- 1 The influence of image masking on object representations during rapid serial visual presentation
- 2
- 3 Amanda K. Robinson^{*a,b,c}, Tijl Grootswagers^{*a,b,c}, Thomas A. Carlson^{a,b}
- 4 *equal contribution
- 5
- ^a School of Psychology, University of Sydney, NSW, 2006, Australia
- 7 ^b ARC Centre of Excellence in Cognition and Its Disorders, NSW, 2109, Australia
- ^c Department of Cognitive Science, Macquarie University, NSW, 2109, Australia
- 9

10 Abstract

11 Rapid image presentations combined with time-resolved multivariate analysis methods of EEG or MEG 12 (rapid-MVPA) offer unique potential in assessing the temporal limitations of the human visual system. 13 Recent work has shown that multiple visual objects presented sequentially can be simultaneously 14 decoded from M/EEG recordings. Interestingly, object representations reached higher stages of 15 processing for slower image presentation rates compared to fast rates. This fast rate attenuation is probably caused by forward and backward masking from the other images in the stream. Two factors 16 17 that are likely to influence masking during rapid streams are stimulus duration and stimulus onset 18 asynchrony (SOA). Here, we disentangle these effects by studying the emerging neural representation of 19 visual objects using rapid-MVPA while independently manipulating stimulus duration and SOA. Our 20 results show that longer SOAs enhance the decodability of neural representations, regardless of 21 stimulus presentation duration, suggesting that subsequent images act as effective backward masks. In 22 contrast, image duration does not appear to have a graded influence on object representations. 23 Interestingly, however, decodability was improved when there was a gap between subsequent images, 24 indicating that an abrupt onset or offset of an image enhances its representation. Our study yields 25 insight into the dynamics of object processing in rapid streams, paving the way for future work using this 26 promising approach.

28 Introduction

29 The human brain processes rapidly changing visual input and can effortlessly extract abstract meaning 30 when stimuli are presented in rapid sequences (Mack, Gauthier, Sadr, & Palmeri, 2008; Mack & Palmeri, 31 2011; Potter, Wyble, Hagmann, & McCourt, 2014; Thorpe, Fize, & Marlot, 1996; VanRullen & Thorpe, 32 2001). Recently, the temporal dynamics of the emerging representation of visual objects have been 33 studied using fast presentation rates and multivariate analysis methods of electroencephalography 34 (EEG) and magnetoencephalography (MEG) (Grootswagers, Robinson, & Carlson, 2019; Marti & 35 Dehaene, 2017; Mohsenzadeh, Qin, Cichy, & Pantazis, 2018). Notably, multiple visual objects 36 represented in different stages of the visual system can be decoded from the EEG signal at the same 37 time (Grootswagers et al., 2019). Object representations persisted for longer when presented at slower 38 presentation rates compared to faster rates (Grootswagers et al., 2019; Mohsenzadeh et al., 2018). 39 Additionally, images presented at slower rates reached higher stages of processing, such that categorical 40 abstraction of animacy was evident for images in 5Hz but not 20Hz RSVP sequences (Grootswagers et al., 2019). The extended neural representations for slower versus faster presentation rates could be 41 42 ascribed to the longer stimulus duration, or the longer stimulus onset asynchrony (SOA) of images in 43 slower sequences.

44

45 Limitations in visual processing during rapid serial visual presentation (RSVP) is likely due to interference 46 from processing multiple images in short succession. Decades of cognitive research have documented 47 limitations in reporting targets during RSVP in phenomena such as the attentional blink (Broadbent & Broadbent, 1987; Raymond, Shapiro, & Arnell, 1992) and repetition blindness (Kanwisher, 1987). Such 48 49 effects are typically studied to investigate high-level cognitive limitations rather than low-level visual 50 processing interference (Raymond et al., 1992; Sergent, Baillet, & Dehaene, 2005). It is important to 51 note, however, that target masking has a large effect on target detection during RSVP; for example, 52 during the attentional blink, masking of the first target attenuates reporting of the second target

53 (Seiffert & Di Lollo, 1997) and masking of the second target is necessary to elicit the attentional blink 54 (Giesbrecht & Di Lollo, 1998). These findings suggest an important effect of low-level visual masking on 55 higher-level processing during RSVP. In fast sequences, images are likely subject to forward masking by 56 the previous image and backward masking by the next image in the sequence. Changing the image 57 presentation rates has the effect of altering the timing of the masks. Backward and forward masking 58 seem to have dissociable effects on perception, with one study showing maximal forward masking for 59 Oms gap between stimuli, and maximal backward masking at 30-90ms gap (Bachmann & Allik, 1976). 60 EEG has shown that backward pattern masking influences processing after approximately 180ms, 61 consistent with recurrent processing rather than feedforward processing deficits (Fahrenfort, Scholte, & 62 Lamme, 2007). Understanding how masking affects the temporal dynamics of image processing during 63 rapid-MVPA can yield important insights about the temporal limitations of the human visual system.

64

65 Studies of periodic visual evoked potentials also provide insights into the effect of image presentation 66 rate on the extent of visual object processing. Faces presented at slower frequencies reach further 67 stages of processing than those at faster rates, such that 15Hz presentations seemed limited to early 68 visual processes, 6Hz showed increased occipitotemporal responses, and 3.75Hz included higher level 69 cognitive effects and frontal responses (Collins, Robinson, & Behrmann, 2018). Retter et al., (2018) 70 showed that SOA and image duration had dissociable effects on the periodic response. Images at 10Hz 71 had larger evoked responses than those at 20Hz, but a 50% on-off image duty cycle (50ms duration, 72 100ms SOA) resulted in larger responses than 100% duty cycle with same SOA (100ms duration, 100ms 73 SOA), a finding attributed to forward masking in the 100% duty cycle condition (Retter, Jiang, Webster, 74 & Rossion, 2018). Taken together, it seems likely that SOA and image duration have separable influences 75 on visual responses, but how these differentially influence the temporal dynamics of individual image 76 processing remains to be seen.

Here, we investigate the effect of image masking on the temporal dynamics of image processing by 78 79 studying the emerging neural representation of visual objects in fast visual streams while separately 80 manipulating stimulus duration and SOA. These factors could be predicted to influence the temporal 81 dynamics of individual image processing in a linear or non-linear fashion. Varying SOA, and thus the 82 amount of time an image can be processed before another image (acting as a mask) appears, could 83 linearly influence the duration of image processing if the length of processing is directly related to the 84 amount of time dedicated to processing the uninterrupted images. Alternatively, there might be a limit 85 on the number of items that can be held in the visual system at once. If SOA influences the dynamics of 86 image processing depending on the stage of processing that is influenced by forward and backward 87 masking, this would predict a non-linear increase in image processing. Our results show that a longer 88 SOA enhances the decodability of the neural representations in a non-linear fashion, regardless of 89 stimulus presentation duration. Our results also suggest that presenting stimuli with no gap between 90 subsequent images (100% duty cycle) delays the processing of each image.

91

92 Methods

93 Stimuli, data, and code are available at: https://doi.org/10.17605/OSF.IO/3RMJ9.

94

95 Stimuli

96 We collected a stimulus set of 24 visual objects spanning 6 categories (Figure 1A). Stimuli were obtained 97 from the free image hosting website www.pngimg.com. The top-level categories were animals and 98 vehicles subdivided into 3 subcategories: birds, dogs, fish, boats, cars, and planes. Each of the 99 subcategories consisted of 4 images each. These images allowed us to investigate visual representations 100 for three different categorical levels: the animal/vehicle distinction (2 categories), object category (6 101 categories, e.g., boats, birds) and image-level (24 images, e.g., yacht, duck). Images were presented using Psychtoolbox (Brainard, 1997; Kleiner et al., 2007; Pelli, 1997) in Matlab. Images were each shown

103 foveally within a square at approximately 3 x 3 degrees of visual angle.

104

105 **Participants and experimental procedure**

106 Participants were 20 adults recruited from the University of Sydney (12 female, 8 male; mean age: 107 25.75, age range 18-52 years) in return for payment or course credit. The study was approved by the 108 University of Sydney ethics committee and informed consent was obtained from all participants. 109 Participants viewed 200 sequences of objects. Each sequence consisted of the 24 stimuli in a random 110 order. To ensure all images were equally masked by other images, the sequences were padded with 12 111 stimuli on both ends, which were excluded from the decoding analysis. The 12 padding stimuli consisted 112 of the same sequence in reverse order, with mirrored versions of the images. Essentially, this meant 113 that each of the 24 experimental images was presented twice per sequence. To keep participants engaged, at the end of each sequence, after a 1000ms blank screen, a random image from the stimulus 114 115 set was presented for 100ms and participants categorised this stimulus as animal or vehicle using a left 116 or right button press (response mappings were alternated between participants). The presentation rates 117 of the sequences were chosen from one of five conditions, which were randomized throughout the 118 study (40 sequences per condition). In conditions 1-3, the presentation duration varied (200ms, 100ms, 119 and 50ms) while keeping the SOA at 200ms. In conditions 3-5, the SOA varied (200ms, 100ms, and 120 50ms) while keeping the presentation duration at 50ms (Figure 1B). This set-up allowed us to use 121 condition 3 as anchor point to compare the effects between varying SOA and duration. In total, 122 participants viewed 9600 presentations, consisting of 80 presentations for each of the 24 images and for 123 the 5 duration/SOA conditions.

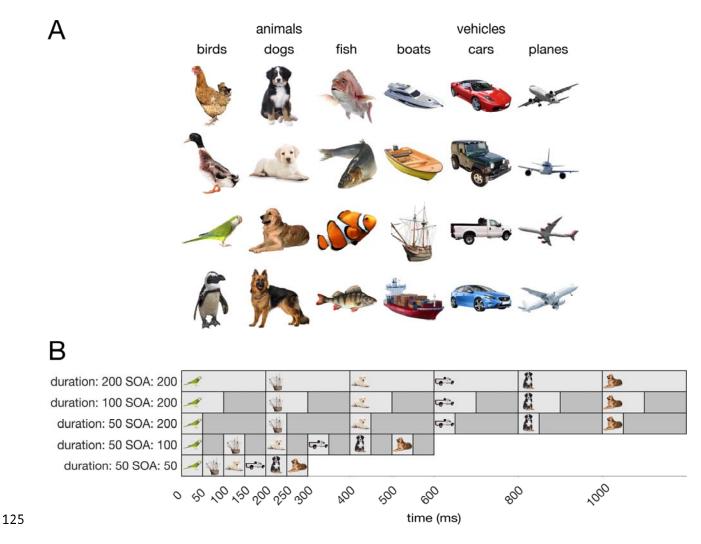


Figure 1. Stimuli and design. A) Experimental stimuli consisted of 24 images of objects organised at three different levels: animal versus vehicle, object category (6 categories e.g., birds, boats) and image (e.g., duck, chicken). B) Example time-lines illustrating the timing of the first six stimuli in a sequence in the different conditions. Images were presented in sequences with image durations of 200ms, 100ms and 50ms, and SOA of 200ms, 100ms and 50ms (5 conditions).

131

132 **EEG recordings and preprocessing**

EEG data were continuously recorded from 64 electrodes (arranged in the international 10–10 system (Oostenveld & Praamstra, 2001)) using a BrainVision ActiChamp system, digitized at a 1000-Hz sample rate. Scalp electrodes were referenced to Cz during recording. EEGlab (Delorme & Makeig, 2004) was used to pre-process the data offline, where data were filtered using a Hamming windowed sinc FIR filter with highpass of 0.1Hz and lowpass of 100Hz. Data were then downsampled to 250Hz and epochs were created for each stimulus presentation ranging from [-100 to 1000ms] relative to stimulus onset. No
further preprocessing steps were applied.

140

141 **Decoding analysis**

142 An MVPA time-series decoding pipeline (Grootswagers, Wardle, & Carlson, 2017; Oosterhof, Connolly, & 143 Haxby, 2016) was applied to each stimulus presentation epoch in the sequence to investigate object 144 representations in fast sequences. Linear discriminant analysis classifiers were trained using an image by 145 sequence cross-validation procedure (Grootswagers et al., 2019) to distinguish between all pairwise 146 groupings within the categorical levels (category, object). This entailed holding out one image from each 147 category in one sequence as test data and training the classifier on the remaining images from the 148 remaining sequences. For pairwise decoding of the non-categorical image-level, we used a leave-one-149 sequence-out cross-validation procedure. The decoding analyses were performed separately for the five 150 duration/SOA conditions. For each condition, this resulted in three decoding accuracies over time (for 151 animacy, category, and image). At each time point, these accuracies were compared against chance 152 (50%), and compared to each other. All steps in the decoding analysis were implemented in 153 CoSMoMVPA (Oosterhof et al., 2016).

154

155 **Statistical inference**

We used Bayes factors (Dienes, 2011; Jeffreys, 1998; Kass & Raftery, 1995; Rouder, Speckman, Sun, Morey, & Iverson, 2009; Wagenmakers, 2007) to determine the evidence for the null and alternative hypotheses. For the alternative hypothesis of above-chance decoding, a uniform prior was used ranging from the maximum value observed during the baseline (before stimulus onset) up to 1 (i.e., 100% decoding). For testing the hypothesis of a difference between decoding accuracies, a uniform prior was set ranging from the maximum absolute difference between decoding accuracies observed during the baseline up to 0.5 (50%). We then calculated the Bayes factor (BF), which is the probability of the data

under the alternative hypothesis relative to the null hypothesis. We thresholded BF>6 as strong
evidence for the alternative hypothesis, and BF<1/6 as strong evidence in favour of the null hypothesis
(Jeffreys, 1998; Kass & Raftery, 1995; Wetzels et al., 2011). BF that lie between those values indicate
insufficient evidence for either hypothesis.

167

To determine onset, offset, and peak time signatures, we defined onset as the second time point where the Bayes factor exceeded 6 and offset as the second-to-last time point where the Bayes factor exceeded 6. Peak decoding time was defined as the latency at which the maximum decoding accuracy was observed in the entire time window. We calculated bootstrap distributions of these latency measures by sampling from the participants with replacement 1000 times and recomputing the abovementioned statistics.

174

175 Results

We examined the temporal dynamics of object processing using rapid-MVPA with sequences of varying image duration and SOA. During the experiment, participants reported whether an image presented after each sequence was an animal or a vehicle. Behavioural performance was high for discrimination of animal (M = 97.10%, SD = 5.23%) and vehicle (M = 98.10%, SD = 3.01%) stimuli.

180

To investigate the temporal dynamics of object processing, we decoded the objects at three levels of categorical abstraction: animacy-level (animate versus inanimate), category-level (birds, dogs, fish, boats, planes, cars), and image-level (24 images; 4 per category). The decoding analyses were performed separately for each SOA and image duration condition. Figure 2 shows the temporal dynamics of all categorical representations varied by SOA and image duration. For the effect of duration (left columns), all durations followed a similar decoding trajectory, but classification was poorer in general for the longest duration, which also happened to be the 100% duty-cycle condition (200ms SOA, 200ms duration). For animacy and category decoding, the first peak (~100ms) was similar across the conditions, but the 200ms duration was lower than both the 50ms and 100ms conditions from 150-200ms, suggesting poorer categorical abstraction for this condition. Additionally, for the individual image decoding analysis, the onset of decoding appeared to be delayed for the 200ms duration relative to the 50ms and 100ms durations.

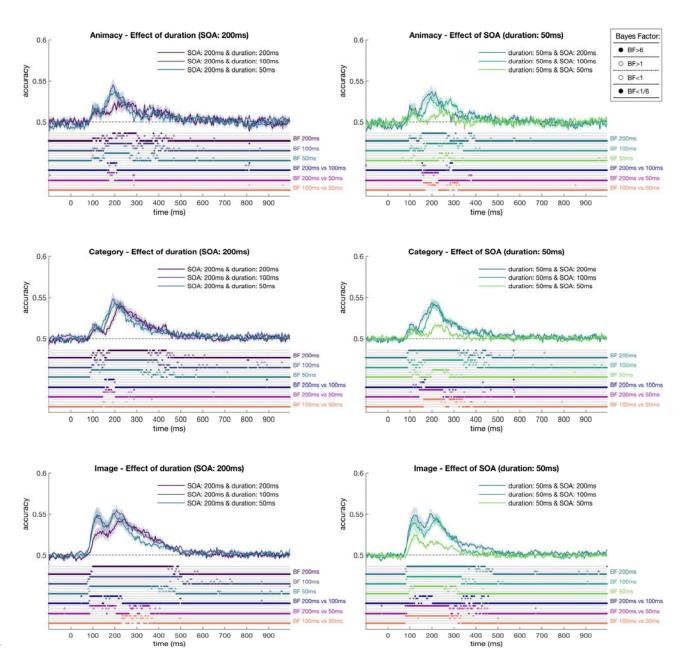
193

194 The right columns of Figure 2 show that for a given image duration (50ms), increasing SOA led to greater 195 neural decoding for all categorical levels. For animacy and category decoding, the initial peak (~120ms) 196 did not differ by SOA, but the larger second peak (~200ms) showed graded responses depending on 197 SOA. At this peak, there was a small but reliable increase in decoding for the 200ms SOA relative to the 198 100ms, and these were both substantially higher than the 50ms SOA. Again, for the image decoding the 199 100% duty-cycle condition (50ms SOA/50ms duration) appeared delayed and had poorer decoding 200 relative to the other conditions. Furthermore, the 200ms SOA had greater decoding than the 100ms 201 SOA condition between 100 and 200ms. Overall, these results imply that longer SOA led to stronger 202 image representations.

203

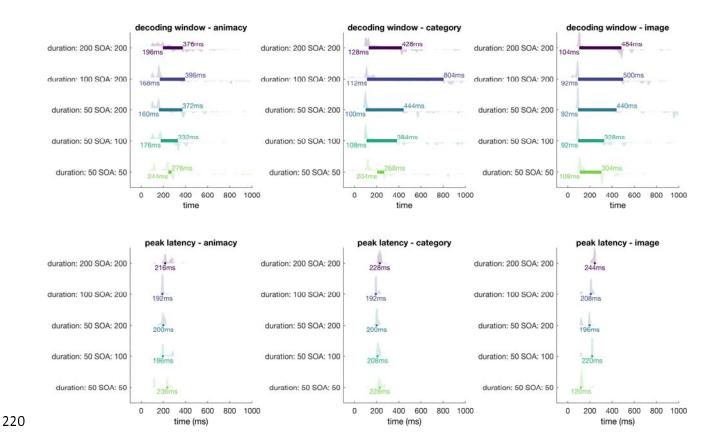
204 To further assess the effect of image duration and SOA on object decoding, we analysed the timing of 205 the decoding window (onset to offset of above-chance decoding) and the latency of peak decoding. 206 Figure 3 shows that the medium duration condition (duration 100/SOA 200) had the longest decoding 207 window for all decoding contrasts. In contrast, the shortest duration and SOA condition (duration 208 50/SOA 50) had delayed onsets and the shortest decoding window. The peak latency results revealed 209 that for animacy and category, the 100% duty-cycle conditions (200/200 and 50/50) had the latest 210 peaks. For the image decoding, however, the 200/200 condition had the latest peak, whereas the 50/50 211 condition had a much earlier peak, suggesting that ongoing processing in the latter condition was 212 limited.

213



214

Figure 2. The effects of duration (left column) and stimulus onset asynchrony (right column) on decoding accuracy at three categorical levels (animacy, category, and image). Dots above the x-axis show the thresholded Bayes factors (see inset). The top three rows show the Bayes factors for above-chance decoding, and the bottom three show Bayes factors for differences between decoding accuracies.



221 Figure 3. Onset, offset, and peak latencies for each condition. Onset was defined as the second time 222 point where the Bayes factor exceeded 6 and offset as the second-to-last time point where the Bayes 223 factor exceeded 6. Top row: For each condition (y-axis), onset and offset are marked by a filled 224 horizontal bar and are annotated at their respective time points. Shaded areas show the onset (above 225 the filled bar) and offset (below the filled bar) latency distributions calculated by bootstrapping 226 participants with replacement 1000 times and recomputing the statistics. Bottom row: Peak latency (time point of peak decoding) for each condition. Shaded areas show the peak latency distribution 227 228 calculated by bootstrapping participants with replacement 1000 times.

229

230 Discussion

231 In this study, we disentangled the effects of duration and stimulus onset asynchrony (SOA) on decoding

232 performance in rapid visual processing streams. Our results showed that shorter SOAs systematically

- reduced the duration of above-chance decoding, as well as the peak decoding accuracy, consistent with
- 234 masking at earlier stages of visual processing. In comparison, there were no graded effects of
- 235 presentation duration on decoding accuracies. Our results also suggest that presenting stimuli without a
- 236 gap (100% duty cycle) leads to delays in visual processing.
- 237

238 Previous work found that fast presentation rates limits visual processing relative to slower presentation 239 rates (Grootswagers et al., 2019). It was however unclear whether this difference was due to shorter 240 stimulus duration or shorter SOA. The results of our study show that stimulus duration and SOA have 241 separable effects on stimulus processing, with the most pronounced effect being that longer SOAs 242 enhance decodability of stimuli relative to shorter SOAs. There also appeared to be an effect of duty 243 cycle, such that neural responses were delayed when images were presented back-to-back (100% duty 244 cycle). These findings are consistent with recent work that investigated the effect of duration and SOA 245 on face response amplitudes in a fast periodic visual stimulation paradigm (Retter et al., 2018). Single 246 unit recordings in temporal cortex of macaques have revealed a similar effect; neural responses to 247 monkey faces in RSVP are stronger and last longer for slower presentation rates (Keysers, Xiao, Földiák, & Perrett, 2001, 2005). Interestingly, Keysers et al. (2001) found that the duration of image 248 249 discrimination coincided with the SOA length plus 60 ms, an effect attributed to neural competition with 250 other images in the sequence (Keysers & Perrett, 2002). Although we found that longer SOA led to 251 longer neural decoding, there was no clear linear relationship between the SOA and length of decoding. 252 Our decoding results utilise whole brain responses, however, which might be one reason for this difference. The current results suggest that SOA influences the degree of masking from subsequent 253 254 images depending on the stage of processing that is disrupted.

255

Our findings suggest that analysing the neural signatures of distractors in RSVP streams can yield insight into the mechanisms underlying visual masking. Without the need of a separate noise mask, a substantial number of presentations or conditions can be tested using rapid-MVPA, increasing the power of such experiments. By shortening the SOA, the masking affected earlier stages of processing, which has significant potential for studying hierarchical processing systems, such as vision (see also McKeeff, Remus, & Tong, 2007). For example, future work could apply rapid-MVPA and varying SOAs to

stimulus sets that vary on orthogonal features that are expected to occur at different stages in the processing streams, such as colour and shape.

264

265 Taken together, our results can be used to guide future visual object representation studies that employ 266 a rapid-MVPA design. Using a 5Hz rate, Grootswagers et al. (2019) obtained 40 epochs for 200 stimuli 267 (8000 epochs in total) in a 40-minute session. Here, we did not observe strong differences between 5Hz 268 and 10Hz presentation rates (200ms and 100ms SOA). Thus, a 10Hz 50% duty cycle presentation 269 paradigm seems to provide a sensitive measure of object decoding accuracy. Notably, this is also a 270 typical frequency used in RSVP paradigms to study target selection processes, which are postulated to 271 involve alpha oscillatory activity (Janson, De Vos, Thorne, & Kranczioch, 2014; Zauner et al., 2012). At 272 10Hz, a 30-minute EEG recording session (excluding breaks) yields 18000 epochs, which has 273 unprecedented potential for studying a large number of different conditions and/or stimuli. It also 274 suggests that it is possible to obtain enough epochs for a small number of conditions in a very short (<5-275 minute) EEG session. This opens up exciting new possibilities to study special populations for whom long 276 experiments often pose significant difficulties, such as children and patients.

277

278 Acknowledgements

The authors thank Ayeesha Dadamia for assisting with data collection, and Denise Moerel for helpful comments on a draft of the manuscript. This work was supported by an Australian Research Council Future Fellowship (FT120100816) and an Australian Research Council Discovery project (DP160101300) awarded to T.A.C. The authors acknowledge the University of Sydney HPC service for providing High Performance Computing resources. The authors declare no competing financial interests.

284

285 References

- 287 Bachmann, T., & Allik, J. (1976). Integration and Interruption in the Masking of Form by Form.
- 288 *Perception*, 5(1), 79–97. https://doi.org/10.1068/p050079
- Brainard, D. H. (1997). The psychophysics toolbox. *Spatial Vision*, *10*, 433–436.
- 290 Broadbent, D. E., & Broadbent, M. H. (1987). From detection to identification: Response to multiple
- targets in rapid serial visual presentation. *Perception & Psychophysics*, 42(2), 105–113.
- 292 Collins, E., Robinson, A. K., & Behrmann, M. (2018). Distinct neural processes for the perception of
- familiar versus unfamiliar faces along the visual hierarchy revealed by EEG. *NeuroImage*, 181,
- 294 120–131. https://doi.org/10.1016/j.neuroimage.2018.06.080
- 295 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),
- 297 9–21. https://doi.org/10.1016/j.jneumeth.2003.10.009
- 298 Dienes, Z. (2011). Bayesian Versus Orthodox Statistics: Which Side Are You On? Perspectives on
- 299 *Psychological Science*, *6*(3), 274–290. https://doi.org/10.1177/1745691611406920
- 300 Fahrenfort, J. J., Scholte, H. S., & Lamme, V. A. (2007). Masking disrupts reentrant processing in human

301 visual cortex. *Journal of Cognitive Neuroscience*, *19*(9), 1488–1497.

- 302 Giesbrecht, B., & Di Lollo, V. (1998). Beyond the attentional blink: visual masking by object substitution.
- 303 Journal of Experimental Psychology: Human Perception and Performance, 24(5), 1454.
- Grootswagers, T., Robinson, A. K., & Carlson, T. A. (2019). The representational dynamics of visual

305 objects in rapid serial visual processing streams. *NeuroImage*, *188*, 668–679.

- 306 https://doi.org/10.1016/j.neuroimage.2018.12.046
- 307 Grootswagers, T., Wardle, S. G., & Carlson, T. A. (2017). Decoding Dynamic Brain Patterns from Evoked
- 308 Responses: A Tutorial on Multivariate Pattern Analysis Applied to Time Series Neuroimaging
- 309 Data. Journal of Cognitive Neuroscience, 29(4), 677–697. https://doi.org/10.1162/jocn_a_01068

- Janson, J., De Vos, M., Thorne, J. D., & Kranczioch, C. (2014). Endogenous and Rapid Serial Visual
- 311 Presentation-induced Alpha Band Oscillations in the Attentional Blink. *Journal of Cognitive*
- 312 *Neuroscience*, *26*(7), 1454–1468. https://doi.org/10.1162/jocn_a_00551
- 313 Jeffreys, H. (1961). *Theory of probability*. Oxford University Press.
- 314 Kanwisher, N. G. (1987). Repetition blindness: Type recognition without token individuation. *Cognition*,
- 315 *27*(2), 117–143.
- Kass, R. E., & Raftery, A. E. (1995). Bayes factors. *Journal of the American Statistical Association*, *90*(430),
 773–795.
- Keysers, C., & Perrett, D. I. (2002). Visual masking and RSVP reveal neural competition. *Trends in*
- 319 *Cognitive Sciences*, *6*(3), 120–125. https://doi.org/10.1016/S1364-6613(00)01852-0
- 320 Keysers, C., Xiao, D.-K., Földiák, P., & Perrett, D. I. (2001). The Speed of Sight. Journal of Cognitive

321 *Neuroscience*, *13*(1), 90–101. https://doi.org/10.1162/089892901564199

- 322 Keysers, C., Xiao, D.-K., Földiák, P., & Perrett, D. I. (2005). Out of sight but not out of mind: the
- 323 neurophysiology of iconic memory in the superior temporal sulcus. *Cognitive Neuropsychology*,

324 22(3–4), 316–332. https://doi.org/10.1080/02643290442000103

- Kleiner, M., Brainard, D., Pelli, D., Ingling, A., Murray, R., Broussard, C., & others. (2007). What's new in
 Psychtoolbox-3. *Perception*, 36(14), 1.
- 327 Mack, M. L., Gauthier, I., Sadr, J., & Palmeri, T. J. (2008). Object detection and basic-level categorization:
- 328 Sometimes you know it is there before you know what it is. *Psychonomic Bulletin & Review*,
- 329 *15*(1), 28–35. https://doi.org/10.3758/PBR.15.1.28
- 330 Mack, M. L., & Palmeri, T. J. (2011). The Timing of Visual Object Categorization. Frontiers in Psychology,

331 2. https://doi.org/10.3389/fpsyg.2011.00165

- 332 Marti, S., & Dehaene, S. (2017). Discrete and continuous mechanisms of temporal selection in rapid
- 333 visual streams. *Nature Communications*, 8(1), 1955. https://doi.org/10.1038/s41467-017-02079-

334

Х

- 335 McKeeff, T. J., Remus, D. A., & Tong, F. (2007). Temporal Limitations in Object Processing Across the
- Human Ventral Visual Pathway. *Journal of Neurophysiology*, *98*(1), 382–393.
- 337 https://doi.org/10.1152/jn.00568.2006
- 338 Mohsenzadeh, Y., Qin, S., Cichy, R. M., & Pantazis, D. (2018). Ultra-Rapid serial visual presentation
- reveals dynamics of feedforward and feedback processes in the ventral visual pathway. *ELife*, 7,
- 340 e36329. https://doi.org/10.7554/eLife.36329
- 341 Oostenveld, R., & Praamstra, P. (2001). The five percent electrode system for high-resolution EEG and
- 342 ERP measurements. Clinical Neurophysiology, 112(4), 713-719. https://doi.org/10.1016/S1388-
- 343 2457(00)00527-7
- 344 Oosterhof, N. N., Connolly, A. C., & Haxby, J. V. (2016). CoSMoMVPA: Multi-Modal Multivariate Pattern
- 345 Analysis of Neuroimaging Data in Matlab/GNU Octave. *Frontiers in Neuroinformatics, 10.*
- 346 https://doi.org/10.3389/fninf.2016.00027
- Pelli, D. G. (1997). The VideoToolbox software for visual psychophysics: Transforming numbers into
 movies. *Spatial Vision*, *10*(4), 437–442.
- Potter, M. C., Wyble, B., Hagmann, C. E., & McCourt, E. S. (2014). Detecting meaning in RSVP at 13 ms
 per picture. *Attention, Perception, & Psychophysics, 76*(2), 270–279.
- Raymond, J. E., Shapiro, K. L., & Arnell, K. M. (1992). Temporary suppression of visual processing in an
- RSVP task: An attentional blink? *Journal of Experimental Psychology: Human Perception and Performance, 18*(3), 849.
- Retter, T. L., Jiang, F., Webster, M. A., & Rossion, B. (2018). Dissociable effects of inter-stimulus interval
- and presentation duration on rapid face categorization. *Vision Research*, 145, 11–20.
- 356 https://doi.org/10.1016/j.visres.2018.02.009
- 357 Rouder, J. N., Speckman, P. L., Sun, D., Morey, R. D., & Iverson, G. (2009). Bayesian t tests for accepting
- and rejecting the null hypothesis. *Psychonomic Bulletin & Review*, *16*(2), 225–237.

- 359 Seiffert, A. E., & Di Lollo, V. (1997). Low-level masking in the attentional blink. Journal of Experimental
- 360 Psychology: Human Perception and Performance, 23(4), 1061.
- 361 Sergent, C., Baillet, S., & Dehaene, S. (2005). Timing of the brain events underlying access to
- 362 consciousness during the attentional blink. *Nature Neuroscience*, *8*(10), 1391.
- 363 Thorpe, S., Fize, D., & Marlot, C. (1996). Speed of processing in the human visual system. *Nature*,
- 364 *381*(6582), 520–522. https://doi.org/doi:10.1038/381520a0
- 365 VanRullen, R., & Thorpe, S. J. (2001). The Time Course of Visual Processing: From Early Perception to
- 366 Decision-Making. *Journal of Cognitive Neuroscience*, *13*(4), 454–461.
- 367 https://doi.org/10.1162/08989290152001880
- 368 Wagenmakers, E.-J. (2007). A practical solution to the pervasive problems of p values. *Psychonomic*
- 369 Bulletin & Review, 14(5), 779–804. https://doi.org/10.3758/BF03194105
- Wetzels, R., Matzke, D., Lee, M. D., Rouder, J. N., Iverson, G. J., & Wagenmakers, E.-J. (2011). Statistical
- 371 Evidence in Experimental Psychology: An Empirical Comparison Using 855 t Tests. *Perspectives*
- 372 *on Psychological Science*, *6*(3), 291–298. https://doi.org/10.1177/1745691611406923
- Zauner, A., Fellinger, R., Gross, J., Hanslmayr, S., Shapiro, K., Gruber, W., ... Klimesch, W. (2012). Alpha
- entrainment is responsible for the attentional blink phenomenon. *NeuroImage*, 63(2), 674–686.
- 375 https://doi.org/10.1016/j.neuroimage.2012.06.075