

Visual short-term memory related EEG components in a virtual reality setup

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Supplementary material

Control analyses

To exclude the possibility that the effects of memory load and eccentricity on the CDA were specific to our selection of the CDA time window (which we gained from the data-driven temporal localizer approach), we conducted a set of supplementary analyses. First, we calculated the mean CDA using the time window reported by Hakim et al. (2019). This alternative time window (400 to 1,450 ms after stimulus onset) started (slightly) later and lasted (substantially) longer than the one yielded by the temporal localizer approach (388 to 1,088 ms). In line with our results, we observed also for this alternative time window that the CDA mean amplitude varied significantly with memory load ($F(1,20) = 14.74, p = .001$) while stimulus eccentricity did not significantly influence mean CDA amplitude ($F(2,40) = 0.45, p = .638$). In contrast to the analysis based on the data-driven time window, there was no significant interaction of eccentricity with the effect of memory load ($F(2,40) = 3.04, p = .059$). Post-hoc paired t -tests, however, revealed the same pattern as found in the main analysis: the difference between trials with low and high memory load was significant for the eccentricities of 4 dva ($\Delta CDA_{\text{low-high}} = 0.47, 95\% \text{ CI } [0.16, 0.78], p = .004$) as well as 9 dva ($\Delta CDA_{\text{low-high}} = 0.39, 95\% \text{ CI } [0.16, 0.62], p = .002$), but not for 14 dva ($\Delta CDA_{\text{low-high}} = 0.14, 95\% \text{ CI } [-0.06, 0.33], p = .155$). To further explore this pattern with a more sensitive analysis approach, we calculated a hierarchical linear mixed-model (using the statistical software package *lme4*; Bates et al., 2015)

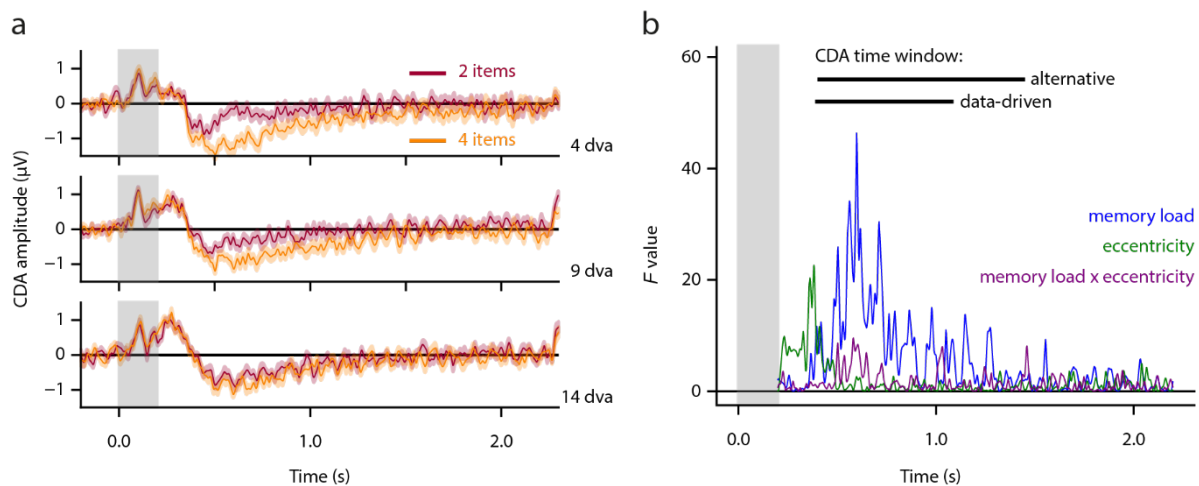
24 which takes single-trial data into account and allows for modeling random intercepts per
25 participant. We modeled the mean CDA amplitude (averaged across channels in our ROI and time
26 points of the respective time window) as a function of memory load (treatment contrast; set size of
27 2 items as a baseline), eccentricity (treatment contrast; 9 dva as a baseline) as well as their
28 interaction. We fitted one of these models separately for the mean CDA calculated for the time
29 window identified with the temporal localizer method and for the alternative time window, and we
30 determined the significance of the model predictors using the R package lmerTest (Kuznetsova et
31 al., 2017).

32 This analysis revealed the same pattern of results for both time windows (Table S1): memory load
33 had a strong influence on the mean CDA amplitude while eccentricity did not have a significant
34 main effect. The interaction effect between memory load and eccentricity was small (compared to
35 the main effect of memory load) but significant for both time windows and was driven by the trials
36 with the largest stimulus eccentricity (14 dva) which exhibited a smaller difference between the
37 memory load conditions as compared to trials with 9 dva or 4 dva. There was no significant
38 difference between the two smaller eccentricity conditions. This is the same pattern that we
39 observed with the (non-hierarchical) rmANOVA paired with post-hoc *t*-tests for the data-driven
40 time window. We suspect that we did not observe a significant interaction effect of memory load
41 and eccentricity with the rmANOVA for the a priori time window due to a lack of statistical power.

42 We will discuss in the following paragraph why the interaction effect is smaller in the alternative
43 time window. The more sensitive mixed-model approach allows to detect also weaker effects.
44 Overall, we conclude that the results gained with the original time window (in accordance with the
45 mixed-model) represent a useful while parsimonious description of the data. The key insight: for
46 the largest eccentricity (14 dva), the load effect on the CDA was substantially weakened as compared
47 to the two smaller eccentricities (Figure S1a).

48 Based on Figure S1a, we assumed that the interaction effect was weaker for the alternative time
49 window as eccentricity might predominantly affect early components of the ERP (like the PNP
50 component). To address this concern, we conducted a time-resolved version of the rmANOVA
51 approach by fitting the according model ($CDA \sim MemoryLoad * Eccentricity$) separately for each
52 sample during the retention interval (Fig. S1b). This revealed different time courses for the main
53 effects of memory load and eccentricity, with the latter peaking substantially earlier, before the
54 onset of the investigated CDA time windows. As expected, the interaction effect only plays a role
55 during the early phases of the CDA time windows and wears off quickly. By using a longer (and/or
56 later) time window to calculate the mean CDA amplitude, the proportion of samples affected by the
57 interaction effect decreases. As the main effect of memory load is stronger and more long lasting, it
58 can also be found more easily in such longer time windows. The post-hoc t -tests, however, did not
59 show differences in CDA mean amplitude for either of the time windows, suggesting that increasing
60 the length of the interval is not sufficient to counteract the effect of the interaction.

61 Supplementary figures & tables



62

63 **Figure S1:** (a) The effect of memory load (2 vs 4 items) on the CDA per eccentricity level. (b) Effects
 64 of a time-resolved rmANOVA, modelling the CDA (i.e., lateralized ERP in our ROI) as a function
 65 of memory load, eccentricity, and their interaction. It is important to stress the exploratory nature
 66 of this supplementary analysis. Therefore, we refrained from performing significance tests.

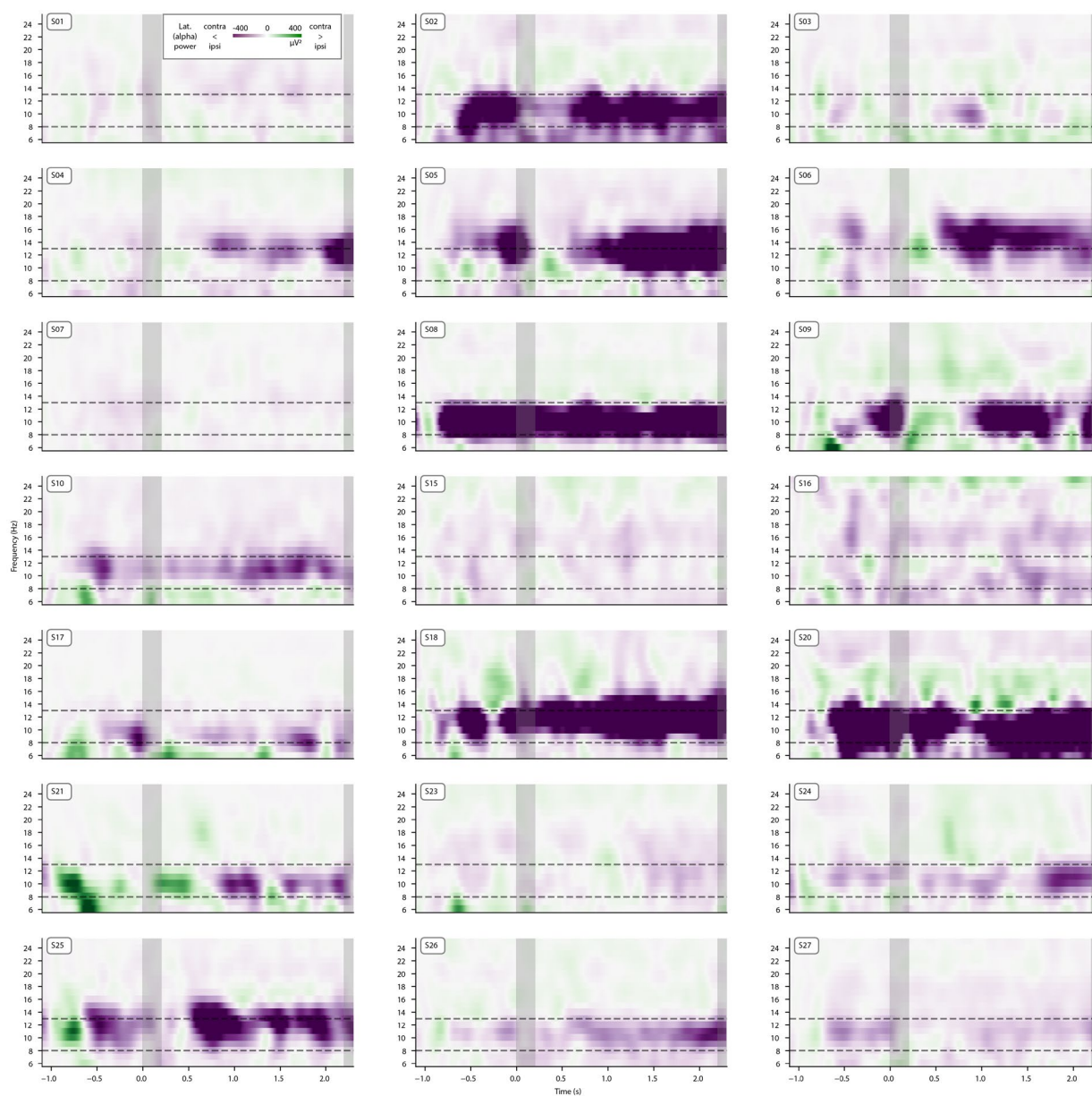
67 **Table S1**68 *Effects in the mixed-model analysis of the mean CDA amplitude for two different time windows*

	predictor	b	t	df	95% CI
CDA time window: 388 – 1088ms	Intercept (2 items; 9 dva)	-0.26	-1.87	34.81	[-0.55; 0.02]
	Memory load (4 vs 2 items)	-0.50	-4.73***	13358.30	[-0.71; -0.29]
	Eccentricity (4 vs 9 dva)	-0.05	-0.45	13358.48	[-0.26; 0.16]
	Eccentricity (14 vs 9 dva)	-0.19	-1.77	13358.38	[-0.39; 0.02]
	Memory load x Eccentricity (4 vs 2 items) x (4 vs 9 dva)	-0.08	-0.54	13358.48	[-0.38; 0.22]
	Memory load x Eccentricity (4 vs 2 items) x (14 vs 9 dva)	0.35	2.37*	13358.37	[0.06; 0.65]
CDA time window: 400 – 1450ms	Intercept (2 items; 9 dva)	-0.17	-1.28	38.38	[-0.44; 0.10]
	Memory load (4 vs 2 items)	-0.44	-4.08***	13358.33	[-0.66; -0.23]
	Eccentricity (4 vs 9 dva)	-0.04	-0.36	13358.55	[-0.25; 0.17]
	Eccentricity (14 vs 9 dva)	-0.19	-1.72	13358.42	[-0.40; 0.03]
	Memory load x Eccentricity (4 vs 2 items) x (4 vs 9 dva)	-0.04	-0.26	13358.55	[-0.34; 0.26]
	Memory load x Eccentricity (4 vs 2 items) x (14 vs 9 dva)	0.32	2.12*	13358.42	[0.02; 0.62]

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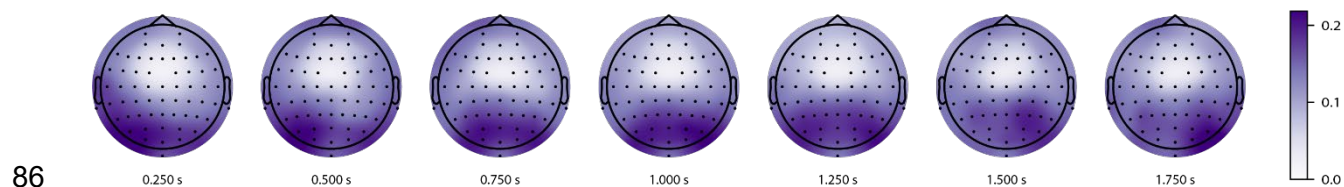
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74 **Figure S2:** Time courses of the decoding performance and the associated spatial patterns for the
75 single participants. Participants are sorted, separately for each subfigure, decreasingly by the
76 maximum average decoding performance in the time window in which the overall decoding
77 performance (across participants) was significantly above chance level. (a) Decoding from the
78 broadband EEG data (ERP). (b) Decoding from time-frequency data (here: alpha frequencies 8–14
79 Hz). As we analyzed induced power, which reflects the non-phase locked signal from an oscillating
80 dipole (i.e., with arbitrary polarity at a given point in time), we show the absolute pattern weights
81 to avoid cancellation across participants and repetitions.

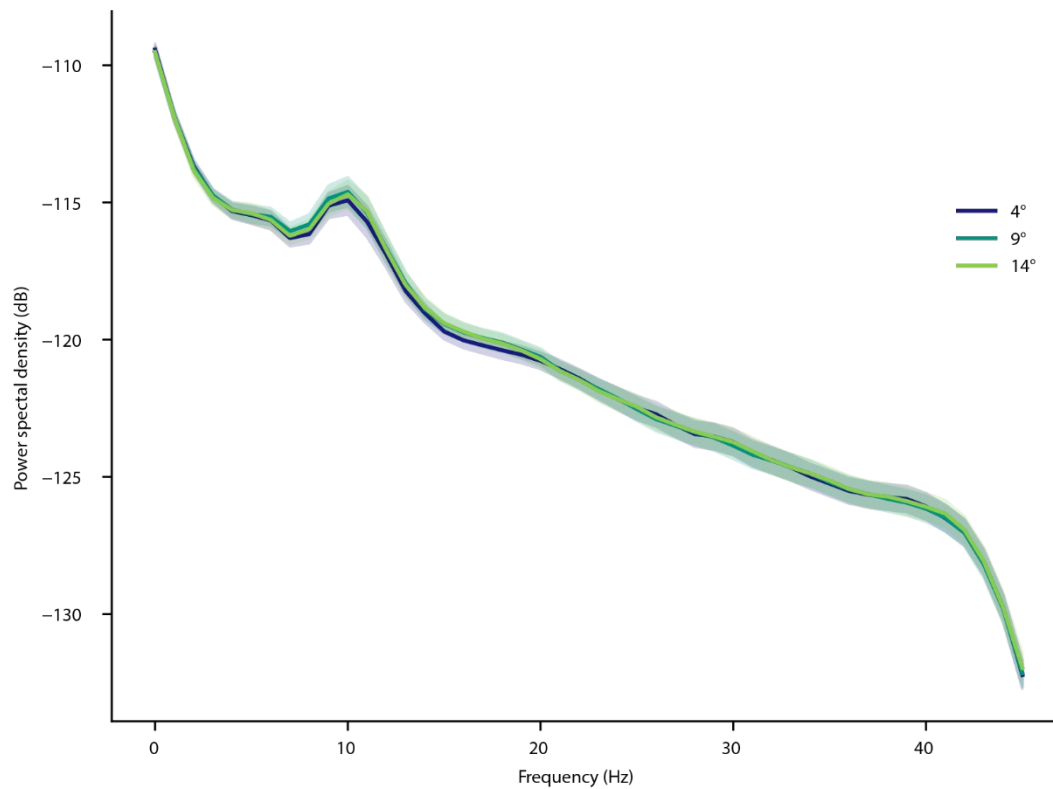


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83 **Figure S3:** Lateralized time-frequency data (difference between contra- and ipsilateral channels) for
 84 each participant. The dotted lines mark the lower and upper limit for the frequency range that we
 85 included in our analyses regarding alpha lateralization.

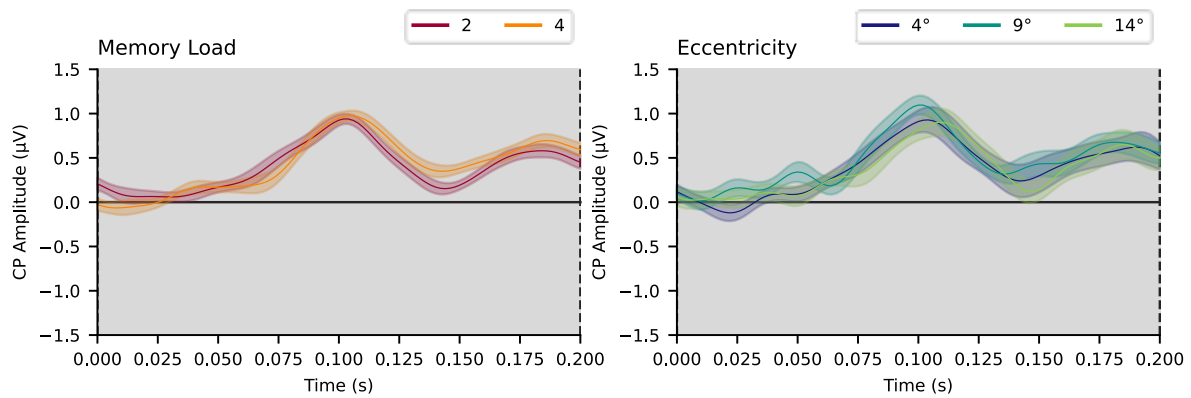


87 **Figure S4:** Averaged absolute pattern weights (gained by multiplying the covariance of the EEG data
88 with the filter weights; Haufe et al., 2014) of the most discriminative CSP component (normalized
89 per time-bin and participant before averaging) for the alpha range. As we analyzed induced power,
90 which reflects the non-phase locked signal from an oscillating dipole (i.e., with arbitrary polarity at
91 a given point in time), we show the absolute pattern weights to avoid cancellation across participants
92 and repetitions.

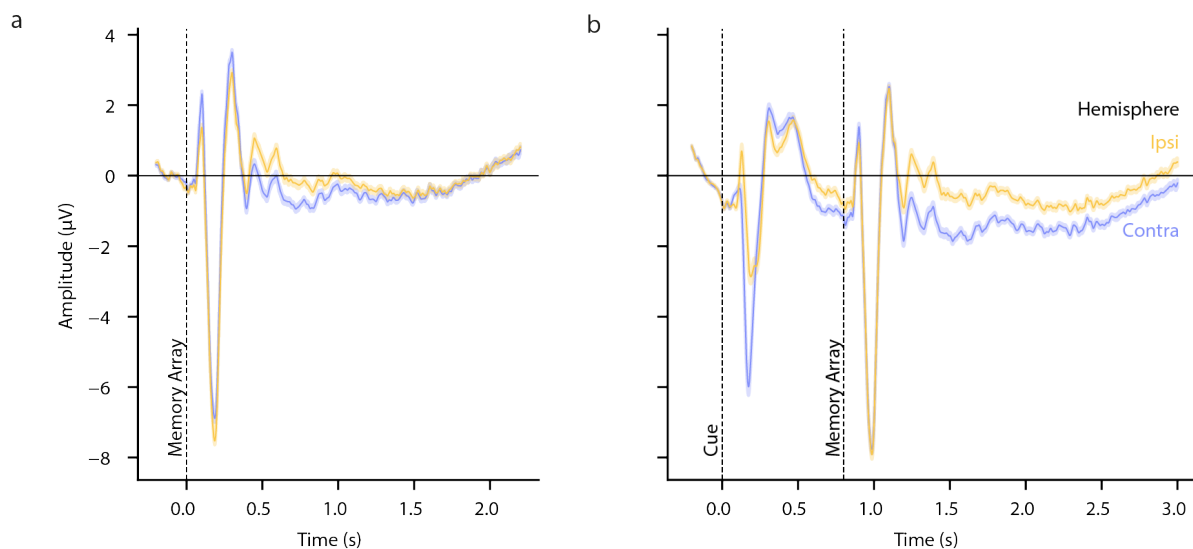


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94 **Figure S5:** Power spectral density (PSD) per eccentricity condition during the retention interval
95 (shaded area: standard error). Data stems from the bilateral channel-pairs which formed the ROI
96 for the calculation of the CDA and lateralized alpha power. We used Welch's method to calculate
97 the PSD with a window size of 512 samples. An alpha peak is recognizable for each of the
98 eccentricity conditions with no evident difference between the conditions.

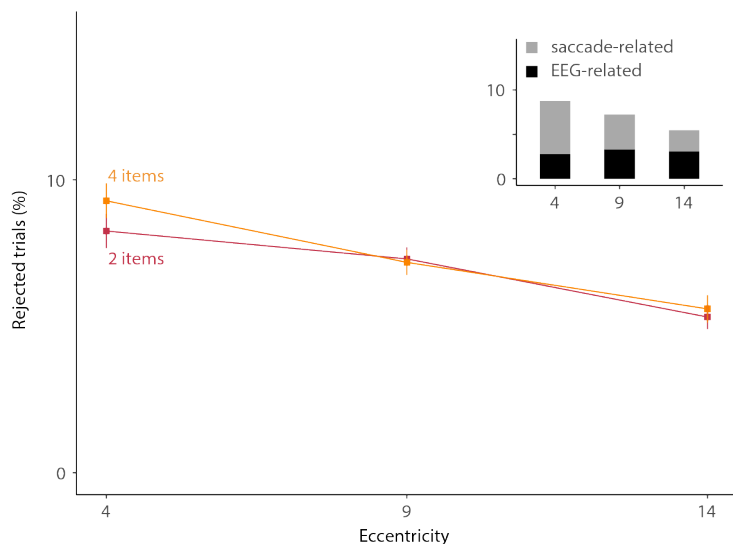


99 **Figure S6:** Early contralateral positivity (zoomed-in version of Figure 2a–b). This early potential
 100 might have been caused by the offset of the asymmetric, arrow-shaped cue which indicated the
 101 relevant hemifield (at time 0, i.e., at the onset of the memory arrays) or indicate enhanced processing
 102 of the stimuli in the cued hemifield (Livingstone et al., 2017). Since in our paradigm, the offset of
 103 the cue and the onset of the memory arrays coincided, it is not possible to clearly determine the
 104 underlying cause of this lateralized potential. However, since it was not modulated by memory load
 105 (in contrast to the CDA) or eccentricity of the memory stimuli (in contrast to the immediately
 106 following PNP component), it is unlikely that its occurrence influenced the effects of primary
 107 interest.



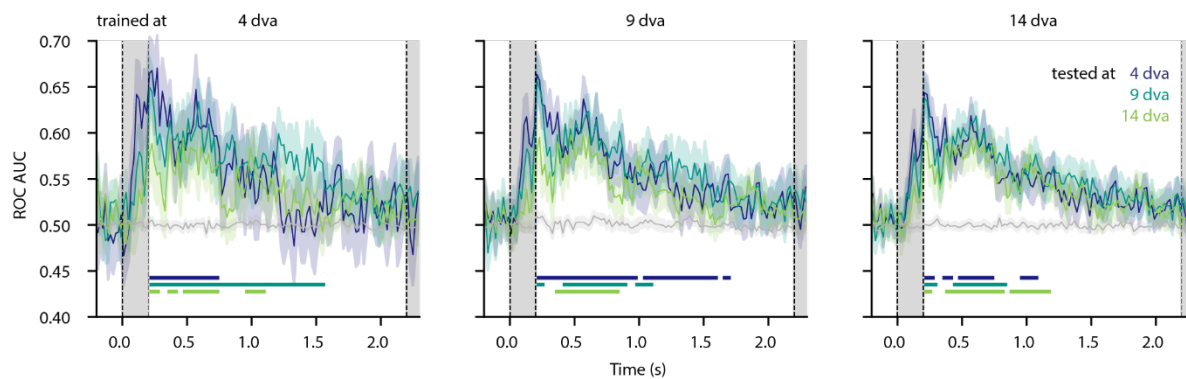
108 **Figure S7:** Event-related potential (ERP) for the hemispheres contra- and ipsilateral to the cued
 109 array (shaded areas: ± 1 SEM corrected for the within-participant measurements). We computed the
 110 average signals from channels that were within the same region-of-interest as those used in the
 111 analysis of the CDA (P3/4, P5/6, PO3/4, PO7/8, O1/2). **(a)** The event-related potential (ERP) time-
 112 locked to memory array onset and normalized by subtracting, on a single-trial basis, the mean
 113 amplitude during a 200 ms baseline period prior to this onset. The analyses regarding the CDA as
 114 well as the ERP-based decoding analyses were conducted on data in this format. **(b)** ERP time-
 115 locked to the onset of the cue (800 ms before onset of the memory stimulus), with the baseline
 116 window being shifted accordingly (i.e., 200 ms before the onset of the cue). The CDA effect is
 117 discernible in both plots. Additionally, the asymmetric cue induced a lateralized response with a
 118 strong positivity during early components (P1, N1), but also another more long-lasting positivity
 119 during later stages of the period between cue and memory array onset (i.e., around 500–800 ms after
 120 cue onset). This period covers the baseline-window chosen for the epochs time-locked to the onset
 121 of the memory array (which formed the basis for our ERP analyses). Therefore, by subtracting the
 122 mean activity in this time-window, we may have introduced an artificial lateralization during the
 123 rest of the ERP. However, this bias would yield a polarity opposite to that of the CDA. Therefore,
 124 such a correction would at most complicate the observation of a CDA, which in turn emphasizes

125 our result that we were able to detect a CDA for all eccentricities. Furthermore, this effect on the
126 baseline-window was (as it occurred before the onset of the relevant stimuli) independent of the
127 experimental manipulations (memory load, eccentricity) and hence impacted all conditions in the
128 same way (i.e., like adding a lateralized constant). Thus, we can rule out concerns of biases in the
129 analysis pipeline as a source underlying our results regarding the CDA or decoding analyses. Finally,
130 we believe that the observed lateralization in the later parts of the cue interval is a concomitant of
131 the spatial shift of attention in response to the cue (Keefe & Störmer, 2021) and is therefore likely
132 to occur in any paradigm which implements spatial cuing preceding the relevant stimulus.
133 Presumably, this effect was also present in various previous studies which used comparable
134 experimental designs/timings. We do not conclude that this is problematic; rather it increases the
135 comparability and compatibility of our findings with previous reports.

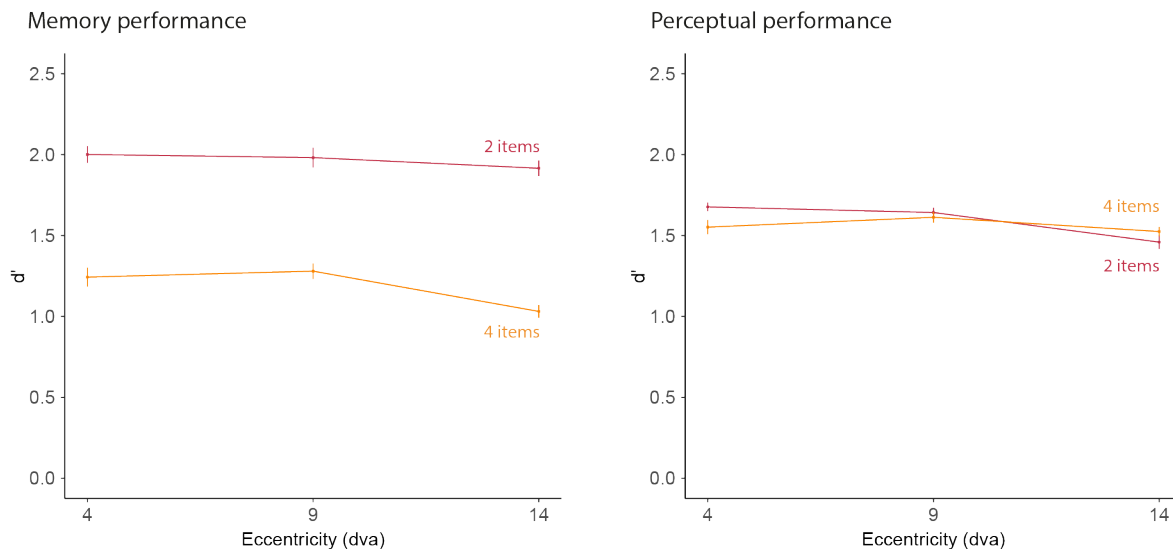


136 **Figure S8:** Average proportion of rejected trials per eccentricity for the ERP analysis. We rejected
 137 trials which contained saccades with an amplitude of at least 2 dva or were classified as noisy by the
 138 cross-validation based trial classification algorithm *autoreject* (Jas et al., 2017). A rmANOVA
 139 identified a significant main effect of eccentricity ($F(2,40) = 8.49, p = .001$) while neither memory
 140 load ($F(1,20) = 0.61, p = .443$) nor its interaction with eccentricity ($F(2,40) = 0.61, p = .550$)
 141 significantly influenced the proportion of rejected trials. Post-hoc t -tests corroborated that in the
 142 largest eccentricity condition significantly less trials were rejected than for 4 dva ($\Delta_{4-14} = 3.31\%$, 95%
 143 CI [1.33, 5.30], $t(20) = 3.48, p = .002$) as well as for 9 dva eccentricity ($\Delta_{9-14} = 1.79\%$, 95% CI [0.46,
 144 3.11], $t(20) = 2.82, p = .011$). There was no difference between the two smaller eccentricities ($\Delta_{4-9} =$
 145 1.53%, 95% CI [-0.13, 3.19], $t(20) = 1.92, p = .070$). Overall, the differences in the number of rejected
 146 trials per condition were rather small. The inset plot shows that the decrease of rejected trials for
 147 the largest eccentricity was the consequence of a smaller number of trials that contained saccades
 148 in the 14 dva eccentricity condition. We suspect that in conditions with smaller eccentricities, there
 149 were more saccadic eye movements, as it is more difficult to suppress reflexive eye movements to
 150 task-relevant stimuli that are located close to the current fixation point. This resulted in a larger
 151 number of involuntary eye movements in these conditions. Regarding the condition with the
 152 highest eccentricity, we excluded the smallest number of trials. Therefore, the absence of a

- 153 significant memory load effect in this condition cannot be attributed to reduced statistical power
- 154 resulting from a smaller number of trials compared to the other eccentricities.



155 **Figure S9:** Results of the cross-decoding analysis for the broadband ERP data. To test whether the
 156 spatial features of the decoding of memory load are substantially different, we trained different
 157 classifiers separately on (80% of) the data from the three eccentricity conditions (see three separate
 158 panels). We then tested each of these classifiers either on the remaining data of the same eccentricity
 159 condition or on 20% of the data from each of the other conditions. The specific parameters of the
 160 classifiers were the same as used in the main decoding analysis on the ERP data (incl. the 100x
 161 repeated 5fold cross-validation). The figure shows the time-resolved decoding performance
 162 averaged across folds, repetitions, and participants and follows the same conventions as Figure 1f.
 163 Classification performance was above chance for each classifier in each condition. However, for
 164 none of the classifiers (trained either on data from the 4, 9, or 14 dva condition), we observed
 165 significant differences between the performances on the different test sets (as tested by a sliding,
 166 cluster-corrected repeated-measures ANOVA).



167

168 **Figure S10:** Behavioral results of the memory task and the perceptual task using d' (i.e., sensitivity)
 169 as a measure of memory performance. Error bars indicate ± 1 SEM, taking into account the repeated
 170 measures design (Baguley, 2012; Morey, 2008).

171 Using d' as a bias-free indicator of memory performance (i.e., sensitivity), we found the same pattern
 172 of results as in the analyses using the proportion of correct responses (i.e., accuracy; see Figure 1b-
 173 c). In the Memory Task, d' was strongly modulated by memory load and varied only little as a
 174 function of eccentricity. Participants showed a significantly higher d' in trials with low ($M = 1.97$,
 175 $SD = 0.26$, 95% CI [1.89, 2.04]), as compared to high memory load ($M = 1.18$, $SD = 0.35$, 95% CI
 176 [1.11, 1.26]; $F(1,20) = 114.14$, $p < .001$). The influence of eccentricity was comparatively small but
 177 significant ($F(2,40) = 3.88$, $p = .029$). Participants' d' was lowest for the eccentricity of 14 dva ($M =$
 178 1.47 , $SD = 0.34$, 95% CI [1.40, 1.54]) and significantly lower than for stimuli presented at 9 dva ($M =$
 179 1.63 , $SD = 0.28$, 95% CI [1.57, 1.69]) or 4 dva ($M = 1.62$, $SD = 0.31$, 95% CI [1.56, 1.69]), as
 180 corroborated by post-hoc t -tests (Δ sensitivity₉₋₁₄ = 0.16, 95% CI [0.02, 0.29], $t(20) = 2.46$, $p = .023$;
 181 Δ sensitivity₄₋₁₄ = 0.15, 95% CI [0.01, 0.29], $t(20) = 2.25$, $p = .036$). There was no significant difference
 182 between the two smaller eccentricities (Δ sensitivity₄₋₉ = -0.01, 95% CI [-0.13, 0.12], $t(20) = -0.14$, p

183 = .893). The interaction between memory load and eccentricity did not affect d' ($F(2,40) = 1.00, p =$
184 .377).

185 Also in the perceptual control task, d' was lowest at the largest eccentricity of 14 dva (**Figure S11**).

186 We corroborated this finding using a two-way rmANOVA and observed a significant main effect of

187 eccentricity ($F(2,40) = 5.91, p = .006$). Post-hoc t -tests revealed lower sensitivity at 14 dva ($M = 1.49,$

188 $SD = 0.18, 95\% \text{ CI } [1.44, 1.54]$) as compared to 9 dva ($M = 1.63, SD = 0.16, 95\% \text{ CI } [1.58, 1.67];$

189 $\Delta\text{sensitivity}_{9-14} = 0.13, 95\% \text{ CI } [0.04, 0.23], t(20) = 2.91, p = .009$) and 4 dva ($M = 1.61, SD = 0.15, 95\%$

190 $\text{CI } [1.57, 1.66]; \Delta\text{sensitivity}_{4-14} = 0.12, 95\% \text{ CI } [0.03, 0.21], t(20) = 2.82, p = .011$). Sensitivity did not

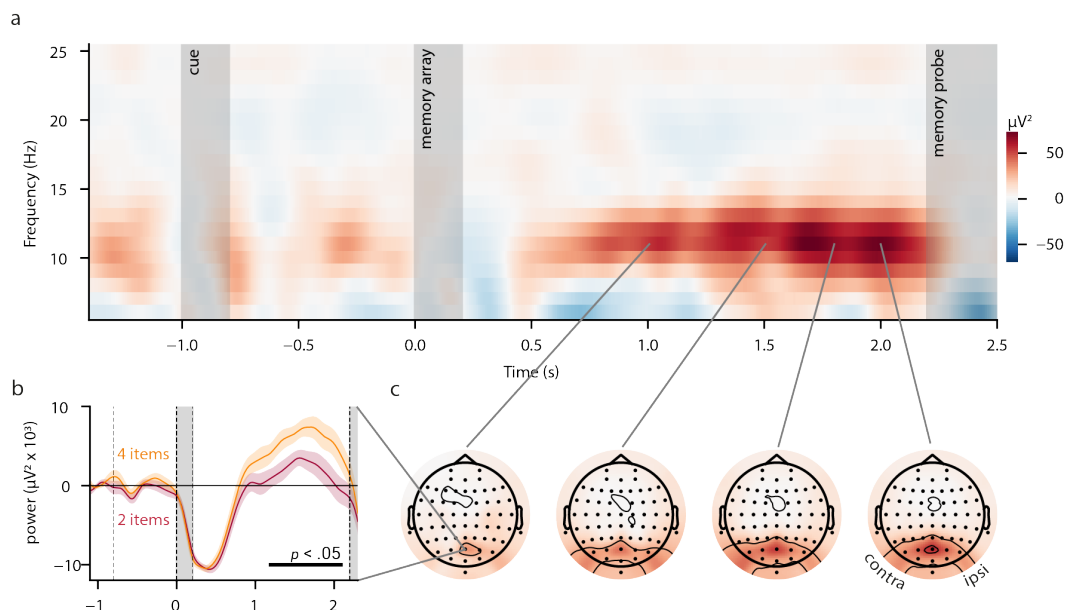
191 differ significantly between the two smaller eccentricities ($\Delta\text{sensitivity}_{4-9} = -0.01, 95\% \text{ CI } [-0.10,$

192 $0.07], t(20) = -0.32, p = .752$). Neither the number of stimuli (low memory load: $M = 1.59, SD = 0.20,$

193 $95\% \text{ CI } [1.53, 1.65];$ high memory load: $M = 1.56, SD = 0.14, 95\% \text{ CI } [1.51, 1.62]; F(1,20) = 0.29, p =$

194 .597) nor the interaction with eccentricity significantly influenced sensitivity in the perceptual task

195 ($F(2,40) = 2.73, p = .077$).



196

197 **Figure S11:** Effect of memory load on global (i.e., non-lateralized) alpha power. Some previous
 198 studies have found that non-lateralized alpha power is modulated by memory load (Pavlov &
 199 Kotchoubey, 2022). In an exploratory analysis, we checked whether this effect is also present in our
 200 data. (a) shows a time-frequency decomposition averaged across all EEG channels (normalized by
 201 subtracting the mean power during a baseline window 300–100 ms before cue onset). During the
 202 retention interval we observed an increase in power in the alpha frequency range which was most
 203 prominent in parieto-occipital sensors along the midline (c). For electrode POz (chosen by visual
 204 inspection of these topographies), we found that this effect was stronger for the high than for the
 205 low memory load condition (b). This was corroborated by a cluster-corrected t -tests comparing the
 206 alpha power values (high vs low memory load) in channel POz during the retention interval, which
 207 yielded one significant cluster (1,168–2,108 ms after onset of the memory array; black line). To test
 208 whether this effect was further modulated by stimulus eccentricity, we modeled the average signal
 209 within the significant cluster with a rmANOVA (factors: memory load, eccentricity). We did not
 210 find a significant main effect of eccentricity ($F(2,40) = 2.32, p = .112$) nor for its interaction with
 211 memory load ($F(2,40) = 2.70, p = .080$). The significant main effect of memory load ($F(1,20) = 4.51,$

212 $p = .046$) just confirmed the result of the cluster-based permutation test which guided the selection
213 of the time-window.

214 **References:**

- 215 Bates, D., Mächler, M., Bolker, B., & Walker, S. (2015). Fitting Linear Mixed-Effects Models Using lme4.
216 *Journal of Statistical Software*, *67*, 1–48. <https://doi.org/10.18637/jss.v067.i01>
- 217 Hakim, N., Adam, K. C. S., Gunseli, E., Awh, E., & Vogel, E. K. (2019). Dissecting the Neural Focus of
218 Attention Reveals Distinct Processes for Spatial Attention and Object-Based Storage in Visual
219 Working Memory. *Psychological Science*, *30*(4), 526–540.
220 <https://doi.org/10.1177/0956797619830384>
- 221 Haufe, S., Meinecke, F., Görgen, K., Dähne, S., Haynes, J.-D., Blankertz, B., & Bießmann, F. (2014). On the
222 interpretation of weight vectors of linear models in multivariate neuroimaging. *NeuroImage*, *87*,
223 96–110. <https://doi.org/10.1016/j.neuroimage.2013.10.067>
- 224 Jas, M., Engemann, D. A., Bekhti, Y., Raimondo, F., & Gramfort, A. (2017). Autoreject: Automated artifact
225 rejection for MEG and EEG data. *NeuroImage*, *159*, 417–429.
226 <https://doi.org/10.1016/j.neuroimage.2017.06.030>
- 227 Keefe, J. M., & Störmer, V. S. (2021). Lateralized alpha activity and slow potential shifts over visual cortex
228 track the time course of both endogenous and exogenous orienting of attention. *NeuroImage*, *225*,
229 117495. <https://doi.org/10.1016/j.neuroimage.2020.117495>
- 230 Livingstone, A., Christie, G., Wright, R., & McDonald, J. (2017). Signal Enhancement, Not Active
231 Suppression, Follows the Contingent Capture of Visual Attention. *Journal of Experimental*
232 *Psychology: Human Perception and Performance*, *43*, 219–224. <https://doi.org/10.1037/xhp0000339>
- 233 Pavlov, Y. G., & Kotchoubey, B. (2022). Oscillatory brain activity and maintenance of verbal and visual
234 working memory: A systematic review. *Psychophysiology*, *59*(5), e13735.
235 <https://doi.org/10.1111/psyp.13735>
- 236