

This file provides documentation for this benchmark and its intended uses. All images, human behavioral data, code to evaluate model performance, and scripts used to visualize results are contained within a [folder at this link](#). Below we outline information about images, data files, and analysis scripts in this folder, as well as information about licensing, hosting, and maintenance plans. We include metadata for this benchmark in croissant format. Finally, we include supplemental figures and descriptions of online/inlab data collection.

IMAGES ARE IN images/

images/

- Folder with all images in this benchmark
- completely flat organizational structure—no nested folders or directories
- images.zip is a zipped version of images/

DATA FILES ARE IN data/

data/benchmark.csv

- data file containing human/model behavioral data central to this benchmark
- Each row contains information for a single trial (i.e., an image triplet)
- Each column contains relevant behavioral and meta data for this triplet, including
 - 'images': list of all images in this trial (which are in /images)
 - 'human_accuracy': averaged human performance on this trial
 - 'human_accuracy_sem': standard error of the mean for trial
 - 'human_rt': averaged reaction time for this trial
 - 'human_rt_sem': standard error of the mean for human reaction times
 - 'human_rt_std': standard deviation of human reaction times
 - 'condition': name of image class (e.g., 'abstract1 contains abstract objects')
 - 'dataset': dataset this trial comes from (e.g., barensen, shapenet)
 - 'trial': name of this trial
 - 'n_subjects': number of participants who completed this specific trial
 - 'odddity_index': location of non-matching (i.e., the answer) object in image list
- Additionally, each row contains results from all models evaluated, e.g.
 - 'dinov2-giant_svm_avg': performance of dinov2-giant using linear
 - 'dinov2-giant_svm_std': standard deviation of dinov2-giant on this trial
 - 'dinov2-giant_svm_sem': standard error of the mean for dinov2-giant on trial
 - 'CLIP_ViT-B-32_svm_avg': as above, but for CLIP
 - ...
 - 'ViT-mae-base_svm_avg': as above, but for MAE

data/df_behavior_subject.csv

- original human data used to estimate performance saved in 'benchmark.csv'
- not averaged across trials, but preserves each participants choice behaviors
- necessary for reliability analyses

data/salienc_maps.pickle

- dictionary with salienc maps for all eye tracking data
- ['population'] [<imagenam>] each image's salienc map averaged across the group
- ['subject'] [<imagenam>] this images salienc maps for each person who saw it

data/df_behavior_wdistance.csv

- Same structure/data as benchmark.csv but with columns for modeling results from distance metrics (instead of a weighted linear readout) using dinov2-base, e.g.:
 - dino_distance_avg: averaged performance of all distance metrics
 - 'dino_distance_std', std of all distance metrics
 - 'dino_distance_sem', sem of all distance metrics
 - 'dino_l1_avg': performance estimated using an l1 distance metric on this trial
 - ...

data/croissant.json

- metadata for images/data in benchmark.csv using croissant format

MODELING AND VISUALIZATION SCRIPTS ARE IN scripts/

scripts/model_evaluation.ipynb

- main script for building the lightweight linear probe used in a model analysis
- requires path to df_behavior.csv
- requires path to images/

scripts/results.ipynb

- Main script to load data + generate all plots
- requires path to all data files

scripts/model_attention_analyses.ipynb

- scripts used to extract attention maps from dinov2 and relate to human gaze

scripts/visualize_data_and_singletrial.ipynb

- simple script to load data and visualize a single trial

scripts/relative_pose_analysis.ipynb

- script used to determine how pose variation relates to human/model performance

scripts/load_croissant.ipynb

- simple demo for loading metadata from croissant.json

LICENSING

The authors bear all responsibility in case of violation of rights, etc., and license these data under CC BY-NC-SA 4.0

HOSTING, LICENSING, AND MAINTENANCE PLAN

To ensure access to this benchmark we intend to host all data and images on Huggingface as a Huggingface dataset and will provide all necessary maintenance.

Upon acceptance, these data will be made accessible through Huggingface as well as a project page hosted on GitHub.

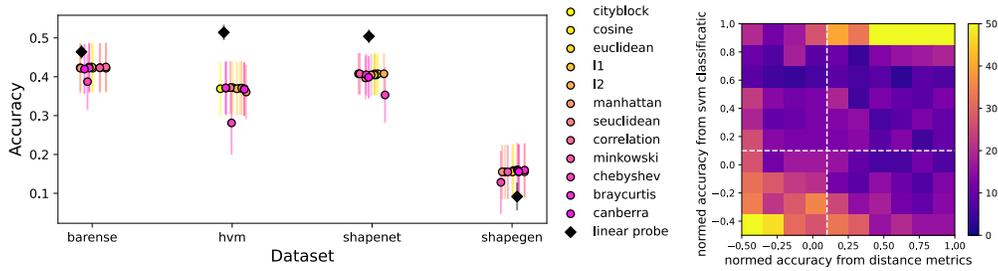


Figure 11:

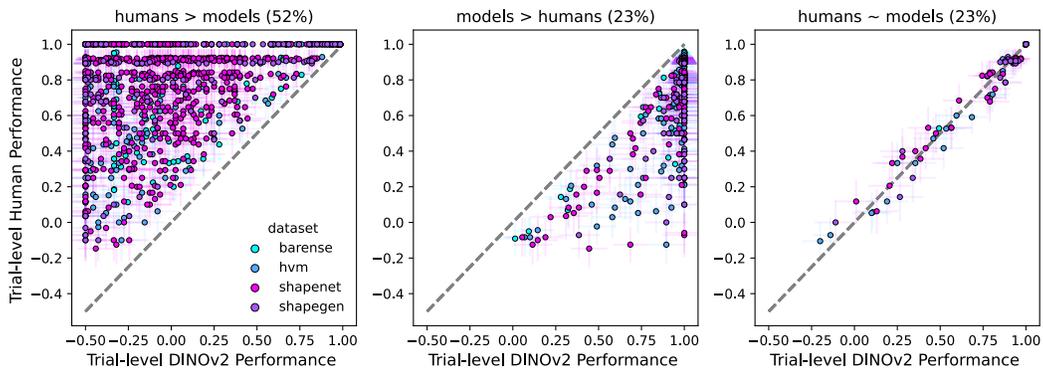


Figure 12: Direct comparison of trial-level human accuracy to the performance of DINOv2

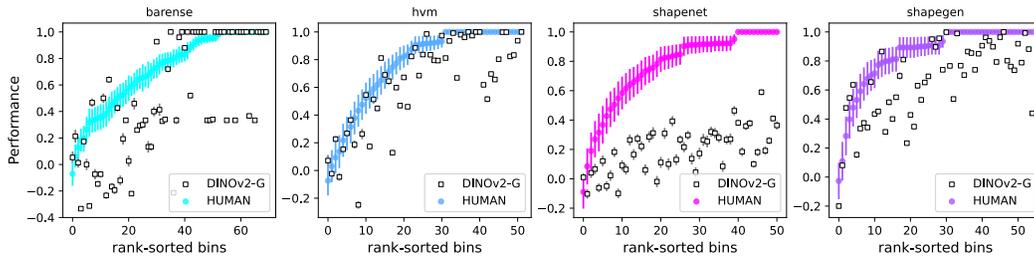


Figure 13:

387 **A.1 Online human data collection**

388 Human experimental data were collected online via Amazon Mechanical Turk and Prolific via
 389 experiments were implemented in JsPsych (De Leeuw, 2015). Each experiment began with an
 390 instruction phase, which introduced them to the task as well as provided 5 practice trials. This
 391 provided an opportunity for participants to acclimate themselves to the task and the controls. Once
 392 the experiment began, participants initiated the beginning of each trial with a button press (spacebar),
 393 such that they can (effectively) pause the experiment whenever they deem appropriate. This was
 394 designed to reduce environmental interference in the experiment. Experiments were designed to
 395 be completed in 10 minutes and participants were paid at a rate of roughly \$16/hour. In addition,
 396 participants were awarded a bonus commensurate with their performance, enabling them to earn up
 397 to twice the base pay. In order to ensure that participants were fairly compensated for their time,

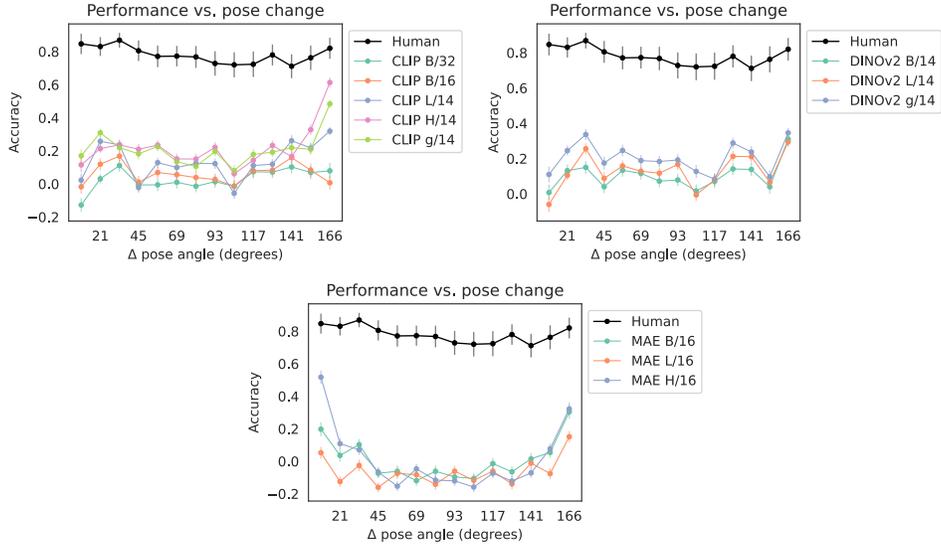


Figure 14: Visualizing viewpoint tolerance in humans and models across stimuli in shapenet

398 even in the case of a crowdsourcing platform errors, trial-by-trial data were collected throughout the
 399 experiment and stored on a custom server built from a Digital Ocean ‘droplet.’

400 We administer two related experimental designs. First, we use a 3-way concurrent visual discrimina-
 401 tion task commonly used to evaluate the role of MTC in perception (Barense et al., 2007; Buckley
 402 et al., 2001; Bussey et al., 2002). This design enables us to determine visual inferences that are
 403 possible with unlimited viewing time, as all stimuli remain on the screen for the duration of the trial.
 404 On each trial, participants are presented with three images and must identify the image that does not
 405 match the other two in terms of object identity (i.e., the ‘oddity’). Participants are given upwards of
 406 ten seconds to complete each trial. At any point in this duration, participants can select the oddity
 407 with a button press (right arrow, left arrow, or down arrow) corresponding to those locations on the
 408 oddity array. After this button press, participants are given feedback related to their performance on
 409 that trial, indicating whether their choice was correct or incorrect. If participants do not press a button
 410 in these ten seconds, the trial is marked as incorrect, feedback is given on the screen encouraging
 411 them to complete each trial within the allotted time.

412 A.2 Eye tracking data collection

413 Eye tracking was performed using an infrared video-based eye-tracker at 1000 Hz (Eyelink 1000; SR
 414 Research). Stimuli were displayed on a 22.5 inch VIEWPixx LCD display (resolution of 1900x1200,
 415 refresh rate of 120 Hz) and responses collected via keyboard. Other sources of light were minimized
 416 during data collection. The stimulus on the sample screen was presented at the central field of view
 417 and spanned up to 10 degrees of visual angle. This stimulus size was selected such that in order to
 418 collect high-acuity visual information from various stimulus locations, participants had to move their
 419 eyes (i.e., make a saccade). Stimuli on the match screen were the same size, but presented side by
 420 side, offset from the horizontal midpoint of the screen by 10 degrees of visual angle. Each experiment
 421 began with gaze calibration, then 5 practice trials to acclimate participants to the experimental setup.
 422 Each trial was initiated by the participants and began with participants maintaining fixation at the
 423 center of the screen (to perform drift correction at the beginning of each trial). Participants completed
 424 each trial at their own pace and there was a brief rest period every 5 minutes. This duration of this
 425 rest period was at the discretion of each participant. After this rest period, there was another gaze
 426 calibration, after which participants again completed a series of trials at their own pace as described
 427 above. For all gaze analyses (e.g., evaluating gaze reliability) we estimate gaze-related events (e.g.

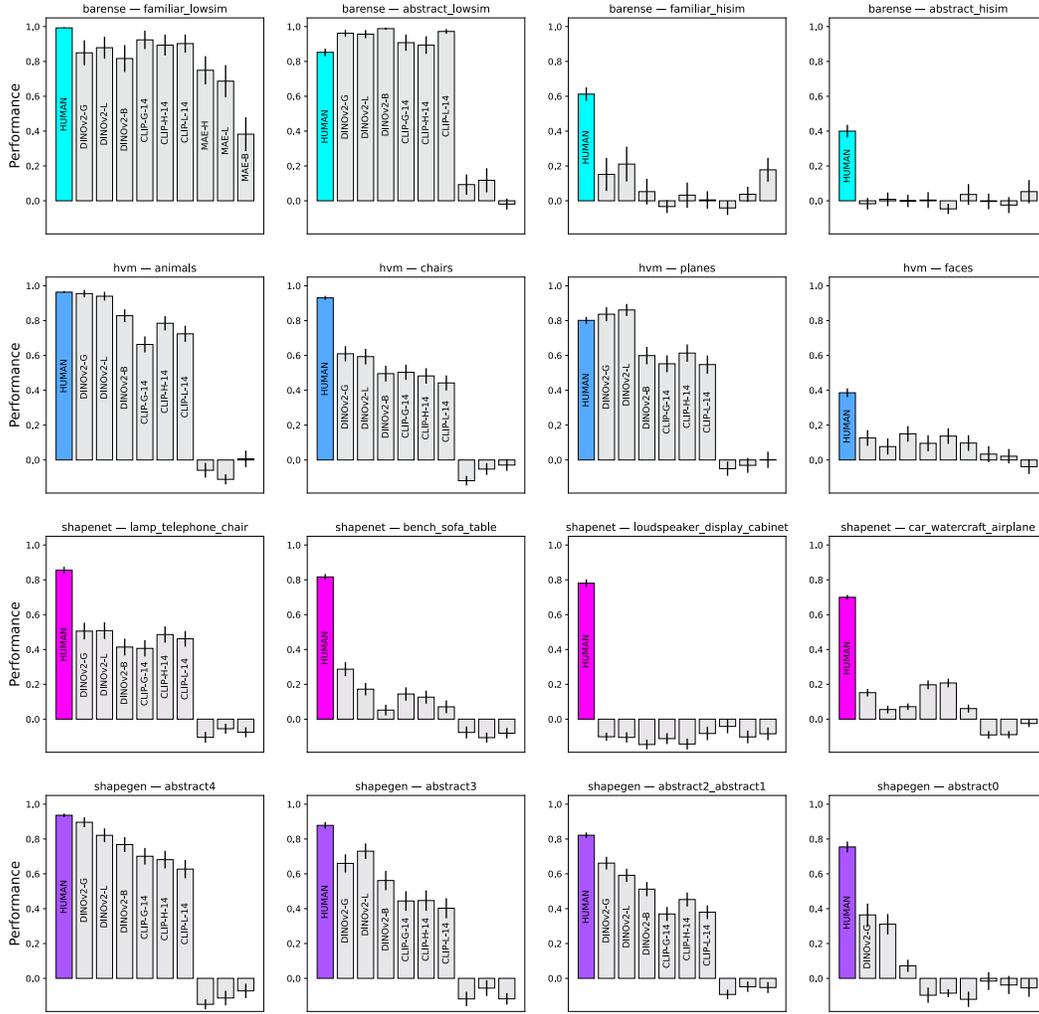


Figure 15: Comparing human performance to multiple vision models across all conditions.

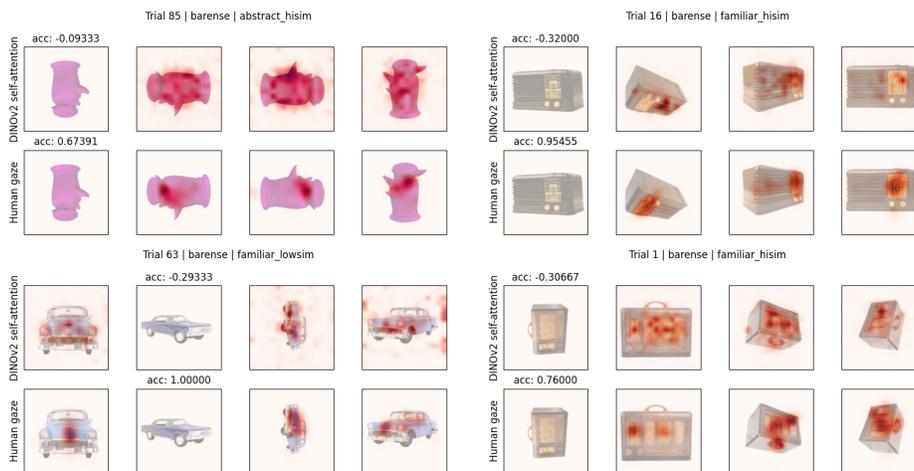


Figure 16: Example comparisons between DINOv2 attention maps and human attention maps.

428 fixations) directly from the raw gaze data using a standard python library (REMoDNaV; Nyström
429 and Holmqvist, 2010).

430 **A.3 Estimating gaze reliability**

431 We estimate the split-half reliability of in-lab gaze dynamics using the following protocol. First, for
432 each trial, a subject-level salience map is generated from the raw gaze behaviors: a 2D histogram
433 is generated from the raw time series, which is then smoothed with a Gaussian kernel. We note
434 that the results reported in this manuscript are robust to the resolution of the 2D histogram and size
435 of the smoothing kernel. This protocol yields a salience map for each image for each subject. We
436 then generate a random split of subjects and partition the salience maps for a given image using this
437 random subject split. We then average across participants in each random split, which results in
438 two salience maps, each corresponding to the random split of participants allotted to that half. We
439 then estimate the correlation between the two (random split-half) salience maps associated with this
440 image. We repeat this protocol for 100 random split-half permutations (i.e., generating a new shuffle
441 of participants each iteration). For each image, we then have a distribution of split-half correlations
442 which enables us to evaluate how similar participants viewed each image. To establish an empirical
443 null we compute the correlation between random splits corresponding the different images within
444 the same trial. Additionally, we estimate the bottom-up salience of each image (Itti et al., 1998) and
445 compute the correlation between this bottom-up salience map and the random splits associated with
446 each permutation of each image.