Top-down specific preparatory activations for Selective Attention and Perceptual Expectations

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Summary

Proactive cognition brain models are mainstream nowadays. Within these, preparation is understood as an endogenous, top-down function that takes place prior to the actual perception of a stimulus and improves subsequent behavior. Neuroimaging has shown the existence of such preparatory activity separately in different cognitive domains, however no research to date has sought to uncover their potential similarities and differences. Two of these, often confounded in the literature, are Selective Attention (information relevance) and Perceptual Expectation (information probability). We used EEG to characterize the mechanisms that pre-activate specific contents in Attention and Expectation. In different blocks, participants were cued to the *relevance* or to the *probability* of target categories, faces vs. names, in a gender discrimination task. Multivariate Pattern (MVPA) and Representational Similarity Analyses (RSA) during the preparation window showed that both manipulations led to a significant, ramping-up prediction of the relevant or expected target category. However, classifiers trained on data from one condition did not generalize to the other, indicating the existence of unique anticipatory neural patterns. In addition, a Canonical Template Tracking procedure showed that there was stronger anticipatory perceptual reinstatement for relevant than for expected contents. Overall, results indicate that preparation during attention and expectation acts through distinguishable neural mechanisms. These findings have important implications for current models of brain functioning, as they are a first step towards characterizing and dissociating the neural mechanisms involved in top-down anticipatory processing.

Introduction

For decades, research in Cognitive Psychology has studied behavior while manipulating external factors, which led to theoretical models that framed cognition mostly from a reactive point of view. Recent years have witnessed a renaissance of proactive cognition, where endogenous top-down mechanisms play a core role in brain functioning. Within this framework, preparation is conceptualized as an endogenous function that takes place in anticipation of incoming inputs or demands and improves subsequent behavior (Battistoni et al., 2017; González-García et al., 2016). Neural preparatory activity has been shown for a plethora of processes, including attention (Battistoni et al., 2017; Kastner et al., 1999; Nobre & Serences, 2018), expectation (Aranda et al., 2010; Kok et al., 2017), working memory (Koshino et al., 2015; van Driel et al., 2017), or cognitive control (Baines et al., 2011; González-García et al., 2016; Hebart & Baker, 2018). Similarly, influential models have proposed different ways in which anticipatory patterns interact with stimulus inputs to guide perception. Examples of this are the Predictive Coding (Auksztulewicz & Friston, 2016a) or the Biased Competition frameworks (Desimone & Duncan, 1995). However, currently these models are silent as to how different preparatory phenomena relate to each other, and whether they reflect common or diverging underlying top-down mechanisms.

Selective Attention and Perceptual Expectation are complex functions that involve top-down and bottom-up elements. Attention refers to the selection of *relevant* information based on specific goals (Nobre & Serences, 2018), while expectation involves predictions based on prior *probability* (Schröger et al., 2015). Studies of selective attention have manipulated the relevance of information using cues that indicate the stimulus or dimensions to respond while ignoring others (Battistoni et al., 2017; Hong et al., 2017; Nobre & Serences, 2018; Stokes et al., 2009). Expectation has primarily (but not only, see e.g. Summerfield & De Lange, 2014) been manipulated with cues that inform about the most probable stimulus (de Lange et al., 2013; Kok et al., 2017; Wyart et al., 2012a). Previous research focusing on the effect of Attention and Expectations on target processing has shown both diverging and overlapping results. Although both lead to behavioral improvements (Ho et al., 2012; Stein & Peelen, 2015), several studies reveal that they can be, at least partially (Rungratsameetaweemana & Serences, 2019; Summerfield & Egner, 2009).

Studies that have tried to overcome frequent confounds between attention and expectation (e.g. Posner, 1980; Schröger, 1996)) have shown separate roles of relevance and probability during target processing (Auksztulewicz et al., 2017; Gordon et al., 2019; Simon et al., 2019; Wyart et al., 2012b; Zuanazzi & Noppeney, 2019). On the other hand, research focused on preparatory activity of either attention or expectation has provided seemingly overlapping results. Cues indicating relevance in selective attention (Battistoni et al., 2017; Nobre & Serences, 2018) preactivate relevant regions of

space processing (Giesbrecht et al.,
2006), specific shape patterns in
visual cortex (Stokes et al., 2009),
patterns in category (Esterman &
Yantis, 2010; González-García et
al., 2018) and object-selective
perceptual regions (Peelen &
Kastner, 2011; Soon et al., 2013).
Similarly, cues providing
probabilistic information lead to the
preactivation of specific templates
of oriented gabors (Kok et al.,
2017), direction (Ekman et al.,
2017), motor patterns (de Lange et
al., 2013) or abstract shapes (Hindy
et al., 2016). However, all these
previous investigations are agnostic
regarding the potential similarities
or differences in such top-down
preparation across relevance and
probability anticipation. Unraveling
the differences in how anticipatory
activity in different contexts reflects
the upcoming information is a
necessary step to understand the
differences between Attention and
Expectation, and is also essential for
theoretical models that explain the
neural basis of these two
phenomena (Auksztulewicz et al.,
2018; de Lange et al., 2018;
Desimone & Duncan, 1995).
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In our study, we employed different
multivariate pattern analyses of
EEG data to directly compare the
representational structure of
anticipated contents for top-down
preparation in contexts capitalizing
on Selective Attention (relevance)
or Expectation (probability). That
is, we examined whether the coding
of anticipated stimulus content
differs on the basis of the type of

Figure 1. Behavioral task and design (A) Behavioral task: example trial. Participants were cued about an incoming target stimulus (a face or a name) with which they performed a sex/gender classification task. **(B)** Block starting screen. In Attention blocks, the cue indicated the relevant stimulus. Participants performed the judgment on top of the screen only if the stimulus matched the cue. In non-relevant trials, participants responded with the "no" key. In Attention blocks the cues were not predictive of the probability of the target (50% faces vs. names) or the response. In Expectation blocks the cue indicated the probability of the stimulus category, which appeared 75% of times. Participants had to perform the judgment regardless of the cue (thus predicted and unpredicted targets were equally task-relevant), which carried no response information either. That is, in both blocks participants responded in all trials and perceptual details were fully equated. **(C)** MVPA classification rationale. In cue-locked within-block analyses, the classifier was trained to differentiate between anticipated (relevant or probable, depending on the block) faces vs. names with one pair of cues and then tested on the other, in either attention or expectation conditions. Note that classifiers could not use block information to differentiate between faces and words, as these were matched in every contrast. Across-blocks, the classifier was trained with one pair of cues within one condition (e.g., attention) and then tested on the other pair of cues within the other condition (e.g., expectation).

anticipation. To do so, we embedded a sex/gender discrimination in a cue-target paradigm. Here,
depending on the block, cues provided information about the upcoming relevance or probability
(Egner et al., 2010; Wyart et al., 2012b) of face or name stimulus categories. In addition, we ran an
independent localizer to study similarities between preparation and perception across contexts. Our
first goal was to study the anticipatory mechanisms during preparation by means of time-resolved
representational similarity analysis (RSA, Kriegeskorte, 2008). Then, we used multivariate pattern
analysis (MVPA, Grootswagers et al., 2017) to examine whether anticipated stimulus categories are
represented with differential fidelity during selected compared to probable targets. Next, we used a
cross-classification approach to directly contrast the patterns of activity underlying the representation
of relevant vs. probable stimuli. Taking into consideration the differences observed during target
processing in these contexts (Jiang et al., 2013; Kok et al., 2012; Wyart et al., 2012a), our hypothesis
was that anticipated relevant vs. probable stimuli would be differently coded. To further understand
the proposed differences, we leveraged a Canonical Template Tracking approach (González-García
et al., 2021; Wimber et al., 2015) to observe the extent to which preparation induces the reinstatement
of specific perceptual information in each condition. Given the dissociations between attention and
expectation observed during target processing (Gordon et al., 2019; Wyart et al., 2012) and the
apparent commonalities reported during their top-down preparatory states (e.g. Battistoni et al.,
2017), our overall hypothesis was that both manipulations would lead to the pre-activation of
anticipated contents but through at least partially different neural mechanisms.
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Methods

136   Methods are reported in accordance with the Committee on Best Practices in Data Analysis and
137   Sharing (COBIDAS) M/EEG (Pernet et al., 2019).
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139   Data and code availability
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141   All original code has been deposited at Github and is publicly available as of the date of submission
142   (github.com/ChemaGP-UGR/AttExpLoc_EEG). Results have been deposited at the OSF framework:
143   . Raw and preprocessed data are not available online due to storage size limitations, but are available
144   under request.
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146   1.   Participants
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Forty-eight participants (mean age = 22.06, range = 18-31; 29 women, 18 men, 1 non-binary) from
the University of Granada were recruited and received from 20 to 25 euros, depending on their
performance. Two additional participants were discarded due to low behavioral accuracy (less than
80%) or excessive noise in the EEG (more than 20% discarded trials). They were all native Spanish
speakers, right-handed with normal or corrected vision, and signed informed consent prior to
participation. Besides, to comply with COVID-19 guidelines, the temperature of participants was
measured upon arrival (always <37ºC), they confirmed to have had no illness symptoms in the days
prior to the experiment and wore a face mask during the whole session. Sample size was calculated
to achieve a statistical power of 80% for an estimated small effect size (Cohen's D = 0.3) and three

independent variables (block x category x cueing). Using PANGEA (Power aNalysis for gEneral ANOVA designs(Westfall, 2016) we obtained a minimum of 32 participants to be able to detect the block x cueing interaction in reaction times and behavioral accuracy, our main behavioral prediction. To fit the counterbalancing scheme, we tested 48 participants. This sample size provides an estimated power of 94% under the described parameters.

2. Apparatus, stimuli, and procedure

Stimulus presentation and behavioral data collection were done with The Psychophysics Toolbox 3 (Brainard, 1997) on MATLAB (v.2020a) in a Microsoft PC. Stimuli were presented on an LCD screen (Benq, 1920x1080 resolution, 60 Hz refresh rate) over a grey background. We employed 160 male and female faces (50% each, with ~6ºx9º visual angle, extracted from The Chicago Face Database(Ma et al., 2015)) plus 160 unique Spanish male and female names (50% each, with ~8ºx2º visual angle). Four different geometrical shapes (circle, square, rain-drop and diamond with thin black outlines, unfilled, ~2ºx2º visual angle) were used as cues in the main task. The sound stimuli employed in the localizer blocks consisted on four different tones (250, 300, 350 and 400 Hz).

The main task was a cue-target paradigm where, depending on the block, cues carried information about either the *relevance* (Attention) or the *probability* (Expectation) of upcoming face or word targets. Each trial started with the presentation of this visual cue. For each participant, and to avoid perceptual confounds, two cue shapes (counterbalanced across participants) were associated with faces and two with names. Importantly, cue pairs (the cue associated with faces and names) changed through the experiment. This way, the first cue for faces (e.g. a circle) appeared in half of the blocks with the first cue for names (e.g. a square) and the other half with the second cue for names (e.g. diamond). The task was to indicate the sex/gender of this target (male or female). Participants pressed one of two keys ("a", "l", counterbalanced across participants) to respond whether or not the target belonged to the gender stated at the beginning of each block. Half of the blocks belonged to the Attention condition, and the other half to the Expectation condition. Participants were verbally instructed to use the cues in the two blocks to respond as fast as possible while avoiding mistakes. At the beginning of each block (Figure 1B), they were informed about the block (Attention or Expectation), the target sex/gender ("Is the target male/female?"), and the two cues (one for faces and one for names). Importantly, and since Attention and Expectation are involved in almost any act of visual perception, we aimed at manipulating one process while keeping the other constant. In Attention blocks, the cue indicated the *relevant* stimulus category to select (either faces or names). Only if the stimulus belonged to the relevant category (50% trials, cued), the participant had to perform the gender discrimination task on the target. Otherwise, participants had to answer '*no*' regardless of the stimulus sex/gender (non-relevant category, uncued). Note that this manipulation of relevance, where further processing has to be applied only to selected stimuli, is similar to that employed in previous literature (e.g. Baldauf & Desimone, 2014b; Saenz et al., 2002; Summerfield et al., 2006; Womelsdorf et al., 2006). Therefore, participants had to give an answer to all Attention trials, and had to be prepared to perform the gender judgment task. Importantly, both relevant and non-relevant targets were matched in expectation, as by design they appeared with a 50% probability after each attention cue. On the other hand, in Expectation blocks the cue indicated the *probable* category of the target, with a 75% likelihood (e.g.(de Lange et al., 2013; Kok et al., 2017) for similar

201 manipulations). Here, participants had to perform the gender discrimination task in all trials, whether
202 or not the target was cued. This way, both the expected and unexpected targets were equally relevant.
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In every trial of the main task, the sequence of events was as follows: a 50 ms cue was followed by
a fixed Cue-Target Interval (CTI) of 1500 ms and then the target appeared for 100 ms. Trials were
separated by 2500 ms intervals. Auditory (tone, 400Hz, lasting for 300 ms) and simultaneous visual
feedback (words "Attention" or "Expectation", depending on the block presented for 500 ms)
appeared in case of a wrong answer 1.3 seconds after target presentation, without altering the trials
duration. Each trial lasted 4.15 seconds and each block 1.23 minutes. This main task was composed
by 32 blocks of 20 trials each, or 640 trials in total.
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In addition, Attention and Expectation blocks were interspersed with Localizer ones, used to measure
target perceptual processing without overt motor activity (adapted from Egner et al., 2010). In these,
the same stimuli as in the main task were presented. They were preceded by auditory cues that
predicted either faces or names with 75% validity. Similar to the main task, there were two tones
predicting faces, and two tones predicting names. One of each type was used per block. This
manipulation was not used for the analyses reported in this work. Participants had to press a key *only*
if the stimulus appeared upside down (rotated 180º, 10% of trials equated in stimulus category and
sex/gender), regardless of cue validity. At the beginning of each block, a screen indicated the type of
block and the two tones for the block. There were 16 localizer blocks of 40 trials each, 640 trials in
total. The cue lasted 200 ms, the visual stimuli appeared after an ISI of 1500 ms and stayed onscreen
for 100 ms. Trials were separated by 1500 ms intervals. Overall, each localizer trial lasted 3.3 seconds
and each localizer block lasted 2.12 minutes.
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225 The three types of blocks (Attention, Expectation and Localizer) appeared in a fully counterbalanced
226 order, so that that they preceded and followed each other an equal number of times. Cues and target
227 stimuli were also fully counterbalanced across participants. In total, the whole experimental session
228 lasted approximately 80 minutes, with additional practice and EEG preparation time.
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230   3.   EEG data acquisition and preprocessing
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232   3.1. Acquisition
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234 High-density EEG was recorded with 64 channels mounted on an elastic cap (actiCap Slim,
235 BrainVision) at the Mind, Brain and Behavior Research Center (CIMCYC) of the University of
236 Granada. Impedances were kept below 10 kΩ as recommended by the amplifiers' manufacturers.
237 EEG activity was referenced online to FCz and recorded at a sampling rate of 1000 Hz.
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239   3.2. Preprocessing
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Data were preprocessed using EEGLAB (Delorme & Makeig, 2004) and custom MATLAB scripts
(López-García et al., 2022). EEG recordings downsampled to 256Hz, digitally low-pass filtered using
FIR filter at 120 Hz and high-pass filtered at 0.1 Hz. A notch bandpass was applied at 50 Hz and 100

Hz to remove line noise and its harmonics. Channels with excessive noise were identified by visual
inspection and removed from the data (1.85% of channels on average, range 0-5). We used different
epochs for cue and target stimuli and in each we split the data into 3 seconds epochs (-1 to 2 seconds
after the onset of each stimulus). Then, Independent Component Analysis (ICA) was carried out to
remove eye artifacts (i.e. blinks and lateral eye movements) using the *runica* algorithm from
EEGLAB. Component rejection was guided by visual inspection of scalp maps, raw activity and
power spectrum. ICLabel was used for further confirmation. In average 1.85 components were
removed per participant (range 2-4). Next, we performed automatic trial rejection to prune the data
from non-stereotypical artifacts. It was based on three factors: 1) abnormal spectra: trials in which
the spectrum deviated from baseline by ±50 dB in the 0–2 Hz frequency window (sensitive to
remaining eye artifacts) or deviated by −100 dB or +25 dB in 20–40Hz (sensitive to muscle activity);
2) improbable data: the probability of occurrence of each trial was computed by determining the
probability distribution of values across trials, with a rejection threshold established at ±6 SD; (3)
extreme values: all trials with amplitudes in any electrode out of a ±150µV range were automatically
rejected (see (Keil et al., 2014; López-García et al., 2020, 2022) for similar preprocessing routines).
The three methods in sum yielded an average of 8% of rejected trials per participant (range 1.8%-
19%). Afterwards, the removed channels were repaired by spherical interpolation. We then applied
common average to re-reference the data, given its widespread use and its optimal adaptation to high-
density recordings(Pernet et al., 2018). Finally, trials were baseline corrected in the -200 to 0 ms prior
to stimulus onset.

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265   4.   Analyses
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267   4.1. Behavioral
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The main task design had three within-subject factors: Block type (Attention vs. Expectation), Cueing
(Cued vs. Uncued) and Stimulus category (Faces vs. Names). We calculated three-way repeated
measures ANOVA separately for behavioral accuracy and reaction times (RTs) employing JASP
(Love et al., 2019). For each participant, trials with longer or shorter RTs than the average ± 2 SDs
were discarded (11.54 % on average). Behavioral results in the localizer were not relevant and were
just considered as exclusion criteria in case of poor performance.

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276   4.2. Representational Similarity Analysis
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Representational Similarity Analysis (RSA) allows relating empirical multivariate measures of brain
activity to theoretical models (Kriegeskorte et al., 2008). We performed RSA using voltage values on
a subject-by-subject basis. Prior to the analyses data were normalized by z-scoring the values across
all trials, regardless of the condition. We then constructed empirical Representational Dissimilarity
Matrices (RDMs) every three time points, which measure the geometrical distances between all
experimental conditions (see MVPA section) and, finally, estimated the relationship between
empirical RDMs and theoretical models.

RDMs were built with data from eight conditions, yielding 8x8 symmetrical matrices. These
conditions were all the possible combinations between the design variables: *cue prediction* (faces and

names); *cue shape* (shapes 1 and 2 for faces, shapes 3 and 4 for names) and *block* (attention and expectation). We employed a Linear Discriminant Contrast (LDC, also known as Crossvalidated Mahalanobis Distance, (see Walther et al., 2016) as measure of distance between conditions, for the following reasons: it is a continuous measure, so it is highly reliable, informative, and lacks a ceiling effect; it includes a crossvalidation loop, which makes it less prone to biases; and it is centered around 0 when the true distance is 0 and therefore it is easier to interpret and more generalizable (Nili et al., 2014). We calculated LDC as described in Bueno & Cravo (2021). For every time point and each pair of conditions, we calculated the mean of each channel. This was done in two different datasets (train and test), to perform two-fold crossvalidation. The distance between the two conditions in the two folds was multiplied by the pseudo inverse covariance matrix between the residuals of the first and the second conditions in the training set, and the distance values were then averaged across the two folds.

Theoretical RDMs (Figure 3a) were built based on the expected distances (assigning values of 0 or 1) between conditions, according to different hypotheses. We built three such model RDMs based on: (1) cue shape (increased similarity between cues with same shape, regardless of the block and predicted target); (2) category (increased similarity between cues that predict the same stimulus category); and (3) block (increased similarity between cues belonging to the same block). The next step was to estimate the share of variance that each of these three model RDMs explained. To do so, at each time point we fitted a linear regression with the model RDMs as regressors and the empirical matrix as dependent variable. As a result, we obtained a t-value for each model, time point and participant that explained a significant *unique* portion of the variance, above that explained by the other regressors.

We used a non-parametric cluster-based permutation method to infer statistical significance at the group level, against empirical chance levels. First, at the single-subject level we randomly permuted the labels of the empirical matrices to generate 100 RSA maps per participant. Then, one map was randomly drawn per subject and the selected maps were averaged. This was repeated 10^5 times to obtain 10^5 permuted group RSA maps. For each time point, the empirical chance distribution was estimated. As expected, this yielded a t-value distribution centered around 0. The above and below thresholds were estimated so that they included the 99% of the distribution. Clusters containing time points with values outside the previously calculated thresholds were measured generating the null distribution of cluster sizes. Finally, we corrected for multiple comparisons using a False Discovery Rate (FDR) approach to determine the smallest cluster size deemed significant for α = 0.01 (López-García et al., 2022). Note that here permuted maps were obtained by randomly exchanging the values of the theoretical matrices.

Besides, to estimate the topography of each model we repeated the analyses for each channel separately, estimating the empirical matrix using all time points. That is, at each channel we fitted a linear regression with the theoretical models as regressors.

4.3. Time resolved Multivariate Pattern Analysis (MVPA)

We used time-resolved MVPA (Grootswagers et al., 2017) to study stimulus category-specific preparation by classifying faces vs. names. Note that we did not intend to directly classify attention vs. expectation, as this approach would be biased by existing block differences between these

conditions. Instead, classifiers were trained and tested to differentiate between faces and names within blocks, and comparisons between attention and expectation were always performed with this base contrast. Also, classifications were done *before* stimulus presentation with cue-locked EEG, where faces and names were anticipated as relevant or probable by the cue (targets were not relevant for this work and thus not analyzed). The steps of the classification were equal for all analyses unless otherwise specified and were performed with both voltage and time-frequency data. Classification was performed using MVPAlab (López-García et al., 2022) running on MATLAB. To maximize observations while reducing computational costs, we performed an MVPA classification every three time points. This way features used for classification were trials by channels matrices of raw voltage/frequency in single time points t_n.

We applied two strategies to increase signal-to-noise ratio. First, the trials used were the result of averaging sets of three trials (randomly selected) in each condition (Grootswagers et al., 2017). Then, we employed a smoothing method, based on a moving average filter with a length of 3 time points. For every t_n the features of the previous and the following data points were averaged, so that $t_n = (t_{n-1} + t_n + t_{n+1})/3$. Then, classes were balanced by subsampling the class with more trials so that the number of trials from the two conditions fed to the algorithm stayed the same (Grootswagers et al., 2017). A five-fold stratified cross-validation loop was implemented (King et al., 2013), which ensures that the proportion of each class stays balanced across folds, thus increasing the classification applicability to unknown data. Data were divided into five parts, which are enough to obtain unbiased results with a relatively small computational cost (Grootswagers et al., 2017; Varoquaux, 2018). The classification algorithm was trained with the first four divisions (training set) and tested on the remaining one (test set). This was repeated five times with the different sets. To improve the performance of the classifiers and the generalizability of multivariate analyses results (Singh & Singh, 2020), we normalized the data. Normalization was carried out within the cross-validation loop (King & Dehaene, 2014). Within each fold, we calculated the mean and standard deviation of each electrode across the training trials. The train set (X_{train}) and testing set (X_{test}) were normalized as follows:

$$X_{train} = \frac{X_{train} - \mu_{train}}{\sigma_{train}} \qquad X_{test} = \frac{X_{test} - \mu_{train}}{\sigma_{train}}$$

Where μ_{train} and σ_{train} are the mean and standard deviation of the training set.

We used Linear Discriminant Analysis as classification algorithm, given its good fit to typical EEG variability and higher sensitivity than similar methods (e.g. Support Vector Machines, see (Grootswagers et al., 2017; Kerrén et al., 2018a)). Time ranges were from -100 ms to 1550 ms. Classification results were estimated with an empirical receiver operative curve (ROC) analysis and reported as the area under the curve (AUC). This method works as an estimate of the true positive rate as a function of the false positive rate. The AUC is a non-parametric criterion-free method, so it does not involve assumptions about of the true distribution of the data (King & Dehaene, 2014). Besides, it is less susceptible to systematic biases and it is especially sensitive to two-class differences. AUC results can be interpreted similarly to classification accuracy, with 0.5 indicating equal probability of true positives and false positives and 1 accounting for perfect discriminability between classes (King et al., 2013). Statistical significance was calculated, as previously, by

369 performing cluster-based permutation analyses. Note that here we randomly permuted the labels of
370 each class, which yielded a distribution centered around chance level (0.5).

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372   4.3.1.   Temporal Generalization Analyses
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To characterize the changes of the signal throughout the temporal window, we employed a temporal
generalization approach (King & Dehaene, 2017). On each time point we trained a classifier following
the process described above. Then, we tested it on all time points of the preparation time window.
This rendered a Temporal Generalization Matrix (TGM) representing the AUC values for each train-
test pair. Statistical significance was then calculated for each TGM following the same rationale as
with time-resolved analyses. The only difference was setting the minimum statistically significant
threshold for cluster sizes to $p<0.001$, to avoid small clusters.

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4.3.2.   Category-specific anticipation within attention and expectation contexts

The process described was applied to the cue-locked EEG separately for Attention and Expectation
blocks, training classifiers to tell apart data from trials in which the cue anticipated faces vs. names.
In a first approximation, we used trials of all cue shapes in each category (Figure 1c). Next, to ensure
that classification results were not biased by perceptual differences between the geometrical shapes
used as cues, we implemented a classification approach across cues. We first trained the classifier
with data from only one pair of cues (e.g., classifying between squares predicting faces and diamonds
anticipating names) and then tested it on the other pair (e.g., testing with circles that predicted faces
and drops that predicted names). Because of the design, the selected training and testing pairs of cues
only appeared together in half of the blocks. This cross-classification ensures that only the common
differences between the two classified pairs will be decoded from the results (Kaplan et al., 2015),
thus removing perceptual confounds. We averaged the results of both directions and their permutation
maps to obtain a greater signal-to-noise ratio and to reduce biases due to the classification of specific
perceptual features. The averaged results were then fed to the same statistical algorithm used
previously to obtain cluster-based thresholds of statistical significance.

Once the results of these classifications were obtained, we compared the scores in Attention vs.
Expectation blocks. To do so, we subtracted the empirical results and the permuted maps of the two
conditions (Attention – Expectation). We used a two-tailed one sample t-test to evaluate differences
in the two directions (Attention > or < Expectation), following the same cluster-based permutation
implementation described above. Note that in this analysis the permuted distribution is centered
around zero.

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404   4.3.3.   Cross-classification between Attention and Expectation
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406 To estimate the degree to which patterns of brain activity are shared for preparation across Attention
407 and Expectation, we employed a cross-classification approach. We trained a classifier with data from
408 one condition and then tested it on the other. We first did this using data from all trials in each
409 condition, incorporating all four types of cues. Again, to rule out perceptual confounds we repeated
410 the analyses following the same rationale described in the previous section (see Figure 1c for a

summary of the condition selection strategies). In addition, we performed a control analysis to ascertain that cross-classification between Attention and Expectation blocks was feasible. Here we trained and tested using cues with the same physical form (e.g. train circle vs. square in attention, test circle vs. square in expectation) to observe whether the classifiers could extract the physical patterns of the cues even across overall changes in block demands.

4.4. Canonical Template Tracking

Finally, we compared the sustained patterns that arose during the preparatory interval with the actual perception of face and name stimuli. To do so, we obtained Canonical Template Patterns (González-García et al., 2021b; Wimber et al., 2015) of brain activity generated by faces and names in the independent localizer blocks. First, we performed an MVPA analysis in localizer trials following the same process as in previous analysis. Then, we selected the time window where the classifiers locked to the target stimuli in the localizer had the highest AUC (i.e., when activity patterns were more dissimilar) across participants (Figure S3a), which was 100 – 300 ms after stimulus onset. Then, and separately for face and name localizer trials, we averaged the raw information of every time-point and trial in the selected window for each channel and category. This resulted in a vector of 64 channel activity values for faces and another one for names in each participant (Kerrén et al., 2018b). Next, these CTP of faces and names were used as regressors in a linear regression where the dependent variable was the raw channel activity for each channel and condition during every time-point in the preparation or target window of the main experimental task. This rendered two t-values per time-point that accounted for the variance explained for each CTP (faces and names). We did this analysis separately in Attention and Expectation, for cues predicting faces and names and for face and name targets. To estimate statistical significance, we used the same cluster-based permutation analysis described above. Here it was not possible to change condition labels, since there were only two categories (faces and names) that were analyzed independently, and later subtracted. Hence, the only feature that allowed for permutations within each condition were channel labels. We shuffled the channel values separately for each CTP.

Results

Figure 2. Behavioral results. (A) Reaction times (in seconds) in Attention and Expectation blocks, for cued and uncued trials. (B) Accuracy in Attention and Expectation blocks, for cued and uncued trials. (C) RTs and accuracy for face and name targets, averaged across conditions. Dots represent individual subject (n=48) scores per experimental condition. Grey lines connect each participant's score in the two conditions of each block. The line inside boxes represents the median, while the limits of the box indicate the first and third quartile. Whiskers indicate the 1.5 inter quartile range for the upper and lower quartiles.

1.   Behavioral

Forty-eight participants completed a cue-target paradigm where, depending on the block, cues carried
information about either the *relevance* (Attention) or the *probability* (Expectation) of upcoming face
or word targets (Figure 1A). Analysis of participants' behavioral results showed that preparation for
the incoming target stimuli affected performance. A three-way repeated measures ANOVA on
reaction times (RTs) showed main effects of Block ($F_{47,1}=56.45$, p<0.001, ηp2 = 0.54), Cueing
($F_{47,1}=5.59$, p=0.022, ηp2 = 0.11) and Category ($F_{47,1}=50.52$, p<0.001, ηp2 = 0.52). Overall, responses
were faster in Attention (M = 569 ms, SD = 0.01) than Expectation (M = 598 ms, SD = 0.01) blocks.
Cueing affected RTs differently depending on the block (Block*Cueing, $F_{47,1}=9.07$, p=0.004, ηp2 =
0.16; Figure 2A). Post-hoc tests showed no effect of Cueing for Attention (t<1) whereas there was an
effect for Expectation ($t_{47,1}=3.6$, p<0.001, Cohen's d = 0.52). Expected (cued) targets elicited faster
responses (M = 592 ms, SD = 0.06) than unexpected (uncued) targets (M = 603 ms, SD = 0.07).
Stimulus Category also induced significant differences. Post-hoc tests showed faster responses for
faces (M = 576 ms, SD = 0.7) than for names (M = 591 ms, SD = 0.75), $t_{47,1}=7.51$, p<0.001, Cohen's
d = 1.08; Figure 2C.

The ANOVA on behavioral accuracy showed results in the same direction albeit less prominent. Both
Cueing ($F_{47,1}=4.33$, p=0.043, ηp2 = 0.01) and Category ($F_{47,1}=7.04$, p=0.011, ηp2 = 0.011) were
significant, and there was no main effect of Block (p>0.05), with both conditions presenting high
accuracy overall (Attention: M = 0.92, SD = 0.07; Expectation: M = 0.93, SD = 0.07). Again, we
found that cueing affected each block differently (Block*Validity, $F_{47,1}=9.67$, p=0.003, ηp2 = 0.15;
Figure 2B). Although there were no differences in Attention (t<1), there was an effect of expectations
($t_{47,1}=3.59$, p=0.003, Cohen's d = 0.52), with cued trials eliciting more accurate responses (M = 0.947,
SD = 0.026) than uncued ones (0.929, SD = 0.035). Post-hoc tests showed less accurate responses to
faces (M = 0.93, SD = 0.4) than to names (M = 0.94, SD = 0.37; Figure 2C), $t_{47,1}=2.65$, p=0.011,

Cohen's d = 0.38; Figure 2C. Overall, these results indicate that, as instructed, cues were used
effectively and also differently across blocks.

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468   2.   Time-resolved profile of preparation
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470 Our first aim was to assess the emergence of specific coding patterns linked to different information
471 content during preparation in Attention vs. Expectation contexts. The perceptual features of the cue

Figure 3. (A) RSA theoretical matrices and (B) results. In (A), black squares represent high similarity between conditions and grey squares represent low similarity. In (B), each line depicts t-values from a linear multiple regression fit for the three theoretical dissimilarity models, showing the unique share of variance explained for each of the factors. The colored straight lines above indicate significance clusters after a cluster-based permutation analysis. Grey shading indicates cue presence onscreen. The cue model (red) indicates dissimilarity between cue shapes, regardless of the block or prediction; the category model (green) indicates similarity between cues predicting faces or names; the Block model (blue) shows overall preparatory differences between Attention and Expectation. (C) Topographies of each model after using time points as features and repeating the analysis on each EEG channel.

472 should be processed, and their contextual meaning extracted to anticipate general target-category
473 information. This anticipated target category should be activated and maintained in working memory

474 to later provide an efficient response. To assess the contribution of each of these sources of
475 information to preparatory activity we employed model-based RSA (Kriegeskorte, 2008).
476

Results of the multiple regression for the three RSA models built (Cue shape, Category and Block,
see Methods section and Figure 3A) revealed the unique variance explained by each of the factors
entered in the analysis. A peak of the Cue shape model appeared first, at 160 ms after cue presentation,
and decayed fast afterwards (Figure 3B, red line). In contrast, the coding of the specific incoming
target category increased progressively along the interval, reaching its peak right before the
presentation of the actual target (Figure 3B, green line). In addition, the variance explained by the
blocks reached its peak around 500 ms after cue onset, slowly decayed along the interval, and fell
below significance levels 100 ms before target onset (Figure 3B., blue line).

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487   3. Time-resolved classification shows increasing AUC throughout the preparation interval.
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We then studied the coding of the anticipated specific information during the preparation interval,
separately for Attention and Expectation. Importantly, we did not compare cued vs. uncued targets,
but cues that predicted (relevant or probable, depending on the block) face vs. word stimuli within
the same block. Using the two cues of each category together for training and testing returned a

classification weighted on the perceptual
features of the cues (see Figure S5 for a
detailed description of the result), similar to
the Cue model in Figure 3 (red line).
Afterwards, we employed a cross-
classification analysis between different
cues to avoid cue perceptual confounds in
the classification (Figure 4). Since two
differently shaped cues coded for each type
of category, we trained the classifier in one
pair and tested it on the other, repeated the
process in the opposite direction and
averaged both. Similar to the Category
model in Figure 3, time resolved cross-
classification showed that the accuracy in
the decoding of the anticipated category
increased as the target onset approached.
We employed the subtraction approach
described in the methods section, but we
found no differences in classification
between Attention and Expectation (all
ps>0.05). We repeated a regression analysis

Figure 4. Raw voltage time-resolved cue decoding results. Result of the time-resolved classification of the category (faces vs. names) to be selected (blue) vs. expected (orange). Horizontal colored lines indicate statistical significance against chance within each block. Grey shading indicates cue presence onscreen. Figure shows the result of training and testing as described in Figure 1C (dotted line). For the results using all the cues together for training and testing see Figure S4.

515 to predict the cross-classification result of each participant based on each RSA model. As expected,
516 the category model (Figure 3, M = 0.35, SD = 2.37) explained the results better than any other. T-

517 values were nominally higher than the cue model (M = 1.04, SD = 1.42) and significantly higher than
518 the block model (p<0.01).
519
520　　　　4.　Relevance- and probability-driven preactivations are stable

Figure 5. Raw voltage temporal generalization results. (**A-B**) Temporal generalization results using the cross-cue decoding scheme, within (**A**) Attention and (**B**) Expectation conditions. Black outlines indicate statistical significance against chance. Black boxes indicate cue presence on screen. (**C-D**) Cross-condition classification results. Classification using the scheme described in Fig 1C, depicting the average of the two train and test directions. (**C**) Time-resolved classification of anticipatory information coding after training and testing the classifier in different conditions yielded no significant decoding. (**D**) Temporal generalization matrix. For visualization purposes, we averaged training and testing in the two directions (train attention and train expectation). The horizontal axis shows the times for training and testing in Expectation, and the vertical axis that of training and testing in Attention. That is, we inverted the axis of one of the directions before averaging. We chose this rationale because activity that is better represented in only one of the conditions should appear only on one side of the diagonal (e.g., above) when training and the opposite when testing (e.g., below). Averaging this way, we can show information that would not be visible doing a standard average. The non-averaged results for both directions can be found in the Supplementary Section (Figure S1). (**E**) A control cross-classification between Attention and Expectation (*Orange: Train Attention – Test Expectation. Purple: Train Expectation – Test Attention*) employing the same pairs of cues to train and test the classifier (e.g. train with circles in Attention, test with circles in Expectation) shows that classification across blocks is feasible. However, whereas the perceptual features of the cues generalize across Attention and Expectation (significance is marked with colored bars), the specific preactivation of contents before target onset does not.

While the previous results provide an initial characterization of neural coding of specific category
information during the preparation interval, they do not allow to explore the extent to which relevant
representations are stable during this time window, since significant decoding on different time points

could be driven in principle by different mechanisms. To investigate this, we employed a cross-time
decoding approach (King & Dehaene, 2014) to compare different patterns of brain activity across the
preparation interval. For this, we trained a classifier in one time point and then tested it on all the
points of the interval (Figure 5A, B). Results showed clear signs of generalization during the
preparation interval in both Attention and Expectation conditions (black outlines). The clusters of
activity grew increasingly larger up until the target's onset, indicating that the underlying patterns
remain relatively stable during the preparation period. We then compared the results of both analyses
following the same rationale as above: we subtracted the result matrices for both conditions and
performed a one-side T-test against 0. This analysis did not yield any significant results, indicating
that the accuracy of anticipatory category decoding was not different between conditions. Altogether,
these results suggest that preparation in both Attention and Expectation leads to a similar level of
discriminability of the anticipated (relevant or probable) category. However, this analysis is agnostic
regarding potential similarities in how anticipated relevant vs. probable information is coded, as
different underlying mechanisms could lead to similar accuracy results.
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539
540   5.   Attention and Expectation induce distinct patterns of preparatory activity
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To examine the mechanisms supporting the classification results, we used multivariate cross-
classification by training the classifier in one condition and then testing it on the other. Similar
patterns of brain activity should generalize between conditions, while differences should provide
chance-level classification. The number of observations employed for this, as well as the inter-block
temporal distance, were also matched. The averaged results (training in attention and training in
expectation) for raw voltage cross-classification are shown in Figures 5C, D (see Figure S1 for the
results split by train and test direction). Common coding between anticipating relevance vs.
probability of stimulus categories was scarce. Small significant clusters appear scattered through the
temporal generalization interval. This result complements the previous analysis by suggesting that
different neural mechanisms support the classification results. In addition, the control analyses
performed using the same physical cues to cross-classify between attention and expectation (see
Figure S2) shows that generalization across blocks is feasible, given that the classifier shows
significant above-chance performance to predict the perceptual shape of cues across attention and
expectation conditions.
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557   6.   Tracking perceptual patterns of brain activity
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We have shown that although Attention and Expectation lead to similar degrees of anticipatory
classification, their underlying neural patterns are partially different. We hypothesized that such
differences could arise due to the extent to which anticipatory representations function as perceptual
reinstatements of the prepared target categories. In this case, the preparatory patterns that allow to
classify anticipated faces and names should be similar to those triggered during actual stimulus
perception. To assess such reinstatement of target representations during preparation, we obtained
canonical template patterns (CTP; Palenciano et al., *in press*) of brain activity associated with faces
and names (as described in the Methods section). We then fit a linear regression with the two CTPs

as regressors to measure the extent to which these explained variance in preparatory activity across
conditions, in each time point (see Figure 8).
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First, to complement the behavioral results regarding face and name processing, we applied the CTP
to EEG activity locked to target onset in Attention and Expectation (Figure S3B). The face CTP
explained better the perception of target faces, while same was true for the name pattern on name
perception. Moreover, face processing was more pronounced, but decreased more rapidly. Name
processing, on the other hand, reached a lower peak but it decreased more slowly over time. As
expected, these results further suggest that faces and names are processed differently.
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Then, we compared how these canonical templates overlapped with preparatory activity in Attention
vs. Expectation contexts. Since both CTPs similarly explained face and name predicting cues, here
we show the average result of the two types of cues separately in Attention and Expectation blocks.
Crucially, both perceptual templates were better predictors of category anticipation in the Attention
compared to the Expectation condition (see Figure 6). Overall, these results indicate that for both
faces and names, regardless of their intrinsic differences in processing, there is an overall stronger
perceptual reinstatement for Attention than for Expectation. Note, however, that this stronger coding
cannot be explained by overall increased difficulty, as responses were faster in attention compared to
expectation blocks (see Behavioral Results section).
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Figure 6. Canonical Template Tracking results. Grey shading represents cue on screen. **(A-B)** CTP results for separate cues for Attention and Expectation. Average of face and name cues **(A)** Name CTP model for cues. **(B)** Face CTP for cues. Horizontal lines depict significance. Black horizontal lines indicate points where the results for Attention were significantly higher than for Expectation. Grey horizontal lines show higher perceptual reinstatement for Expectation.

Discussion

In the present work we studied and contrasted, for the first time, the top-down neural mechanisms
engaged during category-specific preparation in two major information-processing contexts:
Selective Attention and Perceptual Expectations. Our results reveal that the anticipated stimulus
category is active during preparation, regardless of the specific demands. Crucially, we demonstrate
that relevance and probability influence the preparatory patterns of brain activity in a unique manner,
potentially via differences in the perceptual reinstatement of anticipated information. Overall, these
results have important implications for theoretical models that explain how the brain anticipates
relevant vs. probable forthcoming information.
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Our behavioral results show that the paradigm manipulated Attention (relevance) and Expectation
(probability) in an effective manner and as expected (Baldauf & Desimone, 2014b; Kok et al., 2017).
Cued trials in Attention required further processing than those uncued, as participants were required
to evaluate the sex/gender of the cued stimulus category and skip this judgment in uncued ones
(Baldauf & Desimone, 2014a; Saenz et al., 2002; Summerfield et al., 2006; Womelsdorf et al., 2006).
Importantly, for Attention the probability of either category was equal (50%), keeping expectation
constant for relevant and irrelevant trials. The high accuracy in Attention blocks indicates that the
cues were used effectively to select when needed and to respond. On the other hand, both expected
and unexpected trials required selecting the forthcoming target, and thus equated relevance while
leading to predicted and unpredicted stimulus categories. In line with validity effects repeatedly
reported in the literature in expectation contexts, the efficiency of behavior increased in cued trials
(de Lange et al., 2018; Sy et al., 2014). It may be important to note that our results are at odds with
the notion that expected stimuli gain higher task-relevance and thus receive more attention that
unexpected ones (see Rungratsameetaweemana & Serences, 2019), as this interpretation would lead
to expect similar underlying anticipatory patterns for our attention and expectation manipulations.
612
We employed RSA (Carlson et al., 2019; Kriegeskorte, 2008) combined with multiple regression to
study the temporal profile of information coding throughout the preparatory interval. This showed
that anticipatory cues triggered several time-overlapping, yet distinguishable, effects in time (Figure
3B) and spatial arrangement (Figure 3C). The perceptual characteristics of the cue were coded early
in the time interval. Immediately after, and during the majority of the interval and until around 100
ms before target onset, mechanisms pertaining to contexts capitalizing on either selective attention or
expectations were deployed, as revealed by the Block model temporal profile. Evidence of
anticipatory coding of different task sets has been found before (González-García et al., 2021a; Hebart
et al., 2018; Palenciano et al., 2018). Interestingly, this result cannot be explained by motor
preparation, since cues did not predict motor responses; instead, it is most likely due to the control
context in each scenario. Finally, the variance explained uniquely by the Category model on
anticipatory activity increased steadily and peaked before target onset. This pattern of emerging,
ramping-up category representations could reflect two non-exclusive anticipatory mechanisms. Once
the meaning of the cue is extracted and stored in working memory, the actual anticipation of a
stimulus could be facilitated by top-down preactivation of perceptual regions (Auksztulewicz &
Friston, 2016b; Trapp et al., 2016). Simultaneously, the effect of temporal expectations (as the
preparatory interval was fixed) could induce an increasing preactivation of perceptual regions as the
stimulus onset approaches (Jin et al., 2020; Rohenkohl et al., 2012). In either case, these results
suggest that preparation engages a series of mechanisms that act both sequentially and in parallel,

including the processing of bottom-up signals that are transformed into top-down category
anticipation in a context of relevance or probability.

The block model in the RSA suggested that Attention and Expectation led to distinguishable coding
patterns. One mechanistic explanation for this result is that the patterns for the anticipated category
are more robustly coded in one condition over the other. Hence, we employed an MVPA approach
(Grootswagers et al., 2017; Haxby et al., 2014) to classify the prepared category. Crucially, classifiers
did not directly compare Attention vs. Expectation blocks (a discrimination that would be biased by
differential block characteristics) nor cued vs. uncued targets (which would mix reactions to matched
vs. mismatched predictions). Instead, classifiers were trained and tested to differentiate between cues
that predicted (relevant or probable) faces vs. names, within each block. Results matched the RSA
model, pointing to a robust effect of category anticipation. Furthermore, this prediction increased
towards the end of the interval. Again, this resonates with literature on temporal anticipation (Barbosa
et al., 2020; Jin et al., 2020; Ruz & Nobre, 2008). Moreover, prominent theories of attention
(Mongillo et al., 2008; Trübutschek et al., 2017) pose that information can be held in WM by
strengthening the synaptic weights between neurons, allowing for shifts (in this case, steady
increases) in decoding results during the delay period without losing information about the maintained
stimulus. The results revealed no significant differences in AUC between the conditions in either
time-resolved or temporal generalization matrices, and suggest that anticipated targets are coded with
similar fidelity during Expectation and Attention. Importantly, these results were obtained cross-
classifying different pairs of cues, thus avoiding perceptual confounds. However, a similar degree of
accuracy classification does not warrant identical underlying mechanisms. To test this, we performed
multivariate cross-classification between Attention and Expectation. Surprisingly, despite well
above-chance decoding within each type of block, there was little cross-classification between
conditions. Importantly, classification scores were similar when training and testing in different
blocks of the same condition (see Figure S2). In addition, a control analysis using the same cues
showed that generalization across blocks was possible and not prevented by the differential block
context. Classifiers trained and tested in different blocks were able to discriminate the perceptual
nature of the cues, whereas the anticipated category cross-classification was absent (see Fig. 5).
Overall, this set of results suggests that whereas Attention and Expectation both lead to anticipatory
category representations, their top-down mechanisms are partially different, providing further support
to the dissociation of relevance and probability, reported by previous studies, during target processing
(Summerfield & Egner, 2009; Wyart et al., 2012a).

One possible explanation for the differences found is the degree of preparatory perceptual
reinstatement (Kerrén et al., 2018b; Kilner et al., 2007; Muckli et al., 2015; Rose et al., 2016; Smith
& Muckli, 2010; Vetter et al., 2014) in Attention and Expectation. We employed a Canonical
Template Tracking procedure (González-García et al., 2021a; Wimber et al., 2015,Palenciano et al.,
in press) to compare perceptual reinstatement in attention and expectation. We obtained canonical
representations of the two target categories (faces and names) from an independent localizer, and then
estimated the variance explained by these perceptual specific patterns during the anticipatory window.
First, we applied the extracted canonical templates of the actual target processing, with results
supporting that faces are processed differently from words. Importantly, when applied to anticipatory
activity, we found that the canonical templates related differently to the preparation interval,

regardless of the predicted category. Our results showed that the perceptual reinstatement for both
faces and names was higher in Attention than in Expectation. Note that this cannot be due to a higher
difficulty of the attention blocks, as RT showed the opposite pattern (faster responses in Attention).
Instead, this higher reinstatement may be due to Attention directing resources to further activate the
relevant category, making it more relevant than the ignored one. This would be in line with the Biased
Competition model (Battistoni et al., 2017; Desimone & Duncan, 1995; Feldman & Friston, 2010),
that states that during target processing more resources are allocated to relevant stimuli, to enhance
their cortical representation. Crucially, our results extrapolate this finding to the previous top-down
phase, and show that this preparatory reinstatement is larger for relevant than for probable targets. A
further strength of the scope of this finding is that the task presented a large pool of faces and words
instead of repeating a small set of stimuli, and thus we induced proper category preparation, instead
of reinstatements of concrete stimuli.

How Attention and Expectation affect perception is an ongoing debate. Several studies have used
paradigms that combine both processes to study how they interact. Attention has been suggested to
sharpen the differences between expected and unexpected stimuli (Jiang et al., 2013), possibly
changing the oscillatory profile of relevant categories (Auksztulewicz et al., 2017) while reversing
repetition suppression (Kok et al., 2012). Relatedly, Attention boosts signal-to-noise processing by
suppressing noise, while Expectation increases baseline activity in perceptual regions (Wyart et al.,
2012; see also Gordon et al., 2019; Rungratsameetaweemana & Serences, 2019). Predictive coding
accounts propose that Attention increases prediction error of selected stimuli by suppressing noise of
unattended categories, while expectation increases global sensitivity through prediction signals.
Crucially, our results extend this literature by showing anticipatory differences between conditions,
which cannot be accounted for by prediction error differences as targets have not been processed yet.
Speculatively, attention could bias anticipatory neuronal sensitivity by increasing perceptual
differences between relevant categories, coding these changes at least partially in the gamma band
perhaps through anticipatory biasing of error processing units. Expectation could increase sensitivity
to probable categories by increasing excitability of, perhaps, perceptual units.

Although overall our results are a crucial first step to characterize mechanisms across relevance and
probability anticipation, they should be complemented by further studies. First, although the results
are statistically significant after robust cluster-based correction, accuracy values are lower than those
obtained, for example, using target-locked data (e.g., Figure S4). Importantly, decoding accuracies
do not equal effect sizes (Hebart & Baker, 2018). The values we obtained are within typical
classification ranges when studying subtle neural patterns[9,81] (also see [87]) employing non-invasive
human neuroimaging. Future studies may obtain higher accuracies by increasing stimulus repetition
and thus reinforcing specific neural traces of the information anticipated. Second, it could be argued
that the diverging results between the two conditions shown here are not due to differences in stimulus
representation for relevance vs. probability, but to different task demands of the two blocks. However,
as the goal of the study was not to fully differentiate the neural mechanisms of attention and
expectation but to understand how anticipatory information is coded differently in contexts of
relevance vs. probability, we did not merely contrast the two blocks directly. Instead, the analyses
contrasted anticipated faces vs. words in conditions matched within blocks. In the same line, the
arguably potential higher relevance of probable contents could not drive results, as such increased

relevance would equal for predicted probable faces vs. names, and thus the classifier cannot rely on this information. Moreover, and to further show how processes that are similar do generalize across both blocks, we trained the classifier in a pair of cues in one block, and tested it in the same pair for the opposite block (Figure 5E). As expected, significant classification appeared for the cue identity, adding to the idea that block differences do not account for lack of cross-classification of anticipated relevant vs. probable information content.

Third, our scope was limited to the temporal domain, and questions arise regarding potential differences between brain regions. Internal predictions have been generally associated with the hippocampus (Aitken & Kok, 2022; de Lange et al., 2018; Hindy et al., 2016; Stachenfeld et al., 2017), which has a location that challenges EEG sensitivity. Further studies should employ more spatially sensitive techniques. Additionally, we focused on category-based preparation of faces and words, to facilitate having the same task across categories (sex/gender judgments). These two types of stimuli, although frequently combined in the literature (Alm et al., 2016; Amado et al., 2018; Dumas, 2015; Rose et al., 2016; Sperling et al., 2003) have a different spatial layout, which may have generated the anticipation of different spatial templates. Although this is not a confound in our task, as such difference is constant, it may have added a spatial component to the preparation. Finally, we focused our design on visual perception. Studies that have compared the effects of attention and expectation have used auditory and visual stimuli showing promising results (e.g. Jiang et al., 2013). It is likely that the mechanisms found here can be extended to other sensory modalities, but more research is needed to assess this idea.

Altogether, our results show that anticipatory representations are category-specific and also task dependent, and highlight the role of proactive processes that precede stimulus perception. These findings have important implications for current models of brain functioning. We show that instead of single, unitary top-down phenomena, the brain implements distinct modes of content-specific anticipation, which are tailored to task context, operate within partially separate frequency bands and involve different levels of perceptual reinstatement. Predictive coding and attention models that differentiate these processes during target processing should be extended to the preparatory interval, acknowledging the specificity and complexity of these top-down anticipatory phenomena.

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Author Contributions

761 Conceptualization, JMGP and MR; Data Collection, JMGP, DLG, and BAL; Analyses, JMGP and
762 DLG; Resources, JG and MR; Writing – Original Draft, JMGP; Writing – Review & Editing, CGG
763 and MR; Supervision, CGG and MR; Project Administration, MR; Funding Acquisition, MR
764

765 **Declaration of Interests**

766
767 The authors declare no competing interests.

References

Aitken, F., & Kok, P. (2022). Hippocampal representations switch from errors to predictions during acquisition of predictive associations. *Nature Communications*, *13*(1), 1–13. https://doi.org/10.1038/s41467-022-31040-w

Alm, K. H., Rolheiser, T., & Olson, I. R. (2016). Inter-individual variation in fronto-temporal connectivity predicts the ability to learn different types of associations. *NeuroImage*, *132*, 213–224. https://doi.org/10.1016/j.neuroimage.2016.02.038

Amado, C., Kovács, P., Mayer, R., Ambrus, G. G., Trapp, S., & Kovács, G. (2018). Neuroimaging results suggest the role of prediction in cross-domain priming. *Scientific Reports*, *8*(1), 1–10. https://doi.org/10.1038/s41598-018-28696-0

Aranda, C., Madrid, E., Tudela, P., & Ruz, M. (2010). Category expectations: A differential modulation of the N170 potential for faces and words. *Neuropsychologia*, *48*(14), 4038–4045. https://doi.org/10.1016/j.neuropsychologia.2010.10.002

Auksztulewicz, R., & Friston, K. (2016a). Repetition suppression and its contextual determinants in predictive coding. *Cortex*, *80*, 125–140. https://doi.org/10.1016/j.cortex.2015.11.024

Auksztulewicz, R., & Friston, K. (2016b). Repetition suppression and its contextual determinants in predictive coding. *Cortex*, *80*, 125–140. https://doi.org/10.1016/j.cortex.2015.11.024

Auksztulewicz, R., Friston, K. J., & Nobre, A. C. (2017). Task relevance modulates the behavioural and neural effects of sensory predictions. *PLoS Biology*, *15*(12), 1–27. https://doi.org/10.1371/journal.pbio.2003143

Auksztulewicz, R., Schwiedrzik, C. M., Thesen, T., Doyle, W., Devinsky, O., Nobre, A. C., Schroeder, C. E., Friston, K. J., & Melloni, L. (2018). Not all predictions are equal: 'What' and 'When' predictions modulate activity in auditory cortex through different mechanisms. *The Journal of Neuroscience*, *38*(40), 0369–18. https://doi.org/10.1523/JNEUROSCI.0369-18.2018

Baines, S., Ruz, M., Rao, A., Denison, R., & Nobre, A. C. (2011). Modulation of neural activity by motivational and spatial biases. *Neuropsychologia*, *49*(9), 2489–2497. https://doi.org/10.1016/j.neuropsychologia.2011.04.029

Baldauf, D., & Desimone, R. (2014a). Neural mechanisms of object-based attention. *Science*, *344*(6182), 424–427. https://doi.org/10.1126/science.1247003

Baldauf, D., & Desimone, R. (2014b). Neural mechanisms of object-based attention. *Science*, *344*(6182), 424–427. https://doi.org/10.1126/science.1247003

Barbosa, J., Stein, H., Martinez, R. L., Galan-Gadea, A., Li, S., Dalmau, J., Adam, K. C. S., Valls-Solé, J., Constantinidis, C., & Compte, A. (2020). Interplay between persistent activity and activity-silent dynamics in the prefrontal cortex underlies serial biases in working

memory. *Nature Neuroscience*, *23*(8), 1016–1024. https://doi.org/10.1038/s41593-020-
0644-4

Battistoni, E., Stein, T., & Peelen, M. V. (2017). Preparatory attention in visual cortex. *Annals of*
the New York Academy of Sciences, *1396*, 92–107. https://doi.org/10.1111/nyas.13320

Brainard, D. H. (1997). The Psychophysics Toolbox. *Spatial Vision*, *10*(4), 433–436.
https://doi.org/10.1163/156856897X00357

Bueno, F. D., & Cravo, A. M. (2021). Post-interval EEG activity is related to task-goals in
temporal discrimination. *PLoS ONE*, *16*(9 September), 1–19.
https://doi.org/10.1371/journal.pone.0257378

Carlson, T. A., Grootswagers, T., & Robinson, A. K. (2019). *An introduction to time-resolved*
decoding analysis for M/EEG. http://arxiv.org/abs/1905.04820

Christophel, T. B., Cichy, R. M., Hebart, M. N., & Haynes, J. D. (2015). Parietal and early visual
cortices encode working memory content across mental transformations. *NeuroImage*,
106, 198–206. https://doi.org/10.1016/j.neuroimage.2014.11.018

de Lange, F. P., Heilbron, M., & Kok, P. (2018). How Do Expectations Shape Perception? *Trends*
in Cognitive Sciences, *22*(9), 764–779. https://doi.org/10.1016/j.tics.2018.06.002

de Lange, F. P., Rahnev, D. A., Donner, T. H., & Lau, H. (2013). Prestimulus oscillatory activity
over motor cortex reflects perceptual expectations. *Journal of Neuroscience*, *33*(4), 1400–
1410. https://doi.org/10.1523/JNEUROSCI.1094-12.2013

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*(1), 9–21. https://doi.org/10.1016/j.jneumeth.2003.10.009

Desimone, R., & Duncan, J. (1995). Neural Mechanisms of Selective Visual Attention. *Annual*
Review of Neuroscience, 193–222.

Dumas, J. A. (2015). What Is Normal Cognitive Aging? Evidence from Task-Based Functional
Neuroimaging. *Current Behavioral Neuroscience Reports*, *2*(4), 256–261.
https://doi.org/10.1007/s40473-015-0058-x

Egner, T., Monti, J. M., & Summerfield, C. (2010). Expectation and surprise determine neural
population responses in the ventral visual stream. *Journal of Neuroscience*, *30*(49),
16601–16608. https://doi.org/10.1523/JNEUROSCI.2770-10.2010

Ekman, M., Kok, P., & De Lange, F. P. (2017). Time-compressed preplay of anticipated events in
human primary visual cortex. *Nature Communications*, *8*(May), 1–9.
https://doi.org/10.1038/ncomms15276

Esterman, M., & Yantis, S. (2010). Perceptual expectation evokes category-selective cortical
activity. *Cerebral Cortex*, *20*(5), 1245–1253. https://doi.org/10.1093/cercor/bhp188

Feldman, H., & Friston, K. J. (2010). Attention, uncertainty, and free-energy. *Frontiers in*
Human Neuroscience, *4*(December), 1–23. https://doi.org/10.3389/fnhum.2010.00215

Giesbrecht, B., Weissman, D. H., Woldorff, M. G., & Mangun, G. R. (2006). Pre-target activity in
visual cortex predicts behavioral performance on spatial and feature attention tasks.
Brain Research, *1080*(1), 63–72. https://doi.org/10.1016/j.brainres.2005.09.068

González-García, C., Flounders, M. W., Chang, R., Baria, A. T., & He, B. J. (2018). Content-specific
activity in frontoparietal and default-mode networks during prior-guided visual
perception. *ELife*, *7*, 1–25. https://doi.org/10.7554/eLife.36068

González-García, C., Formica, S., Wisniewski, D., & Brass, M. (2021a). Frontoparietal action-
oriented codes support novel instruction implementation. *NeuroImage*, *226*(October
2020). https://doi.org/10.1016/j.neuroimage.2020.117608

González-García, C., Formica, S., Wisniewski, D., & Brass, M. (2021b). Frontoparietal action-
oriented codes support novel instruction implementation. *NeuroImage*, *226*(October
2020). https://doi.org/10.1016/j.neuroimage.2020.117608

González-García, C., Mas-Herrero, E., de Diego-Balaguer, R., & Ruz, M. (2016). Task-specific
preparatory neural activations in low-interference contexts. *Brain Structure and
Function*, *221*(8), 3997–4006. https://doi.org/10.1007/s00429-015-1141-5

Gordon, N., Tsuchiya, N., Koenig-Robert, R., & Hohwy, J. (2019). Expectation and attention
increase the integration of top-down and bottom-up signals in perception through
different pathways. *PLoS Biology*, *17*(4), e3000233.
https://doi.org/10.1371/journal.pbio.3000233

Grootswagers, T., Wardel, S. G., & Carlson, T. A. (2017). Decoding Dynamic Brain Patterns from
Evoked Responses: A Tutorial on Multivariate Pattern Analysis Applied to Time Series
Neuroimaging Data. *Journal of Cognitive Neuroscience*, *29*(4), 677–697.
https://doi.org/10.1162/jocn_a_01068

Haxby, J. V., Connolly, A. C., & Guntupalli, J. S. (2014). Decoding Neural Representational
Spaces Using Multivariate Pattern Analysis. *Annual Review of Neuroscience*, *37*(1), 435–
456. https://doi.org/10.1146/annurev-neuro-062012-170325

Hebart, M. N., & Baker, C. I. (2018). Deconstructing multivariate decoding for the study of
brain function. *NeuroImage*, *180*(August 2017), 4–18.
https://doi.org/10.1016/j.neuroimage.2017.08.005

Hebart, M. N., Bankson, B. B., Harel, A., Baker, C. I., & Cichy, R. M. (2018). The representational
dynamics of task and object processing in humans. *ELife*, *7*.
https://doi.org/10.7554/elife.32816

Hindy, N. C., Ng, F. Y., & Turk-Browne, N. B. (2016). Linking pattern completion in the
hippocampus to predictive coding in visual cortex. *Nature Neuroscience*, *19*(5), 665–667.
https://doi.org/10.1038/nn.4284

Ho, T. C., Brown, S., Abuyo, N. A., Ku, E. H. J., & Serences, J. T. (2012). Perceptual consequences
of feature-based attentional enhancement and suppression. *Journal of Vision*, *12*(8), 1–
17. https://doi.org/10.1167/12.8.15

Hong, X., Wang, Y., Sun, J., Li, C., & Tong, S. (2017). Segregating Top-Down Selective Attention

from Response Inhibition in a Spatial Cueing Go NoGo Task: An ERP and Source
Localization Study. *Scientific Reports*, *7*(1), 1–14. https://doi.org/10.1038/s41598-017-
08807-z

Jiang, J., Summerfield, C., & Egner, T. (2013). Attention sharpens the distinction between
expected and unexpected percepts in the visual brain. *Journal of Neuroscience*, *33*(47),
18438–18447. https://doi.org/10.1523/JNEUROSCI.3308-13.2013

Jin, W., Nobre, A. C., & van Ede, F. (2020). Temporal expectations prepare visual working
memory for behavior. *Journal of Cognitive Neuroscience*, *32*(12), 2320–2332.
https://doi.org/10.1162/jocn_a_01626

Kaplan, J. T., Man, K., & Greening, S. G. (2015). Multivariate cross-classification: applying
machine learning techniques to characterize abstraction in neural representations.
Frontiers in Human Neuroscience, *9*(March), 1–12.
https://doi.org/10.3389/fnhum.2015.00151

Kastner, S., Pinsk, M. A., De Weerd, P., Desimone, R., & Ungerleider, L. G. (1999). Increased
activity in human visual cortex during directed attention in the absence of visual
stimulation. *Neuron*, *22*(4), 751–761. https://doi.org/10.1016/S0896-6273(00)80734-5

Keil, A., Debener, S., Gratton, G., Junghöfer, M., Kappenman, E. S., Luck, S. J., Luu, P., Miller, G. A.,
& Yee, C. M. (2014). Committee report: Publication guidelines and recommendations for
studies using electroencephalography and magnetoencephalography. *Psychophysiology*,
51(1), 1–21. https://doi.org/10.1111/psyp.12147

Kerrén, C., Linde-Domingo, J., Hanslmayr, S., & Wimber, M. (2018a). An Optimal Oscillatory
Phase for Pattern Reactivation during Memory Retrieval. *Current Biology*, *28*(21), 3383-
3392.e6. https://doi.org/10.1016/j.cub.2018.08.065

Kerrén, C., Linde-Domingo, J., Hanslmayr, S., & Wimber, M. (2018b). An Optimal Oscillatory
Phase for Pattern Reactivation during Memory Retrieval. *Current Biology*, *28*(21), 3383-
3392.e6. https://doi.org/10.1016/j.cub.2018.08.065

Kilner, J. M., Friston, K. J., & Frith, C. D. (2007). Predictive coding: An account of the mirror
neuron system. *Cognitive Processing*, *8*(3), 159–166. https://doi.org/10.1007/s10339-
007-0170-2

King, J.-R., & Dehaene, S. (2014). Characterizing the dynamics of mental representations: the
temporal generalization method. *Trends in Cognitive Sciences*, *18*(4), 203–210.
https://doi.org/10.1016/j.tics.2014.01.002

King, J., & Dehaene, S. (2017). *Characterizing the dynamics of mental representations : the
temporal generalization method*. *18*(4), 203–210.
https://doi.org/10.1016/j.tics.2014.01.002.Characterizing

King, J. R., Faugeras, F., Gramfort, A., Schurger, A., El Karoui, I., Sitt, J. D., Rohaut, B., Wacongne,
C., Labyt, E., Bekinschtein, T., Cohen, L., Naccache, L., & Dehaene, S. (2013). Single-trial
decoding of auditory novelty responses facilitates the detection of residual
consciousness. *NeuroImage*, *83*, 726–738.
https://doi.org/10.1016/j.neuroimage.2013.07.013

Kok, P., Mostert, P., & de Lange, F. P. (2017). Prior expectations induce prestimulus sensory
templates. *Proceedings of the National Academy of Sciences*, 201705652.
https://doi.org/10.1073/pnas.1705652114

Kok, P., Rahnev, D., Jehee, J. F. M., Lau, H. C., & De Lange, F. P. (2012). Attention reverses the
effect of prediction in silencing sensory signals. *Cerebral Cortex*, *22*(9), 2197–2206.
https://doi.org/10.1093/cercor/bhr310

Koshino, H., Minamoto, T., Yaoi, K., Osaka, M., & Osaka, N. (2015). Coactivation of the Default
Mode Network regions and Working Memory Network regions during task preparation.
Scientific Reports, *4*(1), 5954. https://doi.org/10.1038/srep05954

Kriegeskorte, N. (2008). Representational similarity analysis – connecting the branches of
systems neuroscience. *Frontiers in Systems Neuroscience*, *2*(November), 1–28.
https://doi.org/10.3389/neuro.06.004.2008

Kriegeskorte, N., Mur, M., & Bandettini, P. (2008). Representational similarity analysis -
connecting the branches of systems neuroscience. *Frontiers in Systems Neuroscience*,
2(NOV), 1–28. https://doi.org/10.3389/neuro.06.004.2008

López-García, D., Peñalver, J. M. G., Górriz, J. M., & Ruz, M. (2022). MVPAlab: A machine
learning decoding toolbox for multidimensional electroencephalography data. *Computer*
Methods and Programs in Biomedicine, *214*, 106549.
https://doi.org/10.1016/j.cmpb.2021.106549

López-García, D., Sobrado, A., Peñalver, J. M. G., Górriz, J. M., & Ruz, M. (2020). Multivariate
Pattern Analysis Techniques for Electroencephalography Data to Study Flanker
Interference Effects. *International Journal of Neural Systems*, *30*(7).
https://doi.org/10.1142/S0129065720500240

Love, J., Selker, R., Marsman, M., Jamil, T., Dropmann, D., Verhagen, J., Ly, A., Gronau, Q. F.,
Šmíra, M., Epskamp, S., Matzke, D., Wild, A., Knight, P., Rouder, J. N., Morey, R. D., &
Wagenmakers, E. J. (2019). JASP: Graphical statistical software for common statistical
designs. *Journal of Statistical Software*, *88*(1). https://doi.org/10.18637/jss.v088.i02

Ma, D. S., Correll, J., & Wittenbrink, B. (2015). The Chicago face database: A free stimulus set of
faces and norming data. *Behavior Research Methods*, *47*(4), 1122–1135.
https://doi.org/10.3758/s13428-014-0532-5

Mongillo, G., Barak, O., & Tsodyks, M. (2008). SynaptiC Theory of Working Memory. *Science*,
319(5869), 1543–1546. https://doi.org/10.1126/science.1150769

Muckli, L., De Martino, F., Vizioli, L., Petro, L. S., Smith, F. W., Ugurbil, K., Goebel, R., & Yacoub,
E. (2015). Contextual Feedback to Superficial Layers of V1. *Current Biology*, *25*(20),
2690–2695. https://doi.org/10.1016/j.cub.2015.08.057

Nili, H., Wingfield, C., Walther, A., Su, L., Marslen-Wilson, W., & Kriegeskorte, N. (2014). A
Toolbox for Representational Similarity Analysis. *PLoS Computational Biology*, *10*(4).
https://doi.org/10.1371/journal.pcbi.1003553

Nobre, A. C., & Serences, J. T. (2018). Building on a Solid Baseline: Anticipatory Biases in

Attention. *Trends in Neurosciences*, *41*(3), 120–122.
https://doi.org/10.1016/j.tins.2018.01.005

Palenciano, A. F., Sedoussi, M., Formica, S., & González-garcía, C. (in press). Canonical template
tracking: measuring the activation state of specific neural representations. *Frontiers in
Neuroimaging.* doi: 10.3389/fnimg.2022.974927

Palenciano, A. F., González-garcía, C., Arco, J. E., & Ruz, M. (2018). Transient and Sustained
Control Mechanisms Supporting Novel Instructed Behavior. *Cerebral Cortex*, 1–13.
https://doi.org/10.1093/cercor/bhy273

Peelen, M. V., & Kastner, S. (2011). A neural basis for real-world visual search in human
occipitotemporal cortex. *Proceedings of the National Academy of Sciences of the United
States of America*, *108*(29), 12125–12130. https://doi.org/10.1073/pnas.1101042108

Pernet, C., Garrido, M., Gramfort, A., Maurits, N., Michel, C., Pang, E., Salmelin, R., Schoffelen, J.
M., Valdes-Sosa, P., & Puce, A. (2018). *Best Practices in Data Analysis and Sharing in
Neuroimaging using MEEG*. 1–64. https://doi.org/10.31219/osf.io/a8dhx

Posner, M. I. (1980). Orienting of attention. *The Quarterly Journal of Experimental Psychology*,
32(1), 3–25. https://doi.org/10.1080/00335558008248231

Rohenkohl, G., Cravo, A. M., Wyart, V., & Nobre, A. C. (2012). Temporal expectation improves
the quality of sensory information. *Journal of Neuroscience*, *32*(24), 8424–8428.
https://doi.org/10.1523/JNEUROSCI.0804-12.2012

Rose, N. S., LaRocque, J. J., Riggall, A. C., Gosseries, O., Starrett, M. J., Meyering, E. E., & Postle, B.
R. (2016). Reactivation of latent working memories with transcranial magnetic
stimulation. *Science*, *354*(6316), 1136–1139. https://doi.org/10.1126/science.aah7011

Rungratsameetaweemana, N., & Serences, J. T. (2019). Dissociating the impact of attention and
expectation on early sensory processing. *Current Opinion in Psychology*, *29*, 181–186.
https://doi.org/10.1016/j.copsyc.2019.03.014

Ruz, M., & Nobre, A. C. (2008). Attention modulates initial stages of visual word processing.
Journal of Cognitive Neuroscience, *20*(9), 1727–1736.
https://doi.org/10.1162/jocn.2008.20119

Saenz, M., Buracas, G. T., & Boynton, G. M. (2002). Global effects of feature-based attention in
human visual cortex. *Nature Neuroscience*, *5*(7), 631–632.
https://doi.org/10.1038/nn876

Schröger, E. (1996). A neural mechanism for involuntary attention shifts to changes in
auditory stimulation. *Journal of Cognitive Neuroscience*, *8*(6), 527–539.
https://doi.org/10.1162/jocn.1996.8.6.527

Schröger, E., Kotz, S. A., & SanMiguel, I. (2015). Bridging prediction and attention in current
research on perception and action. *Brain Research*, *1626*, 1–13.
https://doi.org/10.1016/j.brainres.2015.08.037

Simon, A., Schachtner, J., & Gallen, C. L. (2019). Disentangling expectation from selective

attention during perceptual decision-making. *SSRN Electronic Journal*, *5*(564), 1–19.
https://doi.org/10.4324/9781315853178

Singh, D., & Singh, B. (2020). Investigating the impact of data normalization on classification
performance. *Applied Soft Computing*, *97*, 105524.
https://doi.org/10.1016/j.asoc.2019.105524

Smith, F. W., & Muckli, L. (2010). Nonstimulated early visual areas carry information about
surrounding context. *Proceedings of the National Academy of Sciences of the United States*
of America, *107*(46), 20099–20103. https://doi.org/10.1073/pnas.1000233107

Soon, C. S., Namburi, P., & Chee, M. W. L. (2013). Preparatory patterns of neural activity
predict visual category search speed. *NeuroImage*, *66*, 215–222.
https://doi.org/10.1016/j.neuroimage.2012.10.036

Sperling, R., Chua, E., Cocchiarella, A., Rand-Giovannetti, E., Poldrack, R., Schacter, D. L., &
Albert, M. (2003). Putting names to faces: Successful encoding of associative memories
activates the anterior hippocampal formation. *NeuroImage*, *20*(2), 1400–1410.
https://doi.org/10.1016/S1053-8119(03)00391-4

Stachenfeld, K. L., Botvinick, M. M., & Gershman, S. J. (2017). The hippocampus as a predictive
map. *Nature Neuroscience*, *20*(11), 1643–1653. https://doi.org/10.1038/nn.4650

Stein, T., & Peelen, M. V. (2015). Specific Expectations Enhance Stimulus Detectability by
Increasing Perceptual Sensitivity Content-Specific Expectations Enhance Stimulus
Detectability by Increasing Perceptual Sensitivity. *Journal of Experimental Psychology.*
General.

Stokes, M., Thompson, R., Nobre, A. C., & Duncan, J. (2009). Shape-specific preparatory activity
mediates attention to targets in human visual cortex. *Proceedings of the National*
Academy of Sciences, *106*(46), 19569–19574.

Summerfield, C., & De Lange, F. P. (2014). Expectation in perceptual decision making: Neural
and computational mechanisms. *Nature Reviews Neuroscience*, *15*(11), 745–756.
https://doi.org/10.1038/nrn3838

Summerfield, C., & Egner, T. (2009). Expectation (and attention) in visual cognition. *Trends in*
Cognitive Sciences, *13*(9), 403–409. https://doi.org/10.1016/j.tics.2009.06.003

Summerfield, C., Egner, T., Greene, M., Koechlin, E., Mangels, J., & Hirsch, J. (2006). Predictive
Codes for Forthcoming Perception in the Frontal Cortex. *Science*, *314*(November), 1311–
1314. https://doi.org/10.1017/cbo9781139167291.033

Sy, J. L., Guerin, S. A., Stegman, A., & Giesbrecht, B. (2014). Accurate expectancies diminish
perceptual distraction during visual search. *Frontiers in Human Neuroscience*, *8*(MAY), 1–
18. https://doi.org/10.3389/fnhum.2014.00334

Trapp, S., Lepsien, J., Kotz, S. A., & Bar, M. (2016). Prior probability modulates anticipatory
activity in category-specific areas. *Cognitive, Affective and Behavioral Neuroscience*, *16*(1),
135–144. https://doi.org/10.3758/s13415-015-0373-4

Trübutschek, D., Marti, S., Ojeda, A., King, J. R., Mi, Y., Tsodyks, M., & Dehaene, S. (2017). A
theory of working memory without consciousness or sustained activity. *ELife*, *6*, 1–29.
https://doi.org/10.7554/eLife.23871

van Driel, J., Gunseli, E., Meeter, M., & Olivers, C. N. L. (2017). Local and interregional alpha
EEG dynamics dissociate between memory for search and memory for recognition.
NeuroImage, *149*(October 2016), 114–128.
https://doi.org/10.1016/j.neuroimage.2017.01.031

Varoquaux, G. (2018). Cross-validation failure: Small sample sizes lead to large error bars.
NeuroImage, *180*(April 2017), 68–77.
https://doi.org/10.1016/j.neuroimage.2017.06.061

Vetter, P., Smith, F. W., & Muckli, L. (2014). Decoding sound and imagery content in early
visual cortex. *Current Biology*, *24*(11), 1256–1262.
https://doi.org/10.1016/j.cub.2014.04.020

Walther, A., Nili, H., Ejaz, N., Alink, A., Kriegeskorte, N., & Diedrichsen, J. (2016). Reliability of
dissimilarity measures for multi-voxel pattern analysis. *NeuroImage*, *137*, 188–200.
https://doi.org/10.1016/j.neuroimage.2015.12.012

Westfall, J. (2016). *PANGEA: Power ANalysis for GEneral Anova designs*.

Wimber, M., Alink, A., Charest, I., Kriegeskorte, N., & Anderson, M. C. (2015). Retrieval induces
adaptive forgetting of competing memories via cortical pattern suppression. *Nature*
Neuroscience, *18*(4), 582–589. https://doi.org/10.1038/nn.3973

Womelsdorf, T., Fries, P., Mitra, P. P., & Desimone, R. (2006). Gamma-band synchronization in
visual cortex predicts speed of change detection. *Nature*, *439*(7077), 733–736.
https://doi.org/10.1038/nature04258

Wyart, V., Nobre, A. C., & Summerfield, C. (2012a). Dissociable prior influences of signal
probability and relevance on visual contrast sensitivity. *Proceedings of the National*
Academy of Sciences of the United States of America, *109*(9), 3593–3598.
https://doi.org/10.1073/pnas.1120118109

Wyart, V., Nobre, A. C., & Summerfield, C. (2012b). Dissociable prior influences of signal
probability and relevance on visual contrast sensitivity. *Proceedings of the National*
Academy of Sciences of the United States of America, *109*(9), 3593–3598.
https://doi.org/10.1073/pnas.1120118109

Zuanazzi, A., & Noppeney, U. (2019). Distinct neural mechanisms of spatial attention and
expectation guide perceptual inference in a multisensory world. *The Journal of*
Neuroscience, *39*(12), 2873–18. https://doi.org/10.1523/jneurosci.2873-18.2019

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