Short title: Context-dependent reward-enhanced memory

2 # Can neural correlates of encoding explain the context dependence of

3 reward-enhanced memory?

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Abstract

Selective encoding can be studied by manipulating how valuable it is for participants to

remember specific stimuli, for instance by varying the monetary reward participants receive

for recalling a particular stimulus in a subsequent memory test. It would be reasonable for

participants to strategically attend more to high-reward items compared to low-reward items

in mixed list contexts, but to attend both types of items equally in pure list contexts, where all

items are of equal value. Reward-enhanced memory may be driven by automatic

dopaminergic interactions between reward circuitry and the hippocampus and thus be

insensitive to list context; or it may be driven by meta-cognitive strategies, and thus context-

dependent. We contrasted these alternatives by manipulating list composition, and tracked

selective encoding through multiple EEG measures of attention and rehearsal. Behavioural

results were context-dependent, such that recall of high-reward items was increased only in

mixed lists. This result and aspects of the recall dynamics confirm predictions of the eCMR

(emotional Context Maintenance and Retrieval) model. The power of ssVEPs was lower for

high-reward items regardless of list composition, suggesting decreased visual processing of

high-reward stimuli and that ssVEPs may index the modulation of context-to-item

associations predicted by eCMR. By contrast, reward modulated the amplitude of Late

Positive Potential and Frontal Slow Wave only in mixed lists. Taken together the results

provide evidence that reward-enhanced memory is caused by an interplay between strategic

processes applied when high- and low-reward items compete for cognitive resources during

encoding and context-dependent mechanisms operating during recall.

1   Introduction
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The ability to selectively encode information that has relevance for future goals is a crucial

function of long-term memory. For example, it is more important to remember details of a

medical appointment than a casual conversation with a stranger on the bus to the hospital.

Selective encoding can be studied in the lab by manipulating how valuable it is for

participants to remember specific stimuli, for example by varying the monetary reward

participants receive for recalling a particular stimulus in a subsequent memory test.

Confirming prevalent intuitions, research shows that people remember information that

predicts future reward better than other information (for review see Knowlton & Castel,

2022).

It has been suggested that given our limited encoding capacity, increased memory

performance for reward-related information is due to the propensity to engage prefrontal

metacognitive encoding strategies to selectively encode valuable information and ignore less

important information when both are encountered in the same context (Cohen et al., 2014,

2017). Alternatively, reward-enhanced memory may be driven by automatic dopaminergic

interactions between reward circuitry and the hippocampus (Adcock et al., 2006; Bowen et

al., 2020; Gruber et al., 2016; Shohamy & Adcock, 2010) and occur regardless of whether

high reward and low reward information are encoded in the same or separate contexts. These

two views are not mutually exclusive and may operate together (Cohen et al., 2019), but

evidence for context dependence of reward-enhanced memory cannot be explained by

automatic reward processes alone. The first aim of this study was to investigate if reward-

enhanced memory is context-dependent by using a list-composition manipulation.

The list-composition effect refers to an interaction between how items are processed

during encoding and the composition of the encoding list (McDaniel & Bugg, 2008). List

composition is manipulated by having participants encode two types of items in the same list

context (mixed lists) or in separate list contexts (pure lists). It has been shown that emotion-enhanced memory, i.e. increased memory for items with negative valence compared to neutral valence, is dependent on list context. The effect is present when the items are studied together in mixed lists, but is decreased or disappear altogether in pure lists where negative and neutral items do not compete during encoding and retrieval (Barnacle et al., 2016, 2018; Talmi et al., 2007; Talmi & McGarry, 2012). Of relevance for the present study, we recently provided initial evidence that *reward*-enhanced memory also is context dependent (Hellerstedt & Talmi, 2022; Talmi et al., 2021). In the forementioned studies, we found that reward increased memory performance in mixed lists, where high and low reward items were encoded together, but not in pure lists where they were encoded separately. The first, pre-registered hypothesis (link to pre-registration: https://osf.io/cndzs/?view_only=0bc346eef0e9404a82d526b2db40f3fe) is that we will replicate this pattern of results here, using a larger sample and improved materials.

Strategic, meta-cognitive mechanisms could either operate during *encoding* or during *retrieval*. On the one hand, selective attention, selective rehearsal or semanticization (see for example (Cohen et al., 2014, 2017) could be applied during *encoding* to prioritise encoding of high-reward items. On the other hand, the list composition effect could be explained by competition during *retrieval*. One cognitive computational model that formalizes how list-composition effects could occur is the emotional Context Maintenance and Retrieval Model (eCMR, Talmi, Lohnas, et al., 2019) which has successfully modelled list-composition effects both in the context of emotion (Talmi, Lohnas, et al., 2019) and reward (Talmi et al., 2021). Similarly to previous computational models of list-composition effects (Shiffrin & Steyvers, 1997), this model explains emotion and reward-enhanced memory effects as the consequence of an interaction between encoding and retrieval processes. Specifically, in eCMR the prioritisation of high-reward items at encoding only yields an advantage in a free-

recall test when these items compete with low-reward items for retrieval. eCMR builds on the

Context Maintenance and Retrieval Model (CMR, Polyn et al., 2009), but extends this model

by implementing encoding priority as increased context-to-item associations in a two-layer

neural network. After encoding of a mixed list, when the memory test relies on the context to

cue retrieval (i.e. in a free-recall test, when no other cues are available), these stronger

associations promote the recall of items that were prioritised during encoding. Once a high-

reward item is retrieved, it will reactivate the high-reward context of its encoding, thereby

promoting the recall of items that share the same context (other high-reward items). A second

aim of this study was to test specific behavioural predictions of eCMR. Our second and third

hypotheses are that high-reward items are retrieved first in mixed lists, and that recall would

be clustered according to the reward condition. These hypotheses were not pre-registered, but

are derived directly from the eCMR model (Talmi, Lohnas, et al., 2019).

  A central assumption of the way eCMR simulates encoding of high-reward items is

that they are more tightly-bound to their encoding context, a mechanism considered to be a

computational implementation of context-independent increases in the attention allocated to

prioritised items, and to operate during encoding regardless of list composition. We recorded

EEG during encoding with the aim to test this assumption and more broadly to investigate the

neural mechanisms underlying reward-enhanced memory. We investigated EEG measures

that have been related to attention: steady-state visual evoked potentials (ssVEPs) and event-

related potentials (ERPs), including P3, LPP (Late Positive Potential). Our fourth, pre-

registered hypothesis is that the amplitude of ssVEP and ERP effects will be larger for high

than low reward items regardless of list composition, namely, observed in both mixed and

pure list conditions, reflecting list context-independent increased attention to high reward

items.

The ssVEP technique has been used extensively to study visual attention (for reviews see (Vialatte et al., 2010; Wieser et al., 2016). The technique involves flickering stimuli at a given frequency; increased visual attention is associated with increased ssVEP power in the flicker frequency over occipital electrodes. ssVEP amplitude is increased by reward (e.g. Grahek et al., 2021) and predicts subsequent memory (Martens et al., 2012). ssVEPs and ERPs have traditionally been used as indices of attention in separate literatures, but it is possible to obtain both measures in the same study (e.g. Hajcak et al., 2013; Jiang et al., 2018; Thigpen et al., 2018). The P3 component and the late posterior positivity (LPP) have both been related to increased attention during encoding. The P3 component is related to allocation of attention and reward processing (Polich, 2007; Sato et al., 2005) and has recently been shown to be higher for high- vs. low-reward items during encoding (Elliott et al., 2020). The P3 is a positive peak 250-500ms post stimulus presentation which is maximal over midline parietal electrodes (Polich, 2007). The LPP has traditionally been associated with allocation of attention to emotional compared to neutral stimuli (for review see Hajcak et al., 2010; Schupp et al., 2000, 2006), but it may also more broadly reflect stimulus significance and may thus be sensitive to reward (Hajcak & Foti, 2020). The LPP is a positive slow wave, which is maximal over centroparietal electrodes and onsets around 200-400ms after stimulus presentation and typically lasts as long as the emotional stimulus is presented (Hajcak & Foti, 2020; Schupp et al., 2006). Two studies reported that the emotional modulation of the LPP is insensitive to list composition, and one found that it was related to subsequent memory (Barnacle et al., 2018; Zarubin et al., 2020). In addition to the ERP effects related to attention we also investigated the frontal slow wave (FSW) ERP effect, which has been related to elaborative rehearsal strategies (Fabiani et al., 1990; Mangels et al., 2001). The FSW typically onsets late, around 1000ms post stimulus presentation and has a frontal maximum (Elliott et al., 2020). Crucially, because ssVEPs and ERPs, including the

FSW, are time-locked to stimulus presentation, their modulation by reward suggest that reward influences attention to and rehearsal of valued stimuli during stimulus presentation. Any one of these effects could implement reward-dependent tightening of the association between the context and item layer in eCMR.

The eCMR account of list-composition effects can be seen as an instantiation of the retrieval distinctiveness account (Geraci et al. 2013; McDaniel et al. 2005). Retrieval distinctiveness accounts of list composition effects attribute the effect to prioritization of reward items at encoding, which endows them with properties that render them more distinctive during retrieval, and therefore better retrieved than less-distinctive items during the recall of mixed, but not pure lists. In eCMR, increased context-to-item binding provides a specific mechanism that renders particular items (here, those associated with high reward) 'stronger' than 'standard' items (here, those associated with low reward). eCMR can also be contrasted with two other accounts of list-composition effects. The item-order account (McDaniel & Bugg, 2008) suggests that reward captures encoding resources otherwise dedicated to encoding temporal order, and therefore predicts that temporal order will be less well preserved in the recall of mixed lists and pure, high-reward lists compared to pure, low-reward lists. Although eCMR assumes that reward items are bound more strongly to their context, the portion of context that they are bound more strongly to is the partition that represents reward rather than time. In technical terms, in eCMR high-reward items are more strongly bound to the "source context" (Polyn et al., 2009), rather than the temporal context (Talmi, Lohnas & Daw, 2019), and consequently, it does not predict that the composition of the list or the reward value will influence temporal organisation. Therefore, if temporal organisation scores change as a function of list type and reward, and especially if the direction of change follow that predicted by the item-order account, this result will support the item-order account and challenge eCMR. We have carried out a similar analysis for the

emotion-dependent list-composition effect, which favoured the eCMR account, and therefore,

although we explore this again here, we have not registered this hypothesis. Finally, the

attentional borrowing account attributes list-composition effects to selective rehearsal of

high-reward items. These 'borrows' attention from other items in mixed lists but in pure lists

they cannot do so (Watkins, LeCompte & Kim, 2000; Slamecka & Katsaiti, 1987). Attention

borrowing can take place immediately upon item presentation and shortly after (akin to

massed rehearsal), or during the presentation of other items (akin to spaced rehearsal). While

the former is similar to the extra attention capture that eCMR already simulates, eCMR

cannot account for selective spaced rehearsal of high-reward items. Under natural encoding

conditions it is tricky to pinpoint exactly when people rehearse. However, if selective spaced

rehearsal is the only reason for list-composition effects in free recall, there should be no

effect of reward on EEG marker of attention in pure lists. This prediction contrasts directly

with the eCMR account of the list composition effect, and corresponds to our fourth pre-

registered hypothesis, as described above.

15   The present study was the first to investigate the neural mechanisms underlying the

16   context dependence of reward-enhanced memory. By combining multiple EEG measures of

17   attention and selective rehearsal the study tests the relative contribution of automatic reward

18   signals and meta-cognitive encoding strategies in reward enhanced memory and specifically

19   tests the assumption of increased attention during encoding of the eCMR model.

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21   Methods

Participants

The sample size was determined a priori based on effect sizes in Hajcak et al. (2013). Power

analysis using the WebPower package (Zhan & Mai, 2018) in R (v4.0.5; R Core Team 2021)

indicated that a sample size of 43 is needed to replicate their ssVEP main effect of emotion

($\eta^2_p = .203$) with .90 power at alpha = .05. The data was collected using a pre-registered

stopping rule stating that data collection would continue until the targeted sample size was

reached and that excluded participants based on pre-registered exclusion criteria would be

replaced. Fifty-one participants gave informed consent before participating in the experiment.

The participants were compensated with £25 for their participation plus the performance-

based reward earned in the experiment. The reward was calculated for two randomly selected

lists plus the practice list (the maximum total reward was £16.8). Eight participants were

excluded based on pre-registered criteria: three based on having less than 16 trials in one of

the four conditions after EEG pre-processing, three due to not having an ssVEP at 14.17Hz

when averaging over all four conditions, one participant due to a technical error and another

participant due to performing below the 75% threshold in the distractor task. The final sample

consisted of 43 participants (31 female, 11 male, 1 non-binary; M_{age} = 24.4, SD_{age} = 4.6). The

experiment was approved by the Cambridge psychology research ethics committee.

Materials

The stimuli used in the study were 144 images (585x410 pixels) obtained from a standardized

set of 260 stimuli (Snodgrass & Vanderwart, 1980). The assignment of picture to condition

was randomised. There were three different types of lists: pure high reward (two lists), pure

low reward (two lists) and mixed lists (four lists).

Two coloured frames surrounding the image were used to indicate the reward

condition. We used a photometer to control their luminance. The selected colours were the

following in RGB space, brown 101, 55, 0 with 100% opacity; blue 3,67, 223 with 86%

opacity. The assignment of colour to reward condition was counterbalanced across

participants. The pictures were transformed to grayscale and presented on a 19 inches CRT

monitor with a refresh rate of 85Hz.

Experimental Design

We used a 2 (Reward: High vs Low) by 2 (List composition: mixed list vs pure list) design.

There were nine blocks. The first block was a practice block. Four of the blocks had mixed

lists with both high and low reward items whereas the other four lists were pure and either

only had high reward items (two lists) or only low reward items (two lists). The assignment

of list type (mixed, pure/low reward, pure/high reward) to block was randomised.

Procedure

Each block of the experiment started with an encoding phase. In this phase, the participant

was shown 16 pictures sequentially in the centre of the screen. Each trial started with a

fixation cross with a randomised jittered duration between 3.5, 4 or 4.5 seconds. Next a

picture was flickered at 14.17Hz in the centre of the screen for 2047 ms. The picture had a

coloured frame (blue or brown) which indicated what reward condition it was associated

with. The participants were instructed that they would receive £0.5 for remembering pictures

from the high-reward condition (e.g. pictures with blue frames) and £0.05 for remembering

pictures from the low-reward condition (e.g. pictures with brown frames).

A one-minute distractor task followed the encoding phase. A math problem (e.g. 5 + 9

-3 = 11) was shown on the screen and the participants were asked to indicate via button

presses if the answer was true or false. We told the participants that their responses needed to

be 75% correct in this distractor task to get the monetary reward for the memory task. All

included participants reached this criterion.

A free recall test followed the distractor task. The participants had three minutes to

write down descriptions of the pictures they could recall at any order, using a keyboard. They

were encouraged to "use multiple words or phrases to describe one picture, e.g. 'spotty dog'

and 'cat lying down' as opposed to 'animal'; 'lettuce' and 'broccoli' as opposed to vegetable"

and to "be as specific as possible so we know which picture you are referring to. The

participants were encouraged to continue to try to recall until the end of the response time

window. The last response was on average entered after 111 seconds ($SD = 23$ seconds),

indicating that the participants had enough time respond.

Behavioural data analysis

Accuracy in the final memory test was coded manually by the experimenter (RH) who

compared participants' responses to the pictures shown in each list while being blind to

experimental conditions. Fifty-one percent of the data (22 out of the included 43 subjects)

were coded by a second coder and the percentage of correspondence between the two coders

were 98.2% for this proportion of the data. The pre-registered analysis of memory

performance utilised a generalised mixed effect model using the lme4 package (Bates et al.,

2015) in R. Memory performance was used as a dependent variable and there were two fixed

effects of Reward (high vs low) and List composition (mixed lists vs pure lists) and a random

intercept plus random slope effect of participant. The model formula was:

Memory Performance ~ 1 + Reward * List Composition + (1 + Reward * List

Composition | Participant).

The statistical significance of the fixed effects and their interaction was assessed using

an omnibus χ^2 Wald test as implemented in the car package (Fox & Weisberg, 2019). We

followed recommendations in (Brown, 2021) to handle convergence issues (i.e. we changed

the optimiser and increased the number of iterations for the memory performance GLMM).

According to hypothesis 2, high reward items are more likely to be recalled in the first

output position than low reward items. An analysis of this effect must control for the

expected increase in recall of high reward items. Therefore, for each mixed list, we permuted

the output positions of recalled items 10000 times, to estimate the chance level in our data

when accounting for the increase in recall for high reward items. We then compared the

chances of recalling a high-reward item in the first output position in the observed and the

permuted data with a one sample test (for a similar approach see Healey et al., 2019). A

Wilcoxon signed rank test was used due to that that the Shapiro-Wilk test of normality was

significant ($W = .838$, $p < .001$), suggesting deviation from normality.

Next, we tested whether the participants organised recall based on reward condition

(hypothesis 3). Again, we controlled for clustering that could occur by chance due to higher

memory performance in the high reward than in the low reward condition by permuting the

output order 10000 times. We calculated the reward-based organisation scores for each

participant in the observed and the permuted data using the EMBAM toolbox

(https://github.com/vucml/EMBAM) in MATLAB. Observed reward-based organisation

scores were compared to the mean score in the permuted data using a one sample test. A

Wilcoxon signed rank test was used due to that that the Shapiro-Wilk test of normality was

significant ($W = .935$, $p = .018$), suggesting deviation from normality.

Finally, to contrast eCMR and the item-order account, we used the EMBAM toolbox to

compute temporal organisation scores. We then conducted a one-way repeated measures

ANOVA with temporal organisation score as dependent variable List Type as independent

variable (mixed list, pure list high reward, pure list low reward). Given that eCMR predicts

no difference in temporal organisation scores between list types we complimented the

frequentist ANOVA with a Bayesian one-way repeated measures ANOVA conducted in

JASP (JASP Team 2022, version 0.16.3).

EEG recording and pre-processing

EEG data was acquired using a 128-channel high-density hydrocel Geodesic sensor net,

sampled at 500Hz using a Netamps 300 amplifier. The data was recorded during the covid-19

pandemic and the participants were wearing face masks throughout the experiment which

necessitated excluding the 15 electrodes covering the face. The EEG data was pre-processed

offline using the EEGLAB toolbox in MATLAB with a standard pipeline (see Hellerstedt et

al. 2021 for a study using a similar pipeline) and the steps and settings were pre-registered.

The data was recorded with Cz as reference and was re-referenced offline to the average of

the left and the right mastoid electrodes. Next, the data was high-passed filtered with an FIR

filter with a cut off at 0.1Hz and segmented into epochs starting 2000ms prior to stimulus

onset and ending 3000ms after stimulus onset. Channels and epochs with artefacts were

detected and rejected using customised threshold functions from the FASTER plugin (Nolan

et al. 2010). The criterium for rejection of channels were values outside ±3 for participant-

level z-transformed values. We used the standard settings in FASTER meaning that the

values used for epoch rejection were the amplitude range, variance and deviation from

channels mean and the values used for rejecting channels were each channel's mean

correlation with other channels, variance and hurst exponent, a measure of a signal's long-

range dependence. Next the data was submitted to independent component analysis and

independent components reflecting ocular artefacts were identified by visual inspection of

scalp topographies, time course and activation spectra and were discarded from the data by

back-projecting all but these components to data space. Rejected channels were interpolated

and a low-pass FIR filter was applied with a cut-off at 40Hz. Channels containing artefacts

within single epochs were detected using customised functions from the FASTER plugin.

The criterion for rejecting and interpolating a channel was that z-transformed values exceeded

±3 within a single epoch. We used the standard settings in FASTER meaning that the values

compared against the threshold were variance, median gradient, amplitude range and channel

deviation (Nolan et al., 2010). Epochs with amplitudes exceeding ±100 microvolts were

detected and rejected to clean out any remaining artefacts. Finally, the epochs were sorted

into the conditions. Mean trial numbers ranged between 28.9 and 29.3, out of 32, for the four conditions and all participants contributed with at least 16 trials per condition.

Focal ssVEP analysis

The procedure for the focal ssVEP analysis was pre-registered. The EEG data was baseline-corrected using a pre-stimulus interval of 500ms. Next, the power spectrum was computed for the time-window between 1000-2000ms post stimulus presentation using Fast Fourier Transform with a Hanning taper as implemented in the FieldTrip toolbox (Oostenveld et al., 2011) in MATLAB (ver. 2021a). Amplitude values were extracted between 14Hz and 15Hz, the stimulation frequency, from electrode Oz. The selection of time window and electrode of interest was based on where and when the ssVEP was maximal in time-frequency data when averaging over all four conditions as described in the pre-registration. Extracted ssVEP amplitude data were analysed using linear mixed effects models using the lme4 package (Bates et al., 2015) in R. The extracted ssVEP amplitudes were used as the dependent variable. We started out from a model including a fixed effect of Reward (High/Low) a fixed effect of List composition (Mixed list/Pure list) and random intercept + random slope effects of participant, but needed to reduce the complexity of the random effect structure to only include random intercepts due to convergence problems. The formula for the final model was:

ssVEP amplitude ~ 1 + Reward * List Composition + (1|Participant).

Global ssVEP analysis

Analysing EEG data from a selected time-window and region may overlook effects occurring at other scalp locations and time points. We therefore complemented the focal analysis with a pre-registered global data driven analysis of all time points of the trial, 0-2000ms after

stimulus presentation, and all 113 analysed electrodes and controlled for multiple

comparisons with nonparametric cluster-based permutation tests with the FieldTrip toolbox

(Maris & Oostenveld, 2007).

First, single trial EEG-data was submitted to time-frequency decomposition using

Fast Fourier Transform with Hanning taper. The decomposition was performed between 4

and 30Hz in frequency steps of 1Hz, time steps of 50ms and with a time-window length of

five cycles. Oscillatory power in each frequency band was normalised to dB scale using a

pre-stimulus baseline period between –700 to –200ms to avoid that the estimation of baseline

data points were affected by data in the post-stimulus time window in the stimulation

frequency (as in Allen et al., 2020; Hellerstedt et al., 2021).

Next, nonparametric cluster-based permutation tests were performed for data in the

stimulation frequency between 14-15Hz. In the first step of this analysis t-tests are performed

for every timepoint at every electrode. Significant spatially or temporally neighbouring

samples (including at least two electrodes) at alpha = .05 (uncorrected) were used to form

clusters and their t-values were summed to a cluster-level t-value. The observed cluster-level

t-value was then compared to the maximal cluster-level t-value obtained in 5000

permutations. The Monte-Carlo p-value was calculated as the proportion of the permutated

cluster-level t-values being larger than the observed cluster-level t-value. The exact

spatiotemporal distribution of the clusters should be interpreted with caution given that the

correction for multiple comparisons is applied at the cluster-level, but not at the individual

sample level (Sassenhagen & Draschkow, 2019). We tested the main effect of Reward, the

main effect of List composition and the Reward x List composition interaction.

Global ERP analysis

Two notch filters were applied to filter out the ssVEPs from the EEG data prior to ERP

analysis. The first filter had a lower cut-off at 14Hz and a higher threshold at 15Hz to filter

out the stimulation frequency and a second filter had a lower cut-off at 28Hz and a higher

cut-off at 29Hz to filter out the resonance frequency. The ERP data was baseline corrected

using a pre-stimulus interval between –200ms and 0ms. As for the global ssVEP analysis, the

global ERP analysis used nonparametric cluster-based permutation tests of all timepoints in

the trial from 0-2000ms and included all 113 electrodes and the analysis was pre-registered.

The settings were identical to the global ssVEP analysis.

Results

Behavioural Results

Figure 1A depicts the proportion of items recalled in each of the four list-composition by

reward conditions. We begin by reporting results that correspond to the pre-registered

hypothesis 1 and the pre-registered analysis. There was a reliable main effect of Reward,

($\chi^2(1) = 20.56$, $p < .001$) while the main effect of List composition was not reliable ($\chi^2(1) =$

1.57, $p = .21$). Importantly, the Wald test also indicated that the Reward x List composition

interaction was reliable ($\chi^2(1) = 17.24$, $p < .001$). Consistent with our pre-registered

prediction, follow-up pairwise contrasts showed that memory performance was higher for

high reward than low reward items in mixed lists (z-ratio = –5.200, $p < .001$), but not in pure

lists (z-ratio = –1.716, $p = .086$), indicating that reward enhanced memory depended on list

context. Parameter estimates of the model are presented in Table 1.

Next, we examined two additional hypotheses which were not pre-registered, but

derived from the eCMR model. Participants were more likely to recall high reward items first

in the mixed lists. To test whether this reflected a bias to recall high-reward items first, or

was a consequence of the overall increased recall of high-reward items, we compared

observed and permuted data. A one sample Wilcoxon signed rank test indicated that the

likelihood of retrieving a high reward item in position one was reliably higher in the observed

data ($M = .727$) than in the permuted data ($M = .547$; $V = 760$, $p < .001$; see Figure 1B).

These results support hypothesis 2, suggesting that participants were more likely to retrieve a

high reward item first, even considering the general effect of reward on recall. To test

hypothesis 3 we analysed the likelihood that output order was organised based on reward

condition. This analysis queries whether participants were more likely to retrieve a high

reward item in a given trial if they retrieved a high reward item in the preceding trial.

Confirming hypothesis 3, a Wilcoxon signed rank test showed that reward clustering was

greater than the estimated chance level (M observed $= .589$, M permuted $= .510$; $V = 702$, $p =$

.005; see Figure 1D).

Finally, we assessed the effect of reward and list composition on temporal

organisation scores (Figure 1C). This analysis examines whether participants were more

likely to retrieve items that they have studied consecutively during encoding. A one-way

repeated measures ANOVA indicated that there was no effect of list type (mixed list vs pure

list high reward vs pure list low reward) on temporal organisation scores ($F_{2,84} = .258$, $p =$

.773, $\eta_p^2 = .006$) and a Bayesian one-way ANOVA showed strong evidence for the null

hypothesis over the alternative (BF01 $= 10.792$). These results are in line with predictions

from eCMR and difficult to reconcile with the item-order account.

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Figure 1 Behavioural results. A) Proportion recalled items in the final memory test for the

four conditions.Scatter points indicate means for each participant. The larger black dots

displays the group means and the errorbars reflect the 95% confidence interval of the group

means. B) Proportion of high reward items recalled first in the mixed lists in the observed

data. The dotted line shows the mean proportion of high reward items recalled first in the

permuted data. C) Temporal clustering scores for the three list types. D) Clustering score in

the observed data. The mean clustering score in the permuted data is displayed as a

dotted line.

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ssVEP manipulation check

The frequency spectrum of the EEG was extracted for the whole trial (0-2000ms) averaged

over all conditions to check that the flicker manipulation had induced an ssVEP at 14.17Hz

using a fast fourier transform (FFT) with a Hanning taper. These data confirm that the flicker

manipulation successfully induced increased spectral power at 14Hz and in the resonance

frequency at 28Hz (Figures 2A, 2B and 2C). We next performed a pre-registered time

frequency analysis to detect what time window to use for the focal ssVEP analysis. In line

with previous research, this analysis showed that the ssVEP was maximal over occipital

electrodes and between 1000-2000ms post stimulus presentation (Figures 2C and 2D).

Figure 2 ssVEP Manipulation Check. A) Power spectrum of all conditions combined from

electrode Oz showing increased power in the stimulated frequency at 14Hz and at the

resonance frequency at 28Hz. B) ERP waveform averaged over all trials from Oz showing

the ssVEP at 14Hz. Note that the ssVEP was reducedwith notch filters prior to the statistical

analysis of ERPs. C) Time frequency plot averaged over all four conditions from electrode

Oz showing that the increase in power at 14Hz was maximal around 1000-2000ms. D)

Topographic map over time frequency data between 14-15Hz for the whole epoch averaged

over all conditions and averaged over the whole stimulus time window (0-2000ms).

Focal ssVEP analysis

Figure 3 depicts the pre-registered focal ssVEP analysis. Contrary to predictions, there was

no main effect of Reward, $\chi^2(1) = .60$, $p = .440$, or list composition $\chi^2(1) = 1.32$, $p = .250$,

and the interaction between Reward and List composition, $\chi^2(1) = .08$, $p = .782$ was also not

reliable, indicating that there were no effects of Reward or List composition on ssVEP power.

We had pre-registered a focal ssVEP subsequent memory analysis, to investigate if any

observed effects were related to memory, but we did not conduct this analysis due to low trial

numbers when dividing the EEG trials for the four conditions based on performance.

2   *Figure 3. Focal ssVEP analysis. The scatter dots depict individual ssVEP power between*

3   *1000-2000ms post stimulus presentation from electrode Oz for the four conditions. The*

4   *larger black dots reflects group means and the error bars shows the 95% confidence interval*

5   *of the group mean.*

6
Global ssVEP analysis

Figure 4 depicts the pre-registered exploratory global analysis. There was a reliable negative

cluster in the analysis in the main effect of Reward, showing larger ssVEP power for low

reward than for high reward (p = .003). The cluster was reliable between approximately 750-

1650ms and was maximal over occipital and parietal electrode sites. There was no reliable

main effect of list composition (all ps > .724) and no reliable Reward x List composition

interaction (all ps > .421). Although we have predicted to find a main effect of reward, the

direction of this effect was surprising and contradicted our pre-registered hypothesis, as

discussed in the Discussion section below.

As can be seen in Figure 2C, the experimental task was overall associated with a

reduction in alpha power around 10Hz and in beta power around 20Hz when averaging over

all conditions. We conducted control analyses to investigate if the unexpected reduction in

ssVEP power for high reward items compared to low reward items was driven by a global

reduction in alpha/beta power. If the effect was driven by global alpha/beta suppression, we

would expect to see a similar effect outside the stimulation frequency. To investigate this

possibility, we performed cluster permutation tests including all electrodes and all time points

for the alpha band (8-12Hz, $p = .169$) and the beta band (13-30Hz, $p = .152$), but found no

significant clusters in these frequency bands. We also tried to analyse narrow frequency

bands just below (11-13Hz, $p = .05$, which was not significant given the statistical threshold

of $p=.025$ for the two tailed test) and just above (15-17Hz, $p = .267$) the stimulation

frequency. The results suggest that the effect was most robust in the ssVEP frequency and

therefore, was unlikely to be driven by alpha/beta suppression. Figure 4B illustrates that the

decrease in power for high reward compared to low reward was maximal around the ssVEP

frequency.

Figure 4. Global ssVEP analysis. A) Time frequency data from electrode Oz between 14 and

15Hz for the four conditions. B) Time-frequency plot with the averaged power difference

between high reward and low reward from the five electrodes in the significant cluster in the

global ssVEP analysis. C) Topographic maps of power differences between 14-15Hz for the

time window where there was a reliable negative cluster for the main effect of reward.

Electrodes belonging to a significant cluster which are significant on average in the

displayed time window are highlighted with black stars.

Global ERP analysis

ERP results from the pre-registered global analysis are depicted in Figure 5. There was a

reliable positive cluster for the main effect of Reward (p = .009) between 1050-2000ms with

a right anterior/central distribution, indicating greater ERP amplitude for high reward than

low reward. There was no main effect of List composition (all ps > .130). Mirroring the

behavioural results, there was a reliable Reward x List composition interaction (p = .014)

which was reliable around 860-1320ms and had a posterior and central distribution. Follow-

up analysis showed that there was a reliable positive cluster around 590-1450ms within the

mixed lists, indicating more positive amplitude for high reward items than low reward items

with a widespread topography (p = .001). The effect has a posterior topography around 600-

1000ms fitting with an LPP effect (e.g. Barnacle et al., 2018; Hajcak et al., 2013; Zarubin et

al., 2020) while it has a more anterior topography after 1000ms fitting with the

spatiotemporal characteristics of an FSW effect (e.g. Elliott et al., 2020). The effect is more

long-lived than a typical P3 effect which is a peak rather than a slow wave, although there are

ongoing discussions regarding whether the P3 and the LPP are neural reflections of the same

processes and whether the duration of the effect may be determined by tasks characteristics

(Hajcak & Foti, 2020). There were no reliable clusters when comparing amplitude for high vs

low reward within pure lists (all ps > .837).

Figure 5. Global ERP analysis. A) ERPs from electrodes from two electrodes where the ERP

effects of interest typically are maximal, a frontal electrode (Fz) for the FSW and a

centroparietal electrode (CPz) for the LPP. B) Topographic maps of amplitude differences

from the time windows of the reliable clusters for the different contrasts. The cluster for the

follow up of the List Composition x Reward interaction, the bottom four plots, has been

divided into an early (590-1000ms) and a late (1000-1490ms) part to show the change in

topography over time. Electrodes from reliable clusters are highlighted with black stars if

they are reliable on average in the plotted time window.

Scalp current source densities (CSDs) were computed using the spherical spline method

(Perrin, Pernier, Bertrand & Echallier, 1989) as implemented in FieldTrip to further

investigate if there was support for an anterior and an posterior source for the reward ERP

effect within mixed lists. As depicted in Figure 6 the CSDs provided support for both a

posterior and an anterior source in the 590-1000ms time window (left plot). This is the time

window when the LPP typically is maximal, so this is in line with the interpretation that a

posterior LPP ERP effect and an anterior FSW ERP effect are present in this time window.

The FSW is typically maximal after 1000ms and the LPP is typically reduced then which also

matches the CSD pattern where the posterior source gets weaker while the anterior source

continues to be strong in the 1000-1490ms time window.

590-1000ms   1000-1490ms

Figure 6. Current source density plots from an early and a late time window from mixed lists

where there were significant differences between high- and low-reward ERP amplitudes.

Electrodes from reliable clusters are highlighted with black stars if they were reliable on

average in the plotted time window.

14   **Discussion**

The present study investigated whether reward-enhanced memory is context-dependent, and

the neurocognitive mechanisms underlying reward-enhanced memory. We also tested

predictions of the eCMR computational model of memory. All four behavioural predictions

of eCMR were supported by the data. By contrast, the EEG data challenged our proposed

neural implementation of a key eCMR mechanism. The ERP results showed for the first time

that the modulation of LPP and ssVEP by reward is context dependent, suggesting that they

reflect top-down controlled strategic processes that are applied when high and low reward

items compete for cognitive resources. In contrast to the ERPs, the ssVEPs differentiated

between high and low reward in a context-independent manner, potentially reflecting

increased focus on internal semantic representations relative to external visual perceptual

information when encoding high reward items compared to low reward items. Taken

together, the pattern of the EEG results show that LPP and ssVEPs are functionally

dissociated, suggesting that they measure different aspects of attention.

Reward-enhanced memory was dependent on list context, meaning that recall of items

that predicted reward was increased in mixed lists, but not in pure lists, conceptually

replicating two recent studies from our group (Hellerstedt & Talmi, 2022; Talmi et al., 2021)

despite a number of methodological differences (e.g. nature of the stimuli). The finding that

reward did not enhance recall in pure lists places an important constraint on any explanation

of reward-enhanced memory as solely a function of automatic encoding processes. It aligns

better with other explanations: either the eCMR model and other retrieval-distinctiveness

accounts, where encoding-retrieval interactions play a key role, or the suggestion that

automatic processes operate together with strategic processes to enhance memory for valued

information (Hennessee et al., 2019; Knowlton & Castel, 2022).

Retrieval distinctiveness accounts explain the list-composition effect as a

consequence of prioritised encoding, which is expressed differently during the retrieval of

mixed and pure lists (Gerarci et al. 2013; McDaniel et al 2005). eCMR instantiates this

account with a computational mechanism that makes predictions about recall dynamics.

Findings from the mixed list condition agreed with the recall dynamics that this

computational model predicted. In agreement with the prediction that the high reward items

are more tightly bound to the encoding context and therefore will be more likely to be

recalled first (Talmi, Lohnas, et al., 2019; Talmi et al., 2021), participants recalled high-

reward items earlier than low-reward items. While in previous studies of recall of mixed lists

of 'strong' and 'weak' items, including ours, the propensity to recall 'stronger' items early

was confounded with the increased average recall of such items, here we controlled for this confound yet still observed that high-reward items were recalled earlier than low-reward items. Additionally, we found that recall of mixed lists was organised based on reward condition. This finding confirms a second prediction of eCMR, that recalling a high reward item will change the context of the recall test to overlap more with the high-reward encoding context, and therefore, promote recalling further high-reward over low-reward items. Finally, we obtained evidence that neither reward, nor list composition, alters the temporal organisation of recall. This finding supports eCMR, where high-reward items are bound more tightly to their source context (the context partition that codes reward) but not their temporal context. By contrast, this finding challenges the item-order account of list composition effects (McDaniel & Bugg 2008; Nairne et al, 1991). While recall dynamics qualitatively align with predictions of eCMR, this is not conclusive evidence that this model is correct or detract from a potential role of meta-cognitive strategic encoding processes in reward-enhanced memory. For example, a model that formalises spaced rehearsal processes, e.g. selective rehearsal of high reward items in working memory across the entire presentation of mixed lists, may offer a better fit. EEG data offer additional insight into the contributions of automatic reward processes, meta cognitive processes, and the specific mechanisms of eCMR in reward-enhanced memory.

  ssVEPs offer an unobtrusive measure of the amount of visual attention participants allocate to items during encoding. eCMR embodies tighter associations between the context and highly-attended items regardless of the local context, and we have therefore pre-registered a hypothesis that if ssVEPs indexes this effect, ssVEP power will be increased in the high-reward condition. In contrast, the attentional borrowing account postulates that high-reward items borrow attentional resources or rehearsal time during encoding from low reward items when they compete for attention in mixed lists, but not in pure lists (Watkins,

LeCompte & Kim, 2000). The ssVEP results agreed with the prediction from eCMR in that they showed a context-independent effect of reward. However, surprisingly, we observed *a reduction* in ssVEP power for high-reward items, suggesting decreased visual processing of these stimuli. One plausible interpretation of this result is that participants focused more on rehearsal of semantic information rather than visual perceptual information, perhaps because they needed to describe the pictures verbally in the free recall test, and that semantization and rehearsal were more present for high reward items. If this interpretation is correct, then although the direction of the effect is the opposite of that predicted, the reward-dependent decrease in ssVEP power across both mixed and pure lists may indeed act as a neural marker that matches eCMR's predictions. Exactly how reward modulates ssVEPs may depend on the visual processing demands of the task. In two previous studies reward and subsequent memory increased ssVEP power, but these studies used motion detection (Grahek et al., 2021) and a recognition memory for visual shapes (Martens et al., 2012), both tasks which emphasized visual processing to a greater degree than here, where free recall of simple objects emphasized semantisation. While reward lowered ssVEP power here, it robustly increases ssVEP power in studies that employ negative emotional stimuli (Hajcak et al., 2013; Keil et al., 2008; Talmi, Slapkova, et al., 2019; Wieser et al., 2016), perhaps because such stimuli capture visual processing resources. At present, because the direction of the effect contradicts the pre-registered hypothesis, and its interpretation requires further research, we cannot consider the effect of reward on ssVEP markers of attention to offer strong evidence against the attentional borrowing account of list composition effects.

The ERP analysis showed that high reward items were related to more positive-going ERPs than low reward items, but only in mixed lists. The effect was widespread and included both centroparietal and frontal regions, so the spatial distribution overlaps with both an LPP and an FSW effect while it is more prolonged than a traditional P3 effect. The CSD analysis

provided additional support for the interpretation that there were two separate ERP effects, one posterior (LPP) and one anterior (FSW) that overlapped in time. These late and prolonged ERPs are typically interpreted as indicative of strategic cognitive processing of stimuli. The ERP effects may reflect top-down selective attention to high reward items in mixed lists, as indexed by the more posterior LPP and/or selective rehearsal of high reward items in working memory, as indicated by the more anterior FSW. The similarity between the pattern of the ERP and behavioural results suggests that selective processing of high-reward items in mixed lists is likely to contribute to the reward list-composition memory effect, consistent with the selective encoding account and the attentional borrowing account. The finding that reward is related to an FSW contrasts with a recent study by Elliot et al. (2020) which did not observe an FSW effect during encoding in a value-based memory paradigm. That study used a recognition test which is known to be associated to less strategic encoding compared to free recall tests (Cohen et al., 2017), which may explain the discrepancy in results.

Reward-enhanced memory had previously been related to a posterior P3/LPP like effect, but reward manipulations are associated with a number of potential neural and cognitive processes, which previous work could not distinguish. The ERP results in our study shows that they are only present in mixed lists and is therefore context-dependent. This finding contradicts ERP studies of the emotion-based list-composition effect, which have observed an emotion-dependent increase in LPP amplitudes in both mixed and pure lists (Barnacle et al., 2018; Zarubin et al., 2020). The discrepancy suggests that list-composition effects for reward and emotion rely on at least partially different mechanisms. It is likely that encoding strategies are employed more in reward experiments than in emotion experiments, given that participants are instructed to try to remember high reward information in reward experiments, while there is no instruction to prioritise negative items in emotion experiments,

and evidence that emotion attracts attention obligatorily (e.g. Pourtois et al. 2013; Talmi et al. 2007). Because emotional valence is intrinsic to the stimulus while reward needs to be associated with the stimulus, it is possible that top-down attention processes play a larger role in encoding of stimuli associated with reward. This interpretation agrees with the observation that the subsequent memory effect for emotional stimuli relies on regions associated with the bottom-up attention network, while the subsequent memory effect for neutral stimuli relied on the top-down attention network (Barnacle et al. 2016). The difference between the effect of reward and emotion may be best explained through the construct of 'task usefulness'. Emotional items may be chronically useful because of an evolved sensitivity to threat-related content. By contrast, indication that certain items predict high reward items may only be 'useful' to participants in mixed lists. The effect of context on reward-dependent but not emotion-dependent modulations of the LPP may hint that this component is sensitive to 'task usefulness', rather than reward or emotion *per se*, mimicking recent results where reward did not enhance recall once usefulness was controlled for (Chakravarty et al. 2019).

The ssVEP and the LPP have both been suggested to reflect increased attention to emotional materials in studies of emotional stimuli. Prior research has shown that the two measures have different neural generators. In the context of emotion, the LPP has been suggested to be related to activity in a broad network including both cortical and subcortical regions (e.g. Keil et al., 2002; Liu et al., 2012; Sabatinelli et al., 2007, 2013) while flicker-induced ssVEPs mainly are generated in primary visual cortex and adjacent higher-order cortices (e.g. Russo et al., 2007; Wieser & Keil, 2011). However, this evidence cannot rule out that both LPP and ssVEP are different proximal manifestations of the same distal appraisal of a stimulus as motivationally salient; for example, both scalp measures could be due to downstream effects of amygdala activation (Pourtois et al., 2013). Measuring ssVEPs and LPP simultaneously makes it possible to investigate if these two measures capture the

same aspect of attention. To the best of our knowledge, only one prior study has combined these two measures, but the authors used a very different orienting task of emotional attention, rather than the intentional encoding task in the present experiment (Hajcak et al., 2013). Nevertheless, even though Hajcak and colleagues found that negative emotion increased both the LPP effect and the ssVEP, the two measures were uncorrelated across participants, suggesting that they are measuring different aspects of attention to emotion. Our results provide further evidence for this possibility and extend it to effects of reward. Here, the two measures had a) opposite relation to reward (increased amplitude for the LPP in mixed lists and decreased power for the ssVEP) and b) differed in terms of the influence of list composition, where the LPP was list-context dependent while the ssVEP was not. Taken together, while both components may index increased attention, and may even be due to activations of the same driving influence, we show that they are functionally dissociated.

To conclude, the present study demonstrated that reward enhanced memory depends on list context. This result supports the prediction of eCMR as a computational model of reward-enhanced memory, but they also agree with selective encoding accounts. Going back to the question in the title of this article, our ERP results suggest that at least part of the context dependency of reward-enhanced memory is due to application of strategic processes to selectively encode high-reward items, but only in contexts in which they compete for cognitive resources with low-reward items. The selective encoding account is not a formal model, however, and does not make predictions for the dynamics of free recall. Because the dynamic pattern of recall matched the predictions of eCMR, our results suggest that the formal mechanisms this model embodies are useful to account for some of the effect of reward on free recall. Future computational work may allow retrieved-context models such as eCMR to account for strategic processes, something they are not presently able to simulate. Decreased ssVEP power in the high reward condition may be a neural marker of the tighter

associations eCMR predicts link the context and the high-reward items, but we offer this

interpretation cautiously, given the unpredicted direction of the effect, especially against

previous literature and our own pre-registered hypothesis. Interestingly, the pattern of the

LPP here is different from what have previously been observed in studies of list-context

dependence in emotion and suggest that top-down encoding processes may be more

important to explain list-composition effects for reward than for emotion. To test this

hypothesis, future research could manipulate both emotion and reward, and examine whether

the differences in their effects on the LPP become manifest even when they are measured in

the same experiment. More broadly, the discussion highlights a number of putative

differences between the influence of reward and emotion on the neural mechanism of

intentional encoding. If corroborated, different formal mechanisms may be necessary to

account for recalling different types of motivationally-significant experiences.

References

Adcock, R. A., Thangavel, A., Whitfield-Gabrieli, S., Knutson, B., & Gabrieli, J. D. E. (2006). Reward-Motivated Learning: Mesolimbic Activation Precedes Memory Formation. *Neuron*, *50*(3), 507–517. https://doi.org/10.1016/j.neuron.2006.03.036

Allen, J., Hellerstedt, R., Sharma, D., & Bergström, Z. M. (2020). Distraction by Unintentional Recognition: Neurocognitive Mechanisms and Effects of Aging. *Psychology and Aging*, *35*(5), 639–653. https://doi.org/10.1037/pag0000398

Barnacle, G. E., Montaldi, D., Talmi, D., & Sommer, T. (2016). The list-composition effect in memory for emotional and neutral pictures: Differential contribution of ventral and dorsal attention networks to successful encoding. *Neuropsychologia*, *90*, 125–135. https://doi.org/10.1016/j.neuropsychologia.2016.06.023

Barnacle, G. E., Tsivilis, D., Schaefer, A., & Talmi, D. (2018). Local context influences memory for emotional stimuli but not electrophysiological markers of emotion-dependent attention. *Psychophysiology*, *55*(4), e13014. https://doi.org/10.1111/psyp.13014

Bates, D., Mächler, M., Bolker, B., & Walker, S. (2015). Fitting Linear Mixed-Effects Models Using lme4. *Journal of Statistical Software*, *67*(1). https://doi.org/10.18637/jss.v067.i01

Bowen, H. J., Ford, J. H., Grady, C. L., & Spaniol, J. (2020). Frontostriatal functional connectivity supports reward-enhanced memory in older adults. *Neurobiology of Aging*, *90*, 1–12. https://doi.org/10.1016/j.neurobiolaging.2020.02.013

Brown, V. A. (2021). An Introduction to Linear Mixed-Effects Modeling in R. *Advances in Methods and Practices in Psychological Science*, *4*(1), 2515245920960351. https://doi.org/10.1177/2515245920960351

Cohen, M. S., Cheng, L. Y., Paller, K. A., & Reber, P. J. (2019). Separate Memory-Enhancing Effects of Reward and Strategic Encoding. *Journal of Cognitive Neuroscience*, *31*(11), 1658–1673. https://doi.org/10.1162/jocn_a_01438

Cohen, M. S., Rissman, J., Hovhannisyan, M., Castel, A. D., & Knowlton, B. J. (2017). Free Recall Test Experience Potentiates Strategy-Driven Effects of Value on Memory. *Journal of Experimental Psychology: Learning, Memory, and Cognition*, *43*(10), 1581–1601. https://doi.org/10.1037/xlm0000395

Cohen, M. S., Rissman, J., Suthana, N. A., Castel, A. D., & Knowlton, B. J. (2014). Value-based modulation of memory encoding involves strategic engagement of fronto-temporal semantic processing regions. *Cognitive, Affective, & Behavioral Neuroscience*, *14*(2), 578–592. https://doi.org/10.3758/s13415-014-0275-x

Chakravarty, S., Fujiwara, E., Madan, C. R., Tomlinson, S. E., Ober, I., & Caplan, J. B. (2019). Value bias of verbal memory. Journal of Memory and Language, 107, 25–39. https://doi.org/10.1016/j.jml.2019.03.005

Elliott, B. L., Blais, C., McClure, S. M., & Brewer, G. A. (2020). Neural correlates underlying the effect of reward value on recognition memory. *NeuroImage*, *206*, 116296. https://doi.org/10.1016/j.neuroimage.2019.116296

Fabiani, M., Karis, D., & Donchin, E. (1990). Effects of mnemonic strategy manipulation in

a Von Restorff paradigm. *Electroencephalography and Clinical Neurophysiology*,

75(1–2), 22–35. https://doi.org/10.1016/0013-4694(90)90149-e

Fox, J. & Weisberg, S.(2019). An R compantion to Applied Regression, Third Edition.

Sage.

Geraci, L., McDaniel, M. A., Miller, T. M., & Hughes, M. L. (2013). The bizarreness effect:

Evidence for the critical influence of retrieval processes. *Memory & Cognition*, 41,

1228–1237. http://dx.doi.org/ 10.3758/s13421-013-0335-4

Grahek, I., Schettino, A., Koster, E. H. W., & Andersen, S. K. (2021). Dynamic Interplay

between Reward and Voluntary Attention Determines Stimulus Processing in

Visual Cortex. *Journal of Cognitive Neuroscience*, *33*(11), 2357–2371.

https://doi.org/10.1162/jocn_a_01762

Gruber, M. J., Ritchey, M., Wang, S.-F., Doss, M. K., & Ranganath, C. (2016). Post-

learning Hippocampal Dynamics Promote Preferential Retention of Rewarding

Events. *Neuron*, *89*(5), 1110–1120. https://doi.org/10.1016/j.neuron.2016.01.017

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(7), e13570. https://doi.org/10.1111/psyp.13570

Hajcak, G., MacNamara, A., Foti, D., Ferri, J., & Keil, A. (2013). The dynamic allocation

of attention to emotion: Simultaneous and independent evidence from the late

positive potential and steady state visual evoked potentials. *Biological Psychology*,

92(3), 447–455. https://doi.org/10.1016/j.biopsycho.2011.11.012

Hajcak, G., MacNamara, A., & Olvet, D. M. (2010). Event-Related Potentials, Emotion, and Emotion Regulation: An Integrative Review. *Developmental Neuropsychology*, *35*(2), 129–155. https://doi.org/10.1080/87565640903526504

Healey, M. K., Long, N. M., & Kahana, M. J. (2019). Contiguity in episodic memory. *Psychonomic Bulletin & Review*, *26*(3), 699–720. https://doi.org/10.3758/s13423-018-1537-3

Hellerstedt, R., Moccia, A., Brunskill, C. M., Bowman, H., & Bergström, Z. M. (2021). Aging reduces EEG markers of recognition despite intact performance: Implications for forensic memory detection. *Cortex*. https://doi.org/10.1016/j.cortex.2021.03.015

Hellerstedt, R., & Talmi, D. (2022). Reward does not modulate forgetting in free recall tests. *Learning & Memory*, 29, 430-434. https://doi.org/10.1101/lm.053631.122

Hennessee, J. P., Patterson, T. K., Castel, A. D., & Knowlton, B. J. (2019). Forget me not: Encoding processes in value-directed remembering. *Journal of Memory and Language*, *106*, 29–39. https://doi.org/10.1016/j.jml.2019.02.001

JASP Team (2022). JASP (Version 0.16.3). Computer software.

Jiang, Y., Wu, X., Saab, R., Xiao, Y., & Gao, X. (2018). Time course of influence on the allocation of attentional resources caused by unconscious fearful faces. *Neuropsychologia*, *113*, 104–110. https://doi.org/10.1016/j.neuropsychologia.2018.04.001

Keil, A., Bradley, M. M., Hauk, O., Rockstroh, B., Elbert, T., & Lang, P. J. (2002). Large-scale neural correlates of affective picture processing. *Psychophysiology*, *39*(5), 641–649. https://doi.org/10.1111/1469-8986.3950641

Keil, A., Smith, J. C., Wangelin, B. C., Sabatinelli, D., Bradley, M. M., & Lang, P. J. (2008). Electrocortical and electrodermal responses covary as a function of emotional arousal: A single-trial analysis. *Psychophysiology*, *45*(4), 516–523. https://doi.org/10.1111/j.1469-8986.2008.00667.x

Knowlton, B. J., & Castel, A. D. (2022). Memory and Reward-Based Learning: A Value-Directed Remembering Perspective. *Annual Review of Psychology*, *73*(1), 25–52. https://doi.org/10.1146/annurev-psych-032921-050951

Liu, Y., Huang, H., McGinnis-Deweese, M., Keil, A., & Ding, M. (2012). Neural Substrate of the Late Positive Potential in Emotional Processing. *Journal of Neuroscience*, *32*(42), 14563–14572. https://doi.org/10.1523/jneurosci.3109-12.2012

McDaniel, M. A., & Bugg, J. M. (2008). Instability in memory phenomena: A common puzzle and a unifying explanation. *Psychonomic Bulletin & Review*, 15, 237–255. http://dx.doi.org/10.3758/PBR.15.2.237.

McDaniel, M. A., Dornburg, C. C., & Guynn, M. J. (2005). Disentangling encoding versus retrieval explanations of the bizarreness effect: Implications for distinctiveness. *Memory & Cognition*, 33, 270–279.

Mangels, J. A., Picton, T. W., & Craik, F. I. M. (2001). Attention and successful episodic encoding: an event-related potential study. *Cognitive Brain Research*, *11*(1), 77–95. https://doi.org/10.1016/s0926-6410(00)00066-5

Maris, E., & Oostenveld, R. (2007). Nonparametric statistical testing of EEG- and MEG-data. *Journal of Neuroscience Methods*, *164*(1), 177–190. https://doi.org/10.1016/j.jneumeth.2007.03.024

Martens, U., Gert, A. L., & Gruber, T. (2012). Influences of encoding and retrieval on the steady-state visual evoked potential. *NeuroReport*, *23*(6), 337–341. https://doi.org/10.1097/wnr.0b013e3283516542

McDaniel, M. A., & Bugg, J. M. (2008). Instability in memory phenomena: A common puzzle and a unifying explanation. *Psychonomic Bulletin & Review*, *15*(2), 237–255. https://doi.org/10.3758/pbr.15.2.237

Nairne, J. S., Reigler, G. L., & Serra, M. (1991). Dissociative effects of generation on item and order retention. *Journal of Experimental Psychology: Learning, Memory, and Cognition,* 17, 702–709.

Nolan, H., Whelan, R., & Reilly, R. B. (2010). FASTER: Fully Automated Statistical Thresholding for EEG artifact Rejection. *Journal of Neuroscience Methods*, *192*(1), 152–162. https://doi.org/10.1016/j.jneumeth.2010.07.015

Oostenveld, R., Fries, P., Maris, E., & Schoffelen, J.-M. (2011). FieldTrip: Open Source Software for Advanced Analysis of MEG, EEG, and Invasive Electrophysiological Data. *Computational Intelligence and Neuroscience*, *2011*, 156869. https://doi.org/10.1155/2011/156869

Perrin F., Pernier, J., Bertrand O, & Echallier, J.F. (1989). Spherical splines for scalp potential and current density mapping. *Electroencephalography and Clinical Neurophysiology.* 72, 184-187. https://doi.org/10.1016/0013-4694(89)90180-6.

Polich, J. (2007). *Updating P300: an integrative theory of P3a and P3b. 118*(10). https://doi.org/10.1016/j.clinph.2007.04.019

Polyn, S. M., Norman, K. A., & Kahana, M. J. (2009). A Context Maintenance and Retrieval Model of Organizational Processes in Free Recall. *Psychological Review*, *116*(1), 129–156. https://doi.org/10.1037/a0014420

Pourtois, G., Schettino, A. & Vuilleumier, P. Brain mechanisms for emotional influences on perception and attention: What is magic and what is not. *Biol Psychol* **92**, 492–512 (2013). https://doi.org/10.1016/j.biopsycho.2012.02.007

R Core Team (2021). R: A language and environment for statistical computing. R Foundation for Statistical Computing, Vienna, Austria. URL https://www.R-project.org/.

Russo, F. D., Pitzalis, S., Aprile, T., Spitoni, G., Patria, F., Stella, A., Spinelli, D., & Hillyard, S. A. (2007). Spatiotemporal analysis of the cortical sources of the steady-state visual evoked potential. *Human Brain Mapping*, *28*(4), 323–334. https://doi.org/10.1002/hbm.20276

Sabatinelli, D., Keil, A., Frank, D. W., & Lang, P. J. (2013). Emotional perception: Correspondence of early and late event-related potentials with cortical and subcortical functional MRI. *Biological Psychology*, *92*(3), 513–519. https://doi.org/10.1016/j.biopsycho.2012.04.005

Sabatinelli, D., Lang, P. J., Keil, A., & Bradley, M. M. (2007). Emotional Perception: Correlation of Functional MRI and Event-Related Potentials. *Cerebral Cortex*, *17*(5), 1085–1091. https://doi.org/10.1093/cercor/bhl017

Sassenhagen, J., & Draschkow, D. (2019). Cluster-based permutation tests of MEG/EEG data do not establish significance of effect latency or location. *Psychophysiology*, *56*(6), e13335. https://doi.org/10.1111/psyp.13335

Sato, A., Yasuda, A., Ohira, H., Miyawaki, K., Nishikawa, M., Kumano, H., & Kuboki, T. (2005). Effects of value and reward magnitude on feedback negativity and P300. *NeuroReport*, *16*(4), 407–411. https://doi.org/10.1097/00001756-200503150-00020

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*(2), 257–261. https://doi.org/10.1111/1469-8986.3720257

Schupp, H. T., Flaisch, T., Stockburger, J., & Junghöfer, M. (2006). Emotion and attention: event-related brain potential studies. *Progress in Brain Research*, *156*, 31–51. https://doi.org/10.1016/s0079-6123(06)56002-9

Shiffrin, R. M., & Steyvers, M. (1997). A model for recognition memory: REM—retrieving effectively from memory. *Psychonomic Bulletin & Review*, *4*(2), 145–166. https://doi.org/10.3758/bf03209391

Shohamy, D., & Adcock, R. A. (2010). Dopamine and adaptive memory. *Trends in Cognitive Sciences*, *14*(10), 464–472. https://doi.org/10.1016/j.tics.2010.08.002

Slamecka, N. J., & Katsaiti, L. T. (1987). The generation effect as an artifact of selective displaced rehearsal. *Journal of Memory and Language*, 26,589–607.

Snodgrass, J. G., & Vanderwart, M. (1980). A standardized set of 260 pictures: Norms for name agreement, image agreement, familiarity, and visual complexity. *Journal of Experimental Psychology: Human Learning and Memory*, *6*(2), 174–215. https://doi.org/10.1037/0278-7393.6.2.174

Talmi, D., Kavaliauskaite, D., & Daw, N. D. (2021). In for a penny, in for a pound: examining motivated memory through the lens of retrieved context models. *Learning & Memory*, *28*(12), 445–456. https://doi.org/10.1101/lm.053470.121

Talmi, D., Lohnas, L. J., & Daw, N. D. (2019). A Retrieved Context Model of the Emotional Modulation of Memory. *Psychological Review*, *126*(4), 455–485. https://doi.org/10.1037/rev0000132

Talmi, D., Luk, B. T. C., McGarry, L. M., & Moscovitch, M. (2007). The contribution of relatedness and distinctiveness to emotionally-enhanced memory. *Journal of Memory and Language*, *56*(4), 555–574. https://doi.org/10.1016/j.jml.2007.01.002

Talmi, D., & McGarry, L. M. (2012). Accounting for immediate emotional memory enhancement. *Journal of Memory and Language*, *66*(1), 93–108. https://doi.org/10.1016/j.jml.2011.07.009

Talmi, D., Schimmack, U., Paterson, T. & Moscovitch, M. The Role of Attention and Relatedness in Emotionally Enhanced Memory. *Emotion* **7**, 89–102 (2007). https://doi.org/10.1037/1528-3542.7.1.89

Talmi, D., Slapkova, M., & Wieser, M. J. (2019). Testing the Possibility of Model-based Pavlovian Control of Attention to Threat. *Journal of Cognitive Neuroscience*, *31*(1), 36–48. https://doi.org/10.1162/jocn_a_01329

Thigpen, N. N., Gruss, L. F., Garcia, S., Herring, D. R., & Keil, A. (2018). What does the dot-probe task measure? A reverse correlation analysis of electrocortical activity. *Psychophysiology*, *55*(6), e13058. https://doi.org/10.1111/psyp.13058

Vialatte, F.-B., Maurice, M., Dauwels, J., & Cichocki, A. (2010). Steady-state visually evoked potentials: Focus on essential paradigms and future perspectives. *Progress in Neurobiology*, *90*(4), 418–438. https://doi.org/10.1016/j.pneurobio.2009.11.005

Watkins, M. J., LeCompte, D. C., & Kim, K. (2000). Role of study strategy in recall of mixed lists of common and rare words. *Journal of Experimental Psychology: Learning, Memory, and Cognition*, 26, 239–245. http://dx.doi.org/10.1037/0278-7393.26.1.239.

Wieser, M. J., & Keil, A. (2011). Temporal Trade-Off Effects in Sustained Attention: Dynamics in Visual Cortex Predict the Target Detection Performance during Distraction. *Journal of Neuroscience*, *31*(21), 7784–7790. https://doi.org/10.1523/jneurosci.5632-10.2011

Wieser, M. J., Miskovic, V., & Keil, A. (2016). Steady-state visual evoked potentials as a research tool in social affective neuroscience. *Psychophysiology*, *53*(12), 1763–1775. https://doi.org/10.1111/psyp.12768

Zarubin, V. C., Phillips, T. K., Robertson, E., Swafford, P. G. B., Bunge, T., Aguillard, D., Martsberger, C., & Steinmetz, K. R. M. (2020). Contributions of Arousal, Attention, Distinctiveness, and Semantic Relatedness to Enhanced Emotional Memory: An Event-Related Potential and Electrocardiogram Study. *Affective Science*, *1*(3), 172–185. https://doi.org/10.1007/s42761-020-00012-y

Zhang, Z., Mai, Y. (2018). Webpower: Basic and Advanced Statistical Power Analysis. R package version 0.5.2. https://CRAN.R-project.org/package=WebPower.

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1 Table 1. Model parameters for memory performance GLMM and the targeted ssVEP LMM
2 analysis.
3

	Estimate	SE	z/t^*
Memory Performance GLMM			
Intercept	.75	.11	6.60
Reward	−.27	.06	−4.53
List composition Pure	−.05	.04	−1.25
Reward:List Composition Pure	−.17	.04	−4.15
ssVEP LMM			
Intercept	3.52	.37	9.32
Reward	.04	.06	.77
List composition Pure	−.07	.06	−1.15
Reward:List Composition Pure	−.02	.06	−.28

4
5 *Z for GLMM and t for LMM

6