

# Current Biology

Magazine

symbiotic strategies across the fungal tree of life.

### **DECLARATION OF INTERESTS**

The authors declare no competing interests.

#### **FURTHER READING**

Akhtar, N., Wani, A.K., Dhanjal, D.S., and Mukherjee, S. (2022). Insights into the beneficial roles of dark septate endophytes in plants under challenging environment: resilience to biotic and abiotic stresses. World J. Microbiol. Biotechnol. 38, 79. https://doi.org/10.1007/s11274-022-03264-x.

Bhardwaj, M., Kailoo, S., Khan, R.T., Khan, S.S., and Rasool, S. (2023). Harnessing fungal endophytes for natural management: a biocontrol perspective. Front. Microbiol. 14, https://doi.org/10.3389/ fmicb. 2023.1280258

Bruyant, P., Moënne-Loccoz, Y., and Almario, J. (2024). Root-associated Helotiales fungi: Overlooked players in plant nutrition. Soil Biol. Biochem. 191, 109363. https://doi.org/10.1016/j.soilbio.2024.109363

Gange, A.C., Koricheva, J., Currie, A.F., Jaber, L.R., and Vidal, S. (2019). Meta-analysis of the role of entomopathogenic and unspecialized fungal endophytes as plant bodyguards. New Phytol. 223, 2002–2010. https://doi.org/10.1111/ nph.15859.

Liao, C., Doilom, M., Jeewon, R., Hyde, K.D., Manawasinghe, I.S., Chethana, K.W.T., Balasuriya, A., Thakshila, S.A.D., Luo, M., Mapook, A., et al. (2025). Challenges and update on fungal endophytes: classification, definition, diversity, ecology, evolution and functions. Fungal Diversity 131, 301–367. https://doi.org/10.1007/s13225-025-00550-5

Mesny, F., Miyauchi, S., Thiergart, T., Pickel, B., Atanasova, L., Karlsson, M., Hüttel, B., Barry, K.W., Haridas, S., Chen, C., et al. (2021). Genetic determinants of endophytism in the Arabidopsis root mycobiome. Nat. Commun. 12, 7227. https://doi.org/10.1038/s41467-021-27479-v.

Redman, R.S., Dunigan, D.D., and Rodriguez, R.J. (2001). Fungal symbiosis from mutualism to parasitism: who controls the outcome, host or invader? New Phytol. 151, 705–716. https://doi.org/10.1046/j.0028-646x.2001.00210.x.

Rodriguez, R.J., White Jr., J.F., Arnold, A.E., and Redman, R.S. (2009). Fungal endophytes: diversity and functional roles. New Phytol. 182, 314–330. https://doi.org/10.1111/j.1469-8137.2009.02773.x.

Ruotsalainen, A.L., Kauppinen, M., Wäli, P.R., Saikkonen, K., Helander, M., and Tuomi, J. (2022). Dark septate endophytes: mutualism from byproducts? Trends Plant Sci. 27, 247–254. https:// doi.org/10.1016/j.tplants.2021.10.001.

Saini, H.P., Meena, M., Sahoo, A., and Mehta, T. (2025). A review on fungal endophytes of the family Fabaceae, their metabolic diversity and biological applications. Heliyon 11, e42153. https://doi.org/10.1016/j.heliyon.2025.e42153.

Biosciences Division, Oak Ridge National Laboratory, Oak Ridge, TN 37831, USA. 
<sup>2</sup>Plant-Microbe Interfaces Scientific Focus Area, Genomics, US Department of Energy, Office of Science, Biological and Environmental Research, DE-AC05-00OR22725. 
<sup>3</sup>INRAE, UMR 1136 Interactions Arbres-Microorganismes (IAM), Université de Lorraine, 54000 Nancy, France. 
<sup>4</sup>Department of Biology, Duke University, Durham, NC 27708, USA. 
<sup>5</sup>These authors contributed equally as first authors. 
<sup>\*</sup>E-mail: Claire.Veneault-Fourrey@inrae.fr

### Correspondence

## Visual anagrams reveal highlevel effects with 'identical' stimuli

Tal Boger and Chaz Firestone

A fundamental question in psychology and neuroscience concerns how the mind represents not only lower-level stimulus features such as luminance, contrast, or spatial frequency, but also richer, higher-level properties such as animacy, emotion, or real-world size. Numerous findings suggest that such high-level properties are encoded automatically<sup>1,2</sup>, engage visual attention<sup>3,4</sup>, and organize neural responses<sup>5,6</sup>. However, a critical challenge arises when interpreting such findings: High-level categories systematically covary with lowerlevel features, such that effects attributed to high-level properties may instead be driven by their lowerlevel covariates. Can this challenge be overcome? Here, we introduce a novel approach by leveraging 'visual anagrams' - a diffusionbased technique for generating images whose interpretations change radically with orientation, such as a cow when upright and a mouse when inverted7. Using real-world size as a case study, we generated anagrams depicting a canonically large object in one orientation and a canonically small object in another, and placed them in classic experimental paradigms. Five experiments revealed that many (but not all) effects of real-world size persisted under such conditions. Together, our findings address a longstanding challenge in perception research and establish a broadly applicable tool for psychology and neuroscience.

Consider the rabbit and elephant in Figure 1A. Although they occupy roughly the same amount of space on the page, they differ in their realworld size. An extensive body of research suggests that this high-level difference is actively represented by the mind: Real-world size intrudes on

orthogonal perceptual judgments1,2, drives visual search4, and constrains cortical representation5. But realworld size is not the only feature distinguishing the rabbit and elephant: They also differ in shape, curvature, spatial frequency, viewing angle, and other mid- and low-level properties. Thus, while differences in representation of these objects may arise from differences in real-world size, they may instead arise from correlated lower-level differences (an especially salient possibility given similar findings with distorted, unrecognizable stimuli<sup>2,4,8</sup>). Despite progress on this problem<sup>6,9</sup>, isolating high-level properties from lowerlevel features remains an enduring challenge.

Now consider the rabbit and elephant in Figure 1B. These are actually the very same image, rotated 90°. They are 'visual anagrams', created using a diffusion-based technique that generates static images whose interpretations change radically when rotated<sup>7</sup>. The two images are pixel-wise identical subject to rotation, thus differing in a high-level property (here, real-world size) without differing in features such as curvature, spatial frequency, luminance, contrast, and

Here, we exploit this technique to investigate high-level effects with otherwise 'identical' stimuli, minimizing the lower-level covariation associated with conventional approaches. We generated images depicting a large object in one orientation and a small object in another (for example, rabbit-elephant, butterflybear), placed them in classic paradigms exploring real-world size (https://perceptionresearch.org/anagrams), and asked whether the original findings persist under these conditions.

We first investigated automatic encoding of real-world size using the familiar-size Stroop task<sup>1</sup>. In this task, two images are displayed at different sizes, and subjects must say which is larger on the screen. Despite real-world size being explicitly task-irrelevant, performance is better when displayed size is congruent with real-world size (for example, rabbit-small, elephant-big). Experiment 1 adapted this design to our anagram stimuli. Consistent with previous work, we found a familiar-size Stroop effect



## **Current Biology**

### Magazine



(Figure 1C): Subjects were faster and more accurate on congruent trials than incongruent trials (-21.9 ms, t(50) = 4.75, p < 0.001; +0.8%,t(50) = 3.80, p < 0.001), even when the images were simply rotated versions of one another.

We next explored a connection between real-world size and aesthetic preferences. Previous work suggests that observers prefer canonically small objects to be displayed small, and canonically large objects to be displayed large<sup>8,10</sup>. Consistent with this work, Experiment 2 revealed that subjects preferred canonically large objects to be displayed larger than canonically small objects, even with visual anagrams (+29.3 px, or +9.6%, t(197) = 8.60, p < 0.001; Figure 1D).

Whereas Experiments 1 and 2 included a familiarization phase in which subjects first matched category labels to the anagram stimuli, Experiments 3 and 4 replicated those experiments without this phase. The same patterns emerged (Stroop: -31.7 ms, t(45) = 5.58, p < 0.001; Preferred size: +24.3 px, or +8.2%, t(197) = 8.23, p < 0.001), replicating our results and demonstrating that visual anagrams are readily identifiable without prompting.

Finally, we investigated links between real-world size and attention. Previous work reports that targets are easier to locate when their real-world size differs from distractors4. Using that paradigm, however, Experiment 5 found littleto-no effect with anagram stimuli (11.1 ms advantage, t(48) = 0.51,p = 0.61, BF10 = 0.176; Figure 1E), suggesting that the original findings may indeed be driven by correlated lower-level properties. Importantly, Experiment 5's design replicated earlier search findings using nonanagram stimuli4; those stimuli successfully reproduced previously reported effects (102.6 ms advantage, t(48) = 4.89, p < 0.001), which were significantly stronger than the (nonsignificant) effects with anagrams (91.5 ms difference, t(48) = 3.68,p < 0.001).

Our work confronts the longstanding challenge of disentangling high-level properties from lower-level covariates.

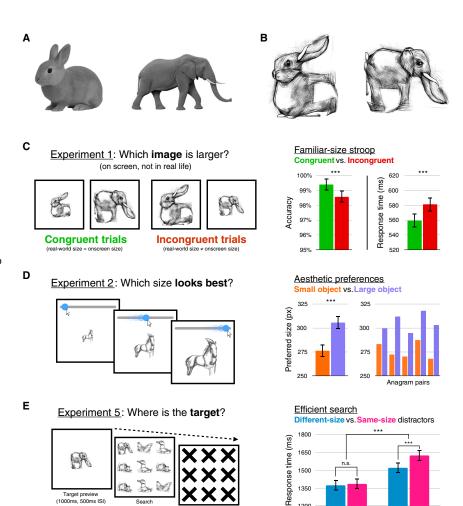


Figure 1. High-level effects with visual anagrams.

(A) This rabbit and elephant differ in a high-level property - real-world size - but also in several mid-level and low-level properties, such as curvature, spatial frequency and contrast. (B) This rabbit and elephant are 'visual anagrams'7; they also differ in real-world size, but contain identical pixels (being the same image rotated 90°). (C) The familiar-size Stroop effect arose with visual anagrams (Experiment 1). (D) Real-world size drove aesthetic preferences with visual anagrams (Experiment 2). (E) Visual search was not facilitated by real-world size when using visual anagrams, although previously reported effects arose with non-anagram

Our results suggest that real-world size per se is represented by the mind: It is encoded automatically and drives aesthetic judgments, in ways that go beyond its lower-level correlates. Not all effects persisted in this way, however, highlighting how this approach can both support and reframe high-level psychophysical effects.

These findings build on previous work showing that many realworld size effects occur even with unrecognizable 'texforms' that preserve mid-level features such as curvature<sup>2,4,8</sup>. That work raises the

question of whether real-world size effects are fully captured by such features or instead go beyond them. Experiments 1-4 suggest that there are indeed effects that go beyond mid-level stimulus features, whereas Experiment 5 suggests that at least some effects are driven mostly or only by such features (in ways that are nevertheless consistent with the original claims).

1350

1200

Anagrams

Ordinary objects

Importantly, our approach is perfectly general. Though we manipulated real-world size, one could generate anagrams of happy faces and sad faces, tools and



# **Current Biology**

Magazine

non-tools, or animate and inanimate objects, overcoming low-level confounds associated with such stimuli3,6. The present work thus serves as a 'case study', yielding concrete discoveries about realworld size and validating a broadly applicable tool for psychology and neuroscience.

### **DECLARATION OF INTERESTS**

The authors declare no competing interests.

### SUPPLEMENTAL INFORMATION

Supplemental information including additional methodological details and analyses and author contributions can be found with this article online at https://doi.org/10.1016/j. cub.2025.08.036.

### REFERENCES

- 1. Konkle, T., and Oliva, A. (2012). A familiarsize Stroop effect: Real-world size is an automatic property of object representation. J. Exp. Psychol. Hum. Percept. Perform. 38,
- Long, B., and Konkle, T. (2017). A familiar-size Stroop effect in the absence of basic-level recognition. Cognition 168, 234-242.
- 3. Levin, D.T., Takarae, Y., Miner, A.G., and Keil, F. (2001). Efficient visual search by category: Specifying the features that mark the difference between artifacts and animals in preattentive vision. Percept. Psychophys. 63,
- Long, B., Konkle, T., Cohen, M.A., and Alvarez, G.A. (2016). Mid-level perceptual features distinguish objects of different real-world sizes. J. Exp. Psychol. Gen. 145, 95-109.
- 5. Konkle, T., and Oliva, A. (2012). A realworld size organization of object responses in occipitotemporal cortex. Neuron 74, 1114-1124.
- 6. Proklova, D., Kaiser, D., and Peelen, M.V. (2016). Disentangling representations of object shape and object category in human visual cortex: The animate-inanimate distinction. J. Cogn. Neurosci. 28, 680-692.
- Geng, D., Park, I., and Owens, A. (2024). Visual anagrams: Generating multi-view optical illusions with diffusion models. In Proceedings of the IEEE/CVF Conference on Computer Vision and Pattern Recognition, pp. 24154-24163.
- Chen, Y.C., Deza, A., and Konkle, T. (2022). How big should this object be? Perceptual influences on viewing-size preferences. Cognition 225,
- Schmidt, F., Kleis, J., Morgenstern, Y., and Fleming, R.W. (2020). The role of semantics in the perceptual organization of shape. Sci. Rep. 10, 22141.
- 10. Konkle, T., and Oliva, A. (2011). Canonical visual size for real-world objects. J. Exp. Psychol. Hum. Percept. Perform. 37, 23-37.

Department of Psychological and Brain Sciences, Johns Hopkins University, Baltimore, MD 21218, USA E-mail: tboger1@jhu.edu (T.B.), chaz@jhu.edu (C.F.)

### Correspondence

## **Coalitionary intra**group aggression by wild female bonobos

Sonya Pashchevskaya1,2,4,\*, Barbara Fruth<sup>1</sup>, Sofia Lunde Kjærland<sup>1</sup>, Leonardo Berton<sup>1</sup>, Linda Vigilant<sup>3</sup>, and Gottfried Hohmann<sup>1,3</sup>

In humans and non-human primates, male aggression and physical violence are common strategies in the struggle over power and are efficient in exerting control over individuals and groups. In contrast, our close relative, the bonobo (Pan paniscus) is often described as an exceptionally peaceful primate due to the lack of lethal aggression or infanticide and the tendency for individuals to reconcile after conflicts. Nonetheless, rates of male aggression are high<sup>1</sup>, but, atypically for primates, bonobo females are observed to confront males without support from others. Despite female exogamy, forming coalitions of unrelated females in response to male aggression appears to be a common strategy, mostly involving charging or chasing and, in some cases, escalating to physical attacks<sup>2-4</sup>. Here, we report on a violent coalitionary attack by resident females against an adult male in a well-studied group of wild bonobos habituated to observation, detailing participants' violent actions and the victim's responses. The assault involved a fraction of the group, while almost everybody was present, and bystanders, including some close maternal kin, did not support the victim. Our observations detail a rare behavior that is not easy to reconcile with the assumed peaceful nature of bonobo society, but which contributes to evolutionary models of aggression<sup>5</sup>.

The following observations lasting two hours were made by S.P., S.L.K., and L.B. on February 18th, 2025, at the research site of the LuiKotale Bonobo Project. At 15:30, a sudden outburst of communal vocalizations indicated an aggressive encounter. The first observer arrived at the spot about two minutes late, the other two about five minutes late. They saw an adult male bonobo, soon identified as Hugo, lying on the ground face down and being continuously assaulted by several adult females: Polly, Tao, Ngola, Djulie and Bella. Polly is a long-term resident, whereas the other four females immigrated into the community between 2012 and 2018 (Table 1). Almost the whole community was quietly observing the scene from a distance of 5-10 meters. The females jumped alternatingly on Hugo's body, stomping on his back and biting his head, legs, neck, fingers and toes. One female bit off a part of Hugo's ear, two others engaged in genito-genital rubbing with each other on top of him. One of the perpetrators bit into his foot and chewed on the removed tissue, then bit his testes. Throughout the attack, Hugo was lying on his belly, covering his head with his hands, emitting monotonous stress hoots. When Hugo's body became more visible, observers noted that his face was disfigured with bleeding marks on lips and eyebrows. He had lost much hair on his head, shoulders and back, and a large chunk of skin was missing from his neck. His hand knuckles were bitten to the bone, several toe phalanges were bitten off, and there were wounds on his testes and penis. After around 25 minutes of constant assaults, the main perpetrators paused and, for the next 90 minutes, were licking blood off the male's body and their own fingers. Throughout that time, other bonobos, including females and their juvenile offspring who had not been involved in the aggression, licked the victim's wounds or fingers of the attackers. Apollo, Hugo's maternal half-brother, approached Hugo and licked his injured scrotum. At 17:30, a part of the group started moving away. Hugo first walked a few steps bipedally and, being pursued by some of the group members, managed to lean on his injured knuckles and run away, pursuers and observers falling behind. Since the event, members of the community have been followed for more than 150 days without seeing Hugo. Given the severity of his injuries, it is likely that the attack was fatal.

Despite no apparent coordination in the violent acts (i.e., females acted simultaneously but independently of each other), the assault against

