

Visual search efficiency is modulated by symmetry type and texture regularity

Rachel Moreau

Department of Psychology,
York University, Toronto, Ontario, Canada
Centre for Vision Research, York University,
Toronto, Ontario, Canada



Nihan Alp

Department of Psychology,
Sabancı University, Tuzla, Istanbul, Turkey



Alasdair D. F. Clarke

Department of Psychology, University of Essex,
Colchester, United Kingdom



Erez Freud

Department of Psychology,
York University, Toronto, Ontario, Canada
Centre for Vision Research, York University,
Toronto, Ontario, Canada



Peter J. Kohler

Department of Psychology,
York University, Toronto, Ontario, Canada
Centre for Vision Research, York University,
Toronto, Ontario, Canada



More than a century of vision research has identified symmetry as a fundamental cue, which aids the visual system in making inferences about objects and surfaces in natural scenes. Most studies have focused on one type of symmetry, reflection, presented at a single image location. However, the visual system responds strongly to other types of symmetries and to symmetries that are repeated across the image plane to form textures. Here we use a visual search paradigm with arrays of repeating unit cells that contained either reflection or rotation symmetries but were otherwise matched. Participants were asked to report the presence of a target tile without symmetry. When unit cells tile the plane without gaps, they form regular textures. We manipulated texture regularity by introducing jittered gaps between unit cells. This paradigm lets us investigate the effect of symmetry type and texture regularity on visual search efficiency. Based on previous findings suggesting an advantage for reflection in visual processing, we hypothesized that search would be more efficient for reflection than rotation. We further hypothesized that regular textures would be processed more efficiently. We found independent effects of symmetry type and regularity on search efficiency that confirmed both hypotheses: Visual search was more

efficient for textures with reflection symmetry and more efficient for regular textures. This provides additional support for the perceptual advantage of reflection in the context of visual search and provides important new evidence in favor of visual mechanisms specialized for processing symmetries in regular textures.

Introduction

As we move through the world, the brain generates our visual experience by rapidly processing a constant stream of visual stimuli. Despite the apparent effortlessness of vision, this process is highly complex. Seminal perception research proposed that visual processing is simplified through a set of fundamental gestalts that provide structural limitations on the interpretation of visual stimuli (Wertheimer, 1938). In the current study, we investigate how one of these fundamental gestalts—symmetry—contributes to the perception of textures and how this impacts the efficiency of visual processing.

Citation: Moreau, R., Alp, N., Clarke, A. D. F., Freud, E., & Kohler, P. J. (2025). Visual search efficiency is modulated by symmetry type and texture regularity. *Journal of Vision*, 25(6):7, 1–13, <https://doi.org/10.1167/jov.25.6.7>.

<https://doi.org/10.1167/jov.25.6.7>

Received June 26, 2024; published May 13, 2025

ISSN 1534-7362 Copyright 2025 The Authors



Symmetries are prevalent in the natural world and can be found in man-made objects throughout human history (Jablan, 2002). Multiple studies have identified a key role for symmetry in scene and object perception (Bertamini, Silvanto, Norcia, Makin & Wagemans, 2018), contributing to behaviors as fundamental as shape perception (Bahnsen, 1928; Machilsen, Pauwels, & Wagemans, 2009) and as sophisticated as judgments about facial attractiveness (Grammer & Thornhill, 1994). Much of this literature has focused on reflection or mirror symmetry, but reflection is only one of four possible symmetry types in the two-dimensional Euclidean plane, with the others being: Rotation, translation, and glide reflection. Although reflection can be seen in the bilateral bodies of many animals and is especially behaviorally relevant for human faces, there are examples of every symmetry type in nature (e.g., rotation symmetry in flower petals, honeycombs, and snowflakes).

This raises the question: How does the visual system process these various symmetry types, and how do they differ from previous findings with reflection symmetry? Since the beginning of symmetry as a research topic in vision research, reflection has been considered unique among the symmetry types (Mach, 1897, eng. translation 1959). Psychophysical studies show that reflection symmetry can be detected pre-attentively, requires less cognitive resources, and allows for faster reaction time than rotation and translation (Wagemans 1995, Wagemans 1997, Olivers & Van Der Helm, 1998; Treder, 2010; Bertamini & Makin, 2014). It has been suggested that the advantage of reflection might be a result of evolutionary pressures to optimize encoding of behaviorally relevant stimuli that have reflection symmetry, such as faces (Grammer & Thornhill, 1994).

Most studies on the role of symmetry in visual behavior have considered one or two axes of symmetry centered on a single location in the image, consistent with the way symmetries would most likely occur over objects in the natural world (Bertamini et al., 2018). However, symmetries can also be found in regular textures known as the wallpaper groups – a set of 17 unique combinations of the four fundamental symmetry types (Fedorov, 1891; Polya, 1924; Liu, Hel-Or, Kaplan, & Van Gool, 2010). The wallpaper name is apt because the textures resemble a Victorian wallpaper or rug (see Figure 1). Regular and near-regular textures are abundant in natural and man-made environments (Liu, Lin & Hayes, 2004), and symmetries in regular textures generate strong responses in the visual cortex of humans (Kohler, Clarke, Yakovleva, Liu, & Norcia, 2016; Alp, Kohler, Kogo, Wagemans, & Norcia, 2018; Kohler, Cottreau, & Norcia, 2018; Kohler & Clarke, 2021) and other primates (Audurier et al., 2022).

The growing literature on wallpaper groups shows that when embedded in regular textures, each of the different symmetry types can give rise to

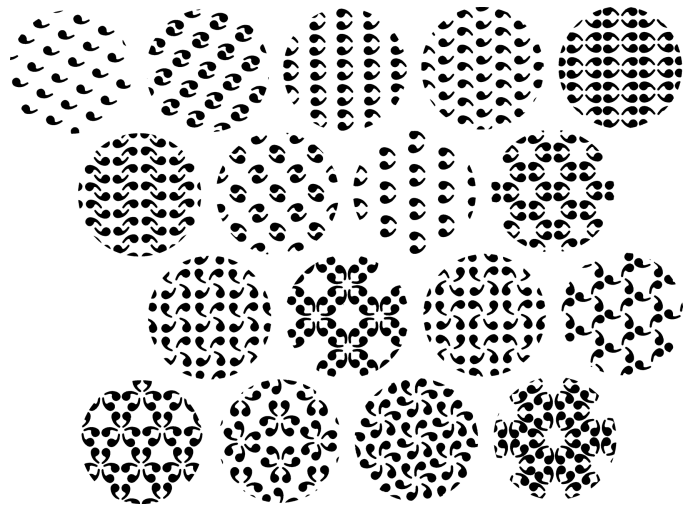


Figure 1. The 17 wallpapers rendered with a comma-like symbol as the repeating element. Illustration based on Wade (1993).

reliable responses in the visual cortex. The behavioral consequences of this have yet to be explored. The current study seeks to address that gap in the literature by investigating the efficiency of processing reflection and rotation symmetries when these symmetries are presented in regular textures and when they are not. This will provide valuable information about how the human visual system handles complex representations of symmetry and regularity, and how these striking patterns may contribute to the perception of natural scenes.

We addressed these questions using a visual search task. Visual search has been used to probe the extent to which a given cognitive process takes place in a serial or parallel fashion. Reaction time and accuracy are measured as participants search for a target. Typically, the target is either presented among distractors or hidden in noise. If information is being processed serially, the observer has to scan through each individual array element until the target is found. Thus adding more distractors to the array will result in longer reaction times, lower accuracy, or both. On the other hand, if information is processed in parallel, the target will “pop-out,” resulting in reaction times and accuracy that are constant across array sizes (Treisman & Gelade, 1980). Previous studies have used visual search to dissociate parallel or serial processing of visual properties such as color and orientation. It is an effective way of differentiating visual properties based on the cognitive resources required for processing (Cavanagh, Arguin, Treisman, 1990; Bundesen, Kyllingsbæk, Larsen, 2003; Kyllingsbæk & Bundesen, 2007). Serial and parallel processing are often presented in binary fashion as two qualitatively distinct types of cognitive processing. However, it is likely more realistic to conceptualize them as endpoints along a spectrum

(Wolfe, 1998; Wolfe, 2016). That is, cognitive processes may operate as strictly serial or parallel but can also fall anywhere between the two. In the current study, we therefore compare conditions in terms of “more serial” or “more parallel.”

Visual search has been used to study symmetry in two main ways: Through within-item symmetry and whole-array symmetry. In within-item studies, individual items in the search array either do or do not have internal symmetry, and participants are asked to either find a symmetrical target among asymmetrical distractors, or an asymmetrical target among symmetrical distractors. This approach has provided some evidence of parallel processing of within-item symmetry (Javadnia & Ruddock, 1988), whereas later work with more diverse and well-controlled stimuli suggested that symmetry detection is a more serial process that requires attention (Olivers & Van Der Helm, 1998). Studies that use the whole-array approach to symmetry in visual search use targets and distractors that do not differ in within-item symmetry but are arranged such that they either do or do not form symmetrical textures across multiple array items. The first to do this was Wolfe and Friedman-Hill (1992), who used oriented lines that were arranged to form symmetrical textures across the search array. Participants were asked to find a target which was oriented such that it disrupted the symmetry of the array. They found that when the distractor arrays were arranged in terms of vertical (mirror) symmetry, finding the target was more efficient than when distractor arrays were arranged in terms of oblique (rotation) symmetry (Wolfe & Friedman-Hill, 1992). Symmetries across the whole array have also been investigated in the context of inter-item symmetries between the target and distractors, and here the findings indicate that when reflection symmetry exists between the target and the distractor, search efficiency is diminished (van Zoest, Giesbrecht, Enns, & Kingstone, 2006). The effect was stronger for vertical than horizontal reflection and is likely the result of shapes that are identical when reflected over the vertical axis being considered as more similar (van Zoest et al., 2006). The current study takes inspiration from both approaches and uses a design that manipulates both within-item and whole-array symmetry in a highly controlled manner.

We used the visual search task to address two research questions: First, how within-item symmetry type (reflection vs. rotation) influences the efficiency of visual processing; Second, how texture regularity, a whole-array property, influences the efficiency of visual processing. We contrasted two wallpaper groups: PMM (Figure 2B) and P4 (Figure 2A). PMM contains fourfold (90°) reflection and twofold (180°) rotation centered at the intersection of the reflection axes. This means that if the PMM pattern is reflected across a horizontal or vertical axis, the resulting pattern is identical to the original. Additionally, a 180° rotation

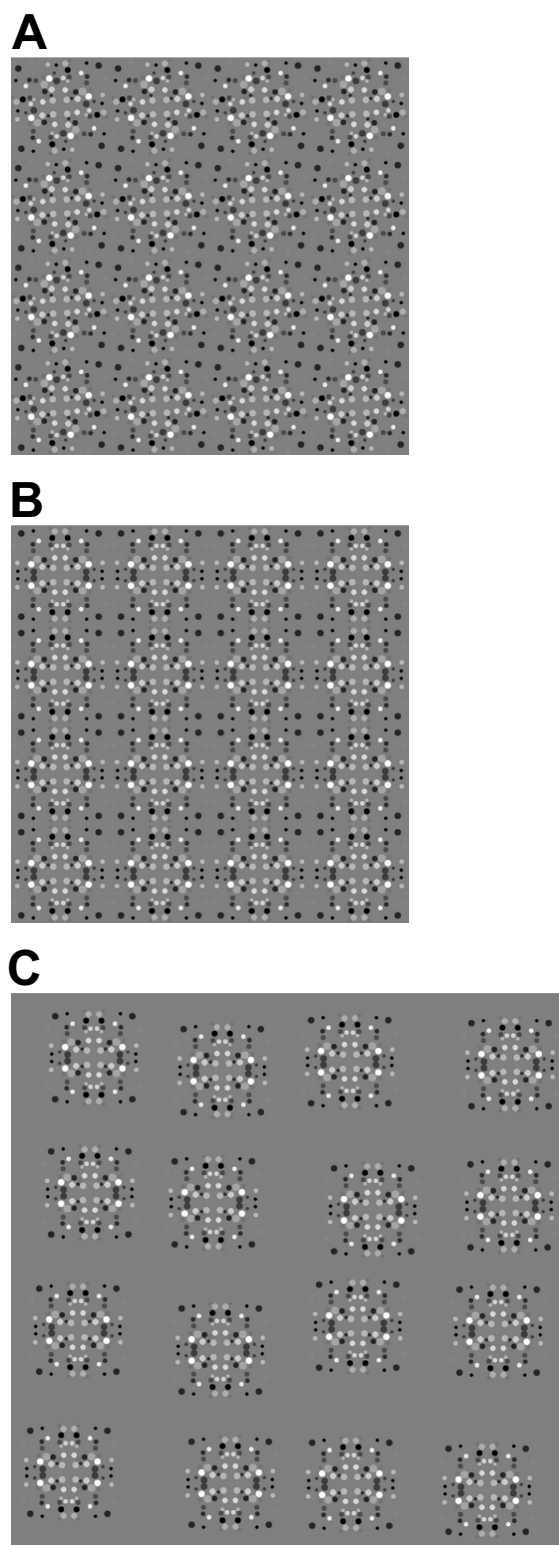


Figure 2. Examples of 4×4 “no jitter” search arrays based on wallpaper group P4 (A) and PMM (B). An example of a PMM in the “jitter” condition is shown in C. Across all stimuli the individual unit cells were the same size. In the jitter stimuli, the overall array was larger, but the unit cell size was the same. For all jitter conditions, the unit cells were presented on a 50% gray background, large enough to contain all possible unit cell positions, to avoid discontinuities along the search array edge.

around its center also preserves the pattern. In contrast, P4 contains fourfold (90°) and twofold rotation but no reflection. The P4 pattern can be rotated by 90° or 180° degrees and be identical to the original, but there is no reflection that preserves the pattern. Both groups (and, in fact, all wallpaper groups) are textures in which a unit cell is repeated on a lattice to tile the image plane, which gives rise to translation symmetry. Wallpaper groups can have various lattice shapes, but the groups used here both had the same lattice and differ only in terms of the symmetries within the lattice. The target stimulus was a random dot pattern that contained no internal symmetry and replaced one of the repeating unit cells when the target was present. The choice of using PMM and P4 as stimuli makes it possible to easily generate exemplars that belong to one group or the other through a very simple image-level operation (see Figure 4 and the Stimuli section of the Methods) and thus manipulate symmetry while controlling for every other image-level attribute (e.g., spatial frequency, contrast, shape of repeating region). Another advantage of these stimuli is that for both PMM and P4, the unit cell contains four repeated and transformed fundamental regions. Our manipulation of wallpaper group allows us to investigate the effect of within-item symmetry. To investigate the effect of regularity across the arrays, we developed “jitter stimuli”, in which gaps were introduced between the repeating unit cells, and unit cell positions were jittered, such that the regularity of the textures was disrupted (see Figure 2). Regularity is by definition a whole-array property and is therefore related to manipulations of whole-array symmetry in previous studies. This means that our design makes it possible to separately measure the effects of within-item symmetry and whole-array regularity. As for the non-jittered displays, the target stimulus was, again, a random dot pattern that replaces one of the array unit cells. Because all wallpaper groups contain translation symmetry, the jitter manipulation also disrupts translation symmetry across the entire pattern. Local translation is preserved, however, between individual unit cells in the pattern. So, although we cannot entirely dissociate the effect of translation symmetry from the effect of regularity, we consider it reasonable to refer to the jitter condition as manipulating regularity and will use that terminology throughout the article.

It is worth noting that across all of our experiments, the visual search task is effectively a search for the absence of symmetry. We will discuss the implications of this in more detail in our General Discussion. Our manipulations of the two dimensions of interest: Symmetry type (PMM vs. P4) and texture regularity (no jitter vs. jitter between unit cells), give rise to a 2×2 design across four visual search experiments, with four array sizes per experiment.

We used the slope of the linear search function to describe how reaction time and sensitivity (d') change with larger search arrays as a measure of processing efficiency. Steeper search function slopes indicate that processing is more serial, whereas shallower slopes indicate more parallel processing, with perfectly flat search function indicating fully parallel processing.

Based on previous results indicating that reflection is processed more efficiently than other types of symmetry, our first hypothesis was that we would find more parallel processing for reflection than for rotation. Our second hypothesis was that regular textures would be processed more efficiently than non-regular textures, because the target would disrupt regularity and perhaps lead to a form of pop-out effect. Our results support the first hypothesis: Across both jittered and un-jittered conditions, reflection symmetries produced more parallel processing. We also confirmed our second hypothesis: Search was more efficient for regular textures across both types of symmetry. There were no interactions between symmetry type and regularity, suggesting that the effect of regularity was independent of the effect of symmetry type. These findings add new evidence to the literature on differential processing of reflection and other types of symmetry, and demonstrate a novel processing advantage for regular textures.

Methods

Stimuli

The stimuli were square arrays of unit cells. Each unit cell was created based on a random dot pattern called a *fundamental region*, that was repeated to form a 2×2 matrix (see Figure 3). Two different sets of transformations were applied to the fundamental region inside the unit cell. In PMM unit cells, the fundamental region is reflected along the vertical axis and then again reflected along the horizontal axis. This produces reflection symmetry along both axes. In P4 unit cells, the fundamental region is rotated 90° clockwise, starting with 0° in the upper left quadrant, then 90° to the upper right, 180° in the lower right, and 270° to the lower left. This produces a fourfold rotation centered at the center of the unit cell (see Figure 3). Importantly, all four experiments used the same random dot patterns as fundamental regions, meaning that image-level properties were matched across conditions. When unit cells are repeated on a lattice to tile the plane, they form regular textures known as wallpaper groups—PMM unit cells produce wallpaper group PMM, and P4 unit cells produce group P4. We used 10 different fundamental regions to create 10 exemplars of each wallpaper type.

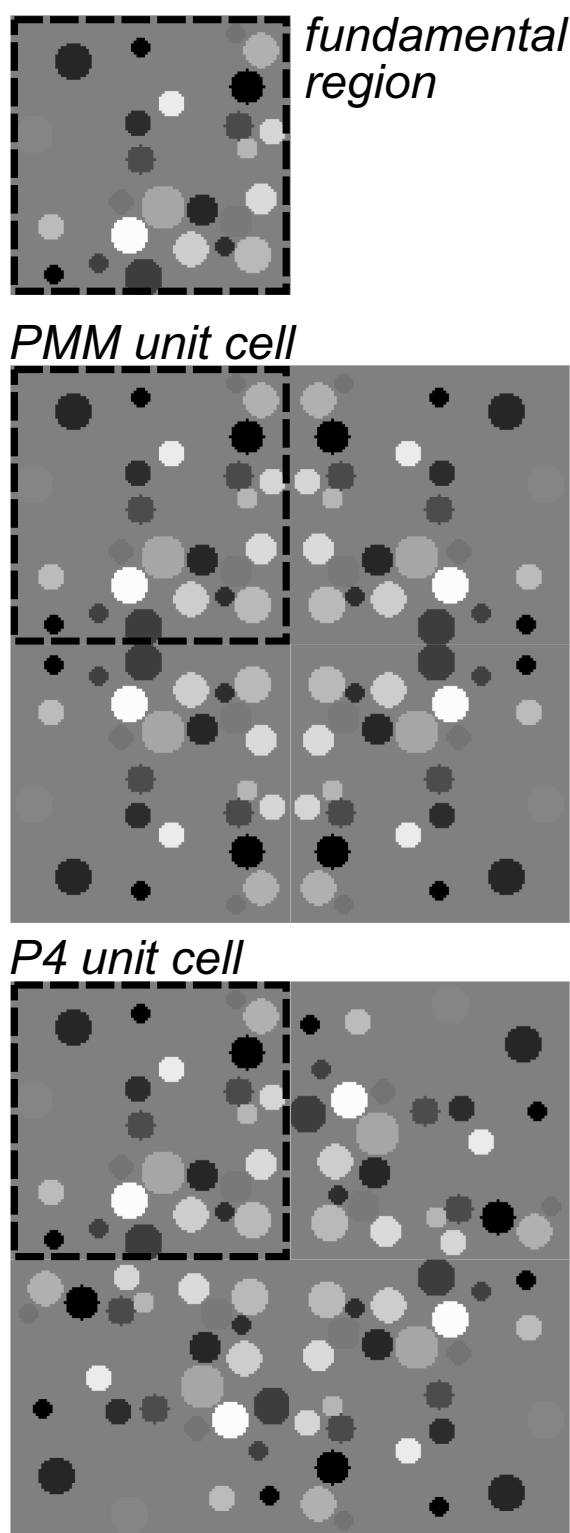


Figure 3. An example of a fundamental region and corresponding PMM and P4 unit cells, with the fundamental region highlighted with a black dotted line. Here, you can see that the random dot pattern from the fundamental region is repeated four times to create a unit cell but using different transformations. This allows for consistency between the two symmetry types regarding the amount of white, black, and shades of gray in each stimulus.

We also created a “random lattice” that contained no symmetry by using four distinct fundamental regions in each quadrant of the 2×2 array. These random unit cells can be embedded in the wallpaper group in place of any of the repeating PMM or P4 unit cells and are matched to the symmetry unit cells in terms of number of dots, contrast and spatial frequency. In our visual search task, the random unit cell serves as the target and wallpaper group unit cells serve as distractors (see Figure 4). As noted in the Introduction, this participants’ ability to identify the absence of symmetry in the search arrays is used as a measure of symmetry processing, across different conditions. We manipulated symmetry type by either using PMM (“reflection”) or P4 (“rotation”) textures. We further manipulated regularity by adding a separate set of conditions in which spacing was introduced around each unit cell corresponding to 20% of the cell width/height, and the position of each cell was jittered randomly between $\pm 15\%$ in both the x and y direction. The jittered conditions were contrasted with un-jittered conditions in which the unit cells were repeated on a lattice with no gaps between the cells, so the distractors formed regular textures. Across all experiments, the unit cells were arranged in 3×3 , 4×4 , 5×5 , and 6×6 wallpapers to create different array sizes, so that the overall pattern size varied between conditions. The size of the unit cell was 100×100 pixels across all experiments.

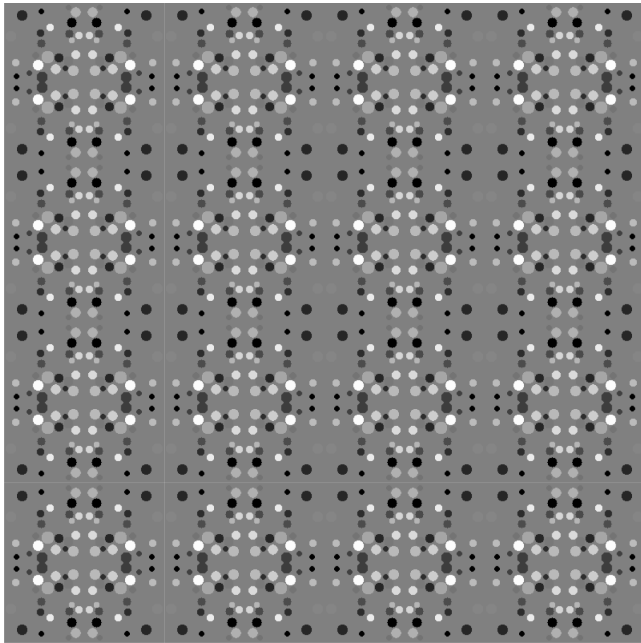
Participants

Participants were recruited through the online participant pool, Prolific. They were compensated £8.50/hr for their participation, and the experiment lasted about 20 minutes. For each of our four experiments, recruited 50 participants, but due to a glitch in our data acquisition pipeline, we ended up with a total of 179 (males = 123, females = 54, other = 2). The average age of all participants was 22.50 ± 3.32 . Across experiments, we removed participants who were unable or unwilling to do the task (Experiment 1 (PMM) = 4; Experiment 2 (P4) = 10, Experiment 3 (PMM jitter) = 2, Experiment 4 (P4 jitter) = 7), based on a criterion explained below. The total number of included participants across the four experiments, was 44, 30, 44 and 38, respectively. Informed consent was obtained before the experiment under a protocol approved by the Office of Research Ethics at York University.

Procedure

All four experiments were written using JsPsych (de Leeuw, 2015), hosted online on Pavlovia.org, and followed the same general procedure. Participants were presented with one block of 24 practice trials,

target absent



target present

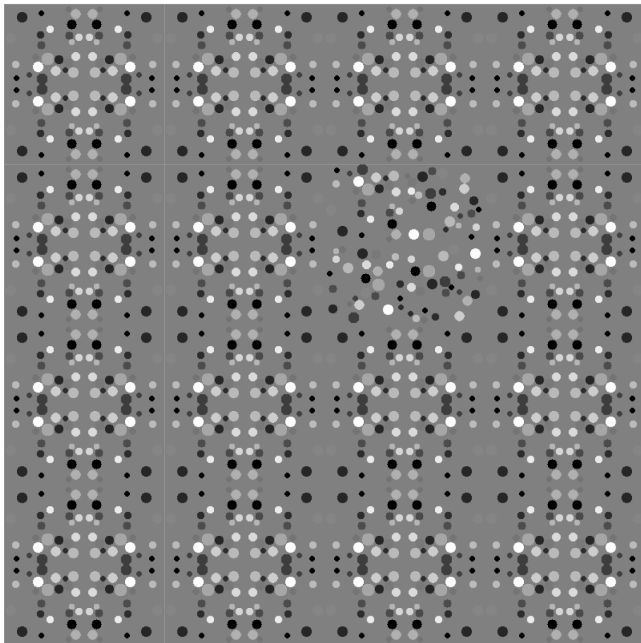


Figure 4. Example images from Experiment 1; PMM target absent versus present example. The “random unit cell” target tile is in the second row from the top, the third column from the left.

followed by 240 experimental trials broken into 10 blocks. The wallpaper array contained a target tile in 75% of trials. Fundamental region exemplars were pseudo-randomly assigned to target-present and target-absent trials so that each of the 10 exemplars was repeated an approximately equal number of times for target-absent and target-present trials, across all array sizes. Trials were shown in random order across exemplars, trial types, and array sizes. When targets were present, their location in the search array was chosen randomly on each trial. Participants were asked to use their keyboard to indicate if a target tile was present or not, pressing the “L” key to indicate that the target was present, and the “D” key to indicate that it was not. Trials only progressed after a selection had been made. After both practice and experiment trials, feedback was provided in the form of the word “Correct!” in green text or “Incorrect!” in red text, presented with a statement indicating how many trials remained in the experiment. Feedback remained on the screen for 1000 ms before the next trial was presented. After each of the 10 blocks, the participants were given the opportunity to take a break before pressing any key to continue. At the end of the experiment, participants were thanked for their participation and redirected to a page where payment could be assigned. A comparative study found that reaction times collected online using the JsPsych package are slightly longer (~25 ms) than those collected in the lab using Psychophysics Toolbox, but that variability is comparable between the two approaches (de Leeuw & Motz, 2016).

Data analysis

We followed the procedure for calculating d' outlined by Macmillan and Kaplan (1985). When individual participants had hit and/or false alarm rates that were 1 or 0, we converted that to usable minimum and maximum rates by adding or subtracting half a trial:

$$\text{minimum} = \frac{0.5}{T}$$

$$\text{maximum} = 1 - \frac{0.5}{T}$$

where T is the total number of target-present (if correcting hit rates) or target-absent (if correcting false alarm rates) trials. This allowed us to use the standard z -score distribution. Participants with $d' < 1$ were considered unable or unwilling to do the task and were removed from further analysis. All statistical analyses were done using JASP (Version 0.18.3).

Results

We computed the median reaction time and d' for each array size, for each participant in each of the four experiments. To test our two hypotheses, we ran a linear mixed models analysis separately on the reaction time and d' data. Symmetry type (wallpaper group) and jitter were between-subject fixed effects, array size, treated as a continuous variable, was a within-participant fixed effect, and the participant was a random effect. For illustration purposes, we also computed the slope of the search function for reaction time and d' individually for each participant (averages across participants are shown in [Figures 5B](#) and [6B](#)). Greater slope values are indicative of more serial processing.

For reaction time, we found a significant main effect of symmetry type ($F(1,152) = 83.745$, $p < 0.001$), indicating that participants were faster for reflection symmetry (PMM) than for rotation symmetry (P4) across all array sizes. There was also a significant main effect of array size ($F(1,152) = 283.191$, $p < 0.001$), indicating that reaction time increased with larger array sizes across conditions. Importantly, the significant main effects were modified by two significant interactions that elucidate the efficiency of the visual search: The first interaction was between symmetry type and array size ($F(1,152) = 6.776$, $p = 0.010$) such that rotation symmetry (P4) produced steeper search functions and thus less efficient search than reflection (PMM). The second interaction was between jitter and

array size ($F(1,152) = 19.258$, $p < 0.001$), such that jittered conditions produced steeper search functions and thus less efficient search than un-jittered conditions. There was no main effect of jitter ($p = 0.345$) but the interaction between symmetry type and jitter approached significance ($F(1,152) = 3.386$, $p = 0.068$). Importantly, we did not find a three-way interaction (symmetry type \times jitter \times array size) ($F(1,152) = 0.165$, $p = 0.685$), suggesting that symmetry type and jitter have separate and independent effects on processing efficiency.

We ran the same analysis with d' values as the dependent variable to determine whether sensitivity was influenced by symmetry type and jitter, and to check for potential speed accuracy tradeoffs. As for the reaction time analysis, we found significant main effects of symmetry type ($F(1,152) = 14.553$, $p < 0.001$), jitter ($F(1,152) = 5.470$, $p = 0.021$), and array size ($F(1,152) = 55.025$, $p < 0.001$). The significant main effects were modified by a significant interaction between symmetry type and jitter ($F(1,152) = 4.317$, $p = 0.039$), but there were no other main effects or interactions (smallest $p = 0.258$). The slopes of the functions relating d' to array size were relatively flat and similar across conditions. The only exception is un-jittered reflection symmetries (PMM), which had a flatter slope than the other conditions (see [Figure 6](#)). This is consistent with the reaction time analysis, which shows that out of all the conditions, un-jittered reflection symmetries (PMM) led to the most efficient processing.

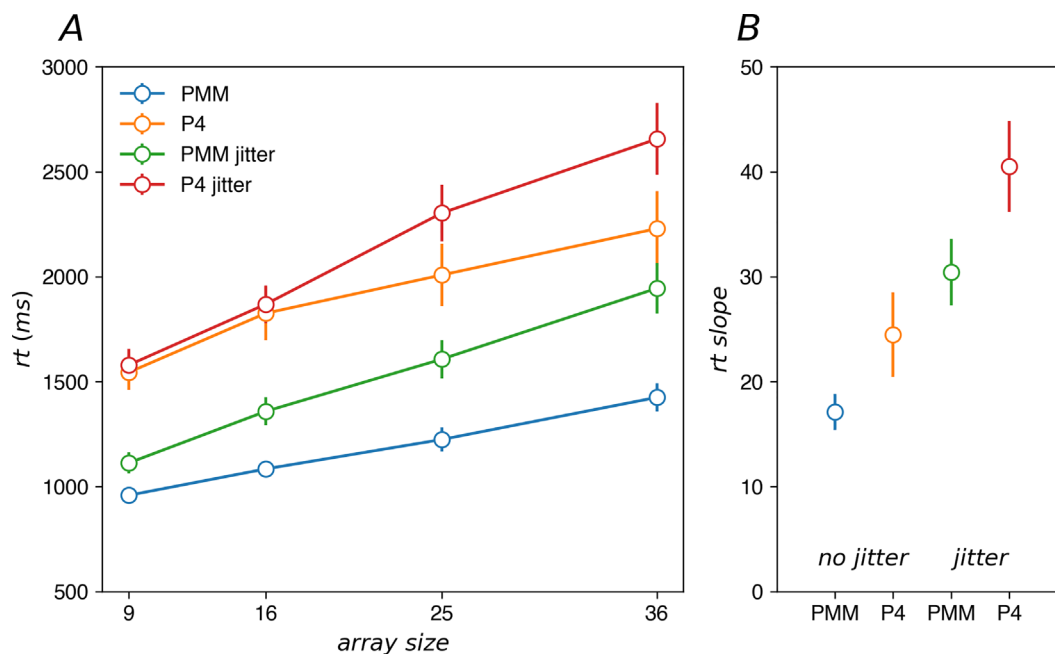


Figure 5. (A). Reaction time data across the four experiments. Error bars reflect the standard error of the mean. (B) Slopes of the corresponding visual search function, averaged across participants, for each of the four experiments. Error bars reflect the standard error of the mean. It is evident that slope values are smaller (more parallel) for PMM than for P4, and for non-jittered compared to jittered conditions.

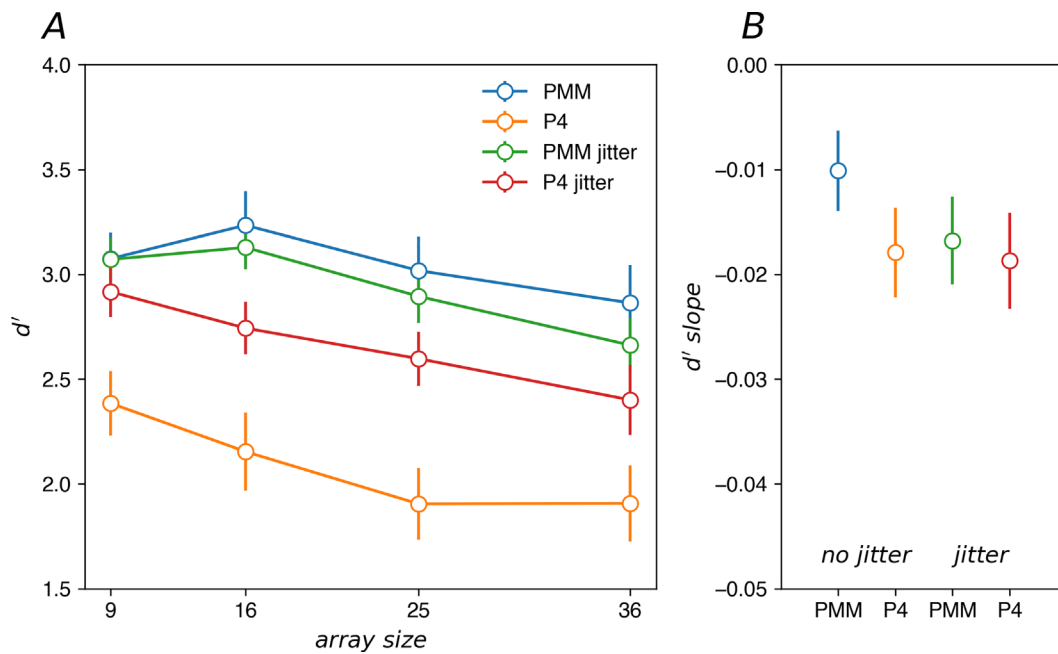


Figure 6. d' plotted in the same way as reaction time, with d' data across the four experiments in (A), and slopes of the corresponding visual search function in (B). Error bars reflect the standard error of the mean. Slopes across array sizes are relatively flat and similar across conditions. The only exception is un-jittered PMM, where the slope is flatter than the others. This implies that this condition was the easiest overall, which is consistent with the reaction time results.

Discussion

Our results identify independent effects of symmetry type and texture regularity on visual search efficiency. This is captured by the interactions between symmetry type and array size, and between jitter and array size, which we observe for reaction time. The interactions show that symmetry type and jitter both influence the slope of the search function, with reflection leading to shallower slopes than rotation, and un-jittered displays leading to shallower slopes than jittered displays. These results can be observed in Figure 5, where the shallowest slope is observed for un-jittered reflection symmetry (PMM) and the steepest for jittered rotation (P4). The absence of a three-way interaction suggests that symmetry type and jitter have separate and independent effects on processing efficiency. The pattern of results for d' allows us to rule out speed-accuracy tradeoffs as an explanation for our reaction time results.

Behavioral literature on symmetry, reflection versus rotation

Previous research generally used symmetry detection tasks when comparing reflection versus rotation symmetries and found that reflection was more perceptually salient than rotation (Mach, 1959; Royer, 1981; Hamada & Ishihara, 1988; Palmer, 1991; Ogden,

Makin, Palumbo, & Bertamini, 2016). We extend these previous findings by demonstrating an advantage for reflection in the context of visual search. We can speculate that the advantage of reflection may be a result of evolutionary pressures because reflection contributes to the identification of bilateral organisms: Members of the same species, predators, and prey. The rapid detection of reflection symmetries would thus be pertinent to survival in hunting, fighting, or mating scenarios. It is easy to imagine that these pressures would apply to rotation symmetry detection to a lesser degree, although rotation symmetry may still facilitate identification of various plants, insects, and marine animals.

Our texture regularity manipulation reveals a novel processing advantage for symmetries when presented in regular textures. This finding is similar to previous demonstrations of whole-array effects of symmetry in visual search, but in our case, we are manipulating regularity rather than symmetry. The ecological relevance of this effect may be related to evolutionary pressures toward detecting disruptions in regular and near-regular textures in the environment, in the context of detecting edible plants, predators, or prey, that are embedded in the background vegetation.

Visual search and symmetry

Studies of symmetry using visual search have identified effects of both within-item symmetry

(Javadnia & Ruddock, 1988; Olivers & Van Der Helm, 1998) and symmetry over the whole array (Wolfe & Friedman-Hill, 1992). The current study builds on previous work in two important ways: First, by making it possible to independently manipulate symmetry type, as a within-item manipulation, and regularity, by definition a whole-array manipulation. Second, to our knowledge, no prior research has examined how rotation symmetry, whether within a target or a distractor, influences performance on a visual search task. Our study allowed us to place each of our conditions along a spectrum of parallel and serial processing and measure the effect of symmetry type and regularity on both.

We did not see evidence of parallel processing for any of our conditions, whereas previous work using both within-item (Javadnia & Ruddock, 1988) and whole-array manipulations (Wolfe & Friedman-Hill, 1992) found evidence of parallel processing of reflection symmetry. We believe there are a few reasons why that may be. First, although our regularity manipulation is a whole-array manipulation in the same class as that used in previous work demonstrating parallel processing (Wolfe & Friedman-Hill, 1992), we are manipulating regularity while previous work manipulated symmetry. When compared to the previous findings, our results thus suggest that whole-array symmetry, but not whole-array regularity, gives rise to parallel processing.

Our within-item manipulation of symmetry, however, is more generally similar to the types of displays used in previous work (Javadnia & Ruddock, 1988; Olivers & Van Der Helm, 1998), especially in our jittered conditions. Should we be surprised to not see parallel processing when within-item reflection or rotation symmetry differed between the target and the distractor? To our knowledge, the evidence of parallel processing in within-item symmetry-based visual search is limited to a study (Javadnia & Ruddock, 1988) that used stimuli based on “textons” (Julesz, 1981). Here, the target and distractor stimuli did differ on symmetry, but also differed in other respects, such as the presence of line junctions of various types, which may have facilitated pop-out of the target among the distractors. A more recent study used more well-controlled stimuli and found no evidence of parallel processing of symmetry across four distinct stimulus types (Olivers & Van Der Helm, 1998). The stimuli used in the current work have targets and distractors that are similarly well-matched for low-level differences, and further allow us to manipulate symmetry type without introducing any low-level differences. Given this high level of control, we are not surprised that we, like Oliver & Van Der Helms (1998), find no evidence of parallel processing of within-item symmetry.

Neuroimaging literature on symmetry, reflection versus rotation

The neuroimaging literature shows that both reflection and rotation produce strong responses in the visual cortex, even when participants are doing an orthogonal task and not paying attention to the symmetry (Kohler et al., 2016), but activity measured using electroencephalography was weaker for rotation than for reflection symmetry (Kohler & Clarke, 2021). A recent direct comparison of responses to different wallpaper groups in the visual cortex of macaque monkeys showed that activation in visual areas (V3 and V4) was approximately 1/3 larger for reflection than for rotation (Audurier et al., 2022). Studies using non-texture stimuli with a single symmetry axis have also consistently found weaker responses for rotation than reflection (Makin, Wilton, Pecchinenda, & Bertamini, 2012; Makin, Pecchinenda, & Bertamini, 2013; Makin, Rampone, Wright, Martinovic, & Bertamini, 2014; Wright et al., 2015). This neural advantage for reflection over rotation is consistent with the behavioral advantage observed in the current study and prior studies discussed above. An important goal for further neuroimaging research will be to directly compare symmetries when presented independently or embedded in regular textures to understand how the behavioral advantage for symmetries in regular textures arises in the brain.

In this study, we found that search was more efficient for PMM compared to P4 patterns. One interpretation is that this is due to a larger difference in neural activity elicited by the random unit cell target within the PMM patterns, compared to when the same type of unit cell is placed within P4 patterns. This would likely occur because the random unit cell produces weaker brain responses than the symmetrical surround. The inverse—that the random unit cell generates stronger responses—is also possible, but unlikely, given the large literature showing that symmetrical patterns generate more brain activity than random patterns. This has been demonstrated for wallpaper groups (Kohler et al., 2018), and for dot patterns with symmetry axes at a single location in the image (see Makin et al., 2022 for a review). Given that the task likely involves finding a region of weaker neural activity, it is interesting to consider the possible involvement of visual area V4. V4 is a topographically organized region on the ventral surface of the occipital cortex, that has been found to respond strongly to symmetry in wallpaper groups (Kohler et al., 2016) and dot patterns (Sasaki, Vanduffel, Knutsen, Tyler, & Tootell, 2005). Ablation of area V4 in macaque monkeys leads to a dramatic weakening of performance on a visual search task when targets were lower brightness or smaller size, but only mild deficits when targets were brighter or larger than the distractors (Schiller & Lee, 1991). This

suggests that our task, which involves symmetry *and* detecting an area of weaker or missing information, may in fact be ideally suited for driving area V4. Testing this hypothesis is an important goal for future electrophysiology or neuroimaging research.

Texture perception

Textures form the patterns that make up the surfaces of objects and environments; they play an essential role in vision in everyday life (Adelson, 2001). An important step toward understanding and analyzing human texture perception was the development of a computational model that made it possible to represent and synthesize visual textures based on joint statistics of the image (Portilla & Simoncelli, 2000). The model has proven highly useful in capturing how texture representations change across the visual field (Balas, Nakano, & Rosenholtz, 2009; Freeman & Simoncelli, 2011) and how natural textures are represented in different areas of the visual cortex (Freeman, Ziemba, Heeger, Simoncelli, & Movshon, 2013; Okazawa, Tajima, & Komatsu, 2015). Importantly, however, this modeling framework is unable to synthesize regular textures like the wallpaper groups used in our experiment (unpublished data) and therefore unlikely to fully explain the regularity effect found in the current data or the brain imaging data mentioned above. The current data offer another piece of evidence suggesting that regular and near-regular textures may play an important role in perception. Our findings highlight the need for the development of models that can describe and synthesize regular textures.

The uniformity illusion (Fukuda & Seno, 2012; Otten, Pinto, Paffen, Seth, & Kanai, 2017) may also be relevant to our results. In this illusion, a regular grid of repeating local elements can undergo illusory completion such that a disruption of the uniformity in the periphery is not detected. The illusion suggests that the visual system has a tendency toward perceiving textures as regular, which implies that visual search should be less efficient when the target is presented in a regular texture, because the target is filled in to give the impression of a uniform pattern. Why do we see the opposite pattern, more efficient search for regular textures? One possibility is that participants rapidly moved their gaze around the search array while performing the task and thereby prevented the grid illusion from occurring. It may also be that the mechanism underlying the uniformity illusion only operates in the far periphery, far enough to not influence our results. Another, related, possibility is that the uniformity illusion requires more tiling of the repeating pattern, than is present in our textures. Otten and colleagues (2017) used relatively large central segments (e.g., $26^\circ \times 14^\circ$ /visual angle), and although they did not systematically test the effect of tiling, the

number of repetitions did in most cases exceed that present in even our largest stimuli. It is worth noting that although these differences could potentially explain the absence of a regularity effect in our results, they do not predict the effect we see in the opposite direction. It is compelling to consider a third possibility, namely that the filling-in mechanism is unable to reproduce the local symmetries, whether reflection or rotation, present in our stimuli. If the visual system has a tendency toward perceiving textures as regular, it is plausible that this failure to fill in the regularity would give rise to a strong error signal, which would in turn make visual search more efficient with a regular search array. Otten and colleagues (2017) have demonstrated that the uniformity illusion can be produced for a wide range of features, and the wallpaper group stimuli used in the current study provide an ideal stimulus set for adding symmetry to that list. Further work should investigate which parameters are required for generating a potential uniformity illusion with repeating elements that have local symmetries.

Potential confounds and limitations

A possible limitation of this study is that although the stimuli were well controlled within participants the data were collected online on participants' own devices, which has the potential for introducing differences between participants. To minimize these differences, participants were required to use a laptop or desktop computer for the experiment (no phones or tablets were allowed). However, we made no attempts to control viewing distance or monitor resolution, which may have led to some variability in the size of the stimuli in degrees of visual angle between participants. In addition, the contrast and luminance of the stimuli may also have varied because of differences in the monitors used by different participants. It is important to note, however, that our effects of interest were measured within-participant, and thus unlikely to be driven by these between-participant differences. Furthermore, any noise added to our measurements by the lack of control is likely compensated for by our ability to get data from a relatively large number of participants, compared to a standard psychophysical experiment.

A second possible limitation is that the target was always asymmetrical, while the distractors were symmetrical. This approach was chosen because it allows us to organize the search arrays into wallpaper groups PMM and P4, thereby connecting our results to the existing literature on visual processing of symmetries in wallpaper groups (e.g., Kohler et al., 2016). It is important to note that the search for asymmetry approach has been taken in previous studies using both within-item (Javadnia & Ruddock, 1988) and whole-array symmetry (Wolfe & Friedman-Hall, 1992). Most importantly, Olivers & Van der Helm

(1998) directly compared search for a symmetrical target among asymmetrical distractors to search for an asymmetrical target among symmetrical distractors and found no differences in search efficiency as quantified by the slope of the search function (Olivers & Van der Helm, 1998, p. 1106). Based on these findings, we conclude that using an asymmetrical target to measure symmetry processing is uncontroversial and well-aligned with the existing literature.

Another possible concern regarding our regularity manipulation is that the regular arrangement of the unit cell on a lattice in no jitter condition may have produced a mid-level visual effect where individual dots in the unit cells are perceptually grouped to form a grid-like pattern. It is possible that this can help guide the visual search task, because the target (a disruption of the grid) becomes easier to spot. We have several responses to this concern: First, it is important to note that such patterns across the whole texture are in a sense inherent to regular textures, and thus difficult to disambiguate from regularity itself. Second, we note that the presence of a grid pattern does not necessarily mean that a disruption of the grid becomes easy to detect, as evidenced by the uniformity illusion (discussed above; see Figure 7 in Otten et al., 2017). Finally, if the grid pattern was driving our regularity effect, we would expect there to be a stronger effect of regularity on PMM, where straight lines are more likely to form across the pattern (see Figure 4). This is not what we see in our results: If anything, the averages for PMM and P4 are more similar in the no jitter condition than in the jitter condition, and the three-way interaction between symmetry type, jitter, and array size is far from significance ($p = 0.685$). Absence of evidence is not evidence of absence, and we should not draw firm conclusions based on a null result, but in the current study our results support the conclusion that symmetry type and regularity have separate and independent effects on processing efficiency.

Finally, we would like to remind the reader that there are 17 distinct wallpaper groups, each with a unique combination of symmetries. We only used 2 out of the 17, which allowed us to target differences between reflection and rotation. Follow-up studies using other wallpaper groups or other types of symmetry such as glide reflection are valuable targets for future research and may produce different results.

Conclusions

The current study used highly controlled stimuli to provide strong evidence that reflection symmetries (wallpaper group PMM) are processed more efficiently than rotation symmetries (wallpaper group P4). Perhaps more surprisingly, the results also show that texture regularity has a significant effect on the processing of

symmetries independent of symmetry type, such that when symmetries are embedded within a regular texture, they are more efficiently processed. Notably, these effects were independent and additive, which suggests that they reflect distinct perceptual mechanisms. As pointed out above, a major distinction of our work from previous work on visual search is our ability to test both within-item and whole-array manipulations in the same display and show that within-item symmetry type as well as whole-array regularity matters for visual search. The findings should prompt further studies of the underlying brain mechanisms and provide a foundation for further research on how symmetries and textures interact during natural, real-world perceptual tasks.

Keywords: symmetry, regularity, serial and parallel processing

Acknowledgments

Supported by the Vision Science to Applications (VISTA) program funded by the Canada First Research Excellence Fund (CFREF, 20162023), by a Discovery Grant from the Natural Sciences and Engineering Research Council of Canada awarded to PJK, and by a grant from the Scientific and Technological Research Council of Türkiye (TUBITAK, 1001 grant 122K922) awarded to NA.

Data availability: The datasets collected for the current study are available in the project wallpaper-search repository, <https://osf.io/b7hq9/>.

Commercial relationships: none.

Corresponding author: Peter J. Kohler.

Email: pjkohler@yorku.ca.

Address: Department of Psychology, Sherman Health Science Research Centre, York University, Toronto, Ontario M3J 1P3, Canada.

References

- Adelson, E. H. (2001). On seeing stuff: The perception of materials by humans and machines. *Human Vision and Electronic Imaging VI*, 4299, 1–12, <https://doi.org/10.1117/12.429489>.
- Alp, N., Kohler, P.J., Kogo, N., Wagemans, J., & Norcia, A. M. (2018). Measuring Integration Processes in Visual Symmetry with Frequency-Tagged EEG. *Scientific Report*, 8, 6969, <https://doi.org/10.1038/s41598-018-24513-w>.
- Audurier, P., Héjja-Brichard, Y., De Castro, V., Kohler, P. J., Norcia, A. M., Durand, J.-B., ... Cottureau,

- B. R. (2022). Symmetry Processing in the Macaque Visual Cortex. *Cerebral Cortex*, 32(10), 2277–2290, <https://doi.org/10.1093/cercor/bhab358>.
- Bahnsen, P. (1928). *Ein Untersuchung über Symmetrie und Asymmetrie bei visuellen Wahrnehmungen*. Princeton, NJ: Lippert.
- Balas, B., Nakano, L., & Rosenholtz, R. (2009). A summary-statistic representation in peripheral vision explains visual crowding. *Journal of Vision*, 9(12), 13, <https://doi.org/10.1167/9.12.13>.
- Bertamini, M., & Makin, A. D. J. (2014). Brain activity in response to visual symmetry. *Symmetry*, 6(4), 4, <https://doi.org/10.3390/sym6040975>.
- Bertamini, M., Silvanto, J., Norcia, A. M., Makin, A. D. J., & Wagemans, J. (2018). The neural basis of visual symmetry and its role in mid- and high-level visual processing. *Annals of the New York Academy of Sciences*, 1426(1), 111–126, <https://doi.org/10.1111/nyas.13667>.
- Bundesen, C., Kyllingsbæk, S., & Larsen, A. (2003). Independent encoding of colors and shapes from two stimuli. *Psychonomic Bulletin & Review*, 10(2), 474–479, <https://doi.org/10.3758/BF03196509>.
- Cavanagh, P., Arguin, M., & Treisman, A. (1990). Effect of surface medium on visual search for orientation and size features. *Journal of Experimental Psychology: Human Perception and Performance*, 16(3), 479–491, <https://doi.org/10.1037/0096-1523.16.3.479>.
- de Leeuw, J. R. (2015). jsPsych: A JavaScript library for creating behavioral experiments in a Web browser. *Behavior Research Methods*, 47(1), 1–12, <https://doi.org/10.3758/s13428-014-0458-y>.
- de Leeuw, J. R., & Motz, B. A. (2016). Psychophysics in a Web browser? Comparing response times collected with JavaScript and Psychophysics Toolbox in a visual search task. *Behavior Research Methods*, 48(1), 1–12, <https://doi.org/10.3758/s13428-015-0567-2>.
- Fedorov, E. (1891) Symmetry in the plane (in Russian). In *Proceedings of the Imperial St. Petersburg Mineralogical Society* (2), 28, 245–291.
- Freeman, J., & Simoncelli, E. P. (2011). Metamers of the ventral stream. *Nat Neurosci*, 14(9), 1195–1201, <https://doi.org/10.1038/nn.2889>.
- Freeman, J., Ziemba, C. M., Heeger, D. J., Simoncelli, E. P., & Movshon, J. A. (2013). A functional and perceptual signature of the second visual area in primates. *Nature Neuroscience*, 16(7), Article 7, <https://doi.org/10.1038/nn.3402>.
- Fukuda, H., & Seno, T. (2012). Healin' Groovy: Movement Affects the Appearance of the Healing Grid Illusion. *Perception*, 41(2), 243–246, <https://doi.org/10.1068/p7132>.
- Grammer, K., & Thornhill, R. (1994). Human (Homo sapiens) facial attractiveness and sexual selection: The role of symmetry and averageness. *Journal of Comparative Psychology*, 108(3), 233–242, <https://doi.org/10.1037/0735-7036.108.3.233>.
- Hamada, J., & Ishihara, T. (1988). Complexity and goodness of dot patterns varying in symmetry. *Psychological Research*, 50(3), 155–161, <https://doi.org/10.1007/BF00310176>.
- Jablan, S. V. (2002). *Symmetry, Ornament and Modularity*. Singapore: World Scientific Publishing Co Pte Ltd.
- Javadnia, A., & Ruddock, K. H. (1988). The limits of parallel processing in the visual discrimination of orientation and magnification. *Spatial Vision*, 3(2), 97–114, <https://doi.org/10.1163/156856888X00069>.
- Julesz, B. (1981). Textons, the elements of texture perception, and their interactions. *Nature*, 290(5802), 91–97.
- Kohler, P. J., & Clarke, A. D. F. (2021). The human visual system preserves the hierarchy of two-dimensional pattern regularity. *Proceedings of the Royal Society B: Biological Sciences*, 288(1955), 20211142, <https://doi.org/10.1098/rspb.2021.1142>.
- Kohler, P. J., Clarke, A., Yakovleva, A., Liu, Y., & Norcia, A. M. (2016). Representation of Maximally Regular Textures in Human Visual Cortex. *Journal of Neuroscience*, 36(3), 714–729, <https://doi.org/10.1523/JNEUROSCI.2962-15.2016>.
- Kohler, P. J., Cottureau, B. R., & Norcia, A. M. (2018). Dynamics of perceptual decisions about symmetry in visual cortex. *NeuroImage*, 167(Supplement C), 316–330, <https://doi.org/10.1016/j.neuroimage.2017.11.051>.
- Kyllingsbæk, S., & Bundesen, C. (2007). Parallel processing in a multifeature whole-report paradigm. *Journal of Experimental Psychology: Human Perception and Performance*, 33(1), 64–82, <https://doi.org/10.1037/0096-1523.33.1.64>.
- Liu, Y., Hel-Or, H., Kaplan, C. S., & Van Gool, L. (2010). Computational Symmetry in Computer Vision and Computer Graphics. *Foundations and Trends® in Computer Graphics and Vision*, 5(1–2), 1–195, <https://doi.org/10.1561/06000000008>.
- Liu, Y., Lin, W.-C., & Hays, J. (2004). Near-regular texture analysis and manipulation. *ACM Transactions on Graphics (ToG)*, 23, 368–376.
- Mach, E. (1959). *The Analysis of Sensations (1897)*. New York: English Transl., Dover.
- Machilsen, B., Pauwels, M., & Wagemans, J. (2009). The role of vertical mirror symmetry in visual shape detection. *Journal of Vision*, 9(12), 11, <https://doi.org/10.1167/9.12.11>.
- Macmillan, N. A., & Kaplan, H. L. (1985). Detection theory analysis of group data: Estimating

- sensitivity from average hit and false-alarm rates. *Psychological Bulletin*, 98(1), 185–199. <http://doi.org/10.1037/0033-2909.98.1.185>.
- Makin, A. D. J., Rampone, G., Wright, A., Martinovic, J., & Bertamini, M. (2014). Visual symmetry in objects and gaps. *Journal of Vision*, 14(3), 12. <https://doi.org/10.1167/14.3.12>.
- Makin, A. D. J., Wilton, M. M., Pecchinenda, A., & Bertamini, M. (2012). Symmetry perception and affective responses: A combined EEG/EMG study. *Neuropsychologia*, 50(14), 3250–3261. <https://doi.org/10.1016/j.neuropsychologia.2012.10.003>.
- Makin, A., Pecchinenda, A., & Bertamini, M. (2013). Visual and emotional analysis of symmetry. *Journal of Vision*, 13(9), 812. <https://doi.org/10.1167/13.9.812>.
- Makin, A. D., Tyson-Carr, J., Rampone, G., Derpsch, Y., Wright, D., & Bertamini, M. (2022). Lessons from a catalogue of 6674 brain recordings. *eLife*, 11, e66388. <https://doi.org/10.7554/eLife.66388>.
- Ogden, R., Makin, A. D. J., Palumbo, L., & Bertamini, M. (2016). Symmetry Lasts Longer Than Random, but Only for Brief Presentations. *I-Perception*, 7(6), 2041669516676824. <https://doi.org/10.1177/2041669516676824>.
- Okazawa, G., Tajima, S., & Komatsu, H. (2015). Image statistics underlying natural texture selectivity of neurons in macaque V4. *Proceedings of the National Academy of Sciences*, 112(4), E351–E360. <https://doi.org/10.1073/pnas.1415146112>.
- Olivers, C. N. L., & Van Der Helm, P. A. (1998). Symmetry and selective attention: A dissociation between effortless perception and serial search. *Perception & Psychophysics*, 60(7), 1101–1116. <https://doi.org/10.3758/BF03206161>.
- Otten, M., Pinto, Y., Paffen, C. L. E., Seth, A. K., & Kanai, R. (2017). The Uniformity Illusion: Central Stimuli Can Determine Peripheral Perception. *Psychological Science*, 28(1), 56–68. <https://doi.org/10.1177/0956797616672270>.
- Palmer, S. E. (1991). Goodness, Gestalt, groups, and Garner: Local symmetry subgroups as a theory of figural goodness. In *The perception of structure: Essays in honor of Wendell R. Garner* (pp. 23–39). Washington, DC: American Psychological Association. <https://doi.org/10.1037/10101-001>.
- Polya, G. (1924). XII. Über die Analogie der Kristallsymmetrie in der Ebene. *Zeitschrift Für Kristallographie-Crystalline Materials*, 60(1), 278–282.
- Portilla, J., & Simoncelli, E. (2000). A Parametric Texture Model Based on Joint Statistics of Complex Wavelet Coefficients. *International Journal of Computer Vision*, 40(1), 49–70. <https://doi.org/10.1023/A:1026553619983>.
- Royer, F. L. (1981). Detection of symmetry. *Journal of Experimental Psychology: Human Perception and Performance*, 7(6), 1186–1210. <https://doi.org/10.1037/0096-1523.7.6.1186>.
- Sasaki, Y., Vanduffel, W., Knutsen, T., Tyler, C., & Tootell, R. (2005). Symmetry activates extrastriate visual cortex in human and nonhuman primates. *Proceedings of the National Academy of Sciences*, 102(8), 3159–3163. <https://doi.org/10.1073/pnas.0500319102>.
- Schiller, P. H., & Lee, K. (1991). The Role of the Primate Extrastriate Area V4 in Vision. *Science*, 251(4998), 1251–1253. <https://doi.org/10.1126/science.2006413>.
- Treder, M. S. (2010). Behind the Looking-Glass: A Review on Human Symmetry Perception. *Symmetry*, 2(3), Article 3. <https://doi.org/10.3390/sym2031510>.
- Treisman, A. M., & Gelade, G. (1980). A feature-integration theory of attention. *Cognitive Psychology*, 12(1), 97–136. [https://doi.org/10.1016/0010-0285\(80\)90005-5](https://doi.org/10.1016/0010-0285(80)90005-5).
- Wagemans, J. (1995). Detection of visual symmetries. *Spatial Vision*, 9(1), 9–32.
- Wagemans, J. (1997). Characteristics and models of human symmetry detection. *Trends in Cognitive Sciences*, 1(9), 346–352. [https://doi.org/10.1016/S1364-6613\(97\)01105-4](https://doi.org/10.1016/S1364-6613(97)01105-4).
- Wertheimer, M. (1938). Laws of organization in perceptual forms. In *A source book of Gestalt psychology* (pp. 71–88). London, UK: Kegan Paul, Trench, Trubner & Company. <https://doi.org/10.1037/11496-005>.
- Wolfe, J. M. (1998). What can 1 million trials tell us about visual search? *Psychological Science*, 9(1), 33–39. <https://doi.org/10.1111/1467-9280.00006>.
- Wolfe, J. M. (2016). Visual search revived: the slopes are not that slippery: A reply to kristjansson (2015). *I-Perception*, 7(3), 2041669516643244. <https://doi.org/10.1177/2041669516643244>.
- Wolfe, J. M., & Friedman-Hill, S. R. (1992). On the role of symmetry in visual search. *Psychological Science*, 3(3), 194–198. <https://doi.org/10.1111/j.1467-9280.1992.tb00026.x>.
- Wright, D., Makin, A. D. J., & Bertamini, M. (2015). Right-lateralized alpha desynchronization during regularity discrimination: Hemispheric specialization or directed spatial attention? *Psychophysiology*, 52(5), 638–647. <https://doi.org/10.1111/psyp.12399>.
- van Zoest, W., Giesbrecht, B., Enns, J. T., & Kingstone, A. (2006). New Reflections on Visual Search: Interitem Symmetry Matters! *Psychological Science*, 17(6), 535–542. <https://doi.org/10.1111/j.1467-9280.2006.01740.x>.