Contextual cuing in the presence of an endogenous cue for attention

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13 Abstract

 $_{14}$ Three experiments explored the interaction between an endogenous cue of attention and

the repetition of a visual search display during contextual cuing. In Experiment 1,

participants readily learnt about repeated configurations of visual search, before being

presented with an endogenous cue for attention towards the target on every trial.

Participants used this cue to improve search times, but the repeated contexts continued to

guide attention. Experiment 2 demonstrated that the presence of the endogenous did not

impede the acquisition of contextual cuing. Experiment 3 confirmed the hypothesis that

21 the contextual cuing effect relies largely on localised distractor contexts, following the

2 guidance of attention. Together, the experiments point towards an interplay between two

23 drivers of attention: after the initial guidance of attention, memory representations of the

²⁴ context are flexibly deployed to facilitate the target detection.

Public significance statement: This study provides a test of how attention is

26 governed by different aspects of the environment. We examine whether the control of

attention by an instructional stimulus (an arrow that directs attention) will interfere with

28 the way in which attention is governed by other learnt visual patterns in the environment.

Keywords: visual search; contextual cuing; endogenous cuing

Word count: 7999

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Contextual cuing in the presence of an endogenous cue for attention

It is well established that the process of visual search is guided by past experience.
When we encounter a scene, the extent to which the configuration of stimuli matches
stored representations in memory will determine the effectiveness of the processing and
subsequent search through the elements of that scene. This cognitive process is studied in
the lab using the contextual cuing (CC) task: participants typically experience a standard
visual search task (i.e., serial processing; slow search), such as searching for a T amongst L
shapes. A set of search configurations is repeated across trials, and response times to
targets are faster compared to those in configurations that do not repeat. Thus, the
repetition of the search configurations leads to a stored representation of (some aspect of)
the configuration in memory, and future processing of the same configuration activates the
memory representation, driving more efficient search behaviour within that scene.

Much work has focused on the nature of the memory and attention processes 43 responsible for contextual cuing. The effect was initially suggested to be implicit in nature, with repeated configurations seemingly guiding search unconsciously: typically participants 45 are unable to articulate their knowledge of the repeated configurations, and show poor ability to recognise configurations in memory tests (e.g., Chun & Jiang, 1998; Colagiuri & Livesey, 2016), although this view of CC has been strongly contested (e.g., Smyth & Shanks, 2008; Vadillo et al., 2016). There are also a number of plausible models of how memory representations of repeated configurations might guide search (e.g., Beesley et al., 2015; Brady & Chun, 2007), with the predominant view being that the memory 51 representations are best characterised as associative in nature, whereby distractors (or 52 groups of distractors, see Beesley et al., 2016) form associations that activate more likely 53 target positions.

The exact nature of how repeated configurations facilitate visual search is also the focus of much debate within the literature. There is a question as to whether CC reflects

enhanced attentional processing of the display, such as by reducing the number of
distractors processed (e.g., Beesley et al., 2018), or whether it facilitates the decision
process once targets have been detected (e.g., Kunar et al., 2007; Sewell et al., 2018). The
current article focuses on the assumed attentional advantage for repeated configurations,
and explores the extent to which this results in an automatic form of attentional bias.
That is, to what extent does the processing of the search configuration control the guidance
of attention, and to what extent does that guidance persist even in the presence of other
top-down control processes that might be driving attention.

A number of studies have explored how flexible the learned behaviours are in 65 contextual cuing. For example, a number of studies have shown that moving the target to a new position within the display will abolish the established CC effect (Makovski & Jiang, 2011; Manginelli & Pollmann, 2009). Notably, Zellin et al. (2013) explored the remapping of target positions over a longer training period, observing that with extended training, 69 new associations will form for these new target positions, though the effects are limited to 70 targets that appear closer to those that are initially trained. This suggests that any 71 relocation effect is driven strongly by a generalisation of the pre-existing associations. 72 Furthermore, strong contextual cuing effects were observed for the initially trained targets 73 in a final "return phase" at the end of the experiment. All of these results point towards CC constituting a fairly inflexible behaviour that is activated somewhat automatically during search. 76

More direct examination of the role of top-down control processes on CC comes from Luque et al. (2017) (Experiment 3). They used a task in which participants were initially given a standard CC experiment (search for a T amongst Ls), before then being told in a second phase that the target would appear in two designated positions along the horizontal mid-line of the screen. Participants were given an explicit instruction to search in these two locations for a new target (a Y); in this phase participants engaged in a new search task requiring controlled attention to specified locations. Yet the underlying configuration of repeated distractors was still present, as was the original target, which
appeared in its trained location for that configuration. Luque et al. found that the
acquired knowledge of the configurations did not affect performance in this second phase:
responses to the new target were comparable when the old target was pointed in either the
same or opposite direction to the new target, suggesting that there was no detectable
processing of the old target (see also Luque et al., 2021). The suggestion is that contextual
cuing can be controlled in the presence of a top-down instruction to search in a new
location - search is not automatic in nature in the CC task.

One potential issue with the studies presented by Luque and colleagues (Luque et al., 2017; Luque et al., 2021) is that participants are instructed to engage in a new search process for a new target object: participants initially search for a T and are later instructed to search for a Y. The role of a prior target template is important for visual search (Vickery et al., 2005; Võ & Wolfe, 2012), and object identities appear to play an important role in the contextual cuing effect (Makovski, 2017, 2018). While it is unclear how dependent CC is on the identity of the target, it is possible that distractor-target associations may well be sensitive to target identity and to the goals of the participant. For this reason, the current study assesses the impact of top-down instruction on CC when participants maintain the same task goal of searching for a single target identity within the display.

The overarching aim of the current study is to explore the interaction between controlled (top-down) attentional processes and the pattern of search behaviour established by the repeated configurations. Specifically we seek to understand whether repeated configurations continue to guide attention even when participants are directed to alter their natural search patterns by the presence of an endogenous cue (an arrow that instructs the participant to direct attention in a specific direction). The experiments explore both the performance aspect of CC in terms of whether it continues to guide behaviour once an endogenous cue is introduced, and also whether the development of the search behaviour is impeded when trained concurrently with the endogenous cue.

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Transparency and Openness

The raw data, analysis scripts, experimental materials, and the manuscript source files, are available at http://github.com/tombeesley/CC_Control. The analyses reported in this manuscript are computationally reproducible from the manuscript source files (using R v4.3.1), which are available at the github repository. The study design and analyses were not pre-registered.

Experiment 1

Experiment 1 sought to examine whether the learnt attentional behaviour that 118 develops during contextual cuing is expressed when participants are directed by an 119 endogenous (instructional) cue to search in a particular region of the visual scene. 120 Participants were first trained with a set of four repeating configurations in phase 1 across 121 5 epochs of 32 trials each. Then prior to phase 2, participants were told that an arrow 122 would appear before every trial indicating the side of the screen on which the target would 123 be located. This arrow was valid on every trial. In phase 2, the repeating configurations were presented in two forms: "consistent", where the target appeared in the same position 125 as it has appeared for that configuration in phase 1; and "inconsistent", where the target 126 appeared in a position in the opposite quadrant of the screen from where it had appeared 127 in phase 1. Random configurations were also presented in this phase. If the contextual cues 128 within the repeated configurations continue to guide attention in the presence of the 129 instructional cue, then we would expect that response times would be faster on consistent 130 trials compared to random trials. In addition, we would also expect that the contextual 131 cues would guide attention away from the (new) target quadrant on inconsistent trials, and 132 so response times should be slower on these trials compared to those on random trials. 133

134 Method

135 Participants

Thirty-one undergraduate students from Lancaster University were recruited (mean age = 20.13, SD = 1.09; 17 identified as female and 14 as female) via the Psychology
Research Participation System in the Department of Psychology at Lancaster University, in return for the opportunity to use the recruitment system for their own research in future years.

Participants were tested individually in a quiet room with a Dell laptop with a

Materials

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15.6" screen, a screen resolution of 1920 x 1080, and a full size external keyboard for 143 participants to use to respond to the task. Participants sat approximately 50 cm from the 144 screen. Stimulus presentation was controlled by MATLAB using the Psychophysics 145 Toolbox extensions (Brainard, 1997; Kleiner, Brainard & Pelli, 2007; Pelli, 1997). 146 Responses to the target stimulus were made by pressing the 'c' or 'n' key on a standard 147 keyboard. All experimental materials are available at the github repository for this study. 148 Distractor stimuli were an 'L' shape (rotated 0°, 90°, 180°, or 270°) while the target 149 stimulus was a 'T' shape (rotated at either 90° or 270°). Stimuli were 8 mm square and 150 arranged in a square grid of 144 evenly spaced cells (12 x 12) which was positioned 151 centrally on the screen and was 170 mm square. The grid itself was invisible to 152 participants. The fixation cross (displayed centrally before each trial) was 4 mm square. 153 The background of the screen was grey (RGB: .6, .6, .6) and the stimuli were presented in 154 black (RGB: 1, 1, 1). There was a small offset in the vertical line of the 'L' distractors, 155 which increased the similarity between the 'L' distractors and the target 'T', making the 156 search task more difficult (Duncan & Humphreys, 1989). 157

Design

Phase 1 employed a within-subjects design with factors of epoch (1-5) and configuration (repeated and random). All configurations contained 16 distractors, equally divided between the four quadrants of the display, and one target. Four repeated configurations were trained. Four target locations were used, with one from each quadrant assigned to each of the repeated configurations. These same four target positions were used for the random configurations throughout the task. Each of these four target positions was chosen at random from one of five locations within each quadrant, that were approximately equidistant from the center of the screen. Distractors could not appear in these target locations.

Phase 2 employed a within-subjects design with factors of epoch (6-10) and 168 configuration (repeated: consistent; repeated: inconsistent; random: consistent; 169 random:inconsistent). On each trial, there was a .5 probability that an "inconsistent" 170 version of the configuration would be presented. This meant that the target was relocated 171 to a diametrically opposed target position such as to maximise the displacement from the 172 trained target position (see Figure 1). This could occur for both the repeated and random 173 configurations, hence creating four unique trial types for this phase. While random 174 configurations did not have a "trained", associated, target position, it is necessary to divide 175 the random trials into consistent and inconsistent trial types in this way in order to assess any target frequency effects that may occur, since the inconsistent target locations used in this phase were novel.

179 Procedure

Participants were tested individually in a quiet testing room. They were given instructions on how to complete the task, including the presentation of an example of a search trial. Participants were shown the two correct responses for the two possible orientations of targets.

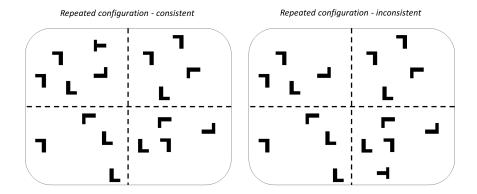


Figure 1
Schematic of the manipulation of target position in consistent and inconsistent trials of phase 2.
The dashed lines show the division of the stimuli into quadrants, but were not present in the task procedure.

Each trial commenced with a fixation cross presented in the center of the screen for 184 500 ms, which was then replaced immediately by the search configuration. Participants 185 searched for the target stimulus and responded with a left or right response depending on 186 its orientation. Reaction times (RTs) were recorded from the onset of the search 187 configuration. Following a valid response (c or n), the configuration was removed from the 188 screen. The ITI was 1000 ms. If participants made an incorrect response to the target 189 orientation, "INCORRECT RESPONSE" appeared in red in the center of the screen for 190 3000 ms, prior to the ITI. If participants did not respond within 6000 ms, "TIMEOUT -191 TOO SLOW" appeared in red in the center of the screen for 3000 ms, prior to the ITI. 192

Each block of eight trials contained each of the four different repeated configurations and four random configurations. These eight configurations could appear in any order with the constraint that the position of the target did not repeat across trials or across consecutive blocks.

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A rest break of 30 seconds was given every 80 trials. Trials started automatically after these breaks.

After 160 trials, prior to phase 2, participants were given an instruction screen
which detailed the arrow that would appear on the screen prior to the configuration. They
were able to ask any questions they had at this stage and then proceeded to phase 2. The
arrow appeared for 1000ms following the fixation cross, before the presentation of the
search configuration. The task was otherwise identical to that used in phase 1.

04 Results

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Our criterion for removing outlier data, at both the participant level and the trial 205 level, was 2.5 standard deviations above or below the mean of the sample. On average, 206 trials ended with a timeout on 1.97% of trials (SD = 2.53). Two participants had an 207 usually high proportion of timeouts and were removed from the analysis. The mean 208 accuracy of participants (not including timeout trials) was 98.10% (SD = 1.65%). One 200 participant had an unusually low proportion of accurate trials and was also removed. The 210 only participant deemed to be an outlier in terms of mean response time (hereafter RT) 211 was also excluded on the basis of the timeout criterion, noted above. 212

For the remaining twenty-eight participants we removed trials with a timeout and inaccurate trials, before removing outliers from the RT data. On average, the proportion of outliers removed was 3.03% (SD = 0.79%). Zero participants had an unusual proportion of trials removed as outlier RTs (greater than 2.5 SDs above the mean).

Figure 2 shows the RT data across the 10 epochs of the experiment. In phase 1 217 (epochs 1-5) a contextual cuing effect emerged, with faster responses to repeated over 218 random configurations. In phase 2, the presence of the guiding arrow led to a clear 219 reduction in the response times. For all participants, the mean RT across epochs 4 and 5 220 was higher than the mean RTs across epochs 6 and 7. Despite the clear evidence for the 221 processing of the endogenous cue, the underlying search configuration continued to play a 222 role in the guidance of attention, with faster response times for (consistent) repeated 223 configurations compared to random configurations. 224

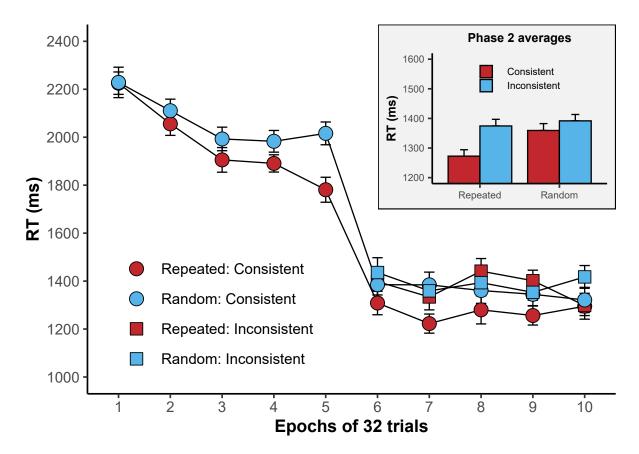


Figure 2

RT data for Experiment 1. The phase 2 averages across the four trial types are shown inset.

Within-subject error bars were computed by a process of normalising the RT data for the sample (Cousineau, 2005).

These data were analysed with a Bayesian ANOVA¹, using the BayesFactor::anovaBF() function in R. All analyses in this study used the default

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¹ The Bayesian analyses here follow the process outlined in Rouder et al. (2017). Briefly, we present the best fitting model evaluated against the null model, and then compare this fit to that of other models. Where the comparison of two models (i.e., A against B) reveals a Bayes Factor of greater than 3, this is taken as support for the components of model A that are not present in model B. Bayes Factors of less than 0.33 are taken as evidence in support of the equivalence of two models. Following Wetzels et al. (2011) we use the terms "substantial" (BF>3; BF<1/3), and "strong" (BF>10; BF<1/10) to reflect the levels of support for the results of the model comparisons.

parameters for the priors, which "places mass in reason-able ranges [of effect sizes] without 227 being overcommitted to any one point" (Rouder et al., 2017, p. 317). First taking the data 228 from phase 1 (epochs 1-5), there was strong support for the model containing the factors of 229 epoch and configuration (repeated vs. random), BF $_{10} = 2.2 \times 10^{12} \pm 0.74\%$. The addition 230 of the interaction term did not improve the model fit, BF = $0.47 \pm 1.86\%$, though there was 231 no evidence for the absence of the interaction. The best fitting model was a better fit than 232 the two models containing only one of the factors, smallest BF = $35.72 \pm 0.98\%$, providing 233 strong support for both the effects of configuration and epoch. Partial eta-squared (n_n^2) 234 effect sizes were calculated using effectsize::eta_squared, giving values of: 0.22 for the effect 235 of configuration; 0.39 for the effect of epoch; and 0.10 for the interaction effect. 236

A Bayesian ANOVA on the data from phase 2 (epochs 6-10) found strong support 237 for the model containing the factors of configuration (repeated vs. random) and target 238 position (consistent vs. inconsistent), $BF_{10} = 47.39 \pm 1.59\%$. The next best fitting model 239 contained these two factors and the interaction term, and was not a substantially worse fit 240 to the data, BF = $0.55 \pm 3.31\%$. The best fitting model (with factors of configuration and 241 target position, but no interaction) was a substantially better fit to the data than the 242 model containing only the factor of configuration BF = $21.26 \pm 1.73\%$ providing evidence 243 that RTs were faster on consistent than inconsistent trials. There was no evidence for a 244 difference between the best fitting model and the model containing only the factor of target 245 position, BF = $2.4 \pm 1.72\%$. The relevant effect sizes (n_p^2) were: 0.14 for the effect of 246 configuration; 0.22 for the effect of target position; and 0.14 for the interaction of these two 247 factors. 248

To further explore responses to the different trial types in phase 2, Bayesian t-tests
were run using BayesFactor::ttestBF (using the default Cauchy prior) for comparisons
between the repeated and random configurations, across the two target position conditions
(consistent and inconsistent). This revealed substantial support for a difference between
the response times on "repeated: consistent" trials and those on the respective random

trials (random: consistent), $BF_{10} = 4.14 \pm 0\%$. There was also substantial evidence to suggest there was no meaningful difference between the response times for the "repeated: inconsistent" trials and the respective random trials, $BF_{10} = 0.24 \pm 0.03\%$.

257 Discussion

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Experiment 1 sought to examine the consequence of an endogenous cue that 258 prompts top-down control of the search process on contextual cuing. In phase 1 we 259 established a robust contextual cuing effect. Following this, participants received 260 instruction that each trial would be preceded by an arrow stimulus that would signal the 261 side of the screen on which the target would appear. This cue was valid on all trials in 262 phase 2. Consistent with these instructions and the processing of this cue, we observed 263 substantially reduced search times in phase 2 compared to phase 1. The same set of 264 repeated configurations were presented in phase 2, but for half of the trials, the target was 265 relocated to the diagonally opposed quadrant of the screen. Therefore, on these "repeated 266 inconsistent" trials, the underlying configuration of distractors predicted the target in a 267 location that opposed that of the (valid) endogenous cue. Across this phase we observed 268 significant contextual cuing for the repeated consistent trials, demonstrating that the 269 underlying configuration of distractors continued to guide attention in the presence of the endogenous cue. However, the repeated inconsistent trials did not lead to an impairment in 271 response times relative to random trials, suggesting that the underlying configuration did not influence search on these trials.

Experiment 2

In Experiment 1 we demonstrated that an established effect of contextual cuing is
maintained even when attention is being guided by the presence of a valid endogenous cue.
That is, we found that the *performance* of an established search behaviour in contextual
cuing is not disrupted by concurrent top-down goals to guide attention in a controlled
manner. In Experiment 2 we wanted to explore whether the *learning* of the contextual cue

itself was affected by the presence of a valid endogenous cue. That is, does the presence of a valid endongenous cue, which leads to a controlled command of attention, limit the 281 development of a contextual cuing effect. To do this, we trained each participant on two 282 sets of repeating configurations. One of these sets was always presented in the presence of a 283 valid endogenous cue, while the other set was always presented in the absence of the 284 endogenous cue. The extent to which there is a "cue-competition" effect between the 285 endogenous cue and the contextual cues can be examined by comparing the contextual 286 cuing effect we observe for the two sets of configurations. Given the clear difference in RTs 287 we observed in Experiment 1 between the trials with the endogenous cue present and the 288 cue being absent, we anticipated the same difference in responding in Experiment 2. 289 Therefore we also included a second phase of Experiment 2 in which we removed the 290 endogenous cue entirely from the task. This second phase therefore allowed us to directly 291 compare the contextual cuing for the two sets of configurations when RTs were at a 292 comparable level. 293

"Cue-competition" effects have been examined previously in contextual cuing. Endo 294 and Takeda (2004) trained participants with a contextual cuing task composed of 295 distractor location configurations and repeating distractor identities. Their experiments 296 suggested that the stronger configural (spatial) cue out-competed the cue provided by the 297 distractor identities. Similarly, Kunar et al. (2014) found that when colour cues and 298 configural cues both predicted the target location, configural cues were dominant and 290 tended to overshadow the weaker colour cue. Beesley and Shanks (2012) looked at the 300 cue-interaction effects within a configuration of distractors. Participants were first trained 301 with half a configuration of repeating distractors that predicted the target (8 out of 16 302 distractors). In a later stage these distractors were paired with a new half-configuration, 303 such that the whole configuration now predicted the same target location. In contrast to 304 the predictions of the vast majority of models of contingency learning, learning about these 305 new predictive distractors was facilitated, rather than impaired in this second phase

(relative to a control condition). Thus, Beesley and Shanks (2012) found that
cue-competition was not observed within a configuration of equally predictive distractors.

Together these studies suggest that the spatial configuration serves as a strong cue for the
target and will out-compete non-configural cues for access to the learning mechanism. The
dominance of the configuration in these situations may therefore lead to the prediction that
the endogenous cue would not "block" the learning of the configuration in the current task.

Method

314 Participants

Thirty-four undergraduate students from Lancaster University were recruited (mean age = 20.74, SD = 5.29; 28 identified as female and 6 as male) via the Psychology Research Participation System in the Department of Psychology at Lancaster University, in return for the opportunity to use the recruitment system for their own research in future years.

319 Materials

Participants were tested in a quiet laboratory testing cubicle, with a standard PC and a 24" monitor set at a resolution of 1920 x 1080 pixels. Since the monitor was larger for this experiment, the dimensions of the presented stimuli had a proportional increase in size: Distractor stimuli were 11 mm square; the search grid was 240 mm square; the fixation cross was 6 mm square. In all other respects, the materials were the same as those detailed in Experiment 1.

Design

Four repeated configurations were created in an identical manner to those used in
Experiment 1. For each participant, two of these configurations were used for the
"cue-competition" condition, in which the arrow cue was presented before the
configuration, while two were used for the "control" condition (no arrow presented). As in
Experiment 1, the four repeated configurations were paired with unique target positions

from each of the four quadrants. We counterbalanced the use of the target quadrants across the factors of configuration type (repeated and random) and cue condition 333 (cue-competition and control). For half of the participants, targets in the top left and 334 bottom right were used for the repeated configurations presented with the arrow 335 (cue-competition) condition, with targets in the top right and bottom left used for 336 repeated configurations in the no-arrow (control) condition. For these participants, random 337 configurations presented with the arrow had targets in the top right and bottom left, and 338 random configurations without the arrow had targets in the top left and bottom right. For 339 the other half of the participants these assignments were reversed (repeated-arrow: 340 top-right and bottom-left; repeated-no arrow: top-left and bottom-right; random-arrow: top-left and bottom-right; random-no arrow: top-right and bottom-left).

Procedure Procedure

The procedure was the same as Experiment 1 with the following differences.

Participants received 320 trials in total. For the first 160 trials, the arrow was presented for
the relevant conditions. For the final 160 trials, the arrow was never presented. Rest breaks
were given every 60 trials.

Results

Our criteria for removing outlier data were identical to Experiment 1. On average, trials ended with a timeout on 2.13% of trials (SD = 1.83). Zero participants had an usually high proportion of timeouts. The mean accuracy of participants (not including timeout trials) was 95.85% (SD = 6.10%). One participant had an unusually low proportion of accurate trials and were removed from the sample. Zero participants were deemed to be an outlier in terms of mean RT.

For the remaining thirty-three participants we removed trials with a timeout and inaccurate trials, before removing outliers from the RT data. On average, the proportion of outliers removed was 2.81% (SD = 1.04%). One participant had an unusual proportion of

trials removed as outlier RTs and were not included in the final analysis.

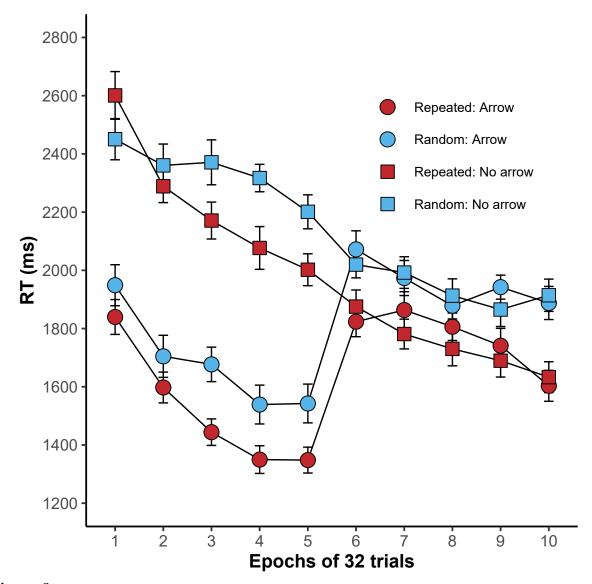


Figure 3

RT data for Experiment 2. Error bars show standard error of the mean on normalised data.

Figure 3 shows the RT data across the 10 epochs of the experiment. Contextual cuing emerged rapidly in both the arrow and no-arrow conditions, with little suggestion that the CC effect was different in the two conditions. The phase 1 data were explored with a Bayesian ANOVA, which revealed that the best fitting model contained the factors of epoch, configuration (repeated vs. random), and endogenous cue (arrow present

vs. arrow absent), with no interaction terms, $BF_{10} = 7.3 \times 10^{100} \pm 1.82\%$. The next best 364 fitting model contained all three factors and the interaction of epoch and configuration, 365 $\mathrm{BF}_{10} = 5.2 \times 10^{100} \pm 3.02\%$, and this model was not a substantially worse fit to the data, 366 $BF = 0.71 \pm 3.53\%$. All other models were substantially worse fits than the best fitting 367 model, largest BF = $0.26 \pm 3.24\%$. Importantly, the interaction term between the factors 368 of endogenous cue and configuration did not improve the fit of the model, providing 369 substantial support for the absence of this interaction, BF = $0.18 \pm 2.17\%$. The relevant 370 effect sizes (n_p^2) were: 0.02 for the effect of epoch; 0.40 for the effect of configuration; 0.85 371 for the effect of endogenous cue; 0.12 for the interaction effect between configuration and 372 epoch; and 0.02 for the interaction between configuration and endogenous cue. 373

When the endogenous cue was removed in the second half of the experiment, RTs 374 were equivalent across the two conditions. An effect of configuration was seen for both 375 cuing conditions, with little discernible difference between the size of the cuing effects. We 376 conducted a Bayesian ANOVA with factors of epoch, configuration and endogenous cue 377 condition (arrow vs. no-arrow). The best fitting model was that with just the factors of 378 epoch and configuration with no interaction between the factors, BF $_{10}$ = 9.6 \times 10^{14} \pm 370 0.8%. There was substantial support for this model over the next best fitting model, BF = 380 $8.97 \pm 1.68\%$. To examine the interaction of the configuration and endogenous cue factors, 381 we compared the model containing those two factors to the model containing the two 382 factors plus the interaction of configuration and endogenous cue, which revealed substantial 383 support for the absence of an interaction, BF = $0.12 \pm 3.63\%$. The relevant effect sizes 384 (n_p^2) were: 0.62 for the effect of configuration; and 0.25 for the effect of epoch. 385

To provide further support for the absence of the interaction between the factors of configuration type and endogenous cue, the data from across the experiment (epochs 1-10) were analysed with a Bayesian ANOVA with only the factors of configuration and endogenous cue. The best fitting model was that with the two factors and no interaction, $BF_{10} = 3.7 \times 10^{51} \pm 1.03\%$. The addition of the interaction term did not strengthen the

model, with substantial evidence for the absence of the interaction, BF = $0.09 \pm 1.82\%$.

The relevant effect sizes (n_p^2) were: 0.77 for the effect of the endogenous cue; and 0.61 for
the effect of configuration.

394 Discussion

Experiment 2 sought to examine whether the presence of a valid endogenous cue 395 would impair the acquisition of a contextual cuing effect. In the first phase, two sets of 396 configurations were trained, one of which was always presented in the presence of the 397 endogenous cue, and one set which was presented without the endogenous cue. Overall 398 there was considerable evidence that the cue was processed and acted upon, as response 399 times to the target were much faster on cued trials. However, there was no evidence to 400 suggest that this initial guidance of attention impaired the acquisition of the configurations 401 on those trials. Furthermore, when the endogenous cue was never presented in the final 402 phase of the experiment, the size of the contextual cuing effect was equivalent between the 403 two sets of configurations; the Bayesian analyses found support for the equivalence of these 404 CC effects. 405

The lack of competition effects seen in Experiment 2 are at odds with some findings in the CC literature (i.e., Endo & Takeda, 2004; Kunar et al., 2014), where competition has been seen by more dominant or salient features of the displays. Instead, the findings point towards a more automatic nature to contextual cuing, whereby associations form ubiquitously, so long as they receive the focus of attention at some point within the search process (e.g., Beesley & Shanks, 2012).

Taken together with the findings of Experiment 1, these data suggest that when
attention is cued in an endogenous manner, the underlying search configuration will still
play a significant role in guiding search for the target. Since the endogenous cue appears at
the point of fixation, the most plausible interaction of these processes is that the guidance
by the endogenous cue is followed by the guidance by the repeated context. The

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equivalence of the CC effects in the two conditions (cued and uncued) would therefore 417 suggest that the guidance by the context was driven largely (or perhaps entirely) by the 418 distractors that appear close to the target. Accordingly, the longer search times in the 419 uncued condition suggest that more distractors are processed in this condition, but that 420 the influence of the repeated distractors on attentional guidance may be limited to those 421 occurring later in the search process, and therefore those nearer to the target. 422 Alternatively, the interaction of these two processes (endogenous and configuration-driven 423 attention) need not interact in this order. It is at least possible that the configuration is 424 processed rapidly at the onset of the trial, before the effects of the endogenous cue on 425 attention are observed. If this is the case, then those repeated distractors that influence 426 search (producing the CC effect) need not be localised around the target. Experiment 3 427 provided a test of these two possible accounts. 428

Experiment 3

Existing data from studies of contextual cuing has pointed towards a localised 430 learning effect for repeated configurations, with those distractors closest to the target being 431 preferentially weighted in the learning process over those located further from the target. 432 For example, Olson and Chun (2002) trained participants with three sets of repeating 433 configurations that differed in terms of which distractors repeated across trials. For one set, the entire global context (all of the distractors) repeated, while for the other two sets 435 only the short-range (those close to the target) or the long-range distractors (those far from 436 the target) repeated across trials. They found no difference between the CC effect in the 437 short-range and global configurations, while the CC effect was not significant for the 438 long-range context. Similar results have been shown by Brady and Chun (2007) which led 439 to the development of the spatial constraints model of contextual cuing, in which 440 distractor-target associations occurring in close proximal space are weighted more heavily 441 in the learning process (over those occurring across greater spatial distance).

It is important to consider how the bias towards local learning may interact with 443 the attentional scanning process during contextual cuing. The analysis of eye-movements 444 during contextual cuing tasks (Beesley et al., 2018; Tseng & Li, 2004) has revealed a 445 characteristic scanning pattern comprising two phases: search initially occurs in a 446 seemingly random manner, as the eyes move between distractors in the central region of 447 the distractor field, before then moving in a more directed manner towards the target 448 position. Contextual cuing appears to result from a cessation of the first (random) search 449 phase at an earlier time point in the entire search process, such that processing of repeated 450 distractors will, on average, result in fewer fixations. With respect to the current study, in 451 Experiments 1 and 2 we have initially directed attention towards the side of the screen that 452 contains the target on cued trials. This may bring about an early cessation of the first 453 phase of the search process. From here, however, it seems that search is still facilitated by 454 the repetition of the context. 455

To test this characterisation of the interaction between the endogenous cue and the 456 repeated context, we exposed participants to the same procedure as used in phase 1 of 457 Experiment 1, which establishes a contextual cuing effect prior to the use of the 458 endogenous cue. In a second phase we then presented the endogenous cue on every trial (as 459 in Experiment 1), but we manipulated the presence of the repeated distractors within the 460 configurations. For each repeated configuration we created two variations: in the 461 "proximal" configurations, only the distractors in the quadrant containing the target match 462 those from the full repeated configuration, while the distractors in the other three 463 quadrants were randomly arranged on each trial; in the "distal" configurations, the 464 distractors closest to the target were randomised, while the distractors in the other three 465 quadrants were the same as those in the full repeated configuration. During this phase we 466 also presented fully repeated configurations and fully randomised configurations. 467 Comparison of the response times across these four trial types will allow us to determine the contribution of proximal and distal distractors to the CC effect when attention is cued

endogenously.

471 Method

472 Participants

Forty-two undergraduate students from Lancaster University were recruited (mean age = 18.64, SD = 2.84; 28 identified as female and 14 as male) via the Psychology
Research Participation System in the Department of Psychology at Lancaster University, in return for the opportunity to use the recruitment system for their own research in future years.

478 Materials

All materials, including stimuli and testing environment were identical to
Experiment 2.

Design

The design of phase 1 was identical to Experiment 1, with four repeated 482 configurations created and presented with random configurations during this phase. For 483 phase 2, each of the four configurations was manipulated to create two alternative 484 conditions. In the "Repeated distal" condition, the four distractors in the target quadrant 485 were randomly arranged on each trial, while the 12 distractors in the other three quadrants 486 were presented in the same positions as had been trained in phase 1. Thus, slower response 487 times for this condition (compared to the fully repeated configurations) would indicate the 488 extent to which participants CC was governed by the distractors closest to the target. For the "Repeated proximal" condition, the four distractors in the target quadrant were presented in the same positions as had been trained in phase 1, while the 12 distractors in the other three quadrants were randomly arranged on each trial. Thus, slower response times for this condition (compared to the fully repeated configurations) would indicate the extent to which CC was governed by the distractors further from the target. Comparison

of the RTs for these different configurations with those of the random configurations would allow for the assessment of whether these subsets of distractors had *any* contribution to the CC effect that had developed during phase 1.

498 Procedure

The procedure was identical to Experiment 1.

500 Results

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Our criteria for removing outlier data were identical to Experiment 1. On average, trials ended with a timeout on 2.81% (SD = 2.25) of trials . Two participants had an usually high proportion of timeouts and were removed from the sample. The mean accuracy of participants (not including timeout trials) was 96.09% (SD = 8.57%). Two participants that had an unusually low proportion of accurate trials and were also removed. Zero participants were deemed to be an outlier in terms of mean RT.

For the remaining thirty-eight participants we removed trials with a timeout and inaccurate trials, before removing outliers from the RT data. On average, the proportion of outliers removed was 3.17% (SD = 0.71%). Zero participants had an unusual proportion of trials removed as outlier RTs.

Figure 4 (main panel) shows the RT data across the 10 epochs of Experiment 3. As 511 in Experiment 1, contextual cuing was readily established in phase 1. These data were 512 subjected to a Bayesian ANOVA which revealed that the best fitting model contained the 513 factors of configuration (repeated vs. random) and epoch, and an interaction between those 514 factors, $BF_{10} = 5.5 \times 10^{24} \pm 1.84\%$. However, the model without the interaction provided 515 a strong fit to the data, $BF_{10} = 5.1 \times 10^{24} \pm 1.09\%$, and a comparison between the two 516 models did not find any evidence in support of the interaction term, BF = $0.94 \pm 2.14\%$. 517 There was strong support for the best fitting model over the remaining models, smallest 518 $BF = 3987.48 \pm 1.86\%$, providing strong support for the factors of epoch and 519

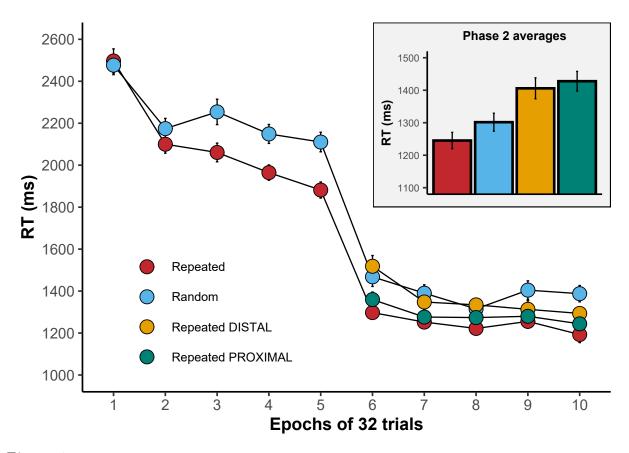


Figure 4 RT data for Experiment 3. Error bars show standard error of the mean on normalised data.

configuration. The relevant effect sizes (n_p^2) were: 0.38 for the effect of the endogenous cue; 520 and 0.47 for the effect of configuration; and 0.08 for the interaction of these two factors. 521

The response times decreased significantly with the presentation of the valid 522 endogenous cue in phase 2. Response times to the fully repeated configurations were 523 somewhat comparable to those when just the proximal repeated distractors were present. Response times for the distal repeated distractors appeared to be slower and comparable to 525 the fully random configurations. The phase 2 data were subjected to a Bayesian ANOVA 526 which found that the best fitting model contained the factors of configuration and epoch but no interaction between the factors, BF $_{10}$ = 1.4 × 10 14 ± 0.47%. This model provided a 528 superior fit to the data compared to the next best fitting model that included the two 529

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factors and the interaction term, BF = $123.41 \pm 0.66\%$, providing strong support for the contribution of the two factors and the absence of an interaction between the two factors.

The relevant effect sizes (n_p^2) were: 0.37 for the effect of configuration; and 0.16 for the effect of epoch.

The inset graph in Figure 4 shows the mean RTs to the four types of configuration, 534 averaged across the 5 epochs of phase 2. To explore the differences in response times, 535 Bayesian t-tests were run for all pairwise comparisons. The response times to repeated and repeated-proximal configurations were both faster than those to random configurations, 537 smallest BF₁₀ = 10313.81 \pm 0%. In contrast, there was no evidence that the response times 538 to repeated-distal configurations were different from those to random configurations, BF₁₀ 539 $=0.39\pm0.04\%$. Response times to repeated configurations were faster than those to 540 repeated-proximal configurations, $BF_{10} = 4.67 \pm 0\%$. Response times to repeated-proximal 541 configurations were faster than those to repeated-distal configurations, $BF_{10} = 31.88 \pm 0\%$. 542

543 Discussion

Experiment 3 explored the localisation of the distractors driving contextual cuing when attention is guided by an endogenous cue. As expected, there was substantial 545 evidence that contextual cuing was present when the distractors close to the target were 546 maintained, but not when these distractors were randomly arranged. These data suggest a 547 particular order to the interplay between the two drivers of attention: initially attention is 548 guided by the endogenous cue towards one half of the screen, and then search is refined by 549 the presence of the valid configural cues (the repeated distractors). Like in Experiment 1, 550 the phase 2 data demonstrate the resilience of the CC effect to changes in the search 551 process. Despite visual search never commencing in a cued manner during the initial 552 acquisition period of phase 1, a CC effect was readily observed in phase 2. Thus it seems 553 that the stored representations of configurations surrounding target positions are very 554 flexibly deployed in visual search. Notably the fully repeated configurations exerted more 555

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of a benefit on search than those containing only the proximal distractors, suggesting that
the repeating distractors beyond the target quadrant have some (but possibly lesser)
influence on search (Brady & Chun, 2007).

These data lend support to the notion that the effect of the repeated configuration is a late process within visual search, and that each trial commences with a random search process that is not guided by the repeated configuration (Beesley et al., 2018; Tseng & Li, 2004). In some ways, these findings represent a paradox of CC: the cuing effect occurs almost at the point at which target detection has been made. One interpretation would be that this exemplifies the role of spatial contiguity in the formation of visual associations (Renaux et al., 2017). Alternatively, it provides support for the proposed "decision threshold" accounts of CC (Kunar et al., 2007; Sewell et al., 2018), which posit that the repeated distractors close to the target ensure a reduced threshold for target detection, resulting in faster response times.

General Discussion

Three experiments explored the impact of a central endogenous cue on the 570 contextual cuing of visual search. In Experiment 1, having established a contextual cuing 571 effect, each trial was preceded by an central endogenous cue of attention in the form of an 572 arrow, directing attention towards the side of the screen in which the target was positioned 573 (this arrow cue was always valid in each of the three experiments). Despite participants 574 clearly using this cue, visual search was still facilitated by the presence of the repeating 575 pattern of visual search. This experiment demonstrated that, once acquired, the activation 576 of the memory representation and its impact on performance of visual search remains 577 intact in the presence of a top-down instruction to guide attention. Experiment 2 examined 578 the storage of these contextual representations, and whether these were impaired by an 579 endogenous cue guiding search. We found equivalent levels of contextual cuing for two sets 580 of configurations, one of which was paired with the cue and one which was not. Together, 581

these two experiments suggest a seamless interplay between these two factors governing 582 attention in visual search: the endogenous cue initially guides attention and the repeated 583 configuration continues to refine and guide attention towards a fixation on the target. In 584 Experiment 3 we therefore explored whether the localised distractors around the target 585 were sufficient to generate CC following the guidance by the endogenous cue. Indeed, there 586 was a significant CC effect in the case of the proximal distractors, albeit slightly weaker 587 than that generated by the entire repeated configuration. Importantly, those repeated 588 configurations that did not contain the proximal distractors failed to generate a CC effect, 589 suggesting that the proximal distractors play a crucial role in search following the guidance 590 of attention by the endogenous cue. 591

The effect of CC on visual search has frequently been characterised as an automatic 592 influence on behaviour (e.g., Chun & Jiang, 1998; Chun & Nakayama, 2000; Geyer et al., 593 2021). This characterisation of CC comes from multiple aspects of the observed effect. 594 Updating of the associations is somewhat slow and seemingly inflexible to changes in the 595 acquired associations (Makovski & Jiang, 2011; Manginelli & Pollmann, 2009; e.g., Zellin et 596 al., 2013), and therefore perhaps reflects a habitual form of behaviour. In addition, 597 contextual cuing has frequently been observed in the absence of above-chance recognition 598 memory for the repeating search configurations (e.g., Colagiuri & Livesey, 2016), which 599 suggests a non-conscious, automatically evoked form of behaviour. Despite this persistent 600 characterisation, the automaticity (or controllability) of CC has rarely been directly tested 601 in the literature. To our knowledge, only the experiments of Luque and colleagues (Luque 602 et al., 2017; Luque et al., 2021) have directly assessed this aspect of CC, by placing the 603 influence of the configuration in competition with top-down goals in the task. Their 604 findings supported the conclusion that CC performance can be controlled and will not 605 guide search for the target when another aspect of the task governs attentional control. In 606 the current study, the repeated configurations continued to have an influence on search performance even when attention had been guided by the endogenous cue. These results

are therefore somewhat at odds with the conclusions of Luque and colleagues (Luque et al., 2017; Luque et al., 2021).

To what extent is this behaviour best characterised as "automatic" in nature? 611 Arguably the clearest demonstration of an automatic effect of a stimulus on behaviour is 612 when the associated behaviour is elicited even when it is counter-productive to the current 613 goals (Moors & De Houwer, 2006). Such a test was constructed in the repeated 614 inconsistent trials of Experiment 1, in which the repeated configuration was associated 615 with a target that was previously located in a position on the opposite side of the screen to the direction indicated by the endogenous cue. If the repeated configuration had an effect 617 on behaviour on these trials, we would have expected to see slower response times compared to random trials. This was not the case: response times were equivalent in these 619 two conditions. As such it is hard to claim here that the configuration is having an 620 automatic effect on behaviour, according to this strict characterisation of such an effect. 621 Nevertheless, the experiments here reveal an interplay between top-down processes and 622 stimulus driven effects on attention in CC. 623

The current data reveal that the influence of repeated contexts has a relatively late 624 control on behaviour in visual search. Previous analysis of eye-movements during CC 625 (Beesley et al., 2018; Tseng & Li, 2004) has shown that contextual cuing (and visual search 626 more generally) has two characteristic components. The first of these is an inefficient 627 search process where search fails to move towards the target in trials with more fixations. This is followed by a phase in which monotonic, positive increments are made toward the 629 target position in the final 3 to 4 fixations. CC reduces the frequency of trials with the initial (random) search period (there are more of such trials for random configurations and 631 fewer for repeated configurations). Thus, the effect of the endogenous central cue in the 632 current study is to eliminate, or considerably reduce, the engagement with this first phase 633 of the search process. The results of this study strongly imply that the positive associative 634 information in the repeating configurations is extracted in the final stages of search and is

localised to the target. This is true both in terms of the performance of an acquired
configuration (Experiments 1 and 3) and the acquisition of the representation for that
configuration (Experiment 2). Perhaps paradoxically, the benefit of repeated configurations
in search occurs shortly before the target is fixated. These data therefore support the view
that late-stage "response threshold" processes may play an important role in the CC of
visual search (Sewell et al., 2018). Notably, the results of Experiment 2 show that the
curtailing of the initial (random) search process, which significantly reduces search times,
does not limit the development of an adequate contextual cuing effect.

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