1	An adaptive perspective on visual working memory distortions
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44 Abstract

45 When holding multiple items in visual working memory, representations of individual items are 46 often attracted to, or repelled from, each other. While empirically well-established, existing 47 frameworks do not account for both types of distortions, which appear to be in opposition. Here, 48 we demonstrate that both types of memory distortion may confer functional benefits under 49 different circumstances. When there are many items to remember and subjects are near their 50 capacity to accurately remember each item individually, memories for each item become more 51 similar (attraction). However, when remembering smaller sets of highly similar but discernible 52 items, memory for each item becomes more distinct (repulsion), possibly to support better 53 discrimination. Importantly, this repulsion grows stronger with longer delays, suggesting that it 54 dynamically evolves in memory and is not just a differentiation process that occurs during 55 encoding. Furthermore, both attraction and repulsion occur even in tasks designed to mitigate 56 response bias concerns, suggesting they are genuine changes in memory representations. Together, 57 these results are in line with the theory that attraction biases act to stabilize memory signals by 58 capitalizing on information about an entire group of items, whereas repulsion biases reflect a 59 tradeoff between maintaining accurate but distinct representations. Both biases suggest that human 60 memory systems may sacrifice veridical representations in favor of representations that better 61 support specific behavioral goals.

62 Memory is a constructive rather than a passive process. For example, people will naturally fill in 63 gaps when recalling a story in an attempt to make the story more coherent (Bartlett, 1932; Loftus, 2005; Roediger & McDermott, 1995). When people study a list of words, they often falsely recall 64 65 or recognize associated words that were not on the original list (Deese, 1959; Underwood et al., 66 1965), and later report these words as actual memories (Schooler et al., 1988). Similarly, visual 67 memory is not analogous to taking a photo - instead, there are many systematic biases in how 68 visual attributes are remembered after a sensory stimulus is no longer available (Alvarez, 2011; 69 Bar, 2004; Brady & Alvarez, 2011; Fischer & Whitney, 2014; Huang & Sekuler, 2010; Koutstaal 70 et al., 2001; Rademaker et al., 2015; Schacter et al., 2011).

71 When people are tasked with remembering a visual item, such memories are often distorted toward 72 existing, learned prototypes (Huttenlocher et al., 1991, 2000; Hemmer & Steyvers, 2009). Such 73 distortion can also occur not toward pre-learned prototypes, but toward the central tendency of a 74 group within a single presentation. For example, when people are asked to remember multiple 75 visual items, these memories are 'attracted' to each other - that is, different objects are remembered 76 as more similar than they really were (Brady & Alvarez, 2011; Dubé et al., 2014; Dubé & Sekuler, 77 2015; Freyd & Johnson, 1987; Huang & Sekuler, 2010; Spencer & Hund, 2002). It has been 78 proposed that this occurs because object-level representations are imprecise, so these unstable 79 representations are constrained by using additional information about the properties of the set of 80 items as a whole (i.e. group-level representation). Thus, inter-item attraction biases may be the 81 result of weighting the representation of each individual object towards the "summary" of the set 82 to achieve a more stable memory at the expense of maintaining distinctions between individual 83 items (Brady & Alvarez, 2011; Huttenlocher et al., 1991).

84 Interestingly, attraction biases are not ubiquitous. Under some conditions, when multiple items are 85 shown at once, memories for individual specific items have been shown to repel each other, being remembered as more different than they really were (Bae & Luck, 2017; Golomb, 2015; O'Toole 86 87 & Wenderoth, 1977; Rademaker et al., 2015; Rauber & Treue, 1998; Suzuki & Cavanagh, 1997). 88 However, far less research has been dedicated to understanding inter-item repulsion biases. 89 Repulsion biases have sometimes been proposed to arise from lateral inhibition, as competition 90 between neurons representing similar feature values may lead to representations that repel away 91 from each other (Johnson et al., 2009; Wei et al., 2012), akin to repulsion resulting from 92 competition during early perceptual processing (Jazayeri & Movshon, 2006; Navalpakkam & Itti, 93 2007; Purushothaman & Bradley, 2005; Regan & Beverley, 1985; Scolari & Serences, 2009; 94 Scolari & Serences, 2010; Smith et al., 2005). However, while providing a possible mechanistic 95 basis, such theories do not straightforwardly explain why repulsion biases sometimes arise and 96 sometimes do not; nor why attraction biases occur for similar stimuli under other circumstances. 97 Despite the importance and pervasiveness of these memory distortions, to date there have been 98 few attempts to understand why memories sometimes attract, while at other times they repel.

99 Because these are rarely studied together, it is still unclear whether these inter-item memory 100 distortions that arise for simultaneously presented items are due to changes in the representations 101 themselves, or if they instead reflect demand characteristics that lead to systematic response biases. 102 For example, repulsion biases can emerge in continuous report paradigms if participants want to 103 actively communicate that they know two items are different, even if participants have access to 104 veridical representations, and most work to date has demonstrated repulsion biases only in such 105 continuous report situations (Bae & Luck, 2017; Golomb, 2015; Rademaker et al., 2015).

106 To establish when attraction biases and repulsion biases arise and whether they are properties of 107 the memory system or a result of stimulus differences or straightforward responses biases that 108 occur only in continuous report tasks, we present a series of experiments. First, we determine 109 whether attraction and repulsion are simply properties of subject's communicative intent in 110 continuous report tasks. Second, we examine whether they arise in predictable circumstances, by 111 manipulating task difficulty and the similarity and distinctiveness of the memoranda. While these 112 are general issues, related to nearly all kinds of memory, we tested these ideas in a well-studied 113 domain – visual working memory for color – where memory representations can be precisely 114 quantified. Task difficulty was increased or decreased by changing how many items must be 115 remembered (set size), how distinctive the colors are from each other (their proximity in color space), and encoding time and memory delay. 116

117 After establishing the empirical phenomena, we adopt the perspective (in the 'Framework' section 118 of the paper) that these inter-item biases for simultaneously presented items may be natural 119 consequences of the memory system attempting to minimize memory error, and that systematic 120 distortion can be adaptive in particular circumstances (Schacter et al., 2011). Specifically, when 121 many items are present and memories for individual items are noisy, attraction biases are known 122 to be optimal for minimizing error (e.g., Brady & Alvarez, 2011). In this case, relying on group-123 level statistics provides an efficient means of retaining at least some information about all items at 124 the expense of precisely representing information about each single item. Repulsion biases can 125 also reduce error in some situations, making them adaptive. In particular, if items would naturally 126 be blended or confused by our memory system (Swan & Wyble, 2014; Oberauer & Lin, 2017) -127 that is, if similar items would interfere with each other – then repulsion can reduce this tendency 128 and reduce error. In this case, the goal is to distinguish highly similar or noisy representations, by

reducing the confusability between memory items. In particular, if items interfere to the extent they overlap in features, then repulsion is adaptive when items overlap in representation. In discussing this framework, we examine whether attraction and/or repulsion occur in the circumstances predicted by this framework, and not in circumstances where biases would be maladaptive to memory performance (i.e. contrary to the adaptive framework).

134 Overall, we find that when distinctiveness between two items goes down, repulsion biases are 135 stronger (up to the point where two items become indistinguishable, and attraction takes over as 136 the dominant force). Repulsion biases also grow stronger with longer delays, suggesting that as 137 memory demands increase and item representations become noisier, memories are biased to keep 138 items individuated. In contrast, we observe attraction biases when individuating items is more 139 difficult due to a higher memory load (in an experiment with 4 instead of 2 memory items), 140 consistent with sacrificing single-item discriminability in order to remember at least some 141 information about ensemble-level features. Importantly, by using a 2-alternative forced-choice 142 paradigm we were able to test the role of demand characteristics: the results imply that repulsion 143 biases are not the result of participants trying to communicate that they can distinguish two targets 144 in a continuous report task. Collectively, these studies suggest that, given task-imposed constraints, 145 attraction or repulsion biases may help to improve behavioral performance even though these 146 biases may lead to non-veridical memories.

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Experiment 1: Memory distortion vs. response strategy

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152 Do memory items truly 'repel' each other when people hold in mind a small number of similar 153 items? In Experiment 1 we sought to replicate this basic repulsion effect and to determine if 154 previously reported biases (e.g., Bae & Luck, 2017; Golomb, 2015) are more likely to reflect 155 memory distortions, or if they are a result of changes in response strategy to communicate an 156 understanding of the continuous reproduction task. That is, when participants remember a pair of 157 colors, they can communicate their awareness of the colors being distinct from one another by 158 exaggerating the difference between the two. When cued to report one of the two remembered 159 items on a continuous color-wheel, this strategy would result in an answer repelled away from the 160 uncued non-target item – mimicking a repulsion bias. We directly addressed this possible response 161 strategy by having participants remember two colored items over a brief delay (Figure 1a), after 162 which they perform a 2-alternative forced-choice (2-AFC) task comparing the correct (cued target) 163 color to an incorrect (distorted foil) color (Figure 1a, b). By presenting participants with the correct 164 answer on every trial, such response biases are discouraged as they are detrimental to task 165 performance, and an understanding of the task is best communicated by picking the correct color. 166 To distinguish between attraction and repulsion in this 2-AFC paradigm, the incorrect foil color 167 was distorted by 6° relative to the correct target color, and the distortion was either towards the 168 non-target (i.e. 'attracted' to the non-target) or away from the non-target (i.e. 'repelled' from the 169 non-target). If memories for the two colors were repelled from each other, a foil color that was 170 distorted toward the non-target would be less often confused with the correct answer (have a higher 171 accuracy) than a foil color that was distorted *away* from the non-target (have a lower accuracy).

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173 Methods

174 The datasets from all of the current studies (plus the code used to generate the stimuli and analyze175 the data) are available in the OSF repository

176 https://osf.io/qp6xk/?view_only=0559769c587c4c8294288451e8af239e

Participants. 45 naïve participants were recruited from Amazon Mechanical Turk. In this and all other experiments reported, all experimental procedures were approved by the UCSD Institutional Research Board, all online participants provided written informed consent, and all reported normal or corrected-to-normal vision without color-blindness. Participants were naïve to the purpose of the study and received payment (\$6 per hour) for their time.

182 Stimuli & Procedure. All stimuli were drawn on a 500×500 pixels white background with a 183 black border around it (1 pixel wide). The fixation cross was in the middle of the canvas, and 12 184 small circular placeholders were shown around fixation, each centered at a distance of 120 pixels. 185 Each placeholder had a radius of 20 pixels, and the inter-placeholder distances were 62 pixels 186 (center-to-center). Placeholders were positioned such that six of them were on the left, and the 187 other six were on the right side of fixation. Furthermore, two placeholders were always presented 188 directly to the left and right of fixation, centered at 35 pixels from fixation. Memory items were 189 colors selected from a subset of CIE L*a*b color space (L = 70, a = 20, b = 38, radius =60). Note, 190 while one of the memory items was always selected randomly from this color space, the second 191 item always differed from the first by 45°. The location probe, cueing participants which memory 192 item to report on, was a small equilateral black triangle, 20 pixels wide and 20 pixels tall.

193 Participants were shown two memory items for 150ms at two randomly selected placeholders in 194 the display (out of 12 possible placeholders), with the restriction that there were always at least 2 195 empty placeholders between the two memory items. After a 750ms delay, a location cue (arrow)

196 indicated which of the items was the memory target, and two response options appeared in the 197 placeholders directly to the left and right of fixation. One of the response options was always the 198 correct color (i.e. identical to the color that was cued), while the other option was always a foil, 199 and participants made a 2-AFC judgment between the two response options. The foil always 200 differed from the correct color by 6° in color space, either in the direction towards (50% of trials) 201 or away (50% of trials) from the non-target memory item. The positions (left or right of fixation) 202 of the correct and foil response options were completely randomized. Participants had to press "z" 203 or "m" to select the choice presented on the left or right of fixation, respectively, before proceeding 204 to the next trial. There were 60 trials per condition (a total of 120 trials per participant).

205 Results

As predicted by an account where repulsion is a genuine memory phenomenon, participants were better at rejecting a foil color that was distorted toward the non-target memory item than rejecting a foil color that was distorted away from the non-target memory item – an indicator of repulsion bias ($t_{(44)}=3.98$; p<0.001; Figure 1c). In other words, performance was higher when a foil was distorted towards the non-target memory. This shows that repulsion biases occur even in a 2-AFC format with an objectively correct answer vs. an objectively incorrect answer, implying that repulsion is not merely the result of this particular a priori plausible response strategy.

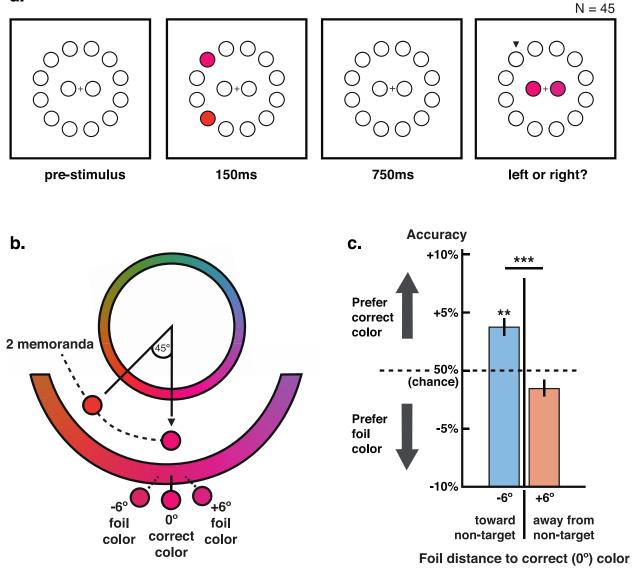


Figure 1. Task and results from Experiment 1. a.) Participants remembered two memory items that were always 45° apart in color space. Memory items were briefly presented for 150ms at two randomly chosen placeholder locations. After a 750ms delay, participants reported the color of the target item (cued with an arrow) by choosing between two options, one always being the correct color, and the other always being an incorrect foil color that was distorted from the correct color by 6° in a direction either toward or away from the non-target color. In the example trial shown here, the correct response is shown on the left, while the foil on the right is distorted in a direction

221 away from the non-target color. b.) Two memory colors were selected to lie within 45° of each 222 other in color-space (at any possible position on the color wheel). The target color (cued after the 223 delay) was always one of the response options during the 2AFC phase of the trial (i.e. the "correct 224 color"). The other response option was a foil color. The foil color always differed 6° from the 225 correct target color and could be distorted towards (-6°) or away $(+6^{\circ})$ from non-target color. c.) 226 Participants preferred the correct color to the foil when the foil was distorted toward the non-target 227 color, as indicated by above-chance performance (blue bar; $t_{(44)}=3.73$; p=0.006). This differed 228 significantly from trials on which the foil was distorted away from the non-target color (compare 229 blue and red bars; $t_{(44)}=3.98$; p<0.001), with a trend towards participants preferring the incorrect 230 foil over the correct answer, as indicated by numerically below-chance performance (red bar; 231 $t_{(44)}$ =-1.76; p=0.08). This is the expected result when memory for the target is distorted away from 232 the non-target (i.e. when there is a repulsion bias). Error-bars represent + 1 within-subject SEM. 233 Double and triple asterisks indicates p < 0.01 and p < 0.001 respectively.

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Experiment 2: Memory distortion vs. response strategy, and the role of task engagement.

237 We next replicated and extended Experiment 1 with additional foil colors that were 25° away from 238 the correct memory target. We added 25° foils in this second experiment to test the possibility that 239 participants simply favored all colors distorted away from the non-target color by way of a 240 response strategy, even though such a strategy would result in objectively incorrect performance 241 in this task. After all, if participants meant to communicate their awareness of the two memory 242 colors being distinct, they would prefer any foil away from the non-target over the correct answer. 243 In this were the case, 25° foils would be favored even more than 6° foils, because they are more 244 clearly away from non-target color. This hypothesis is schematically shown in Figure 2a (top panel; 'Prediction 1'). By contrast, if memories of the two colors were truly repelled from one another, and participants remembered the target item as further from the non-target than it actually was, performance should depend on the degree of foil distortion. Specifically, participants should be more likely to choose the foil (and give an incorrect answer) when it closely matches their distorted memory (e.g. the $+6^{\circ}$ foil), but more likely to choose the correct color when the distortion of the foil becomes irreconcilable with their memory (e.g. the $+25^{\circ}$ foil). This hypothesis is also schematically shown in Figure 2a (bottom panel; 'Prediction 2').

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253 Methods

Participants. 45 new naïve participants were recruited from Mechanical Turk for Experiment 2.
For the control experiment replicating Experiment 2 (Appendix Figure 1) we recruited another
independent set of 45 participants from Amazon Mechanical Turk.

257 Stimuli & Procedure. The stimuli and task were identical to Experiment 1, except that in 258 Experiment 2 the foil could differ from the correct color by either 6° (45% of trials), 25° (45% of 259 trials), or 180° (10% of trials). As in Experiment 1, on half of these trials the foil was in the 260 direction *toward* the non-target in color space, while on the other half of trials the foil was *away* 261 from the non-target in color space. Given how easily distinguishable the 180° foils were from the 262 correct color, these trials served as catch trials. For the control experiment replicating Experiment 263 2 (presented in Appendix Figure 1), the foil could differ from the correct color by either 6° (90%) 264 of trials), or 180° (10% of trials). In Experiment 2, there were 30 trials per main condition (total of 4 main conditions, i.e. 6° vs. 25° foils, crossed with distortion away vs. toward non-target) plus 12 265 266 catch trials (a total of 132 trials per participant). In the replication study of Experiment 2, there 267 were 60 trials per condition (6° foils, with distortion away vs. toward non-target) plus 12 catch trials (a total of 132 trials per participant).

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270 Results

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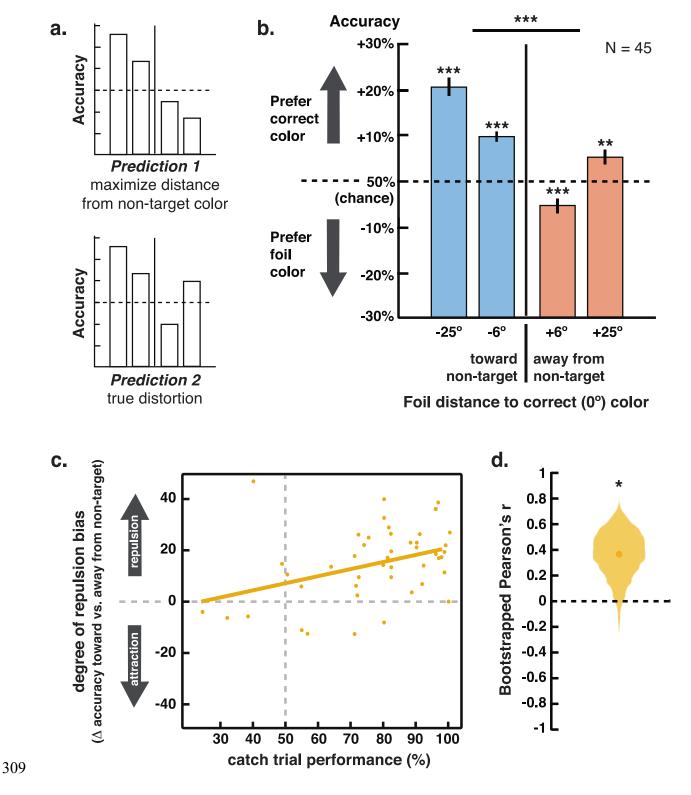
272 We replicated Experiment 1, as participants were again better at rejecting a foil color that was 273 distorted *toward* compared to *away* from the non-target memory item ($F_{(1,44)} = 49.2$; p<0.001; 274 Figure 2b, compare blue and red bars). Interestingly, subjects more often selected foils that were 275 6° away from the non-target color compared to the correct target color, resulting in below-chance 276 level performance in this condition ($t_{(44)}=3.41$; p=0.001; Figure 3b, compare +6° bar against chance 277 accuracy). This is consistent with a strong degree of memory distortion, where participants prefer 278 a repelled foil color relative to the correct answer. In contrast, subjects successfully rejected all 279 other foils resulting in above-chance level performance ($t_{(44)}=8.70, 7.70, and 3.05; p<0.001,$ 280 <0.001, and 0.004 for foils that were -25° , -6° , and $+25^{\circ}$ relative to the non-target color, 281 respectively; Figure 2b). Thus, participants showed a clear repulsion bias that cannot be easily 282 explained by response strategy. Instead, the data are consistent with a target memory that was truly 283 distorted away from the non-target item by several degrees.

284

In addition to replicating Experiment 1 and bolstering the case in favor of a true repulsion bias (and not a response strategy), we wanted to know if the degree of repulsion bias was related to the level of task engagement from our participants. To this end, Experiment 2 included foils that were 180° away from the cued memory target on 10% of the trials. We termed trials with a 180° foil "catch trials", as subjects should rarely, if ever, confuse these foils with the correct color. Thus, performance on catch trials provides a useful measure of overall task engagement and effort. Critically, if the repulsion bias is adaptive and improves memory, one would expect the degree of 292 repulsion to positively correlate with overall performance. In contrast, if biases arise due to lack 293 of effort or some other non-task related factor like response strategy, we might expect repulsion 294 bias to be negatively correlated with performance (or uncorrelated). We quantified the degree of 295 repulsion as performance on trials with foils distorted towards the non-target (both by 6° and 25°), 296 minus trials with foils distorted away from the non-target (both by 6° and 25°). This metric will be 297 larger for participants with stronger repulsion. We found a moderate positive correlation between 298 the degree of repulsion bias and overall task engagement (Pearson's r=0.37; p=0.013; Figure 2c) 299 supported by a bootstrapping analysis (bootstrapped mean Pearson's r=0.37, two-tailed p=0.048; 300 Figure 2d). This positive correlation between repulsion bias and overall task engagement was 301 replicated in an independent set of 45 naïve subjects (Pearson's r=0.39; p=0.009; Appendix Figure 302 1). Thus, repulsion biases do not appear to arise solely in participants putting in low or moderate 303 effort, instead, they are strongest in participants with the highest levels of task engagement.

304

305 Overall, Experiments 1 and 2 provide evidence for a repulsion bias that cannot be explained by
306 these straightforward, a priori reasonable communicative strategies resulting in simple response
307 biases, or a lower amount of effort.



311 Figure 2. Experiment 2 hypotheses and results. a.) The two panels show our predictions if 312 participants were trying to strategically avoid non-target colors leading to a response bias 313 (prediction 1, top) vs. if participants had memories that were truly distorted away from one another 314 (prediction 2, bottom). b.) Data from 45 subjects showed a pattern consistent with true memory 315 distortions as in prediction 2. Participants performed significantly below chance (i.e. preferred the 316 foil over the correct response option) only when the foil was distorted 6° (but not 25°) away from 317 the non-target color. This is in line with a true distortion of the remembered color and is indicative 318 of participants finding that the foil more accurately reflected their memory representation. 319 Presented with any other foil (foils distorted towards the non-target, or a foil distorted farther away 320 from the non-target), participants chose the correct answer more often than chance. Error-bars 321 represent ± 1 within-subject SEM. c.) Degree of repulsion bias (indexed as accuracy differences 322 between all trials with foils distorted toward and away from the non-target color) plotted against 323 general memory performance (indexed by performance on catch trials). Each dot represents a 324 single participant. We found stronger repulsion biases in participants with better general memory 325 performance (Pearson's r = 0.37, p=0.013). Note that the position of the dots are slightly 326 independently jittered by random noise (+/- 5%) to aid visualization of all 45 data points. The solid 327 yellow line represents the best fit to the unjittered data. d.) Distribution plot of bootstrapped 328 Pearson's r between repulsion magnitude and general memory performance (5000 iterations of 329 resampling with replacement). Single, double and triple asterisks indicate p < 0.05, p < 0.01 and p 330 < 0.001 respectively.

It is still possible that the repulsion bias is the result of a response strategy whereby the participantis trying to signal not only an understanding of the task (leading to repulsion), but also wants to

communicate which of the two items was being recalled (leading to repulsion only for the probeditem).

335

336 Such an account would naturally predict a disappearance of the repulsion bias when not one, but 337 both memory items were probed. To investigate this possibility, we reanalyze an existing open 338 data set (Adam, Vogel, & Awh 2017) where participants were required to reproduce the colors of 339 two memoranda in a random order. To quantify the repulsion bias, we took the absolute difference 340 between the two stimulus colors presented and compared this to the absolute difference between 341 the two responses participants made. In case of repulsion, response errors will be further apart in 342 color space than the actual stimuli were. Indeed, we found that differences between the response 343 errors were significantly larger than the stimulus differences ($t_{(1,16)} = 3.11$, p<0.01). This suggests 344 that also in a whole report task, items at set size 2 repel each other systematically.

345

346 Overall, while it is never possible to rule out all possible response strategies. Some aspects of these 347 effects could still be happening at response stages, even if they are not explainable by the response 348 strategies we test here and that are most plausible a priori. However, we have shown they apply 349 not only in continuous report where a single item is probed, but also in continuous report where 350 both items are probed, and in two kinds of forced-choice tasks, including one where there is a 351 single objectively correct answer and a single objectively incorrect answer. While different 352 response strategies could be at work in each task, giving rise to this pattern, this work provides 353 significant evidence in favor of a mnemonic shift account.

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Experiment 3: Attraction vs. repulsion.

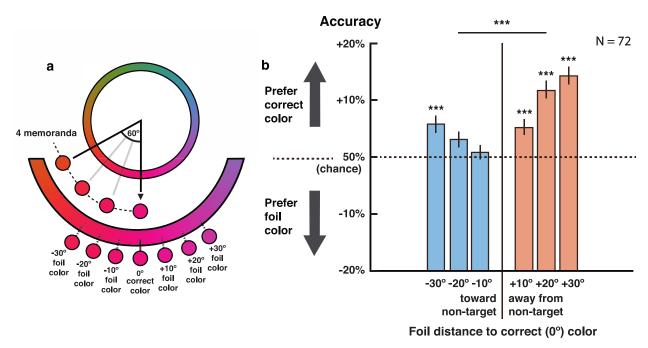
357 We next sought to manipulate task factors to test if we could systematically flip distortions from 358 repulsion to attraction, even for the same kind of stimuli. We used the same experimental paradigm 359 as in Experiments 1 and 2, but increased the number of colors participants had to remember from 360 2 to 4 items (see also Appendix Figure 2). Given well-documented limits on the amount of 361 information that can be retained in working memory (e.g., Bays, 2015; Bays et al., 2009; Luck & 362 Vogel, 1997; Ma et al., 2014), remembering 4 items should be quite challenging for the majority 363 of participants. Our adaptive framework suggests that when it is challenging to maintain 364 individuated representations of all memory items, a partial reliance on group-level statistics (Brady 365 & Alvarez, 2011) or partial blending between items (Swan & Wyble, 2014; Oberauer & Lin, 2017) 366 is optimal, because it supplements the noisy information available about each of the individual 367 items with information from the other items. In this context, when participants are presented with 368 a foil that is distorted *towards* the colors of the other items in the set (Figure 3a), they should be 369 more likely to confuse the foil with the correct (cued) target color (i.e. show an attraction bias) – 370 the exact opposite of the repulsion bias observed in the previous experiments. To test this, in this 371 experiment the four to-be-remembered colors spanned 60° of color space (in 20° steps), and we 372 always cued one of the colors on the "edge" of this set. There were 6 possible foil conditions, of 373 which 3 were distorted towards the other non-target items, and 3 were distorted away from the 374 other non-target items (Figure 3a).

375

376 Methods

377 Participants. A total of 72 naïve participants were recruited from Amazon Mechanical Turk. This
378 is more than in Experiments 1 and 2 due to the increased difficulty of the task associated with the
379 higher set size (thus requiring more power). Participants received \$8 per hour for their time.

380 Stimuli & Procedure. Stimulus and task presentation was identical to Experiments 1 and 2 with 381 the following exceptions: Participants were shown four color items for 800ms, memory item 382 locations were random (could be any 4 placeholders out of the possible 12) with the restriction that 383 there was always at least 1 empty placeholder between each of the memory items. The four items 384 were remembered over a 1000ms delay. The four colors were within 60° from each other in color 385 space, and all colors were equally spaced from one another (i.e. the shortest possible color distance 386 between two items was 20°; see also Figure 3A). The memory target probed at the end of the delay 387 was always one of the colors at the edge of the set. Again, the correct color was always included 388 as one of the response options, while the foil color differed by either 10°, 20°, or 30° from the 389 correct target color option. The foil color could be either toward the colors of the other memory 390 items (note how a -20° foil is identical to one of the other colors in the display, and a -30° foil is 391 exactly the mean of all 4 colors), or it could be *away* from the other colors. There were 20 trials 392 per main condition (total of 6 conditions, 10° vs. 20° vs. 30° foils, and distortion away vs. toward 393 non-target) which means a total of 120 trials per participant.



395 Figure 3. a.) A set of four colors were selected to lie within 60° of each other in color-space (all 396 separated by steps of 20°) and were presented at random spatial positions (chosen from 12 possible placeholders; see Appendix Figure 2). The cued memory target color (to be reported after the 397 398 delay) was always one of the colors on the edge of the set. In this diagram, the target is the 399 memorandum with the arrow pointing at it. After a 1000ms delay, participants performed a 2-AFC 400 memory test. One of the options was always the correct (cued) target color, while the other choice 401 was an incorrect foil of which the color differed by either 10°, 20° or 30° from the correct target 402 color. The foil could be distorted towards $(-10^\circ, -20^\circ, \text{ or } -30^\circ)$ or away $(+10^\circ, +20^\circ, \text{ or } +30^\circ)$ from 403 the center of the four colors in the memory set. b.) Accuracy was lower when the task was more 404 difficult: When subjects had to choose between the correct color and a foil color that was very 405 similar to the correct color (for example, differed by 10°) accuracy was closer to chance compared 406 to when subjects had to choose between the correct color and a foil that differed more from the 407 correct color (for example, differed by 30°). Importantly, performance was worse when the foil 408 color was distorted toward the other memory colors in the set (i.e. the blue bars are lower overall 409 than the red bars). This indicates an attraction of the cued item towards the other non-target items. 410 Error-bars represent + 1 within-subject SEM. Asterisks represent significance levels of differences 411 between foils that were toward vs. away from non-target, with the triple asterisks indicating 412 p<0.001.

413

414 **Results**

We performed a 3x2 repeated-measure ANOVA, and found a significant main effect of the distances of the foils from the target ($F_{(2,142)} = 13.14$; p<0.0001), and a significant main effect of the direction of the foil ($F_{(1,71)} = 15.48$; p<0.0001). There was no significant interaction ($F_{(2,142)} = 1.93$; p=0.15). Specifically, we found that participants were more accurate when the foil

419 colors were more dissimilar from the correct color, making discrimination easier: Accuracy was 420 53%, 57.4% and 60.1% correct for foils that were 10°, 20° and 30° away from the correct color, 421 respectively (Figure 3b, compare bars with smaller versus larger target-foil distances). 422 Importantly, participants were also better at choosing the correct answer when the foil color was 423 distorted away from the other non-target colors in the set (60.4% correct) compared to when the 424 foil color was distorted toward the other non-target colors (53.2% correct; Figure 3b, compare blue 425 and red bars). This implies an attraction bias towards the remembered non-target items, and stands 426 in contrast to the repulsion bias found with set size 2 (in Experiments 1 and 2). Previous work has 427 demonstrated that attraction biases in visual working memory arise from slight shifts toward the 428 gist, and not solely from swaps or guesses based on the average color (e.g., Brady & Alvarez, 429 2011). Consistent with this, we found little evidence for swaps and guesses in our data as well: In 430 particular, the -20° foil was the same color as one of the non-target items; and the -30° foil was the mean of all colors in the set. Nevertheless, neither the -20° nor the -30° foils were selected as often 431 432 as -10° foil – indicative of only a slight attraction toward the other colors.

433

Experiment 4: Biases depend on the degree of distinctiveness between items.

434 In a fourth experiment (Figure 4a) we sought to determine if reducing the distinctiveness between 435 items (by making items increasingly similar or noisy) impacts the amount of repulsion bias in a 436 manner consistent with our framework. In particular, if the memory system naturally blends 437 together similar items (as in the models of Swan & Wyble, 2014; Oberauer & Lin, 2017), then two 438 items that are recognizably distinct (i.e., can still be told apart) but still similar enough to likely be 439 blended, repulsion should arise (see 'Framework' section). To this end, we asked participants to 440 remember 2 colors, and we independently manipulated both memory encoding time (50, 150 and 500ms) and distance in feature space between the two colors (0°, 20°, 45°, 90° and 135°). If less 441

442 easily distinguishable colors need to be differentiated from one another in order to improve 443 behavioral performance, a higher degree of similarity between the 2 memory items should result 444 in a stronger repulsion bias – but critically, there should be an exception for colors that are so 445 similar that they are perceived as the same color and are thus put into a single 'chunk' or group. 446 Furthermore, the color distance that creates maximal repulsion should depend on how precise the 447 representations are: Two very precise representations at a given color distance may not require 448 repulsion to be differentiated, while two more imprecise representations at that same color distance 449 could be more easily differentiated with repulsion. In other words, when two memory 450 representations are not too similar or too distinct, the magnitude of repulsion bias will depend on 451 the precision of the memories. Repulsion bias might be necessary if the memory representations 452 are relatively less precise. Representational precision should vary with encoding time (i.e. memory 453 should be more precise at longer encoding times). Since Experiments 1 and 2 suggest that repulsion 454 biases reflect changes in encoding and memory as opposed to response strategy, here we used a 455 continuous report task where subjects had to report the remembered color by choosing from a 456 continuous 360° color-wheel. The use of a continuous report task allowed us to generalize our 457 findings beyond the 2-AFC paradigm, and to gain insight into how memory biases manifest in 458 response error distributions.

459 Methods

460 **Participants.** 24 healthy volunteers (15 female, mean age of 19.75 ± 1.52) from the University of 461 California San Diego (UCSD) community participated in the experiment in person. All procedures 462 were approved by the UCSD Institutional Research Board and all participants provided written 463 informed consent, and reported normal or corrected-to-normal vision without color-blindness. 464 Participants were naïve to the purpose of the study and received partial course credit for their time.

465 Stimuli & Procedure. Stimuli were rendered on a CRT monitor with a 60-Hz refresh rate and a 466 screen size of 40 x 30 cm. Stimuli were generated using MATLAB and the Psychophysics toolbox 467 (Brainard, 1997; Pelli, 1997). Participants were instructed to maintain fixation throughout, aided 468 by a white central fixation dot (0.5° diameter) presented on a dark-gray background of 2.37 cd/m2. 469 Memory items were colors randomly selected from a subset of CIE color space (L = 70, a = 20, b =470 38, radius =60), as was done in the previous three experiments. Sixteen white placeholders (4.3°) 471 radius, 0.2° thick line) were positioned around the fixation point (centered at 10.5° from fixation). 472 The locations of the two memory targets were selected at random with the exception that (1) they 473 were always presented in the same hemifield to maximize inter-item competition (Alvarez & 474 Cavanagh, 2005; Cohen et al., 2016; Störmer et al., 2014) and (2) there were always 2 empty 475 placeholders between the two memory items (i.e. they were spaced $\sim 4^{\circ}$ apart, center-to-center).

476

477 On each trial (Figure 4a), two colored stimuli were presented for either 50ms, 150ms or 500ms 478 and participants had to remember the colors as precisely as possible. The colors of the two memory 479 items could be either 0° , 20° , 45° , 90° or 135° apart in color space (with $\pm 3^{\circ}$ random jitter). After 480 a 750ms delay, one of the two colors was probed via a spatial cue (the rim of the placeholder in 481 one location got thicker). Along with the spatial probe, a randomly oriented color-wheel (with 10° 482 radius, 1° wide) was presented around fixation, and a crosshair appeared at the fixation point. 483 Participants used the mouse to move the crosshair to the hue on the color-wheel that most closely 484 resembled the remembered color at the probed location. The next trial began ~1s after participants 485 clicked the mouse and this procedure was repeated 96 times per experimental condition (i.e. a total 486 of 1440 trials per participant). Presentation of the 5 different color distances and 3 different 487 encoding times was fully counterbalanced.

488 Analyses. We calculated the difference between the cued target color and the reported color 489 (reported^o – target^o) on each trial. To investigate the systematic relationship between the cued color 490 and the non-target color, we flipped the sign of the error such that the non-target color was always 491 counter-clockwise to the cued target in the error distribution. The circular standard deviation was 492 used to quantify subjects' response precision (i.e. larger deviations indicate less precision). Biases 493 in subjects report were quantified by computing the proportion of responses on the "clockwise" 494 side of the error distribution (i.e. the side opposite to that of the non-target). We centered this bias 495 onto 0 to get a percentage score for the bias as follows (see also Figure 4b):

496
$$bias = \frac{responses \ away * 100}{total \ responses} - 50$$

497 We expect this bias metric to be roughly 0% if no biases exist, >0% if there is repulsion away from 498 the non-target, and <0% if there is attraction toward the non-target. Note that this metric reflects 499 relative repulsion/attraction biases rather than being an absolute metric, since potential "swap" 500 errors (where the target and non-target colors are confused, and a subject mistakenly reports the 501 non-target) would be counted as "attraction". Thus, this metric is conservative to the extent that 502 potential swap errors would inflate attraction biases and underestimate repulsion biases. To 503 benchmark our model-free metrics of memory precision and bias, we also fit a von Mises (circular 504 analogue of a normal distribution) to our error distributions using 2 parameters: standard deviation 505 (*vmSD*) and bias (μ). We used repeated-measures analysis of variance to evaluate the impact of 506 encoding time and color similarity on both the model-free (circular standard deviation and 507 percentage bias metric) and estimated (*vmSD* and μ) parameters.

508 Results

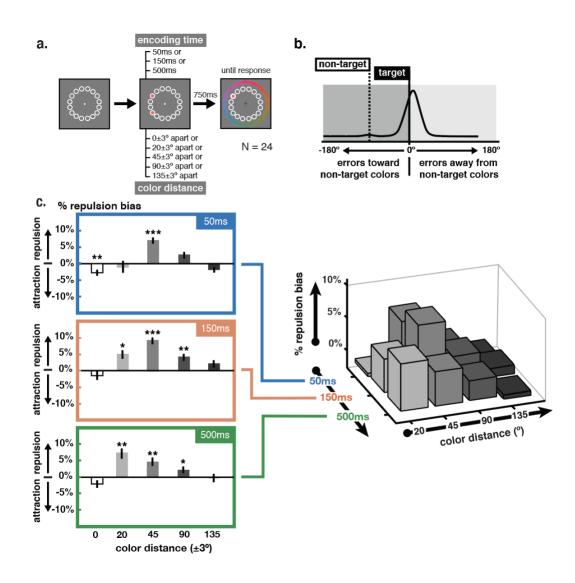
We confirmed that memory precision was higher at longer encoding times, with circular standard deviations of 26.5°, 25.0°, and 22.0° for encoding times of 50, 150 and 500ms, respectively $(F_{(2,46)}=65.17, p<0.001)$. Memory precision also differed as a function of color distance, with circular standard deviations of 17.4°, 21.4°, 25.3°, 29.1° and 29.4° for color distances of 0°, 20°, 45°, 90° and 135°, respectively ($F_{(4,92)}=69.49$, p<0.001), showing increasingly noisy responses as two colors differed more.

515 To quantify the repulsion bias, we used our model-free bias metric (as discussed above), where 516 values >0 indicate repulsion, and values <0 indicate attraction. We found differences in repulsion 517 at longer encoding times, with biases of 0.8%, 3.4% and 2.4% for encoding times of 50, 150 and 518 500ms, respectively ($F_{(2,46)}$ =9.19, p<0.001; compare the 3 panels on the left in Figure 4c). The 519 amount of repulsion also differed as a function of distance in color space between the two memory 520 items, with biases of -2.2%, 3.8%, 7%, 3% and -0.6% for color distances of 0°, 20°, 45°, 90° and 521 135°, respectively ($F_{(4,92)}=13.14$, p<0.001; compare values along the x-axis in the left panels in 522 Figure 4c).

523 Importantly, there was an interaction between encoding time and color distance ($F_{(8,184)}=3.78$, 524 p<0.001; Figure 4c). For example, the strongest repulsion bias shifted from 45° at the shortest 525 encoding time (50ms) to 20° at the longest encoding time (500ms). This is in line with the idea 526 that the maximum amount of repulsion depends on both color distance and representational 527 precision. Note how two very similar colors presented at very short encoding times show a 528 decreasing amount of repulsion (with repulsion disappearing when two items were 20° apart and 529 shown for only 50ms). This pattern likely emerges because people are no longer able to individuate 530 the two items, as shown in a control experiment (Appendix Figure 3). Interestingly, the repulsion 531 of two memory representations away from one another is not a simple lateral shift, but instead

532 leads to significantly skewed response distributions (Appendix Figure 4).

Together, these results are consistent with our framework and suggest that representations are biased to become more distinctive in order to maintain individuated representations (although in the limit people need to be able to dissociate item colors during encoding before any repulsion can occur). This means that with shorter encoding times we see maximal repulsion when two items are sufficiently distant in feature space (i.e. at 45° but not 20°). It also means that when longer encoding time leads to representations that are more precise, items must be very similar (i.e. differ by 20° in color space) to achieve maximum repulsion.



541 **Figure 4.** a) In Experiment 4, participants remembered two memory items that were either 0° , 20° , 45°, 90° or 135° apart in color space (each with \pm 3° of jitter), and that were briefly presented for 542 543 either 50ms, 150ms or 500ms. Participants reported the color of the cued item (indicated by a 544 thicker outline at one of the placeholder locations) by choosing the remembered color on a color-545 wheel. b) While non-targets could have a color that was either counterclockwise or clockwise in 546 feature-space relative to the cued color, error distributions were constructed (for each subject and 547 condition) by always plotting the non-target color as counter-clockwise from the cued color. This 548 cartoon depicts one such error distribution. Attraction and repulsion biases were operationalized 549 as the difference in the percentage of responses that were toward (dark gray shading) vs. away 550 from (light gray shading) the non-target color. c.) The 3D bar plot (right) shows repulsion as a 551 function of both encoding time (z-axis) and inter-item distance in color space (x-axis). Repulsion 552 at each encoding time is replotted in the three sub-panels (left) to show the within-subject standard 553 error (+1 SEM) for each condition, and to show the data from trials with a 0° inter-item difference 554 (not shown in the 3D bar plot) where no repulsion or attraction should exist. Overall, repulsion 555 biases were more prevalent when the two memory colors were more similar. Especially when 556 encoding time increased, and responses become more precise, did the remembered colors need to 557 be very similar to observe maximal repulsion. Single, double and triple asterisks indicate $p \le 0.05$, 558 p < 0.01 and p < 0.001 respectively (tested against no-bias; uncorrected for multiple comparisons).

Note that the above analyses, based on non-parametric quantifications of precision and bias, were confirmed with an additional analysis based on the standard deviation and bias parameters of a von Mises distribution fit to the error distributions (Appendix Figure 5).

563 In Experiment 2 we had found that the degree of repulsion bias was related to the level of task 564 engagement (Figure 2c). This indicated that a lack of effort was not the source of the repulsion 565 biases found in that experiment. To make sure this finding was not due to the specific 2AFC or 566 online nature of Experiment 2, we also analyzed the data from the current experiment, which was collected in the lab using a continuous report paradigm. Here, baseline performance was quantified 567 568 by the circular standard deviation of each subject (with lower circular standard deviation indicating 569 better performance), while bias was quantified by the percentage of responses away from non-570 target color (values >0 indicating repulsion). We found strong negative correlation between 571 circular standard deviation and bias (Pearson's rho = -0.81, p<0.001, Bayes factor = 3872; Figure 572 5) supporting and extending our findings from Experiment 2. In the current analysis, the 573 correlation is very prominent, possibly owing to the high number of trials (1440) per subject. 574 Participants with better performance (smaller circular deviation) tended to have stronger repulsion 575 bias (responses away from non-target colors were higher than 0), showing that repulsion biases 576 are strongest in participants with the highest levels of task engagement.

577

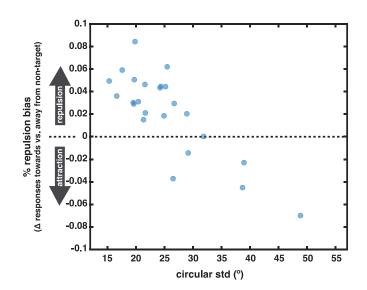


Figure 5. A strong negative correlation between bias (y-axis) and memory performance (as indexed by the circular standard deviation on the x-axis) demonstrates that repulsion is stronger in participants whose performance is better. This replicates the correlation between task performance and magnitude of repulsion biases in Experiment 2 (Figure 2c; and see also a replication experiment in Appendix Figure 1), and clearly demonstrates that a lack of effort cannot explain repulsion biases.

585

586

Experiment 5: Repulsion biases grow with longer delays

587

588 Finally, we tested whether repulsion biases become stronger with increasing memory noise. In 589 Experiments 1-4, biases emerging during encoding cannot be dissociated from those emerging 590 during the delay. Therefore, here we focus on memory noise that arises during the delay. To 591 manipulate memory noise, we compared performance across different memory delay durations. 592 Note that while some have argued that memory noise does not change as a function of delay 593 interval (e.g., Huang & Sekuler, 2010; Magnussen & Greenlee, 1992; Regan & Beverley, 1985; 594 Zhang & Luck, 2009, 2011), subsequent studies have since demonstrated that, with adequate 595 power, representations do become noisier over time (Rademaker et al., 2018; Shin et al., 2017). 596 We reasoned that if repulsion bias functions to keep two memory representations distinct, then this 597 repulsion bias should grow stronger as the memory delay (and thus memory noise) increases. 598 Alternatively, it is possible that when the two representations become increasingly noisy over time, 599 responses may instead become biased toward the average of the two colors, and thus repel less, or 600 even attract. We tested these predictions in an experiment where we manipulated delay duration 601 (250ms, 750ms, or 5000ms; see Appendix Figure 6 for stimulus presentation details) as 602 participants remembered two items. Encoding time was fixed at 250ms, and color distance between

the two items was fixed at 45° (i.e., values that yielded the largest repulsion bias in Experiment 4).
Subjects recalled the target color using a continuous report paradigm. We quantified bias in a
model-free manner as in Experiment 4.

606

607 Methods

608 **Participants.** A total of 60 naïve participants were recruited using Amazon Mechanical Turk. For 609 the control experiment (presented in Appendix Figure 7), an additional 50 naïve participants were 610 recruited from Amazon Mechanical Turk. All participants provided their informed consent, and 611 were paid approximately \$8 per hour for their time. Five participants out of 60 were excluded 612 because of poor baseline performance (mean circular standard deviation more than 70° which was 613 > 2 SD of the group). For the control experiment, 3 participants were excluded for the same reason.

614

615 Stimuli & Procedure. Stimuli and task procedures were identical to Experiments 1-2 (i.e., two 616 stimuli at a 45° color distance were briefly shown at two of 12 placeholders on the screen and 617 remembered over a delay before responding) with the following exceptions: There were no 618 placeholders next to fixation, instead, there was always a light gray circle visible (237 pixel radius, 619 2 pixels wide, #d3d3d3 hex color) outside of the placeholders (see Appendix Figure 6). This grey 620 circle turned into a randomly rotated color wheel during the response period (color wheel of the 621 same dimensions as the grey circle). The two memory stimuli were presented for 250ms and 622 participants remembered the color of each stimulus for a 250ms, 750ms, or 5000ms delay period. 623 After the delay, one of the two colors was probed, and participants reported the cued color by 624 moving a white circle along color wheel (i.e. via a continuous recall procedure as in Experiment 625 4). This procedure was repeated 60 times for each of the 3 delay period conditions (i.e., 180 trials

626 per participant in total). For the replication experiment (Appendix Figure 7), the procedure was 627 identical, with the exception that stimuli were only presented for 150ms (instead of 250ms).

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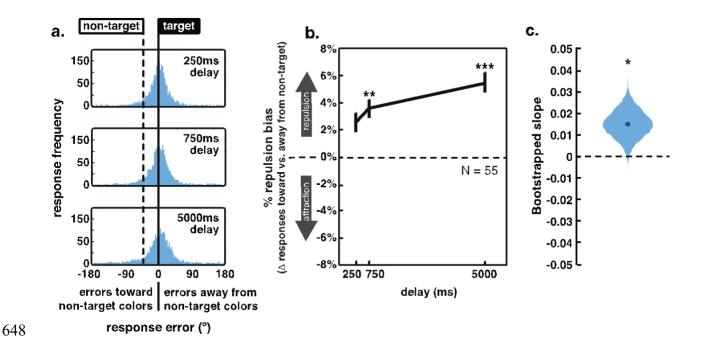
629 Results

630

631 First, we found that the width of recall error distributions significantly differed across the three 632 memory delays (Figure 6a), with circular standard deviations of 33.8°, 34.5° and 38.6° for delays 633 of 250ms, 750ms and 5000ms, respectively ($F_{(1,54)}=38.33$, p<0.001). This is consistent with the 634 notion that there is an increase in memory noise as items have to be remembered over longer 635 delays. We also found that the repulsion bias grew monotonically with delay duration, from 2.5%, 636 to 3.6%, and 5.6% for delays of 250ms, 750ms and 5000ms, respectively (Figure 6b and 6c; $F_{(1,54)}$ 637 =5.36, p=0.025), suggesting larger repulsion biases with increasing delay duration. This effect was 638 replicated in a control experiment using an independent set of subjects (Appendix Figure 7) and 639 cannot be explained by changes in swap rate with delay (i.e., swaps happen when subjects 640 mistakenly report the non-target color instead of the target color; see Appendix Figure 8).

641

Thus, when two similar (but dissociable) items have to be remembered, we observe repulsion. As the items are held in memory for increasingly longer durations, they repel further apart as they become noisier (we do not observe a switch to attraction biases). The increase in repulsion with longer delays suggests that the repulsion bias is at least partly related to the storage of information in memory, and is not purely due to perceptual factors or response strategies.



649 Figure 6. a.) At increasing delays, error distributions become wider (larger circular standard 650 deviation), indicating increasing memory noise. The distributions also reveal a high number of 651 responses biased away from the non-target. b.) The proportion of responses biased away from the 652 non-target, when quantified for the three delay-duration conditions, revealed a repulsion bias that 653 grew monotonically stronger as the delay time increased. Error-bars represent + 1 within-subject 654 SEM. Double and triple asterisks indicate $p \le 0.01$ and $p \le 0.001$ respectively (tested against no-655 bias; uncorrected for multiple comparisons). c.) To assess the increase of repulsion bias with delay, 656 one can fit a line through the three points in (b) and calculate the slope – a positive slope indicating 657 an increasing repulsion. Shown here is a distribution plot of bootstrapped slopes (5000 iterations 658 of resampling with replacement). The single asterisk indicates p<0.05. This confirms a statistically 659 robust effect, with repulsion bias growing as a function of delay duration.

660

647

An adaptive framework

661 In five main experiments (and three control experiments), we found that memory representations 662 were repelled from each other when the memoranda were highly similar (Experiments 1–2), when 663 memory representations were noisier (Experiment 4), and more when representations were 664 remembered over longer delay intervals (Experiment 5). We confirmed that these effects do not 665 simply reflect straightforward demand effect or straightforward response biases, and they hold 666 across different experimental paradigms. Moreover, we showed that participants with excellent 667 performance and task engagement showed large repulsion biases, suggesting that these biases do 668 not simply reflect a lack of effort to precisely remember the colors. Finally, when memory load 669 increased and it was harder for participants to maintain individuated representations, memory 670 biases reversed from repulsion to attraction (Experiment 3).

In this framework, we focus specifically on memory biases between amongst two or more simultaneously presented memory stimuli – which is different from categorical biases and the serial dependence effect. Overall, the experiments we presented here argue against the idea that some studies find attraction biases and some find repulsion biases purely as an artifact of using different stimuli. They also argue against the idea that such biases arise primarily from some form of motor-response strategies.

We instead suggest they these inter-item biases can be thought of as adaptive distortions by our memory system, designed to reduce error. The broad framework we adopt is that visual working memory faces at least two distinct problems. First, the capacity of working memory is limited, and when more items must be stored, they are stored with more noise (Bays & Husain, 2008; Ma et al., 2014; Zhang & Luck, 2008). In such cases, summary statistics or other ways of blending across items can be used to somewhat improve memory of individual items (Brady et al., 2011; Brady & Alvarez, 2015; Lew & Vul, 2015). The second problem is that access to memories is not automatic and not independent of cues and context. Instead, there can sometimes be confusion between items that arises when items are similar in context and features. Indeed, prominent process models of working memory that focus on feature-location binding predict that items are automatically blended if they are similar (Swan & Wyble, 2014; Lin & Oberauer, 2017). Avoiding such confusion is important to reducing error when such blending is not optimal (e.g., when item representations are not noisy, but are similar and so likely to be blended).

We do not attempt to make a precise quantitative model that could be fit to performance on our tasks. However, it is useful to formalize these ideas to see if it is plausible that reducing error is the overall goal of attraction and repulsion, and to ask whether the factors that affect the magnitude of each problem determine when we should expect attraction and repulsion to be strong or weak. We do that here.

Attraction. For the purposes of considering attraction, we assume that the information subjects have about the display is (1) information about the entire set of colors (i.e., participants know if the items were all red), and (2) information about each specific item, with, for now, the simplifying assumption that there is no confusion as to which color goes with which item (i.e., when a subject remembers the color of the i^{th} item, they never mistakenly retrieve the color of the j^{th} item). Given these assumptions, we can predict if memory distortions would be optimal to minimize error if subsequently asked to report the feature associated with an individual item.

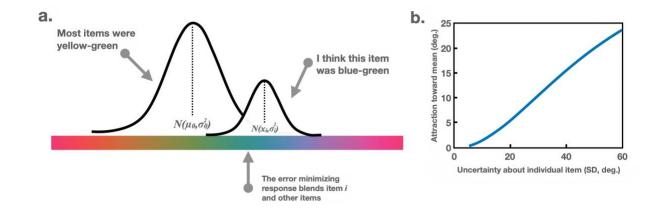
In general, the observer has an estimate of the mean (μ_0) and the uncertainty (σ_0) about the color of the entire set of colors – i.e., the ensemble – and a noisy estimate of the color of a given item (with mean x_i , and uncertainty σ_i). This gives rise to a hierarchical situation because the color of each item is part of the overall set of colors. Given this hierarchy, the optimal error-minimizing color to assign to an item follows from hierarchical Bayesian models, which for the simplest caseof two nested normal distributions is:

708
$$optimal = \frac{\sigma_i^2}{\sigma_i^2 + \sigma_0^2} \mu_0 + \frac{\sigma_0^2}{\sigma_i^2 + \sigma_0^2} x_i$$

709

721

710 That is, remembering and reporting colors according to this rule results in less error on average 711 than reporting based only on your memory of an individual item (i.e., reporting only x_i). However, 712 the cost for this increased accuracy is distortion: following this rule results in attraction toward the 713 mean color of the set. Intuitively, this distortion actually increases performance because if there is 714 a noisy sample of a given color that is green-ish blue, but the mean of the entire set of colors is 715 yellow-ish-green, it is more likely the sample was inaccurate by being too blue as opposed to being 716 too green (Figure 7a). Thus, when taking into account information from both levels, the optimal 717 color to report is slightly greener than the actual sample associated with that one color alone. That 718 is, reporting colors in this way is actually more accurate – resulting in less error on average – than 719 reporting the color you believe an item to be without pulling it toward the average of the set (Brady 720 & Alvarez, 2011; Huttenlocher et al. 2000).



722 Figure 7. a) Attraction is the error-minimizing thing to do when you have uncertainty about an 723 individual item, but know how that item related to the entire set. Blending the information about 724 the individual item with the information about the other similar items improves performance in 725 this circumstance. Intuitively, this distortion actually increases performance because if there is a 726 noisy sample of a color that is green-ish blue, but the mean of the entire set of colors is yellow-727 ish-green, it is more likely the sample was inaccurate by being too blue as opposed to inaccurate 728 by being too green. Thus, when taking into account information from both individual item and 729 group levels, the optimal color to report is slightly greener than the actual sample. b) The amount 730 of attraction that is optimal depends on several factors, but it most clearly depends on the 731 uncertainty about the individual item you are probed on: The more uncertain you are about its 732 color (the wider the normal distribution associated with it), the more attraction is optimal.

Three aspects of the optimizing equation above are relevant for attraction in a typical working memory tasks. For a given set size, more uncertainty about each item will lead to a greater reliance on information about the entire set as opposed to information about the specific item (as σ_i goes up, you weigh x_i less and μ_0 more, Figure 7b). Thus, in general, manipulations that increase uncertainty about individual items, such as decreasing encoding time or increasing delay time (Rademaker et al., 2018; Shin et al., 2017; Schurgin et al., 2020), should result in more attraction if all else is held equal.

The second relevant factor is related to the clustering of individual item values in feature space.Consider a display with a single well-formed cluster of colors that are all some shade of yellow-

ish green, as illustrated in Figure 7¹. If all the items are part of this single cluster, then as the colors get more similar to each other, the uncertainty (σ_0) associated with the group mean will go down and the group color will have a bigger influence on the optimal decision. When σ_0 gets very small, as would happen if the colors were all very similar, this factor assigns nearly all the weight to the group color and none to individual items, regardless of the uncertainty associated with the individual items.

748 A final relevant factor for attraction is that increases in memory set size don't just increase the 749 uncertainty associated with each item (i.e., drive up σ_i , which would increase attraction). Instead, 750 larger set sizes also lead to more precise estimates of the mean and less uncertainty about the entire 751 set of colors (μ_0 and σ_0), since there are more samples to constrain these values. Thus, if the items 752 are relatively tightly clustered on the color wheel at all set sizes, then, as set size goes up, your 753 certainty about the color of the whole set (the ensemble color) goes up (in the same way that having more trials would decrease the standard error of your estimate of the mean in a typical experimental 754 755 setting). This decreases σ_0 , exacerbating the attraction effect even more than just increasing σ_i 756 alone.

As a result, at larger set sizes, and particularly when the items are tightly clustered in feature space, this framework predicts a stronger attraction effect than at smaller set sizes, even with similar clustering. This follows because there are two factors driving attraction – as set size goes up, certainty about the average color of the set goes up, and the item representations themselves get noisier. In contrast, for small set sizes, only in very noisy individual-item conditions or in

¹ Of course, more complex scenarios exist: i.e., if 3 items are red-ish and 3 are blue-ish on a display of 6 items, participants may form 2 clusters and items may be selectively attracted toward the cluster they are part of (Chunharas & Brady, 2019), but we set that aside here for simplicity.

conditions where the set of items are so similar that σ_0 is much smaller than σ_i – would the framework predict any appreciable attraction effects, even though such attraction effects should be robust in displays at higher set sizes when there is clustering of the features.

765 **Repulsion.** In contrast to attraction effects, which should be amplified at large set sizes, our 766 framework suggests that repulsion biases should be error-reducing primarily at small set sizes 767 when items are highly confusable.

768 When considering attraction biases, our model assumed that when subjects seek to retrieve 769 information about color *i*, they can successfully retrieve only information about color *i* (i.e., x_i 770 reflects only color i). However, human memory in general is based on cued-retrieval: content-771 based access rather than direct access (Gallistel & King, 2011). That is, unlike a computer, which 772 stores an item in a given spot in RAM and then accesses that exact address again later, human 773 memories are retrieved by matching operations based on content. As a result, more similar 774 memories are more likely to be confused at retrieval or to interfere with each other. While widely 775 recognized in long-term memory (e.g., Criss, Malmberg & Shiffrin, 2011), this aspect of memory 776 retrieval is typically also present in models of visual working memory when they focus on cued-777 retrieval (Swan & Wyble, 2015; Oberauer & Lin, 2017).

Importantly, such models of memory blend together the representation of different items *all the time* because of interference between memory representations, as a natural consequence of cued retrieval. For example, when storing just two item similar items, the "binding pool" model of Swan and Wyble (2015) predicts that the two items will be attracted to each other significantly (see Figure 8). As we have seen, however, this is not in any way optimal: with strong memories, and few items to give rise to a tight ensemble distribution, attraction will not reduce error. 784 Thus, in this scenario, an adaptive system must balance the need to avoid overlap between item 785 representations and the need to maintain an accurate memory. If the representations are encoded 786 veridically, they will have significant interference and be blended inappropriately. If they are 787 represented as more distinct from each other than they really were, this will come with its own 788 reduction in accuracy although it will also reduce inappropriate blending. The memory system 789 must strike a balance, with systematic repulsion to offset the blending that would otherwise occur, 790 but not so much repulsion that it impairs accuracy overall. We can simulate this in the Binding 791 Pool model (with all of the default parameters) simply by adding an attraction or repulsion step to 792 the encoding process, and seeing what happens to (1) the resulting bias, and (2) error. In the binding 793 pool model, the error minimizing amount of repulsion for storing 2 items that are 15 degrees apart 794 in color space is $\sim 5 \text{ deg}$ (Figure 8). More repulsion is required to minimize error when items are 795 more similar and/or when items are represented with more uncertainty.

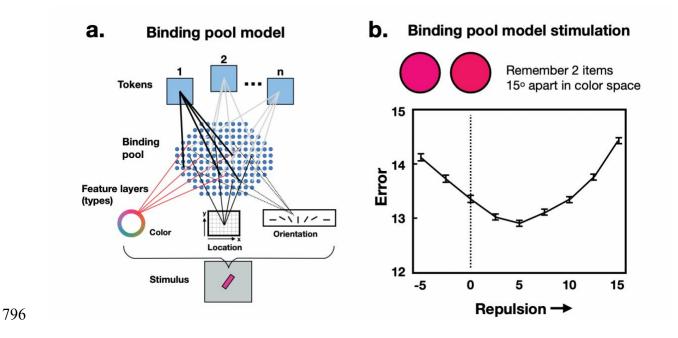


Figure 8. a) Schematic of the Binding Pool model, reproduced from Swan and Wyble (2014). In
this model, each stimulus evokes an activation in a set of feature layers (here: color, location, and

799 orientation). These features are then encoded into a shared binding pool layer and tied to a 800 particular 'token'. When provided a cue at test (like the item location) the associated token can be 801 activated and the color or orientation retrieved. Notably, the binding pool layer, which is shared 802 between all items and the source of the capacity limits of the system, also results in the items 803 features being necessarily blended (e.g., Swan & Wyble, 2014, Figure 11), even when only 2 items 804 are represented. Thus, by default, this model, like many others, always predicts attraction between 805 memory representations. b) We simulated what would happen if repulsion was added to the 806 encoded information in the binding pool model, to provide a concrete case study for how repulsion 807 could be used to overcome the blending inherent in a model such as the Binding Pool model, and 808 reduce overall error. In particular, we asked the model to store and recall 2 items that were 15° 809 apart in color space. As part of the encoding stage, we added an additional step that introduced 810 repulsion of the colors of the two items before they were put in the binding pool. In 100 simulations 811 of the model at each of 9 levels of additional attraction or repulsion added at encoding, we 812 calculated the model's error. We found that error was minimized when the items were repelled 813 away from each other by $\sim 5^{\circ}$ before being entered into the binding pool layer.

814 Summary: attraction and repulsion. Our adaptive framework holds that attraction biases (when 815 memory is very noisy) can be understood as optimal using a straightforward hierarchical Bayesian 816 integration model. Effectively, attraction biases arise because integrating summary statistic 817 information results in reduced error even if it results in systematic distortion (Brady & Alvarez, 818 2011; Huttenlocher et al., 2000). This framework makes a clear set of predictions about when 819 attraction should occur: when items are clustered in color space and individual items are associated 820 with a higher degree of uncertainty than the ensemble color. In practice, this ends up happening 821 primarily when set size is high, or when set size is low but items are very similar relative to the 822 item-level uncertainty. In contrast, repulsion bias can be understood as balancing the avoidance of 823 overlapping representations with the need for accurate representations. Insofar as overlap is present 824 and attraction is not adaptive, this model predicts that items should repel from each other. At low 825 set sizes, this means that repulsion is expected whenever items are similar enough, and uncertainty 826 high enough, that the memory representations overlap substantially. At high set sizes, the extent 827 to which repulsion will be useful in lowering error is severely reduced by the crowding of the 828 feature space with other items, and the fact that attraction and repulsion pull in different directions, 829 with attraction likely being dominant. Overall, we believe this adaptive framework can providing 830 a guiding theory for conceiving of when attraction and repulsion arise in memory.

831

General Discussion

832 Our memory is susceptible to systematic distortions. Even across short periods of time, specific 833 memories become affected by the overarching categories that memory items belong to (categorical 834 biases) or by information viewed in the immediate past (serial dependence). The research 835 presented here focused on a different kind of distortion: inter-item distortions that occur in memory 836 when we try to hold multiple items in mind. When encoding and remembering multiple items at 837 once, mnemonic representations can be subject to systematic distortions that can make items either 838 more separable (repulsion biases) or more similar (attraction biases). While both types of inter-839 item distortion are well documented, it is not clear when repulsion or attraction will occur as a 840 function of the type of information being remembered and current task demands. Here, we 841 examined when each type of bias arises. We found that memory representations were repelled 842 away from each other when the memoranda were highly similar (Experiments 1–2), when memory 843 representations were noisier (Experiment 4), and when representations were remembered over 844 longer delay intervals (Experiment 5). We confirmed that these effects do not simply reflect

straightforward response strategies, and occur in many distinct tasks, and we showed that highperforming participants showed larger repulsion biases which suggests that these biases do not simply reflect a lack of effort to precisely remember the colors. Finally, when memory load increased and it was harder for participants to maintain individuated representations, memory biases reversed from repulsion to attraction (Experiment 3).

850 Past work has found evidence for attraction biases (Brady et al., 2011; Brady & Alvarez, 2011, 851 2015; Dubé et al., 2014; Dubé & Sekuler, 2015; Huang & Sekuler, 2010; Lew & Vul, 2015; Lorenc 852 et al., 2018; Utochkin & Brady, 2020), repulsion biases (O'Toole & Wenderoth, 1977; Rauber & 853 Treue, 1998; Suzuki & Cavanagh, 1997; Scotti et al., 2021), or both (Bae & Luck 2017; Rademaker 854 et al 2015; Golomb 2015). Our model and empirical work identifies several key factors that drive 855 these effects and provides evidence that both can arise even in similar paradigms. This is important, 856 as using highly comparable paradigms and memory for a single feature (color) argues against the 857 more mundane explanation that differences in stimulus features (such as orientation in Bae & 858 Luck, 2017; Dubé & Sekuler, 2015; Huang & Sekuler, 2010; Lorenc et al., 2018; Utochkin & 859 Brady, 2020, spatial location in Lew & Vul, 2015; Suzuki & Cavanagh, 1997, motion direction in 860 Kang & Choi, 2015, or color in Brady & Alvarez, 2015; Golomb, 2015) lead to attraction in some 861 studies and repulsion in others.

Here, we tested the general account that when subjects were trying to encode items in a memory display, repulsion and attraction were driven largely by the inter-item relationship between memoranda. We proposed a way to conceive of these biases and when they arise based on adaptive framework. In particular, we suggested that these biases may be natural consequences of the memory system attempting to minimize memory error, if systematic distortion is adaptive in particular circumstances (Schacter et al., 2011). When many similar items are present and so 868 memories for individual items are noisy, attraction biases are known to be optimal for minimizing 869 error (e.g., Brady & Alvarez, 2011). Repulsion biases can also reduce error in some situations, 870 making them adaptive. In particular, if the items would naturally be blended or confused by our 871 memory system (Swan & Wyble, 2014; Oberauer & Lin, 2017), then repulsion can reduce this 872 tendency and reduce error when we have strong and distinct item memories. Importantly, these 873 biases are not simply inherited from perceptual processing: as noise accumulates in memory over 874 time (reducing the signal-to-noise if memory items), and the need to keep memoranda distinct 875 grows, a corresponding increase in the repulsion bias is observed. Importantly, very recent work 876 (performed since the first presentation of the experiments in the current paper) has confirmed 877 various key aspects of our framework: As memories get weaker, biases switch from repulsion to 878 attraction (Lively, Robinson, & Benjamin, 2021), and repulsion biases increase with longer 879 memory delays (Scotti et al., 2021).

880 Based on these results, the degree and type of bias likely depends on the overall discriminability 881 of a stimulus feature under investigation (such as color, space, orientation, etc.): If features are 882 very readily discriminable, then repulsion will only occur when two items are very similar. Poorly 883 discriminable features will need to differ more before they are susceptible to repulsion. In other 884 words, the data suggest that the extent and type of bias will directly map onto the just-noticeable-885 differences (JND) of a given stimulus feature (and of individual subjects). Using JND as a standard 886 unit might be an interesting approach that allows us to compare the various effects previously 887 reported. Even though we tried to use the same stimulus feature and investigate various task 888 manipulation in this paper, it is still not easy to compare the results with previously reported 889 findings where many interesting inconsistencies are waiting to be explored.

890 Even though our experiments were designed to rule out specific forms of response strategy, it is 891 still possible that our findings could be explained by other response strategies that closely resemble 892 the framework proposed here. For one example, it is possible that foils in Experiments 1 and 2 893 were too similar to the true answer, and that subjects might choose between the two response 894 options by the process of elimination (i.e., "I did not know which one was the target color so I am 895 going to choose the one that is less similar to the non-target"). In this hypothetical case, it is still 896 unclear how the subject would know which response option is less similar to non-target without 897 knowing which one is more similar to the target – making it a possible but implausible strategy. 898 We would like to note how recent neuroscience studies have demonstrated that memory 899 representations drift over time (Compte, Brunel, Goldman-Rakic, & Wang, 2000) - a process 900 which is not likely to be susceptible to response strategies.

901 Mechanisms of memory biases. Attraction biases can occur both in absolute stimulus space (e.g., towards particularly salient colors (Bae et al., 2015)) or arise from the similarity between items in 902 903 an individual display (as in the current work). These attraction biases are straightforwardly 904 explained as arising from gist-based or ensemble-based representations, and a combination of these 905 global representations with item specific representations. Many models claim that attraction biases 906 are the result of weighting the representation of each object towards the "summary" of the set to 907 achieve a more stable memory at the expense of maintaining distinctions between individual items 908 (Brady & Alvarez, 2011), or via blending items together if they are similar (e.g., Swan & Wyble, 909 2014; Oberauer & Lin, 2017). The category learning literature has carefully demonstrated that this 910 is in general an adaptive strategy that serves to minimize error (Huttenlocher et al., 2000).

Repulsion biases have traditionally been more difficult to understand. Previous studies have shownthat repulsion biases occur when two items are task-relevant and proximal in feature space

913 (Rademaker et al., 2015; Bae & Luck, 2017, Golomb 2015). However, the benefits of repulsion 914 biases are still unclear. Here, we suggest that repulsion biases serve to maximize distinctiveness 915 between items, when individual item representations are strong but items are similar enough to be 916 more difficult to distinguish. This helps reduce blending between items that naturally occurs in the 917 memory system (Swan & Wyble, 2013; Oberauer & Lin, 2017). Any factor that affects 918 distinctiveness in memory should thus impact the degree of repulsion biases (e.g., encoding time, 919 feature similarity, memory delay). Interestingly, previous work has frequently found repulsion not 920 only between items, as in the current work, but in absolute terms as well. For example, when asked 921 to remember an orientation that is near, but not quite at, vertical, people will systematically report 922 the orientation as further from vertical than it really was (Jastrow, 1892; Smith, 1962). One 923 framework that has been useful to understand these absolute biases is to dissociate the physical 924 space of the stimuli (e.g., absolute orientation) from the psychological representation of the stimuli 925 (e.g., people may over-weight certain values in a systematic manner). A clear example of a warped 926 psychological space is the massive overrepresentation of vertical and horizontal orientations, 927 presumably to efficiently code environmental regularities (Girshick et al., 2011; Wei & Stocker, 928 2015). Accounting for this selective over-representation of certain stimulus values in 929 psychological space can explain biases like repulsion from cardinal axes, and the reason why these 930 biases tend to arise in parts of stimulus space where discrimination thresholds are lowest (e.g., the 931 most overrepresented stimulus values) (Wei & Stocker, 2015, 2017).

This conception of psychological space is designed to address long-term biases that are likely crystalized in the neural architecture of the visual system, whereas the biases we examined in the current work are more dynamic. Despite the apparent disconnect, a common mechanism such as the warping of psychological space may be at play in both stable long-term phenomena and in 936 more dynamic short-term regimes. In the current work, this would mean that a strong 937 representation of an item "stretches" the psychological representation of stimulus space near that 938 item, resulting in repulsion of other items in a manner similar to how cardinal orientations repel 939 nearby items. This is consistent with other short-term effects: For instance, spatial judgments are 940 distorted by top-down factors such that there is repulsion bias away from currently attended 941 locations (Suzuki & Cavanagh, 1997). Attention, which leads to well-documented changes in 942 visual sensitivity (i.e., lower discrimination thresholds, see Carrasco, 2011), may also adaptively 943 bias perception and memory on demand, as biases typically manifest when discrimination 944 thresholds are low across a variety of visual features such as orientation, motion direction, spatial 945 frequency, and visual speed (see Zhang & Luck 2011 for a summary). Thus, attention amplifying 946 discrimination at a single color may strengthen the representational space there, resulting in 947 repulsion. In sum, conceptions of psychological space, and how it is distorted when particular sets 948 of stimuli are over-represented, may be a useful framework for considering biases at all possible 949 time scales (see also Schurgin et al., 2020, for details on the widely applicable utility of this 950 concept).

951 What might be the neural substrates of biased representations? When a task requires focal attention 952 to a small set of items to remember – as is the case in paradigms that create repulsion bias – the 953 discriminability of the relevant items can be improved by biasing responses in early visual cortex 954 to maximize the separability of their corresponding neural representations. For example, attention 955 to highly similar features, akin to remembering two highly similar colors in Experiments 1, 2, and 956 5, has been shown to modulate neurons tuned just away from the attended features. This 'off-957 target' gain can improve performance because neurons tuned away from the attended features 958 undergo the largest change in firing rates because the two features fall along the steepest part of 959 their bell-shaped tuning curves. In turn, this off-target gain gives rise to systematic biases in 960 behavioral reports such that people see stimuli as repelled from the actual feature values (Jazayeri 961 & Movshon, 2007; Navalpakkam & Itti, 2007; Scolari & Serences, 2009). Such repulsion would 962 be expected if the off-target gain happening in early visual cortex was interpreted as a veridical 963 representation of the world at higher stages of processing. While previous work in this domain has 964 focused on selective attention to continuously present stimuli, a similar type of modulation in the 965 domain of working memory might give rise to repulsive biases in mnemonic representations. 966 Indeed, repulsion biases grow with delay only when a memory is actively held in mind (but 967 disappears when an attention-demanding task is performed during the delay), suggesting that the 968 repulsion bias is not a product of some passive process, but instead requires active maintenance 969 (Scotti et. al., 2021). While speculative, this type of adaptive neural modulation may map onto the 970 psychological space framework, such that changes in the discriminability of stimuli in early visual 971 cortex - either due to a lifetime of experience or to dynamic changes in the focus of attention -972 lead to a warping of perception and memory.

974 **References**

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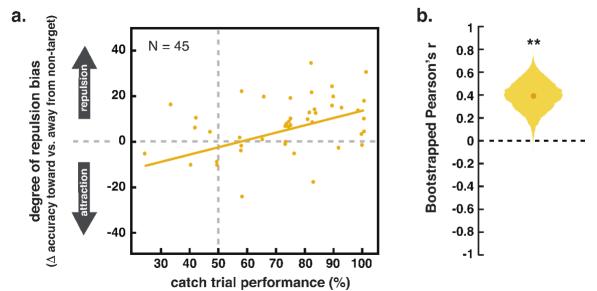
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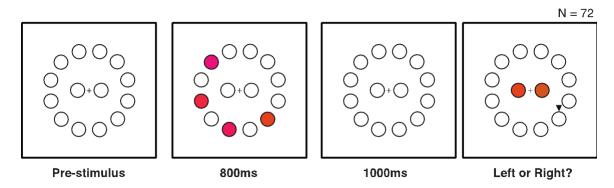
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1161	An adaptive perspective on visual working memory distortions
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1169	This PDF file includes:
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1171	Appendix Figures 1–8
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1173	Other supplementary materials for this manuscript include the following:
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1175	Data, code
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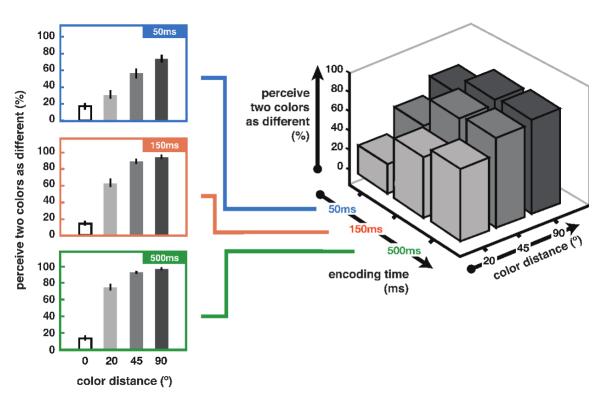


1191 1192 Appendix Figure 1. Results from a control experiment (N=45) replicating Experiment 2: In this experiment, only 1193 foils distorted by 6° relative to the correct color were used (towards and away from the non-target – similar to Exp. 1194 1), while we also included 10% of catch trials (similar to Exp. 2). Participants were an entirely new and independent 1195 set of 45 naïve Amazon mechanical Turk workers. a.) The degree of repulsion bias (indexed as the difference in 1196 accuracy between trials with foils distorted toward, and trials with foils distorted away from the non-target color), 1197 plotted against people's general level of engagement with the memory task (indexed by performance on catch trials). 1198 Each dot represents a single subject. These data demonstrate stronger biases away from the non-target color in 1199 participants with higher levels of task engagement. b.) We bootstrapped the data in (a.) 5000 times: on each bootstrap 1200 we sampled 45 subjects with replacement, and re-calculated the correlation between repulsion bias and general task 1201 engagement. This gives a distribution of bootstrapped Pearson's r, which is depicted in the violin plot. The dot in the 1202 middle indicated the mean bootstrapped correlation (r = 0.39). The double asterisks indicate a p-value of p<0.01. 1203



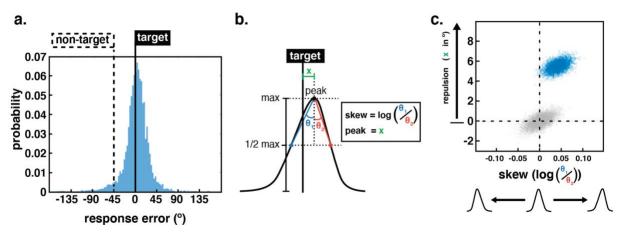
Appendix Figure 2. Task progression in Experiment 3: Participants had to remember a set of four colors (shown at randomly selected locations from a set of 12 possible locations, with at least one empty placeholder between items). The four colors were presented for 800ms, after which participants remembered them during a 1–second memory delay. Subsequently, participants saw a location cue (triangle) indicating which memory item to respond to, as well as two response options presented directly left and right of fixation. Participants chose between the correct (cued) color and a foil color.





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Appendix Figure 3. Results from a same-different color discrimination task as a control for Experiment 4. This control experiment probed whether two colors can or cannot be perceptually discriminated at various encoding times 1223 and color distances: Two colors that were either exactly the same (50% of trails) or differed by 20°, 45°, or 90° in CIE 1224 1*a*b* color space (50% of trials) were simultaneously presented for either 50ms (blue), 150ms (orange), or 500ms 1225 (green). Participants on Amazon mechanical Turk (18 in total) reported whether the two colors were the same or 1226 different. Each participant completed 90 trials in total. The 3D bar plot (right) shows accuracy as a function of 1227 encoding time and color distance. Repeated-measures ANOVA's demonstrate both main effects of encoding time 1228 $(F_{(2,34)}=36.7, p<0.001)$, color distance $(F_{(3,51)}=212.5, p<0.001)$ and an interaction $(F_{(6,102)}=9.32, p<0.001)$. This means 1229 that participants could not tell two colors apart when they were presented very briefly and were very similar to one 1230 another (i.e., encoding time of 50ms and color distance of 20°). The inability of subjects to tell two very similar colors 1231 apart at very short encoding times explains why repulsion biases were not found in these extreme cases.



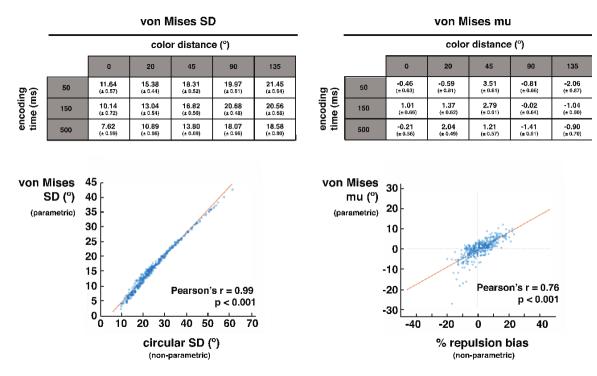
1236 1237 Appendix Figure 4. Asymmetry of Experiment 4 error distributions: a) An example error distribution from all 24 1238 participants combined, in the condition showing the strongest repulsion bias (i.e. encoding time of 150ms and color 1239 distance of 45°). First, note how the peak of the error distribution is not aligned with the cued color, but instead is 1240 shifted away from the non-target color. Second, note how the shape of the distribution is asymmetrical, with the side 1241 away from the non-target being steeper. b) Due to the possible presence of non-target responses (i.e. where a subject 1242 mistakenly reports the color of the non-target instead of the target), we did not wish to measure skewness using circular 1243 skewness measures on the raw response distribution. Instead, we first derived a kernel density estimator (KDE). The 1244 peak of the distribution (x) was defined as the degree of error with maximum probability. The skewness was defined 1245 by the log ratio between the angle toward (θ_1) vs. away (θ_2) from the non-target color at half maximum height of the 1246 KDE $(\log(\theta_1/\theta_2))$. c) A scatter plot showing the relationship between skew and peak. Each dot represents skew and 1247 peak on one bootstrapping iteration (of 5000 total iterations) calculated by randomly resampling the data from 24 1248 participants with replacement (data from the condition shown in a.). The horizontal zero line represents scenarios with 1249 no shift in the distribution peak, while the vertical zero line represents scenarios without any skew (thus, the 0.0 point 1250 represents a perfectly symmetrical distribution). We found both a systematic shift of the peak (p<0.001 from 1251 bootstrapping) as well as skew (p<0.01 from bootstrapping). Furthermore, the shape of the dot cloud shows that 1252 stronger repulsion is associated with a stronger skew (r=0.45; p<0.001). To test the validity of the metrices, we 1253 reanalyzed the same data with randomized signed errors and plotted in grey color. The randomized signed errors 1254 distribution centers at zero in both skew (x-axis) and bias (y-axis) suggesting that the significant bias and skew were 1255 not spurious.

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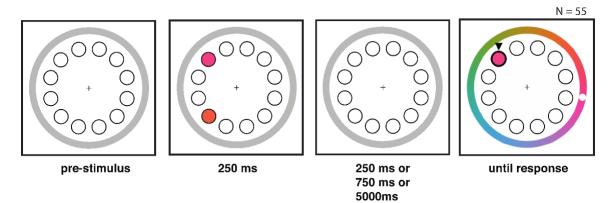
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1263 Appendix Figure 5. Parametric vs. non-parametric quantifications of memory precision and bias in Experiment 4. 1264 For this experiment we non-parametrically quantified memory precision as the circular standard deviation (with 1265 smaller standard deviations indicating higher precision) and we quantified biases as the difference in the percentage 1266 of responses that were toward vs. away from the non-target color (with a negative bias indicating attraction, and 1267 positive bias indicating repulsion). To validate these measures, we also parametrically fit the data using a von Mises 1268 distribution with two independent parameters to reflect memory precision (vmSD) and bias (mu). We found a high 1269 agreement between parametric vs. non-parametric measurements (Pearson's r = 0.99 and 0.76, for precision and bias, 1270 respectively; both p<0.001). The correspondence between these measures is shown in the scatter plots at the bottom 1271 of this figure. Furthermore, we repeated our statistical analyses with the parametric von Mises parameter estimates 1272 (tables in the top of this figure), showing significant differences in memory precision as a function of encoding time 1273 $(F_{(2,46)}=13.7, p<0.001)$, color distance $(F_{(4,92)}=21.09, p<0.001)$, and an interaction $(F_{(8,184)}=3.76, p<0.001)$. The 1274 repulsion bias is marginally impacted by encoding time (F_(2,46)=3.08, p=0.056), significantly impacted by color 1275 distance ($F_{(4,92)}=9.54$, p<0.001), and there is a significant interaction ($F_{(8,184)}=2.66$, p<0.01). Note that the mixture 1276 modelling assumes that the error distribution follows a symmetric circular distribution. However, the true error 1277 distributions were skewed which makes it less accurate in estimating the true biases and the memory strengths.

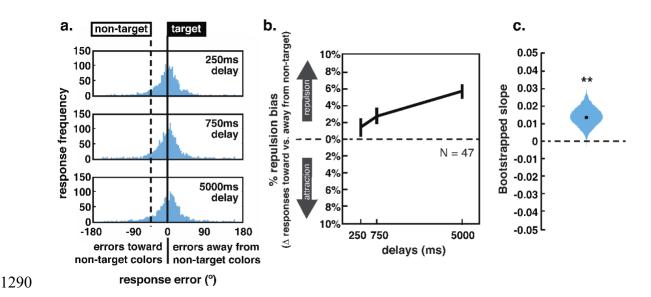
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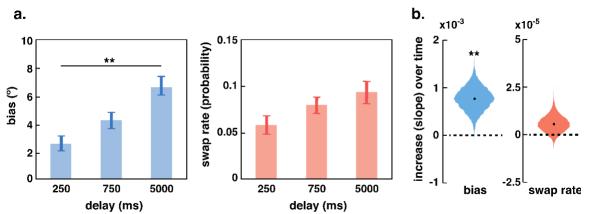




1291 Appendix Figure 7. Results from a control experiment (N=47) replicating the finding from Experiment 5 that memory 1292 biases increase with longer delays. Here, we collected 36 trials per condition per subject (a total of 108 trials per 1293 subject) a.) Error distributions at each delay, revealing a high number of responses biased away from the non-target. 1294 b.) The quantified repulsion bias (i.e. percentage of responses away from the non-target color) shows that repulsion 1295 grew monotonically stronger as the delay duration increased (1.4%, 2.7%, and 5.6% for delays of 250ms, 750ms and 1296 5000ms respectively; $F_{(1,46)}=6.62$, p=0.013). Error-bars represent ± 1 within-subject SEM. c) To assess the increase 1297 of repulsion bias with delay, one can fit a line through the three points in (b) and calculate the slope – a positive slope 1298 indicates repulsion bias growing as a function of delay duration. Shown here is a distribution plot of bootstrapped 1299 slopes (5000 iterations of resampling with replacement). The double asterisk indicates p<0.01 confirming a 1300 statistically robust effect.

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 $\begin{array}{c}1303\\1304\end{array}$ Appendix Figure 8. a.) Fitting a mixture model with swap errors to the data in Experiment 5 confirms that 1305 repulsion bias grows stronger with longer delay intervals (blue; F_(1,54)=10.2; p=0.002), confirming what we found 1306 with our non-parametric repulsion bias measure. The frequency of swap errors did not significantly change 1307 across time (red; $F_{(1,54)}=1.87$; p=0.178). b.) We computed slopes of bias and swap errors as a function of time – 1308 positive slopes indicating an increased repulsion or swap rate over time. We evaluated significance by 1309 resampling with replacement 10,000 times. Repulsion bias grew significantly stronger as the delay interval 1310 increased (blue), replicating our findings using a non-parametric bias measure. Swap errors did not increase 1311 significantly as the delay interval increased. These results suggest that the increase in repulsion bias that we 1312 found when using either parametric or non-parametric methods cannot be explained by a reduction in swap errors 1313 (if anything, swap errors increase with delay, numerically).