

1 Relationships between individual differences in dual process and electrophysiological
2 signatures of familiarity and recollection during retrieval

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15 **Abstract**

16 Our everyday memories can vary in terms of accuracy and phenomenology. According to one
17 theoretical account, these differences hinge on whether the memories contain information
18 about both an item itself as well as associated details (remember) versus those that are devoid
19 of these associated contextual details (familiar). This distinction has been supported by
20 computational modeling of behavior, studies in patients, and neuroimaging work including
21 differences both in electrophysiological and functional magnetic resonance imaging. At
22 present, however, little evidence has emerged to suggest that neurophysiological measures
23 track individual differences in estimates of recollection and familiarity. Here, we conducted
24 electrophysiological recordings of brain activity during a recognition memory task designed
25 to differentiate between behavioral indices of recollection and familiarity. Non-parametric
26 cluster-based permutation analyses revealed associations between electrophysiological
27 signatures of familiarity and recollection with their respective behavioral estimates. These
28 results support the idea that recollection and familiarity are distinct phenomena and is the
29 first, to our knowledge, to identify distinct electrophysiological signatures that track
30 individual differences in these processes.

31 **Introduction**

32 We have all had the experience of running into someone we know walking down the
33 street. Sometimes just seeing that person can help us recollect their name and where and when
34 we last encountered them. At other times, we might feel confident that we have met that
35 person because the face seems so familiar, even if we are otherwise unable to recover any
36 details about that person. A long history of memory research has suggested that recollection
37 and familiarity vary in terms of retrieved information (i.e., item vs. context information),
38 vividness (Cooper and Ritchey, 2019; Jacoby and Dallas, 1981; Roediger and Blaxton, 1987;
39 Tulving, 2002; Woroch and Gonsalves, 2010), and subjective experience (Leynes and
40 Nagovsky, 2016; Souchay et al., 2013). Moreover, studies using electroencephalography
41 (EEG; Addante et al., 2012b; Curran, 2002; Diana et al., 2011; Duarte et al., 2004; Düzel et
42 al., 1997; Leynes and Phillips, 2008; Rugg and Curran, 2007; Tsivilis et al., 2001; Wilding et
43 al., 1995; Woroch and Gonsalves, 2010), magnetoencephalographic (Evans and Wilding,
44 2012), studies of patients with brain damage (Addante et al., 2012a; Aggleton et al., 2005;
45 Aly et al., 2011; Bowles et al., 2010; Duarte et al., 2005, 2004; Wang et al., 2014, 2014), and
46 functional magnetic resonance imaging (fMRI; Diana et al., 2012, 2007; Eldridge et al., 2000;
47 Vilberg et al., 2006; Vilberg and Rugg, 2008; Yonelinas et al., 2005) are consistent with the
48 idea that recollection and familiarity depend on different neural substrates. Accordingly, many
49 theories (Jacoby, 1991; Mandler, 1980; Tulving, 1985; Yonelinas, 1999) and computational
50 models (Elfman et al., 2014; Norman and O'Reilly, 2003; Selmecky and Dobbins, 2014) have
51 proposed that recollection and familiarity are driven by different processes (see Eichenbaum
52 et al., 2007; Ranganath, 2010a, 2010b; Ranganath and Rainer, 2003; Reagh and Ranganath,
53 2018; Wilding and Ranganath, 2012; Yonelinas, 2002 for extensive reviews of this literature),
54 though this idea remains somewhat controversial (e.g., Wixted, 2007).

55 One class of models (Park and Donaldson, 2019; Yonelinas, 2002, 1999, 1994;
56 Yonelinas et al., 2010) particularly emphasizes the idea that recollection and familiarity can
57 be distinguished on the type of associated details recovered, as well as phenomenology.
58 According to this view, familiarity is typically associated with strong memory for an item
59 with minimal information about associated contextual details, whereas recollection is
60 accompanied by retrieval of information about the item and about the context in which it was
61 encountered (Addante et al., 2012a; Diana et al., 2008; Eichenbaum et al., 2007; Ranganath,
62 2010b; Yonelinas, 2002, 1999, 1994; Yonelinas et al., 2010).

63 Many studies have used recordings of event-related potentials (ERPs) to differentiate
64 between neural correlates of recollection and familiarity based on sensitivity to particular
65 experimental variables, temporal dynamics, and scalp topography (Addante et al., 2012b;

66 Bridger et al., 2012; Curran, 2004, 2000; Park and Donaldson, 2019; Rugg et al., 1998a,
67 1998b; Rugg and Curran, 2007; Wilding and Herron, 2006, 2006; Wilding and Ranganath,
68 2012), but there is significant controversy over how these results should be interpreted (Hou
69 et al., 2013; Lucas et al., 2010; Nie et al., 2014; Paller et al., 2007; Thakral et al., 2016; Voss
70 et al., 2012; Voss and Paller, 2016, 2006; Yovel and Paller, 2004). Many studies have
71 reported ERP differences between old and new items (i.e., an ERP “old-new effect”) around
72 300-600 ms post-stimulus, with a mid-frontal or fronto-central scalp topography (Curran et
73 al., 2006; Friedman and Johnson, 2000; Mecklinger, 2006, 2000; Rhodes and Donaldson,
74 2007; Tsivilis et al., 2001). The topography and latency of this “mid-frontal old-new
75 effect”(Rugg et al., 1998a) resembles the N400 ERP component reported in psycholinguistic
76 studies (Kutas and Hillyard, 1984), although the mid-frontal old-new effect often has a more
77 anterior distribution (Wilding and Ranganath, 2012). The mid-frontal old-new effect has often
78 been contrasted with a late-onsetting old-new effect that is maximal at parietal sites. This
79 latter effect has been called the “parietal old-new effect” or the late positive component (LPC)
80 (Friedman and Johnson, 2000; Olichney et al., 2000; Paller and Kutas, 1992; Smith, 1993;
81 Wilding and Ranganath, 2012), and is more left-lateralized for words and more widespread
82 for pictures and actions (Leynes et al., 2017).

83 Results from a number of studies have been used to support the idea that the mid-
84 frontal old-new effect may be a neural correlate of familiarity, whereas the parietal old-new
85 effect may be a neural correlate of recollection (Addante et al., 2012b; Friedman and Johnson,
86 2000; Griffin et al., 2013, 2013; Leynes et al., 2005; Olichney et al., 2000; Paller and Kutas,
87 1992; Park and Donaldson, 2019; Rhodes and Donaldson, 2007; Rugg et al., 1998b; Speer
88 and Curran, 2007; Wilding and Ranganath, 2012, 2012; Wynn et al., 2020, 2020) for reviews
89 see (Curran et al., 2006; Friedman, 2013; Friedman and Johnson, 2000; Rugg and Curran,
90 2007), although results from other studies have argued against this idea (Bridger et al., 2012;
91 Greve et al., 2007; Kelley and Wixted, 2001; Leynes et al., 2017; Paller et al., 2007; Voss and
92 Federmeier, 2011; Voss and Paller, 2006; Yovel and Paller, 2004). One complication in
93 interpreting studies differentiating between the mid-frontal and parietal old-new effects is that
94 the exact timing and topography of these effects can differ considerably across studies, and it
95 is possible that recollection and familiarity might be differentiated by ERP modulations that
96 do not exhibit the typical characteristics of these two old-new effects (Friedman et al., 2005;
97 Ranganath and Paller, 2000; Tsivilis et al., 2001). Data-driven ERP analysis methods (Maris
98 and Oostenveld, 2007) might offer a way to reconcile these views and identify the extent to
99 which recollection and familiarity can be differentiated.

100 It is also notable that most ERP studies have focused on old-new effects in group
101 averages, and little is known about whether ERP modulations track individual differences in
102 memory performance. If different ERP measures are predictive of putatively different
103 memory processes, there should be unique correlations between the behavioral and ERP
104 measures of each memory process. Recently, several studies have focused on correlating
105 ERPs with individual differences in recognition memory performance (Amico et al., 2015;
106 Angel et al., 2010; Chen et al., 2014; MacLeod and Donaldson, 2017), but these studies have
107 not revealed a clear picture. Angel et al. (2010) correlated overall recognition memory
108 performance (corrected recognition rate) with the magnitude of the parietal old-new effect,
109 but this study was performed on a small sample (14 participants), and it focused only on the
110 parietal old-new effect within an a priori time window. More recently, MacLeod and
111 Donaldson (2017) correlated the magnitude of the left parietal old-new effect with recognition
112 performance. Across three tasks, the authors found significant old/new effects in the left
113 parietal ERP (Hit>CR; R-CR > K-CR), but correlations between the ERP and behavioral
114 measures were inconclusive. This may have arisen because, although the total number of
115 participants was high (122), only 20 participants were included in the correlation of R/K
116 effect magnitude with behavioral data. In addition, these analyses focused only on the late left
117 parietal effect, estimated as the mean ERP difference within an a priori time window, averaged
118 across three parietal electrodes and did not attempt to differentiate correlates of recollection-
119 and familiarity-based recognition. Chen et al. (2014) correlated FN400 (mid-frontal old-new
120 effect) magnitudes with recognition performance in a large sample (64 participants), but this
121 study used only overall recognition discriminability (d') and response time as behavioral
122 indices of memory performance, which does not distinguish recollection from familiarity.
123 Whereas the above studies focused specifically on previously identified ERP old-new effects,
124 Amico et al. (2015) used data-driven non-parametric analyses to characterize individual
125 differences in overall recognition performance (Hit and FA rates, the sensitivity index d' , the
126 decision criterion c , and the mean RT for Hit trials). However, this study had a relatively
127 small sample size (18 participants) and did not attempt to separately estimate familiarity and
128 recollection.

129 To summarize, prior studies have focused on relationships between ERP components
130 and recognition memory performance, but no previous study, to our knowledge, has shown a
131 relationship between individual differences in behavioral estimates of recollection and
132 familiarity and the putative ERP correlates of these processes described above. Furthermore,
133 given the known variability in timing and topography in these EEG signatures with different

134 types of stimuli, there is no agreed upon standard definition that can be applied consistently
135 across studies.

136 The present study seeks to address these limitations by testing the hypothesis that
137 individual differences in recognition memory performance are related to variability in
138 electrophysiology in terms of two recognition memory processes: familiarity and recollection.
139 In order to test whether individual differences in electrophysiological signatures of familiarity
140 and recollection were related to differences in their dual process estimates, we measured
141 ERPs while participants made recognition memory judgments using the Remember/Know
142 method (Tulving, 1985). To identify ERPs related to individual differences in familiarity and
143 recollection in an unbiased manner, we used a data-driven analysis approach in which ERP
144 differences were correlated with individual differences in dual process estimates of familiarity
145 and recollection using non-parametric cluster-based permutation analysis. If dual process
146 estimates of familiarity and recollection are associated with particular electrophysiological
147 signatures, we should observe correlations between familiarity-related ERPs and familiarity
148 dual process estimates and between recollection-related ERPs and recollection dual process
149 estimates.

150

151 **Methods**

152 *Participants*

153 49 participants took part in the study. They were right-handed and had normal or
154 corrected-to-normal visual acuity and no history of neurological or psychological disorders.
155 Participants were excluded due to contamination of mastoid channels (N=1), incidental MRI
156 finding (N=1), not completing the task (N=2), and not fulfilling the inclusion criteria of at
157 least 30 recollection trials (Boudewyn et al., 2018; Cohen, 2014; Luck, 2014, 2005), N=7).
158 Thus, 38 participants ($N_{\text{female}} = 26$, mean age = 25.7 ± 4.2 years) were included in the final
159 analyses. A post-hoc sensitivity analysis of a bivariate normal model correlation using
160 G*Power 3.1 software (Faul et al., 2009, 2007) with a two tailed alpha value of 0.05 showed
161 that a sample size of 38 with a power of 0.8 could detect a medium to large effect size of
162 0.436 (Cohen, 1988) with a correlation interval of ± 0.32 . The study was approved by the
163 Institutional Review Board of the University of California at Davis and all participants
164 provided informed consent prior to participation. Participants were compensated \$20/hr for
165 their time.

166

167 ***Stimuli***

168 Study materials came from the Bank of Standardized Stimuli (BOSS) developed by
169 Brodeur and colleagues (Brodeur et al., 2014, 2010). Using the normative data provided by
170 the BOSS creators, we selected the images that were well-agreed on for the name (>29%),
171 category (>29%), object (>2 on a 1-5 scale), and viewpoint (>2 on a 1-5 scale), and rated as
172 familiar (>2 on a 1-5 scale), and simple (<2 on a 1-5 scale for complexity). Duplicate items
173 were removed as well as items in any of the animal ("Crustacean", "Mammal", "Reptile",
174 "Bird", "Insect", "Canine", "Feline", "Sea mammal", "Fish"), body part, and war weapon
175 categories. In addition, any potentially emotional, disturbing, or unpleasant images were
176 removed (e.g., syringe, hunting knife) as well as objects that a research assistant deemed hard
177 to recognize (e.g., contact lens) or redundant with another object. Images were re-sized in
178 Adobe Photoshop to 500 x 500 pixels. Objects from this list of 644 remaining objects were
179 then used to generate study and lure lists. Object categories used in the stimuli represent the
180 general makeup of the original BOSS stimuli (see Brodeur et al., 2014, 2010 for details).
181 Original stimuli used in the paradigm (<https://osf.io/4s7uy/>) as well as the original source
182 code for stimulus presentation

183 (https://github.com/hallez/eetemp_eeg_pub/tree/main/experiment-scripts) are available online.

184 270 objects were uniquely drawn for each subject such that each participant saw a
185 distinct set of items although, by chance, some objects necessarily overlapped between
186 participants. Of the 270 objects selected for a participant, 180 images were used as study
187 items during the encoding phase and 90 were used as lures during the retrieval phase. 180
188 study items were divided into five encoding lists of 36 items each. One of four encoding
189 questions ("Would you find this item in a supermarket/convenience store?", "Would this item
190 fit in a fridge/bathtub?") was paired with each item. In each of the five encoding lists, each
191 question was paired with 9 different items such that each question was used the same number
192 of times in each encoding list. Lists for object retrieval were constructed by randomly
193 selecting 30 studied and 15 lure items. The proportion of objects from each encoding list (1-5)
194 and encoding question (fridge/bathtub/supermarket/convenience store) was not constrained in
195 the construction of the retrieval lists. This manipulation was used to ensure the participants
196 were paying attention to individual items and was not included in the analyses described in
197 this work. For a retrieval list, the 180 studied items and 90 lure items were randomly
198 intermixed, using the *numpy.random.shuffle* tool in Python (Version 2.7.14). Participants
199 were given a practice phase with four objects. Objects used in the practice phase were drawn
200 from a separate stimulus set of computer-generated images (previously used in (Dimsdale-

201 Zucker et al., 2018) so that they would not be confused with any of the studied items. Four
202 practice objects were randomly selected for each subject from a set of 12 objects.

203

204 ***Procedure***

205 The experiment comprised of practice, encoding, and retrieval phases. EEG data were
206 collected during the retrieval phase. Each trial during the retrieval phase included object
207 recognition and source memory test components. The present study focuses on the EEG
208 signature that differentiates recollection and familiarity during item recognition as measured
209 with respect to the object recognition judgment.

210 For practice, a research assistant walked the participant through each phase of the task.
211 The practice for encoding was followed by familiarizing participants with the response scales
212 that were used for the object recognition and source judgments. Participants were told that the
213 difference between familiarity and recollection is that familiarity is feeling like they know or
214 have seen this item but being unable to recall from where. In contrast, recollection is
215 accompanied by memory for both the item itself as well associated details (e.g. an experience
216 or association from when the item was originally studied, the source or context of the item).
217 This description was supplemented with an example of meeting someone in the grocery store
218 and either feeling like you know them but not having access to their name or how you know
219 them (“familiar”) versus running into an individual and knowing their name or where you’ve
220 encountered them previously (“remember”). New responses were explained as being
221 analogous to the experience of meeting a stranger or a totally novel person. The participants
222 were given practice trials along with feedback from the experimenter and detailed instructions
223 to ensure they understood how to use this response scale.

224 During the encoding phase, objects and questions (“Would this item fit in a fridge?”,
225 etc.) appeared on the screen for 250 ms. Within each list, all four questions were presented in
226 a random order, an equal number of times across items. The participant made their response to
227 the question during this interval. A short presentation time was used in order to eliminate
228 potential eye movements during encoding. Response key mappings for the yes/no judgments
229 were randomized across participants. Responses were recorded but accuracy was not analyzed
230 as the purpose of this task was simply to orient participants to the objects and their respective
231 sources. Between lists, there was a 30 second break before the text label (e.g. “List 1”) for the
232 next list was shown.

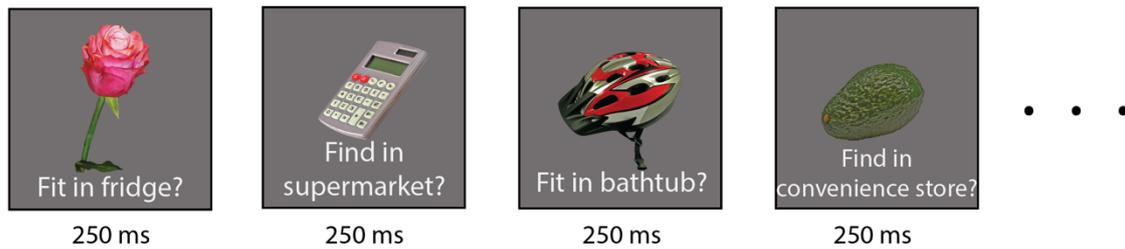
233 After studying all 180 objects, participants were then setup for EEG recording. Details
234 of the EEG setup can be found in the *EEG acquisition and processing* section. This cap setup

235 served as the delay period (approximately 45-60 min) between the encoding and retrieval
236 phases. During the retrieval phase, participants saw 270 objects (180 old and 90 novel) split
237 across six blocks of 45 trials (30 old and 15 novel objects). Each block began with a 23
238 second warm up period that walked the subject through getting in a comfortable position,
239 blinking, and preparing to begin responding while allowing the EEG recordings to stabilize.
240 Each trial started with a white central fixation cross on top of a gray background that
241 remained in the foreground of the screen throughout the entire trial. Next, an object appeared
242 and remained on the screen for 700 ms (Figure 1). Participants knew to withhold their
243 response at this time. A capital white “T” (“think cue”) came on screen for 1700 ms. This
244 timing was based on a similar paradigm (Gruber et al., 2008). Again, no responses were made
245 while the think cue remained on the screen to minimize movement-related artifacts in the
246 EEG data in the time window of interest.

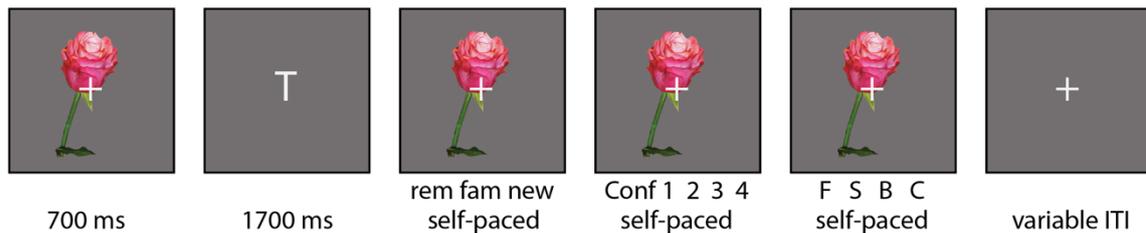
247 Finally, the object reappeared, and the participant was allowed to make a self-paced
248 object recognition judgment (Remember, Feels Familiar, or New; response order
249 counterbalanced). After making their memory judgment, the object remained on the screen
250 but the response scale updated to a confidence judgment (1=highly, 2=moderately,
251 3=somewhat, 4=not at all, Figure 1). This was followed by a four-option forced-choice
252 question about the item’s original encoding question (again, the order of the four options was
253 randomized for each subject but remained constant throughout the experiment). Trials were
254 separated by 2 second ITIs with a white fixation cross. For EEG timing precision, the timing
255 of all screens was taken as the ceiling of the expected duration multiplied by the monitor’s
256 frame rate.

257

A. encoding



B. item recognition (EEG)



258

259 **Figure 1.** Paradigm design. During the encoding phase (A) participants studied items and
260 answered one of four questions: “Would you find this item in a supermarket/convenience
261 store?”, “Would this item fit in a fridge/bathtub?”. Items were presented on top of a gray
262 background with questions written in white text. During (B) participants saw old and new
263 items, while undergoing EEG recordings. Items were presented on a gray background with a
264 white fixation cross overlaid. Each item was presented briefly (700 ms). Initial item
265 presentation was followed by a think cue (white letter “T”). Following this, participants made
266 their memory response to the item (rem =Remember, fam =Familiar, new =New item),
267 followed by a memory confidence level judgment and a question source memory judgment
268 (“fridge”, “supermarket”, “bathtub”, “convenience store”; abbreviated above in figure as “F S
269 B C”). ERPs were recorded during the item recognition phase. Black outlines are for figure
270 visualization purposes only and did not appear during the task to participants.

271

272

273 **Behavioral measures**

274 Behavioral measures of recollection and familiarity were calculated using dual process
275 estimates, which are intended to give an independent measure of these two processes, while
276 accounting for response bias (Yonelinas, 2002). Recollection was estimated as the difference
277 between the Remember hit rate and the Remember false alarm rate ($[count\ of\ all\ remembered$
278 $trials / count\ of\ all\ old\ items] - [count\ of\ all\ remembered\ false\ alarms / count\ of\ all\ new$
279 $items]$) and familiarity was estimated as the difference between the Familiar hit rate corrected
280 by the inverse Remember hit rate and Familiar false alarm rate corrected by the inverse
281 Remember false alarm rate ($[Familiar\ hit\ rate / (1 - Remember\ hit\ rate)] - [Familiar$
282 $false\ alarm\ rate / (1 - Remember\ false\ alarm\ rate)]$), (Yonelinas, 2002). Source judgments to
283 old items were computed to determine whether participants could retrieve information about

284 the question. We expected that accurate retrieval of the orienting task used to encode each
285 item (source memory) should be more likely for items judged as recollected (MacKenzie et
286 al., 2018; Yonelinas, 2002) than for items associated with “familiar” responses.

287

288 ***EEG acquisition and analysis***

289 EEG data were recorded in a sound attenuated chamber at a rate of 512 Hz using a
290 BioSemi (<http://www.biosemi.com>) ActiveTwo system with 64 active Ag/AgCl scalp
291 electrodes embedded in an elastic cap in an extended version of the international 10/20
292 system. Additional electrodes were placed at the left and right mastoids to be used for offline
293 re-referencing. In addition, the electrooculogram (EOG) was recorded with an additional four
294 electrodes – bipolar vertical channels located approximately 1cm above and below the
295 subject’s left eye and horizontal ocular channels located approximately 1 cm lateral to the
296 outer canthus of each eye. The EEG was recorded relative to a common mode sense active
297 electrode near Cz for online referencing. Participants were instructed to blink normally while
298 maintaining focus at the center of the screen and while minimizing muscle tension and any
299 large movements.

300 EEG data preprocessing and analyses were performed using EEGLab (Delorme and
301 Makeig, 2004), ERPLab (Lopez-Calderon and Luck, 2014), and custom code implemented in
302 MATLAB r2014b (www.mathworks.com) for EEG processing, available in a public github
303 repository (https://github.com/hallez/eetemp_eeg_pub/tree/main). Data intended for ERP
304 analyses were downsampled offline to 128 Hz, re-referenced to the average of the mastoid
305 channel signals, and high-pass filtered at 0.1 Hz (IIR Butterworth filter, half-amplitude
306 cutoff=0.2 Hz, slope=12 dB/octave). The data were separately high-pass filtered for ICA
307 (Winkler et al., 2015) at 1Hz (IIR Butterworth filter, half-amplitude cutoff=1.60 Hz, slope=12
308 dB/octave), based on Makoto Miyakoshi's preprocessing pipeline
309 (https://scn.ucsd.edu/wiki/Makoto's_preprocessing_pipeline). Next, bad channels were
310 detected using the *trimOutlier* function (lower standard deviation threshold=2; upper standard
311 deviation threshold=200) and then epoched (epoch start = -500ms; epoch end = 1000ms).
312 After epoching, outlier epochs were automatically rejected using a thresholded approach
313 implemented with *pop_jointprob* (local channel threshold = 6; global threshold = 2).
314 Independent component analysis (ICA) was performed with the *runica* algorithm (Delorme et
315 al., 2007) holding out mastoid and outlier channels, and the SASICA toolbox (Chaumon et
316 al., 2015) was used to manually review and identify eyeblink-related components for removal
317 from the data. At manual review, any additional epochs or channels that were determined to

318 be outliers were identified for removal or interpolation, respectively. Bad channels were
319 interpolated from the ICA corrected data (mean number of interpolated channels was 1.3 ± 1.9
320 per block). At this point, all six blocks were merged into a single file. Baseline correction was
321 applied to each trial using a pre stimulus period from 200 ms prior to the onset of the first
322 image in a trial. Average number and standard deviation (with the range provided in
323 parentheses) of trials included in the analysis was: 66 ± 12 (34-83), 54 ± 15 (33-94), and 79 ± 24
324 (30-127) for Correct Rejection, Familiar, and Remember trial types, respectively. Raw and
325 preprocessed files can be found at <https://osf.io/4e3pq/>.

326 Given that ERPs related to long-term memory are known to shift their latency and
327 distribution with different types of stimuli (Addante et al., 2012b; Busch et al., 2004; Taylor,
328 2002; Yonelinas, 2002), we performed non-parametric cluster-based permutation analysis
329 (Maris and Oostenveld, 2007) implemented in the FieldTrip Matlab toolbox (Oostenveld et
330 al., 2011), which corrects for the multiple comparisons problem (MCP) arising from the fact
331 that the effect of interest (i.e. a difference between experimental conditions) is evaluated at
332 large number of data points, here: (channel, time)-pairs. The approach combines neighboring
333 values that are likely to be correlated (e.g., neighboring time points and/or spatial locations) to
334 reduce the problem of multiple comparisons. Therefore, this method allowed us to compare
335 ERPs between trial types (Familiar, Remember, and Correct Rejection) for each (channel,
336 time) data pair and identify statistically meaningful differences. Under the null hypothesis of
337 exchangeability, assuming averages from Familiar, Remember, and Correct Rejection trials
338 are drawn from the same probability distribution, cluster alpha $p = 0.05$ and 0-700 ms time
339 window were used. In order to isolate familiarity and recollection electrophysiological
340 estimates while controlling for other cognitive processes that are not related to memory, ERPs
341 from the following trial types were compared: Familiar as compared to Correct Rejection
342 trials, and Remember as compared to Familiar trials, respectively. Such comparisons are
343 widely used in the studies of old-new effect, including Remember/Know procedures (Duarte
344 et al., 2006, 2004).

345 The comparisons between each pair of trial types were performed via a two-tailed t-
346 test (alpha = 0.05) using *depsamplesT* function, which clustered samples whose t-value was
347 larger than a priori threshold ($P=0.05$) on the basis of temporal and spatial adjacency. Cluster-
348 level statistics were calculated by taking the sum of the t-values within every cluster and the
349 maximum of the cluster-level statistics was taken. Monte Carlo correction for the MCP with
350 1000 draws from the permutation distribution was used. Channel neighbors for spatial
351 clustering were found based on the template method, using 'Biosemi64_neighb' template.

352 In order to test our critical question of whether ERP signatures associated with
353 recollection and familiarity are associated with dual process estimates, we computed
354 correlations between these measures. This was also performed using a non-parametric cluster-
355 based permutation test. In this case, *Ft_statfun_correlationT* function and Pearson *r*
356 coefficient were used in order to test if there was a relationship between familiarity and
357 recollection dual process estimates per subject (quantitative independent variable) and their
358 (channel, time) EEG data (dependent variable). The correlations of both difference waveforms
359 (Familiar – Correct Rejection, and Remember – Familiar) with behavioral estimates of
360 familiarity and recollection, as generated from dual process estimates, were tested.

361

362 **Results**

363 ***Behavioral results***

364 Participants were highly accurate at discriminating studied from unstudied objects
365 (Table 1). To account for response bias, we also computed dual process estimates of
366 familiarity and recollection (Yonelinas, 2002). Mean familiarity and recollection estimates are
367 presented in Table 1. Each studied item was scored according to the item recognition
368 judgment and accuracy for the question type the item had been paired with (Table 1). The
369 accuracy of source memory was significantly above chance for both familiarity ($t_{37} = 7.0, P <$
370 0.001) and recollection ($t_{37} = 10.3, P < 0.001$), but participants were significantly more likely
371 to correctly retrieve source information for Remember than Familiar trials ($t_{37} = 6.0, P <$
372 0.001). These findings are in agreement with other work showing that accurate source
373 judgments can be made on the basis of both recollection and familiarity (Addante et al.,
374 2012b; Diana et al., 2011, 2008), but that retrieval of contextual details should be more likely
375 when an item is recollected, as compared with familiarity-based recognition (for a
376 comprehensive review, see Yonelinas, 2002).

377

378 **Table 1.** Behavioral results of object recognition and source memory performance. (A)
379 Proportion of “remember” and “familiar” responses to old and new items presented as hit
380 rates, false alarm rates, and source memory rates are presented (with standard deviations)
381 separately for “remember” and “familiar” responses. (B) Mean behavioral estimates of
382 recollection and familiarity derived from the dual process model are presented (with standard
383 deviations).

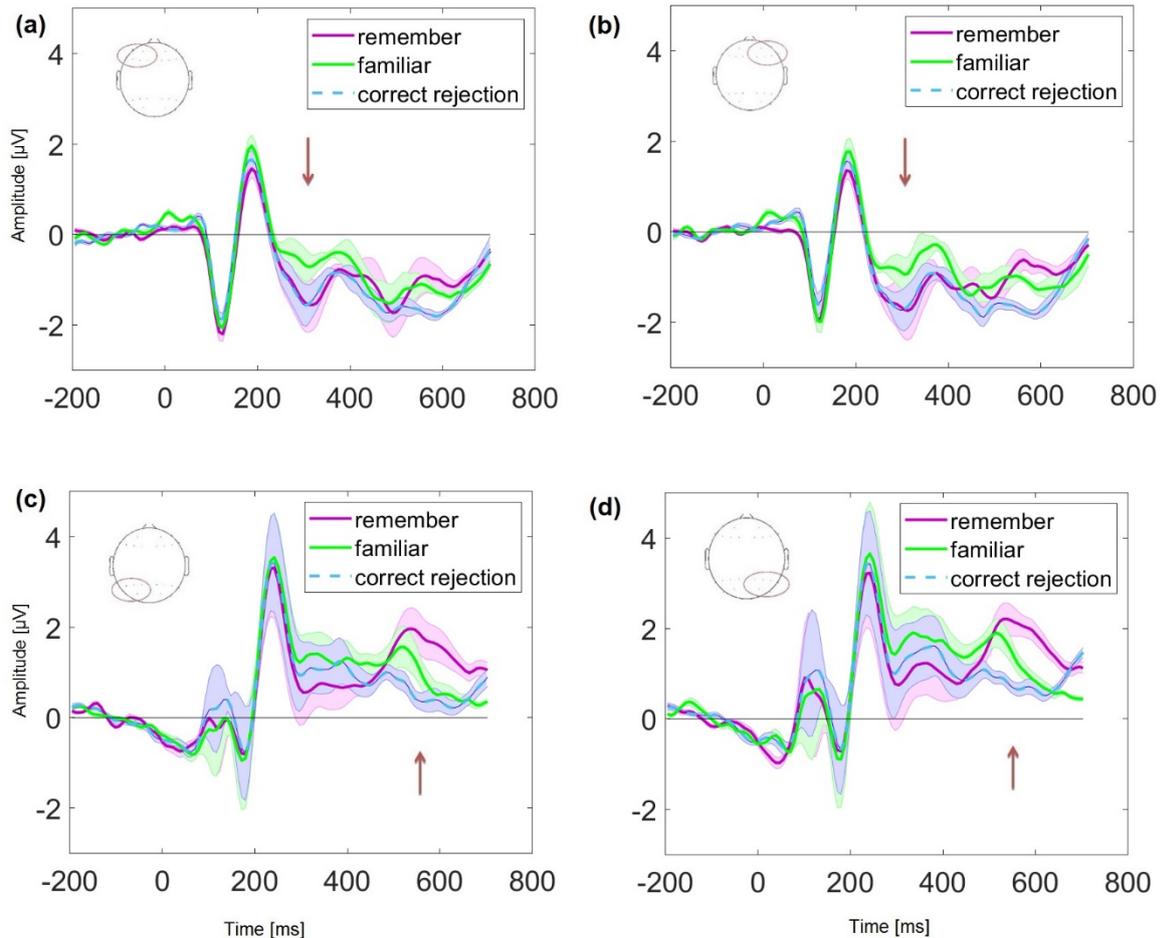
(A)		Remember	Familiar
Object recognition performance	Hit rate	0.47±0.14	0.32±0.08
	False alarm rate	0.03±0.05	0.18±0.10
Source memory accuracy		0.47±0.13	0.33±0.07
(B)		Recollection	Familiarity
Dual process estimate		0.44±0.12	0.45±0.15

384

385 ***ERP results***

386 Before analyzing individual differences in ERP correlates of recognition, we
387 conducted analyses to examine overall ERP old-new effects in order to be able to compare our
388 results to previous reports of ERP differences between recollection and familiarity. We
389 separately averaged ERPs for successfully recognized items associated with Remember
390 responses, for recognized items associated with Familiar responses, and for Correct Rejection
391 responses (Figure 2). These averages were done solely for visualization purposes. We report
392 statistical comparisons between conditions in the following section. Averaged ERPs revealed
393 a sustained negative deflection for all trials types, beginning approximately at 220 and lasting
394 until around 400 ms, (Figure 2a-b) and a positive deflection from approximately 500 to 700ms
395 after stimulus onset (Figure 2c-d).

396



397

398 **Figure 2.** ERP correlates of recollection and familiarity. Grand averaged ERPs on Remember
399 (purple), Familiar (green) and Correct Rejection (dashed blue) trials are separately averaged
400 for four groups of channels split by frontal and parietal for each hemisphere (Woodruff et al.,
401 2006). Arrows are meant to delineate time periods of interest, but do not indicate statistical
402 comparisons: (a) left frontal (F1, F3, F5, F7, AF3, AF7), (b) right frontal (F2, F4, F6, F8,
403 AF4, AF8), (c) left parietal (P1, P3, P5, P7, PO3, PO7), and (d) right parietal (P2, P4, P6, P8,
404 PO4, PO8). Shaded areas represent standard deviation of the mean. Note that these average
405 traces from electrode groups are presented for visualization purposes, but electrodes were
406 analyzed separately in the data-driven statistical analyses.

407

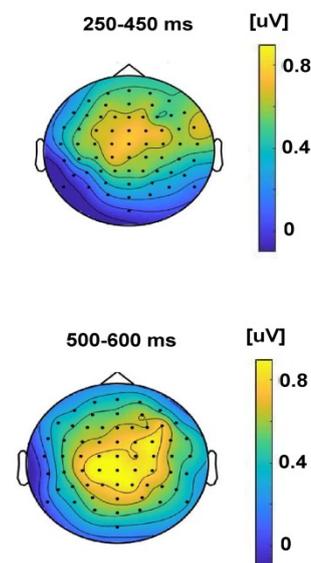
408 In addition, to better visualize the familiarity and recollection ERP effects, we also present
409 difference waveforms: Familiar-minus-Correct Rejection and Remember-minus-Familiar
410 (Supplemental figure 3).

411

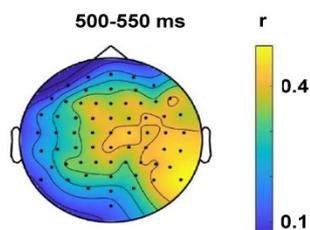
412 To quantify ERP correlates of familiarity at the group level, we contrasted ERPs
413 associated with familiar responses against ERPs associated with correct rejection responses.
414 As described in the *Methods*, these, and all subsequent ERP analyses, were done using a data-
415 driven non-parametric cluster-based permutation analysis procedure (Maris and Oostenveld,
2007). This method allows us to identify statistically significant differences between

416 conditions, though it does not permit specific conclusions about the precise temporal or
417 spatial extent of these differences (Sassenhagen and Draschkow, 2019). We can, however,
418 identify the cluster extent in time and location as descriptive information about the observed
419 data. As such, this analysis identified two spatiotemporal clusters that corresponded to the
420 significant difference ($p=0.001$ cluster corrected for both clusters) in the observed data: from
421 approximately 250 to 450 ms at frontal and fronto-central scalp sites, and from approximately
422 500 to 600 ms over central, centro-parietal, and right fronto-central sites. Figure 3a shows
423 topographic distributions of the ERP differences between familiar and correct rejection trials
424 corresponding to these time windows.

(A) Overall familiar-correct rejection ERP



(B) ERP-behavior correlation



425

426 **Figure 3.** ERP correlates of familiarity. (A) Topographic maps illustrate distributions of mean
427 ERP differences between familiar and correct rejection trials corresponding roughly to the
428 significant clusters identified in the data-driven permutation analysis. (B) A topographic map
429 illustrates a distribution of correlations between familiar - correct rejection ERP differences
430 with dual process estimates of familiarity. Color bars show ERP voltage difference (panel A)
431 or Pearson's r correlation coefficient values (panel B).

432

433 Next, we conducted data-driven analyses to identify ERP correlates of individual
434 differences in familiarity-based recognition. This analysis revealed a significant cluster in the
435 observed data ($p=0.001$ cluster corrected), extending approximately from 500 to 550 ms
436 (Figure 3b). This correlation was most pronounced over right central and centroparietal areas.
437 For completeness, we also analyzed correlations between familiar – correct rejection ERP
438 differences and behavioral estimates of recollection. These analyses revealed no significant
439 clusters. To summarize, we found that ERPs were sensitive to familiarity-based recognition,
440 both at the overall group level and at the level of individual differences.

441
442 Next, to quantify ERP correlates of recollection, we contrasted ERPs associated with
443 remember hits against ERPs associated with familiar hits. Analyses at the group level
444 revealed significant differences corresponding to two spatiotemporal clusters ($p=0.001$ cluster
445 corrected for both clusters). The first cluster extended from approximately 250 to 450 ms
446 during which ERPs for remember trials were more negative than ERPs for familiar trials and
447 had widespread scalp topography (Figure 4a upper panel). The second cluster extended from
448 approximately 550 to 700 ms, and manifested as an enhanced positivity for remember trials
449 compared to familiar trials. This latter effect had a centro-parietal scalp topography, largely
450 consistent with prior reports of recollection-related ERP effects (Addante et al., 2012b;
451 Curran, 2000; Duarte et al., 2004; Ranganath and Paller, 2000; Rugg et al., 1998b, 1998a;
452 Rugg and Curran, 2007; Wilding, 2000; Wilding and Ranganath, 2012). Figure 4a shows
453 topographic distributions of the ERP differences between remember and familiar trials during
454 these time windows.

455 Having established significant remember – familiar differences at the group level, we
456 next conducted a data-driven analysis to determine whether ERP differences between these
457 trial types were positively correlated with dual process estimates of recollection. This analysis
458 revealed a significant cluster ($p=0.001$ cluster corrected) in the observed data extending from
459 600 to 650 ms with a broad scalp distribution, particularly over central and fronto-central
460 scalp sites (Figure 4b), where ERP amplitudes were positively correlated with recollection
461 estimates. For completeness, we also analyzed correlations between remember – familiar ERP
462 differences and behavioral estimates of familiarity. These analyses revealed no significant
463 clusters. To summarize, these analyses revealed significant ERP correlates of recollection at
464 the group level and at the level of individual differences.

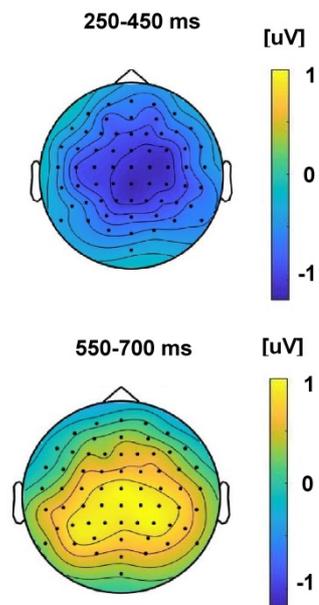
465 To exclude a possible explanation of the results hinging on signal-to-noise ratio
466 differences as a function of the number of trials contributing to the ERPs, we reran the

467 analysis after equating the number of trials, which resulted in obtaining the same pattern of
468 results. In supplemental figures 4 and 5 we also present topographic maps illustrating
469 correlates of familiarity and recollection for the whole analyzed time window with
470 highlighted electrode clusters on the basis of which the null hypothesis was rejected.

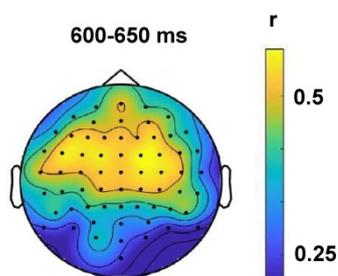
471

472

(A) Overall remember-familiar ERP



(B) ERP-behavior correlation



473 **Figure 4.** ERP Correlates of recollection. (A) Topographic maps illustrate distributions of
474 mean ERP differences between remember and familiar trials during time windows that
475 correspond roughly to the significant clusters identified in the data-driven permutation
476 analysis. (B) A topographic map illustrates a distribution of correlations between Remember –
477 Familiar ERP differences with dual process estimates of recollection. Color bars show ERP
478 voltage difference (A) or Pearson's r correlation coefficient values (B).

479

480 **Discussion**

481 This study was designed to test the hypothesis that the neural correlates of recollection
482 and familiarity-based recognition are predictive of individual differences in episodic memory

483 performance. Data-driven analyses of ERPs during memory retrieval revealed overall effects
484 broadly consistent with previous ERP studies of recognition memory, with an early mid-
485 frontal ERP modulation that was enhanced for familiarity-based recognition, and a late
486 posterior ERP modulation that was enhanced for recollection-based recognition. Critically,
487 our data-driven analyses only revealed significant relationships between Familiar– Correct
488 Rejection ERPs and individual familiarity estimates (Figure 3b), whereas we only found
489 significant relationships between Remember – Familiar ERPs and individual recollection
490 estimates (Figure 4b). ERP-behavior correlations were seen at a relatively late latency
491 (>500ms post-stimulus) for both familiarity and recollection. These findings suggest that
492 ERPs can provide useful markers of individual differences in recognition memory processes.

493 Although the central goal of the study was to look at individual differences in behavior
494 and electrophysiology, we first wanted to determine the extent to which our results concurred
495 with results from previous ERP paradigms. Consistent with a large body of evidence from
496 ERP studies of recognition memory, we observed an early ERP old-new effect related to
497 familiarity-based recognition and a late old-new effect related to recollection-based
498 recognition. It is notable, however, that many other recognition memory correlates have been
499 reported with time courses and scalp distributions that vary across paradigms (c.f., Wilding
500 and Ranganath, 2012). This variance makes sense, because processes that support recognition
501 memory can occur within 200 ms of the onset of a word or picture, and there is likely be
502 extensive parallel, feedforward, and feedback processing throughout different brain networks
503 (Clarke et al., 2011; Clarke and Tyler, 2014; Halgren et al., 2006; Marinkovic et al., 2003),
504 resulting in field potentials that overlap in space and time at the scalp. Thus, the use of
505 different types of stimuli (e.g., words, pictures, etc.) or different kinds of memory decision
506 procedures across studies could likely engage different subprocesses that could impact the
507 timing or topography of ERP correlates of memory (Bader et al., 2020; Busch et al., 2004;
508 Taylor, 2002; Yonelinas, 2002).

509 In order to address this concern in an unbiased manner, we adopted a statistical
510 technique, non-parametric cluster-based permutation testing (Maris and Oostenveld, 2007),
511 that relies on the data to determine both significant time windows and electrode clusters. This
512 was also adapted when identifying spatiotemporal clusters that correlate with behavioral
513 memory measures. Our analyses revealed a rich picture, such that two different
514 spatiotemporal clusters were associated with familiarity, and two different clusters were
515 associated with recollection. Although our analysis methods do not permit precise inferences

516 about the timing of these effects, it is notable that recollection and familiarity were each
517 associated with clusters in relatively early and late time windows. This analysis enabled us to
518 identify reliable effects without relying on assumptions from previous work, and, in turn, may
519 have enabled us to uncover the relationships between electrophysiology and memory
520 measures that have been previously mixed in other reports in the literature (Curran et al.,
521 2006; Friedman and Johnson, 2000; Mecklinger, 2006, 2000; Olichney et al., 2000; Paller and
522 Kutas, 1992; Rhodes and Donaldson, 2007; Rugg et al., 1998a; Smith, 1993; Tsivilis et al.,
523 2001; Wilding and Ranganath, 2012). For instance, we identified different neural correlates of
524 familiarity and recollection, but these results did not conform to the expectation (Curran et al.,
525 2006) that familiarity-related neural processes should always precede those related to
526 recollection.

527 A second key finding from this study is that ERPs also tracked individual differences
528 in familiarity- and recollection-based recognition. Again, the use of data-driven approaches
529 revealed results that might not have been obtained by assuming that individual differences in
530 behavior should correlate with the magnitude of well-known ERP old-new effects. As shown
531 in Figure 3, ERP correlations with familiarity estimates were seen over right posterior sites
532 approximately 500-550 ms post-stimulus, a window which overlapped with the time window
533 during which a significant group-level ERP familiarity effect was observed over central sites.
534 Likewise, as shown in Figure 4, ERP correlations with recollection estimates were seen
535 approximately 600-650 ms at frontal scalp sites, a time window that overlapped with a
536 significant group-level ERP recollection effect with a centro-parietal topography.

537 One way to think about these results is that those who had higher behavioral estimates
538 of recollection or familiarity showed ERP effects that were larger in magnitude than those
539 who had lower recollection or familiarity estimates. To explore this possibility, we calculated
540 grand averaged ERPs on remember, familiar and correct rejection trials for low and high
541 (median split) familiarity estimate performers (Supplemental Figure 1) and low and high
542 (median split) recollection estimate performers (Supplemental Figure 2). Both the familiarity
543 (familiar – correct rejection difference within 500-600 ms time window) and recollection
544 effects (remember – familiar difference within approximately 600-700 ms time window) were
545 more pronounced in high than in low performers. Alternatively, it is possible that group-level
546 ERP recollection and familiarity effects overlapped in time from separate ERP components
547 that differentiated between those with high versus low performers. Although the latter
548 possibility is less parsimonious, we cannot conclusively differentiate between topographic

549 changes driven by increases in the strength of activity in the same configuration of neural
550 sources vs. topographic changes driven by the involvement of different neural sources
551 (Urbach and Kutas, 2002).

552 Our analyses were guided by models which propose that recollection and familiarity
553 independently contribute to successful recognition memory. These models align with a vast
554 body of evidence from lesion, intracranial EEG, and functional neuroimaging evidence
555 demonstrating that familiarity disproportionately depends on representations of item-related
556 information by the perirhinal cortex, whereas recollection disproportionately depends on
557 binding of item and context information by the hippocampus (Davachi, 2006; Eichenbaum et
558 al., 2007; Ranganath, 2010b; Ranganath and Ritchey, 2012). Other researchers, however,
559 have been more agnostic about memory *content*, instead focusing on the idea that all retrieved
560 information is summed together to provide an overall sense of the *strength* of a memory
561 (Kelley and Wixted, 2001; Wixted, 2007). According to this view, “remember” and “familiar”
562 responses reflect different points along a single one-dimensional continuum of memory
563 strength (Kelley and Wixted, 2001). It is important to note that such single-process models do
564 not attempt to characterize memory per se, but rather to account for the way decisions are
565 made on a memory task—for instance, it is possible that there are qualitatively different
566 neural signals for different kinds of memory content, and that the information is integrated
567 into a single strength of evidence signal when making a behavioral response (Gold and
568 Shadlen, 2007, 2001).

569 Our study was not designed to conclusively adjudicate between single- and dual-
570 process models, but it is not clear that a single memory strength process would be sufficient to
571 fully account for our results. If one were to assume that remember and familiar responses vary
572 along a single memory strength continuum, and if ERPs reflect an aggregated measure of
573 memory strength, then we would expect any ERP old-new effect to be larger for remember
574 responses than for familiar responses. However, as we can see from the raw traces in Figure 2,
575 prior to approximately 500 ms, there is an enhanced positivity for familiar trials that is
576 virtually absent for remember trials. This might seem counterintuitive, but it aligns with the
577 dual process model. According to models that assume independent contributions of
578 recollection and familiarity to recognition (Aggleton and Brown, 1999; Atkinson and Juola,
579 1973, 1974; Atkinson et al., 1974; Eichenbaum et al., 1994; Jacoby, 1984, 1991, 1983; Jacoby
580 et al., 1992; Jacoby and Dallas, 1981; Mandler, 1980; Norman and O’Reilly, 2003; Yonelinas,
581 2002, 2001a, 2001b, 1999, 1997, 1994), a familiar response is made only when familiarity is

582 very high and recollection has failed. A remember response, in turn, happens when
583 recollection is successful, even if the item's familiarity is relatively low. Thus, the model
584 would predict that an ERP correlate of familiarity can be very large on familiar trials and
585 attenuated, or even absent, on remember trials (see also Diana et al., 2011). Moreover, if we
586 solely consider behavioral performance, we can look at the associated, or source, information
587 that can be retrieved when an item is successfully remembered. Although there is evidence
588 that both familiarity and recollection can support accurate source memory (Addante et al.,
589 2012b; Diana et al., 2011, 2008; Yonelinas, 2001a), recollection-based responses are more
590 closely associated with retrieval of contextual details (Diana et al., 2012; Dimsdale-Zucker et
591 al., 2018; Ranganath, 2010b, 2010a; Ranganath and Rainer, 2003). We observed significantly
592 above chance source memory performance for items correctly given both familiar and
593 remember responses. However, source memory performance on familiar trials was
594 significantly lower than for remember trials. This fits with the dual-process account of
595 recognition memory phenomenology (Park and Donaldson, 2019; Yonelinas, 2002, 1994;
596 Yonelinas et al., 2010). Another key point is that recollection dual process estimates only
597 correlated with remember – familiar ERPs and familiarity estimates only with familiar –
598 correct rejection ERPs. Moreover, neither correlation effects overlapped in time and
599 topography. The findings are compatible with a dual-process account. However, we
600 acknowledge that the present non-parametric cluster-based permutation test methods do not
601 allow us to make strong conclusions about the topography and timing of the familiarity and
602 recollection ERP effects.

603 Another controversy in prior ERP studies of recognition memory has focused on the
604 functional significance of the mid-frontal ERP old-new effects (Bridger et al., 2012; Paller et
605 al., 2012, 2007; Voss et al., 2012; Voss and Federmeier, 2011; Voss and Paller, 2006; Yovel
606 and Paller, 2004). Results from many studies have supported the idea that this old-new effect
607 is enhanced during familiarity-based recognition, and that it is relatively insensitive to factors
608 that influence recollection (Addante et al., 2012b; Bridger et al., 2012; Curran, 2004, 2000;
609 Friedman and Johnson, 2000; Park and Donaldson, 2019; Rugg et al., 1998a; Rugg and
610 Curran, 2007; Wilding and Herron, 2006; Wilding and Ranganath, 2012). However, a number
611 of findings also support the idea that the mid-frontal old-new effect could instead reflect
612 conceptual priming, which refers to more fluent processing of conceptual information that has
613 been recently encountered (Guillem et al., 2001; Jelicic, 1995; Levy et al., 2004; Mitchell and
614 Bruss, 2003; Nessler et al., 2005; Olichney et al., 2000; Paller et al., 2012, 2007; Ullsperger et

615 al., 2000; Voss and Paller, 2007, 2006). For instance, thinking about the meaning of the word
616 “banana” might make it more likely to come into mind when asked to generate the names of
617 fruit words. Yovel and Paller (2004) used photographs of faces never seen before the
618 experiment as stimuli, to isolate a pure familiarity effect. The authors found no association
619 between familiarity and N400s and suggested that familiarity with faces may arise by a subset
620 of the neural processing responsible for recollection, while the N400 reductions observed in
621 the literature may reflect verbally mediated conceptual priming effects instead of familiarity.
622 Another study (MacKenzie and Donaldson, 2007) obtained similar posterior old/new effect
623 indexing familiarity for faces. However, in contrast to Yovel and Paller, the old/new effects
624 associated with familiarity and recollection were topographically dissociable, consistent with
625 a dual process view of recognition memory.

626 MacLeod and Donaldson (2017) also investigated the functional utility of the left
627 parietal old/new effect using verbal stimuli. Their results revealed that ERP measures (defined
628 as the mean ERP old/new difference within 500-800 ms post-stimulus averaged across left
629 parietal electrodes: P1, P3, and P5) of retrieval were not related to behavioral performance.
630 The authors concluded that the relationship between the left parietal effect and recollection is
631 more complex than previously thought in the sense that the variation in the magnitude of the
632 left parietal old/new ERP effect does not always reliably predict variation in episodic
633 recollection between participants. However, the paper does not fully address the relation
634 between ERPs and behavioral estimates of familiarity and recollection for several reasons: (1)
635 the ERP effect was restricted only to the late left parietal effect, (2) the behavioral measures
636 of recollection used in this study does not dissociate recollection from familiarity, and (3)
637 only 20 participants were included in the correlation of R/K effect magnitude with behavioral
638 data.

639 More recently, Wang et al. (2020) employed conceptually impoverished items
640 (kaleidoscope images) as stimuli in a recognition memory test with a modified
641 Remember/Know paradigm and they also observed that ERPs for Know hits were more
642 positive than those for Correct Rejection items within 500-800 ms. Putting all these results
643 together, there is considerable evidence suggests that the N400-like ERP effects are
644 modulated by conceptual priming and familiarity (Bader and Mecklinger, 2017; Nessler et al.,
645 2005; Wolk et al., 2004).

646 The controversy over ERP correlates of familiarity and conceptual priming relates to
647 the broader question regarding the relationship between fluent processing and familiarity-
648 based recognition memory. Substantial evidence exists to suggest that neural processes
649 associated fluent processing of conceptual information are also related to familiarity—for
650 instance, N400-like potentials occur in the perirhinal cortex during both conceptual priming
651 and recognition memory (Nobre and McCarthy, 1995, 1994; Staresina et al., 2012).
652 Moreover, damage to the left perirhinal cortex impairs both conceptual priming and
653 familiarity-based recognition memory (Bowles et al., 2007; Wang and Yonelinas, 2012a,
654 2012b; Wang et al., 2010). Finally, fMRI studies have shown that activity in the left perirhinal
655 cortex during encoding predicts both conceptual priming and familiarity-based recognition,
656 and perirhinal activity has been correlated with behavioral performance on both conceptual
657 priming and familiarity-based recognition measures (Dew and Cabeza, 2013; Diana et al.,
658 2010; Haskins et al., 2008; Heusser et al., 2013; Ranganath et al., 2004; Voss et al., 2009;
659 Wang et al., 2015, 2014). Although these findings do not rule out the possibility that
660 conceptual priming and familiarity can be dissociated (Paller et al., 2012), they are consistent
661 with the broader idea that fluent processing of an item’s conceptual features can contribute to
662 one’s subjective sense that the item is familiar (Mecklinger and Bader, 2020; Taylor and
663 Henson, 2012; Wang and Yonelinas, 2012b).

664 In summary, the current study presents evidence to suggest that ERPs can be used to
665 identify neural correlates of recollection and familiarity, both at the group level, and at the
666 level of individual differences. The present findings provide support for the idea that ERPs
667 can be used as biomarkers of underlying memory processes in healthy individuals, patient
668 populations, or specific populations, like older adults or children (MacLeod and Donaldson,
669 2017). The combined use of behavioral and ERP measures, as in the present study, might be
670 especially useful in the identification of those who are at risk for disorders such as
671 Alzheimer’s disease (Xia et al., 2020). Additionally, the present results also highlight the
672 potential value of data-driven analysis methods as a means to identify neural correlates of
673 cognitive processes, complementing approaches that focus on specific, well-characterized
674 components.

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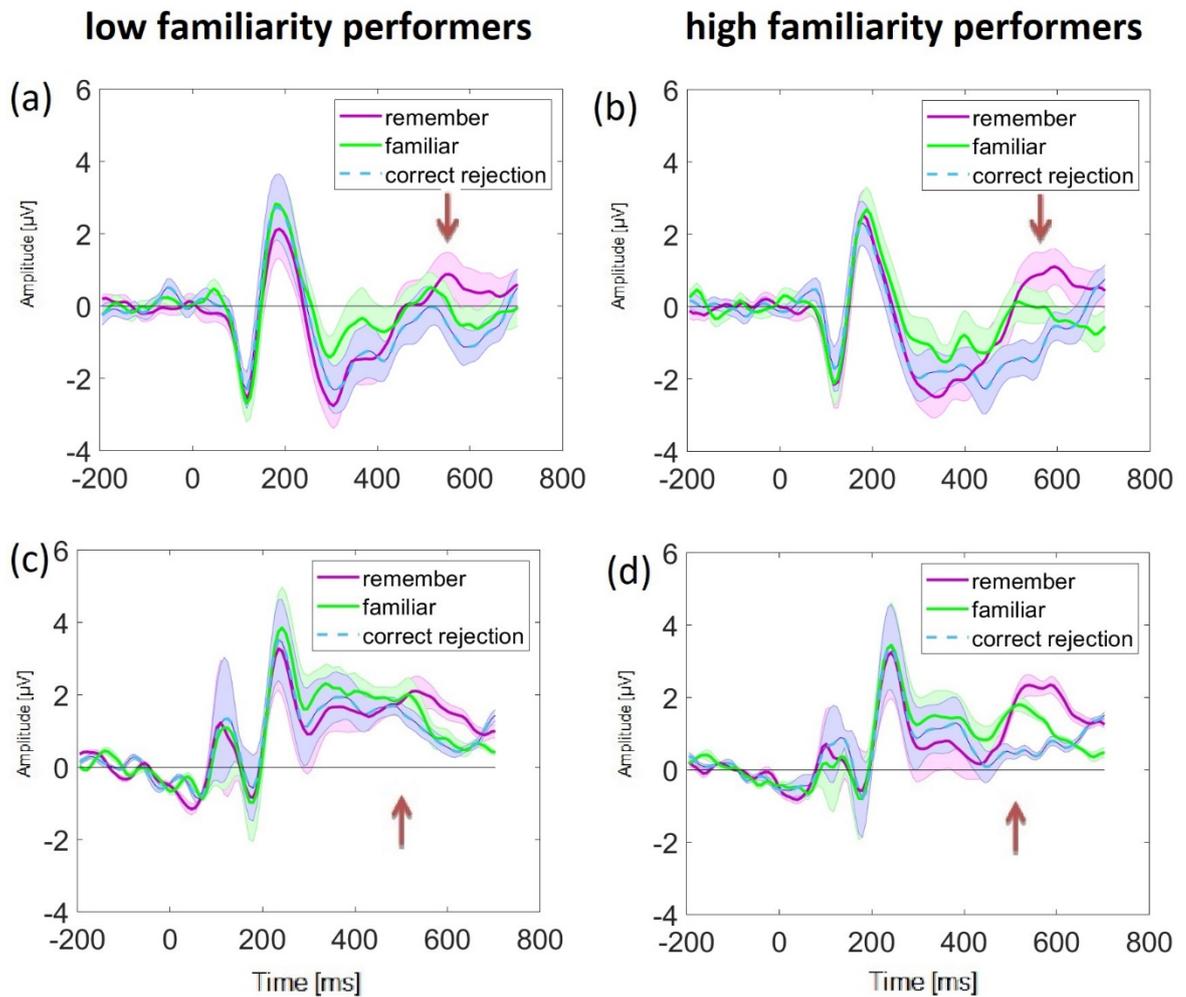
678 Acknowledgement

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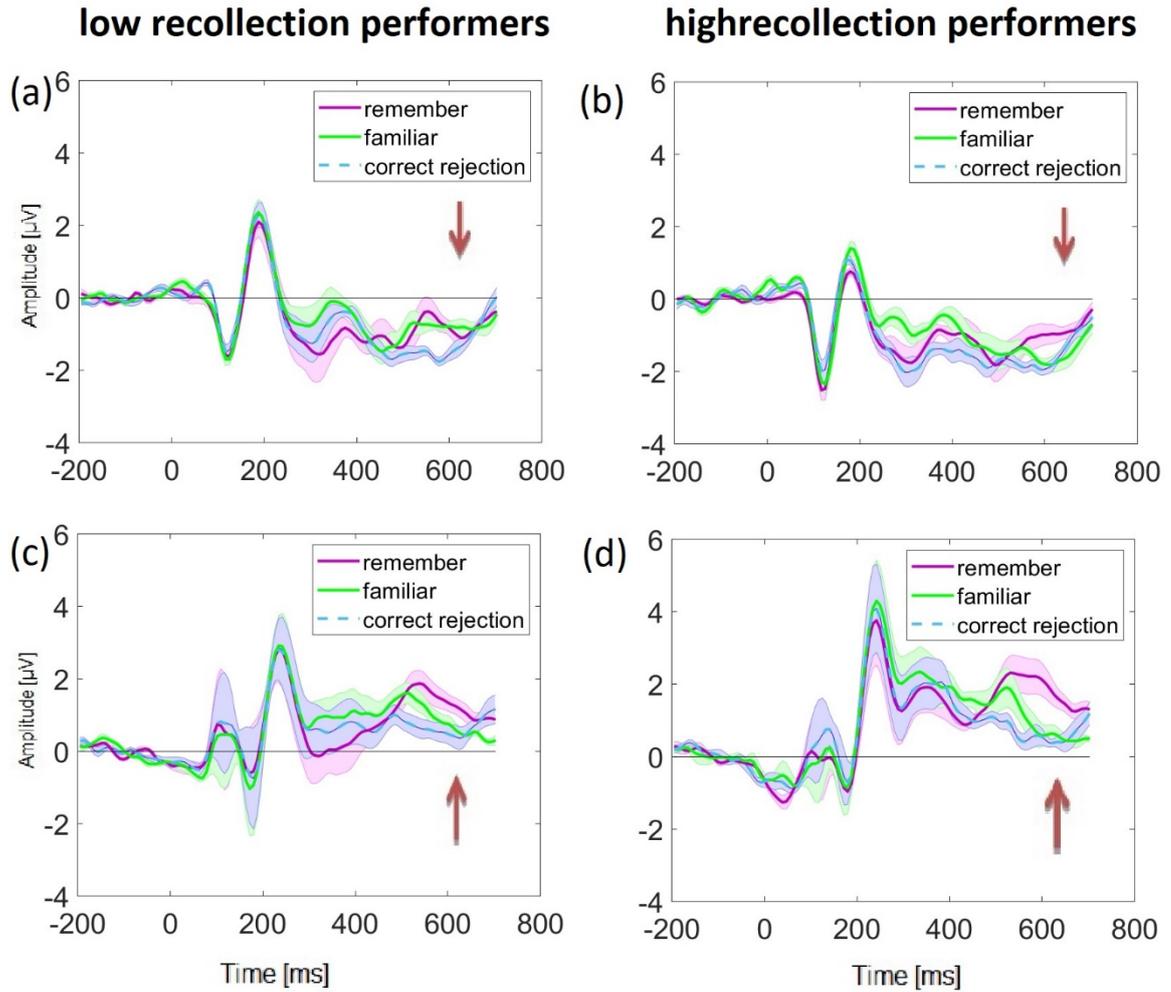
688 Supplementary materials



689

690 **Supplemental Figure 1.** ERP correlates of familiarity. Grand averaged ERPs on Remember
691 (purple), Familiar (green) and Correct Rejection (dashed blue) trials for low and high
692 performers based on median split. Shaded areas represent standard deviation of the mean.
693 Arrows are meant to delineate time periods of interest, but do not indicate statistical
694 comparisons: (a) low familiarity estimate performers at Cz, (b) high familiarity estimate
695 performers at Cz, (c) low familiarity estimate performers at right parietal (P2, P4, P6, P8,
696 PO4, PO8), and (d) high familiarity estimate performers at right parietal (P2, P4, P6, P8,
697 PO8). Note that the average traces from electrode groups are presented for visualization
698 purposes, but electrodes were analyzed separately in the data-driven statistical analyses.

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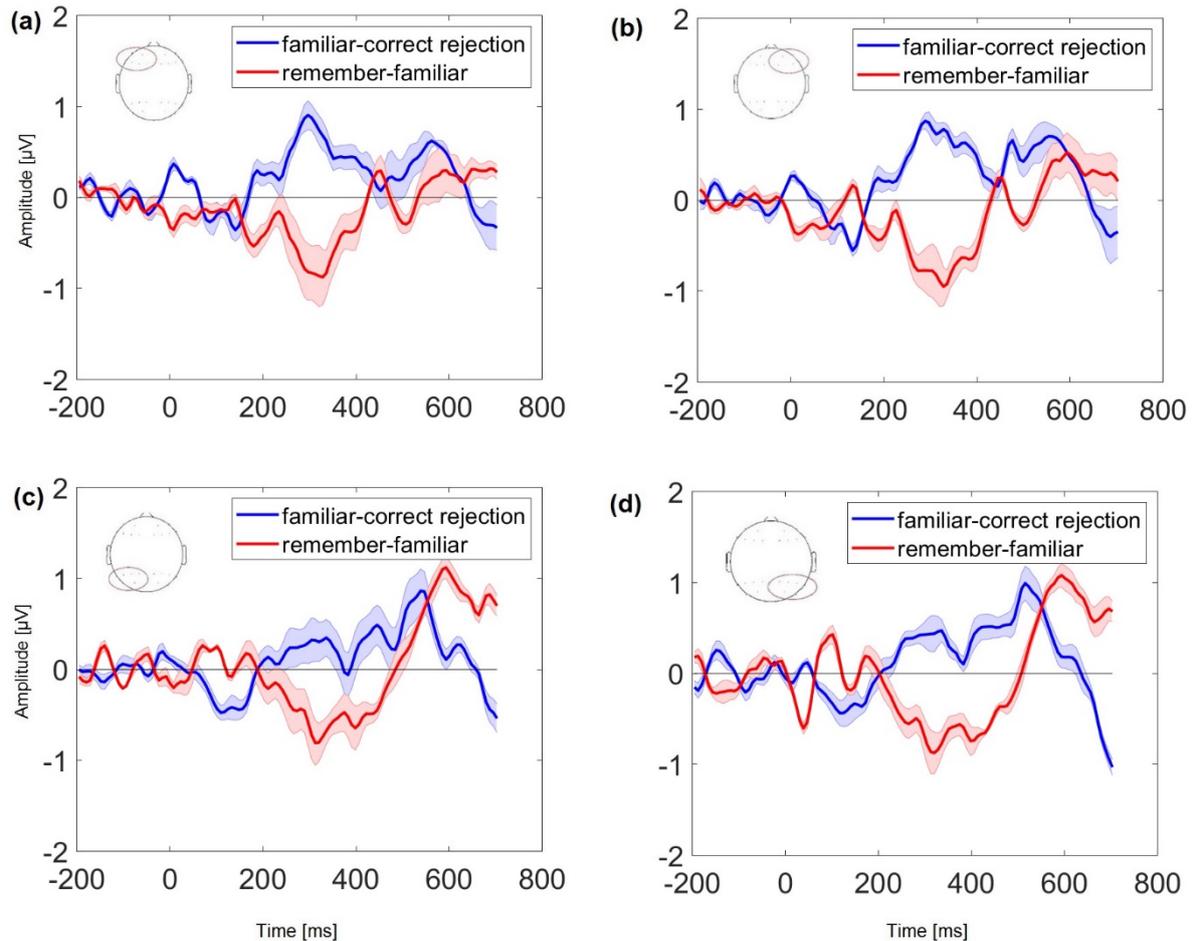


700

701 **Supplemental Figure 2.** ERP correlates of recollection. Grand averaged ERPs on Remember
702 (purple), Familiar (green) and Correct Rejection (dashed blue) trials for low and high
703 performers based on median split. Shaded areas represent standard deviation of the mean.
704 Arrows are meant to delineate time periods of interest, but do not indicate statistical
705 comparisons: (a) low recollection estimate performers at frontal (F1, F3, F5, F7, AF3, AF7,
706 F2, F4, F6, F8, AF4, AF8), (b) high recollection estimate performers at frontal (F1, F3, F5,
707 F7, AF3, AF7, F2, F4, F6, F8, AF4, AF8), (c) low recollection estimate performers at parietal
708 (P1, P3, P5, P7, PO3, PO7, P2, P4, P6, P8, PO4, PO8), and (d) high recollection estimate
709 performers at parietal (P1, P3, P5, P7, PO3, PO7, P2, P4, P6, P8, PO4, PO8). Note that the
710 average traces from electrode groups are presented for visualization purposes, but electrodes
711 were analyzed separately in the data-driven statistical analyses.

712

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714

715 **Supplemental figure 3.** ERP correlates of recollection and familiarity. Grand averaged ERP

716 difference waveforms: Familiar-minus-Correct Rejection (red) and Remember-minus-

717 Familiar (blue), separately averaged for four groups of channels split by frontal and parietal

718 for each hemisphere (Woodruff et al., 2006): (a) left frontal (F1, F3, F5, F7, AF3, AF7), (b)

719 right frontal (F2, F4, F6, F8, AF4, AF8), (c) left parietal (P1, P3, P5, P7, PO3, PO7), and (d)

720 right parietal (P2, P4, P6, P8, PO4, PO8). Shaded areas represent standard deviation of the

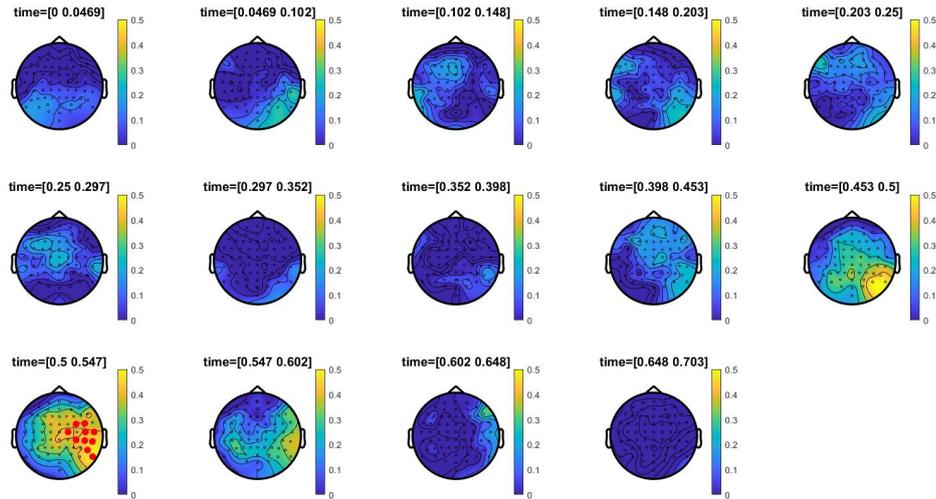
721 mean. Note that these average traces from electrode groups are presented for visualization

722 purposes, but electrodes were analyzed separately in the data-driven statistical analyses.

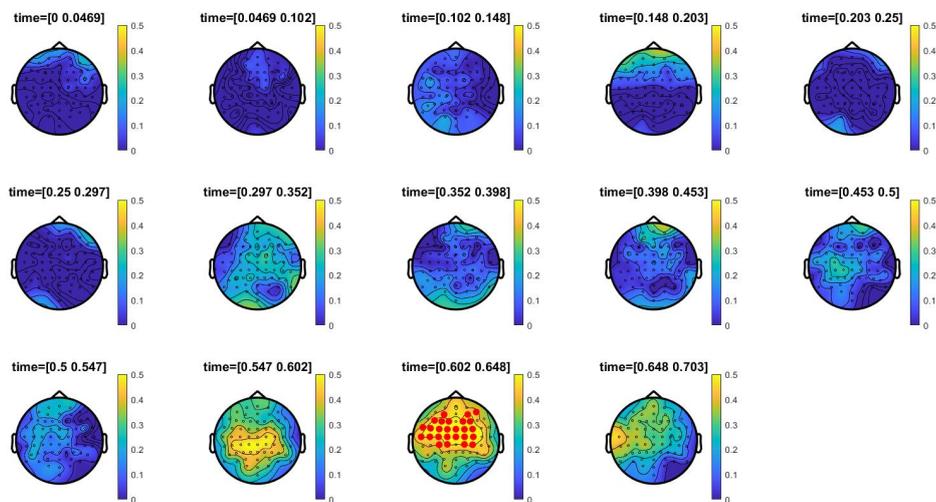
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727 **Supplemental figure 4.** ERP correlates of familiarity. Topographic maps illustrate a
728 distribution of correlations between Familiar - Correct Rejection ERP differences with dual
729 process estimates of familiarity for 50 ms time bins. Electrode clusters on the basis of which
730 the null hypothesis was rejected are highlighted with red asterisks. All timepoints and all 64
731 electrodes were included in the permutation test within specified 0-700 ms time window, at P
732 < 0.05 , cluster corrected. Color bars show Pearson's r correlation coefficient values.
733
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736 **Supplemental figure 5.** ERP Correlates of recollection. Topographic maps illustrate a
737 distribution of correlations between Remember – Familiar ERP differences with dual process
738 estimates of recollection for 50 ms time bins. Electrode clusters on the basis of which the null
739 hypothesis was rejected are highlighted with red asterisks. All timepoints and all 64 electrodes
740 were included in the permutation test within specified 0-700 ms time window, at $P < 0.05$,
741 cluster corrected. Color bars show Pearson's r correlation coefficient values (B).
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Item response (scored)	Highly confident	Moderately confident	Somewhat confident	Not at all confident
Recollect	66.36 (27.18)	20.91 (14.21)	4.19 (3.25)	1.33 (0.58)
Familiar	19.06 (14.06)	18.42 (6.33)	16.97 (8.84)	3.76 (3.07)
Miss	9.93 (9.18)	12.53 (8.25)	9.97 (5.88)	2.89 (2.35)
Correct rejection	28.91 (19.59)	24.39 (12.25)	13.33 (7.88)	4.11 (2.83)

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Supplemental Table 1. Mean and standard deviation counts of confidence level by scored item response.

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