Explaining the Human Visual Brain 2019 competition and workshop

Romuald A. Janik

Jagiellonian University Kraków

RJ 1907.00950 [q-bio.NC]

Outline

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The Algonauts Project The competition fMRI MEG

My approach RDM peculiarities Effective receptive field Surrogate features Conclusions

The workshop Other approaches Some interesting talks

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The Algonauts Project

algonauts.csail.mit.edu

The quest to understand the nature of human intelligence and engineer more advanced forms of artificial intelligence are increasingly intertwined. The Algonauts Project brings biological and artificial intelligence researchers together on a common platform to exchange ideas and advance both fields...

Researchers at MIT, Freie Univ. Berlin, Singapore University of Technology and Design

Idea: Organize a competition on the borderline of neuroscience/ML and a subsequent workshop@MIT...

Another edition is planned for 2020...

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- ... apparently a very active research field...
- ... also slightly controversial however for higher levels of human visual processing, DNN features seem to be better than anything else

Ties in exactly with our Research Goal #11:

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- ▶ The idea is to show a set of images to a human/DNN
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- Use Spearman's correlation coefficient
- Take the off-diagonal components of each RDM and flatten them into a vector...
- Substitute the values by their ranks in each vector
- Compute an ordinary correlation coefficient of the rank vectors...
- Take the square... $\longrightarrow R^2$
- In the competition we have 15 human subjects, so 15 human RDM's

$$score(RDM_{DNN}) = \frac{1}{15} \sum_{i=1}^{15} R^2(RDM_{DNN}, RDM_i)$$

- The human RDM's differ between themselves so one cannot hope to get a perfect score...
- Normalize by a noise threshold

$$\frac{score(RDM_{DNN})}{score\left(\frac{1}{15}\sum_{i=1}^{15}RDM_{i}\right)}$$

How to compare different RDM's?

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fMRI and MEG tracks of the competition



N=15



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from Yalda Mohsenzadeh lecture

Data Structure



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from Yalda Mohsenzadeh lecture

General Linear Model: Constructing BOLD signals



from Yalda Mohsenzadeh lecture

11 / 28

Visual Recognition in the Brain



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from Yalda Mohsenzadeh lecture

12 / 28

MEG

Magnetoencephalography (MEG) / Electroencephalography (EEG)



from Yalda Mohsenzadeh lecture

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MEG – digression: an interesting application

Possible Neural Architectures



(King et al., 2016)

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(see also King, Dehaene 2014)

from Yalda Mohsenzadeh lecture

MEG - digression: an interesting application

A Neural Architecture with Recurrent Interactions



(Rajaei, Mohsenzadeh, Ebrahimpour, Khaligh-Razavi, 2019)

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from Yalda Mohsenzadeh lecture

Training datasets



image_09.jpg







image_10.jpg

image 03.jpg

image_11.jpg

image_19.ipg

image 77.ipg



image_18.jpc



mage 75.ipg









image 76.ip





image 28.ipg

image 78.ipp



image_017.jpg image_025.jpg

image_009.jpg



image_026.jpg

image_027.jpg



- The competition results were
- After the end of the 16 / 28







image_011.jpg

mage_019.jpg



image_003.jpg



image_010.jpd

image_018.ipg

Training datasets

image 02.ipc

image_10.jpg











image_17.jpg



image 25.ipg



Test dataset





image 03.jpg

image 11.ip



27 image 28.ipg

image 04.ipg

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image_009.jpg

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mage_002.ipg

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All 3 datasets are distinct

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image_59.jpg





mage 68.jp image 76.ip

image 52.jpg

image_60.jpg

image 77.ipg







image 78.ipp

Å. image_61.jpg

image 53.jpg









image_62.jpg





Training datasets

image 02.ipc

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image 26.ipc

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image_009.jpg

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image 53.ipc





image 04.ipg

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mage 70.ip



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mage_019.jpg



image_026.jpd

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image_027.jpg

mage_028.jpd

- All 3 datasets are distinct
- The competition results were based on the test dataset
- After the end of the competition the participants had to give predictions on a hidden test set (very similar to the test dataset) to check for overfitting 16 / 28

- I got 2^{nd} place in the MEG track and 3^{rd} in the fMRI track
- On the hidden test dataset, I got 2^{nd} place in the MEG track and 1^{st} in the fMRI track
- A requirement of the competition was to post a report on the arXiv (or biorxiv)
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Representational Dissimilarity Matrices (RDM) by construction have two rather unexpected and somewhat unwelcome features:

- They can miss a very strong discriminative signal (if correlated)
- ▶ They are influenced by irrelevant uninformative features...

$$1 - R(x, y) = 1 - \frac{(x - \langle x \rangle)(y - \langle y \rangle)}{\sigma_x \sigma_y}$$

 $x_i = +1 \qquad y_i = -1$

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This behavior (insensitivity to the global signal) can be countered by adding uninformative features...

▶ This effectively transforms Pearson RDM into cosine dissimilarity

$$1 - \frac{x \cdot y}{|x||y|}$$

 This modification significantly increases the scores... (average of NN activations is relevant for describing brain RDM's)

- ▶ To some extent, the constant level matters...
- The above suggests another (apart from cosine) possible modification of RDM definition:

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resnet50

block1 $256 \times 56 \times 56$ block2 $512 \times 28 \times 28$ block3 $1024 \times 14 \times 14$ block4 $2048 \times 7 \times 7$

Use adaptive_max_pool2d to reduce each layer to k imes k

IT: use 2×2 EVC, EARLY, LATE: 5×5

average pooling

- NN convolutional features partition the image into various resolutions
- At the same time, features become more higher level...

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Results **much worse** with average pooling

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Feature selection

- ▶ RDM is a "global" measure. Features cannot be assessed in isolation... → first pick some reference set...
- **Erase** or add a NN feature \longrightarrow see how the score changes.
- Try to avoid overfitting...

(choosing feature weights to maximize score on training dataset does not generalize...)

A For each of the 15 subjects individually evaluate the reference score and the modified score (with an added or erased feature). Then take the mean/z-score of the 15 differences.

B Randomly choose 30 subsets of 1/4 images and use these for the reference and modified scores. Take the mean/*z*-score of the 30 differences.

- For feature pruning use option A on CV test folds as well as on predictions on other dataset...
- For adding features, we used also a modification of B, with 10 different splits into 5 parts, and requiring positivity on both 118 and 92 datasets...
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- ► RDM is a "global" measure. Features cannot be assessed in isolation... → first pick some reference set...
- ▶ **Erase** or **add** a NN feature → see how the score changes..

Try to avoid overfitting...

(choosing feature weights to maximize score on training dataset does not generalize...)

A For each of the 15 subjects individually evaluate the reference score and the modified score (with an added or erased feature). Then take the mean/z-score of the 15 differences.

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B Randomly choose 30 subsets of 1/4 images and use these for the reference and modified scores. Take the mean/z-score of the 30 differences.

For feature pruning use option A on CV test folds as well as on predictions on other dataset...

 For adding features, we used also a modification of B, with 10 different splits into 5 parts, and requiring positivity on both 118 and 92 datasets...
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This feature selection procedure (option A) can also be used to study the importance of parts of receptive fields (maxpool2 of vgg19 on the 118 image dataset) (positive values bad)

We erase corners in EARLY and EVC... The score increases also on the test dataset..

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We erase corners in EARLY and EVC...

Solutions for EVC and EARLY MEG are very simple...

EVC

- 1. block2 of resnet18 4.26
- **2.** reduce to 5×5 ; extend by 0.2 6.41/2
- eliminate 1/4 of worst features (algorithm B) 25.23
- 4. eliminate corners; $0.2 \rightarrow 0.0$ 26.90/
- 5. add best features (enhanced 2×) 28.29
- add best features from maxpool2 of vgg19 (shrunk 0.5×)

Score: 28.40

Erronously adding worst features from other layers instead of $\mathbf{5.+6}$, gave the best score: 32.68

EARLY

- 1. maxpool2 of vgg19
- **2.** reduce to 5×5 , extend by 0.5
- eliminate bad features (z > 0.15 on either dataset, algorithm A)
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6.41/24.01

- **3.** eliminate 1/4 of worst features
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Solutions for IT and LATE – surrogate features

Question: What (abstract) features would reproduce the given brain RDM (averaged across subjects)? (as measured by Spearman's...)

Use Multidimensional Scaling (MDS):

 $\overline{\textit{MDS}}_{118\times118}^{4}\longrightarrow \mathbb{R}^{118\times10}$

repeat with 10 random seeds

using the constructed 100 features gives a score around 77% for the same dataset

General procedure:

- 1. Fit the resulting 100 features with NN featuresfit for each layer individually, then combine fits using ridge regression
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Question: What (abstract) features would reproduce the given brain RDM (averaged across subjects)? (as measured by Spearman's...)

Use Multidimensional Scaling (MDS):

$$\overline{\textit{MDS}}^4_{118\times 118} \longrightarrow \mathbb{R}^{118\times 10}$$

repeat with 10 random seeds

using the constructed 100 features gives a score around 77% for the same dataset

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- 1. Use resnet50, convolutional features reduced to 2×2
- For 118 dataset MDS features: ridge regression; OMP(6) For 92 dataset MDS features: OMP(7)
- **3.** Concatenate to get 300 features
- 4. Prune bad features imposing positivity on 118 dataset
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- Add in 75+75 ICA from block1, block3 of resnet34

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 - Iots of NN features versus small number of images
 - The three datasets were quite distinct...
- Sometimes CV, as well as assessment of feature importance, was not reliable
- Try to stick to simple models...
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