Guderian, S., Frey, J. U., Heinze, H. J., & Duzel, E. (2005). Reward-related fMRI activation of dopaminergic midbrain is associated with enhanced hippocampus-dependent long-term memory formation. Neuron, 45, 459–467. https://doi.org/10.1016/j.neuron.2005.01.010, PubMed: 15694331
- Wu, Y., & Zhou, X. (2009). The P300 and reward valence, magnitude, and expectancy in outcome evaluation. *Brain Research*, 1286, 114–122. https://doi.org/10.1016/j.brainres.2009.06.032, PubMed: 19539614
- Yin, H., Wang, Y., Zhang, X., & Li, P. (2018). Feedback delay impaired reinforcement learning: Principal components analysis of reward positivity. *Neuroscience Letters*, 685, 179–184. https://doi.org/10.1016/j.neulet.2018.08.039, PubMed: 30170042