Anatomically Constrained ResNets Exhibit Opponent Receptive Fields; So What?

Abstract

Primate visual systems are well known to exhibit varying degrees of bottlenecks
in the early visual pathway. Recent works have shown that the presence of a bot-
tleneck between 'retinal' and 'ventral' parts of artificial models of visual systems,
simulating the optic nerve, can cause the emergence of cellular properties that have
been observed in primates: namely centre-surround organisation and opponency.
To date, however, state-of-the-art convolutional network architectures for classifica-
tion problems have not incorporated such an early bottleneck. In this paper, we ask
what happens if such a bottleneck is added to a ResNet-50 model trained to classify
the ImageNet data set. Do the emergent properties observed in simpler models
still appear in these considerably deeper and more complex models? Does the
bottleneck affect performance or robustness? Does the bottleneck bias the model
in some way? or, does something else happen?

1 Introduction and Motivation

Convolutional Neural Networks (CNNs) with a bottleneck designed to mimic the anatomical con-
straint imposed by the optic nerve have recently been shown to learn representations which more
closely align with those found in primate visual systems. Specifically, Lindsey et al. [2019] showed
that a reduction in the number of channels or convolutional neurons in the second layer of a CNN
gives rise to centre-surround receptive fields in the first two layers followed by orientation selectivity
in subsequent layers. Lindsey et al.'s experiments were based on a relatively simple model of the
visual system (aka 'Retina-Net') constructed with a model of the retina (the first two layers up to
the bottleneck), a model of the ventral stream (stacked convolutional layers) and a pair of fully
connected layers to produce a classification. These models were trained on a greyscale version of the
CIFAR-10 data set [Krizhevsky et al., 2009]. Harris et al. [2019, 2020] subsequently showed that
the same network architecture (albeit with colour input) learns a simple opponent colour code in the
bottleneck layer followed by a reduction in colour opponency in later layers. Harris et al. [2020]
further demonstrated that the general trend that the bottleneck induces opponency was true across a
range of data sets from CIFAR-10 to ImageNet [Russakovsky et al., 2015].

Modern neural networks for solving large-scale classification problems like ImageNet look very
different to the aforementioned 'Retina-Net' and its colour variant, and do not incorporate any form
of bottleneck near the input layer. At the same time, other recent research has demonstrated that there
may be benefits to better modelling of the visual system in state-of-the-art models. For example,
Geirhos et al. [2019] show that CNNs trained on ImageNet are biased towards texture information in
contrast to Human observers who make classifications predominantly according to shape information.
They subsequently show that promoting a shape bias in trained models can improve both accuracy
and robustness. Dapello et al. [2020] show that simulating the function performed by the primary
visual cortex over the input increases robustness to adversarial attacks in which images which have

Submitted to 2nd Workshop on Shared Visual Representations in Human and Machine Intelligence (SVRHM).

Figure 1: Anatomically constrained ResNet-50 model diagram. The only modification to the ResNet-50 architecture is that it expects an input with N_{BN} channels. We trained such models on the ImageNet data set for $N_{BN} \in \{1, 2, 4, 8, 16, 32\}$.

been imperceptibly (to a human) adjusted such that the classification given by the trained network is no longer correct. Their model for simulating V1 (V1Block) is based on the classical observations (for example by Hubel and Wiesey [2004]) that the receptive fields of cells in V1 are orientation selective (modelled as Gabor filters) and that these cells can be described as either simple or complex (modelled with different acitvation functions). Parameters for the Gabor filters in the V1Block are sampled according to known distributions from the neuroscience literature rather than being learned. The model also assumes that the input to the V1Block is just the normalised pixel values (subtract 0.5 and divide by 0.5 for each RGB colour channel) and eschews any notion of processing performed by the retina itself, or by the Lateral Geniculate Nucleus (LGN).

In particular, since convolutional neurons already reflect many of the functional properties of cells found in the early visual system, it is prudent to ascertain whether more simple modifications, such as the bottleneck explored by Lindsey et al. [2019], Harris et al. [2019, 2020], can be used to the same effect. This paper starts to ask what happens if we place a bottleneck at the beginning of a performant classification network trained on the ImageNet data set. Does the bottleneck induce similar emergence of primate-like cell types? How does the bottleneck affect classification accuracy of the networks? Does a bottleneck in this setting increase robustness of the model to adversarial attacks or does it induce some kind of bias in the function performed by the network?

2 Early Visual Representations in Anatomically Constrained ResNets

In order to add a bottleneck to an existing model, we propose pre-pending the 'Retina' portion of the model from Lindsey et al. [2019]. This consists of two convolutional layers with 32 channels and N_{BN} channels respectively, where N_{BN} is the bottleneck width. Unlike the model of Lindsey et al. [2019] which uses a kernel size of 9, we choose the kernel size to ensure a large enough receptive field whilst preserving consistency with the first layer of the target model where possible (e.g. 7 for a ResNet). Padding is chosen such that the resolution of the output is the same as the resolution of the input. In this way the bottleneck acts as a kind of parameterised pre-processing of the input. We trained constrained variants of a ResNet-50 (depicted in Figure 1) for the range of bottleneck widths used by Lindsey et al. [2019] ($N_{BN} \in \{1, 2, 4, 8, 16, 32\}$) on the ImageNet Large Scale Visual Recognition Challenge (ILSVRC) [Russakovsky et al., 2015] data set. We trained three repeats of each model variant, giving 18 models in total. The training regime is described in Appendix D.

We first computed the distribution of opponent cell types following the approach from Harris et al. [2019, 2020] in the second retina layer (Retina-2) and first ResNet layer (ResNet-1). This approach consists of presenting a set of stimuli which vary in hue to the cell in order to obtain a hue response curve. The cell is colour opponent if the response curve crosses the baseline response of the cell to a zero stimulus. Interestingly, our results showed that networks with $N_{BN} = 1$ have no colour opponent cells in either of the layers we studied. This contrasts with the finding of Harris et al. [2019, 2020], who showed that almost all cells in the retina layers of their more constrained networks were colour opponent. We presume that the range of colour stimuli in ImageNet is sufficiently broad that when restricted to one channel, better performance can be obtained with a luminance opponent encoding than a colour opponent one. To test this, we additionally present a set of stimuli which vary

(a) Retina-2 opponent cell distribution

(b) ResNet-1 opponent cell distribution

Figure 2: Colour and luminance opponency distributions in the Retina-2 and ResNet-1 layers of anatomically constrained ResNet-50 models.

Figure 3: Super stimuli for various classes of fruit for networks with $N_{BN} = 1$ and $N_{BN} = 32$ obtained by gradient ascent in Fourier space following Olah et al. [2017]. Even when the networks are restricted to a single channel ($N_{BN} = 1$), they still learn a strong representation of colour.

in luminance (i.e. full field grey-scale stimuli with values ranging from zero to one) and perform the same process of comparing to the baseline (response of the cell to constant stimulus of 0.5 for this experiment) to infer luminance opponency. Note that any cell whose response is linear in the sense that extremes of input value correspond to extremes of response will be classified as luminance opponent. It is therefore appropriate to say that any cells which are not luminance opponent are to some degree non-linear.

Figures 2a and 2b give the results for this experiment in the Retina-2 and ResNet-1 layers respectively. The results show that the vast majority of colour opponent cells are also luminance opponent, particularly in the more constrained networks. In the Retina-2 layer, there is an increase in cells which are luminance opponent, but not colour opponent, for networks with $N_{BN} < 4$. This in turn suggests an increase in the linearity of cell response for these networks, which accords with the findings of Lindsey et al. [2019], Harris et al. [2020]. Almost all cells in the first ResNet layer are luminance opponent. In both layers, approximately half of the cells are also colour opponent in networks with all but the tightest bottlenecks, where the percentage of colour opponent cells drops to zero. Ultimately, the pattern in the colour opponent cell distribution shown by Harris et al. [2019, 2020] is replaced with a similar pattern in luminance opponent cell distribution in these networks. We explore these cells further and find some evidence that they exhibit centre-surround organisation in Appendix A.

In the neuroscience literature, the distinction between luminance opponency and colour opponency is often unclear. For example, it has been argued that many of the cells which preferred achromatic stimuli in Monkey V1, described by Hubel and Wiesel [1968], may be better described as colour opponent but with imbalanced cone response such that they are also responsive to luminance [Johnson et al., 2001, Lennie et al., 1990, Schluppeck and Engel, 2002, Shapley and Hawken, 2011]. We therefore must ascertain whether the luminance opponent cells are still able to convey valuable information regarding the colour of the input. To do this, we propose performing gradient ascent

Figure 4: The shape bias (following Geirhos et al. [2019]) across bottleneck widths.

Figure 5: Top-1 accuracy on the ILSVRC validation set across different bottleneck widths.

on a random input in Fourier space, following Olah et al. [2017], to obtain super stimuli which the networks classify with high confidence as a range of classes which are characterised by a particular colour. Figure 3 shows these stimuli for a range of fruit classes in the ImageNet data set for models with $N_{BN} = 1$ and N_{BN}. From the figure we can see that the networks with a single ($N_{BN} = 1$) luminance opponent channel in the bottleneck layer still make use of colour to inform their decisions, learning the characteristic red of a strawberry, yellow of a lemon, green of an apple, and so on.

It is clear from Figure 3 that neither of the models is particularly biased towards shape in that the super stimuli have appropriate textures and colours but not shapes. To quantify this, we now measure the shape bias of the models. Following Geirhos et al. [2019], we evaluate the models on the 'StylizedImageNet' data set [1], which contains images with a texture-shape cue conflict, requiring networks to recognise objects based on shape rather than texture. Figure 4 shows that networks with tighter bottlenecks exhibit a small increase in shape bias. The model with a bottleneck width of 1 trained on ImageNet achieves the highest shape bias, outrunning a standard (i.e. without our modification) ResNet-50 model. Note, however, that the shape bias of all of the models is far lower than the models trained with the approach from Geirhos et al. [2019].

3   What Effect Does a Bottleneck Have on Performance?

In this section we explore whether the changes in function induced by the bottleneck correspond to any tangible improvement in performance. The most obvious first step is to consider the effect of the bottleneck on the accuracy of the model. Figure 5 shows that networks with bottleneck width of 4 or greater all achieve validation set performance that is within the margin of error (one standard deviation) of the others and that networks with tighter bottlenecks show a small but clear drop in performance. Importantly, all of the networks perform within 2% of the ResNet-50 without our modification compared to the pre-trained version provided in torchvision (76.15%), which has minor differences in the training procedure that might explain this. This contrasts with the networks considered by Dapello et al. [2020] which show a larger reduction of around 4.5% reaching an accuracy of 71.7% (although again note that there are minor differences in the training procedure).

Robustness to natural adversarial examples   We now consider robustness to the ImageNet-A data set [Hendrycks et al., 2019], which contains 7500 naturally occurring images which are consistently mis-classified by pre-trained models. The images correspond to 200 ImageNet classes in which normal models (ResNet50, VGG16, etc) have a top-1 accuracy of more than 90% using the ImageNet validation data set. The ImageNet-A data set typically results in a top-1 accuracy of around 2-3% on the same models. Figure 6 shows the top-1 accuracy of our models on the ImageNet-A data set. Figure 11 from the appendix shows the Area Under the Response Rate Accuracy curve (AURRA) which measures how calibrated the models are (how a model's confidence in its prediction relates to its accuracy). Both of these figures illustrate that the tighter bottlenecks do induce slightly better performance on this task.

Robustness to L_∞ constrained artificial attacks   To study a more general notion of adversarial robustness, we follow the guidelines given by Carlini et al. [2019]. We use FoolBox [Rauber et al., 2017, 2020] to compute the worst case performance of the models under a range of attacks. We

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[1] Available from `https://github.com/rgeirhos/Stylized-ImageNet`

Figure 6: Robustness to natural adversarial examples from the ImageNet-A data set [Hendrycks et al., 2019] as a function of bottleneck width.

Figure 7: Robust accuracy of different bottlenecks (line styles) against L_∞ adversarial attack budgets.

explore a range of perturbations using different L_∞ thresholds between 0 and 1.0. The attacks include Project Gradient Descent, which was utilised by Dapello et al. [2020] to demonstrate improved robustness of their model. A full description of the specific attacks is given in Appendix C. Figure 7 shows the robust accuracy, the worst-case performance following all attacks, of the models across different L_∞ perturbation budgets. Results were computed by sampling 10000 images from the ImageNet validation set uniformly for each class. The results show that the increased opponency and small increase in shape bias actually correspond to a decrease in robustness to targeted adversarial examples. In Appendix B, we further show that the CIFAR-10 trained models from Harris et al. [2019, 2020] show no significant difference in adversarial robustness as a function of bottleneck.

4 Discussion

In this paper we have explored the addition of a bottleneck to ImageNet trained ResNet-50 models. We have shown that cells in the bottleneck layer of models with tight bottlenecks learn opponent receptive fields in accordance with the CIFAR-10 results of Harris et al. [2019, 2020]. In contrast to those results, the opponent cells we find here are best described as luminance opponent rather than colour opponent. That said, we have shown that these luminant opponent cells still exhibit some form of spectral response and that, even in the extreme case of $N_{BN} = 1$, the model is able to learn about colour. Whether the cells in networks with tight bottlenecks have centre-surround receptive fields of the kind observed by Lindsey et al. [2019] is unclear, although there is some evidence that they do. It is, however, clear that the Retina-2 cells respond more linearly in networks with tight bottlenecks, echoing the finding of Lindsey et al. [2019]. We have further shown that these differences in function correspond to a small but consistent increase in the shape bias of the networks.

Our investigation into the performance of the different bottlenecked models is surprising in the sense that it perhaps asks more questions than it answers. At the outset our hypothesis was that the bottleneck might encourage both shape bias as well as adversarial robustness; the typical nature of centre-surround opponent cells observed in prior studies would for example suggest that the early network would perform isotropic band-pass filtering (like a Laplacian of Gaussian) which one presumes would naturally bias to towards shape representations[2]. In terms of accuracy, the different bottlenecks only have a mild effect with the tightest bottlenecks (below 8) having slightly worse performance. The results of testing the models on natural adversarial examples indicates that the tightest bottlenecks do perform better on the ImageNet-A data set (both in terms of accuracy and calibration). However, it is unclear whether this increase corresponds to an increase in robustness or merely indicates that the type of function learned differs for tighter bottlenecks. In any case, the worst case accuracy, computed after a set of targeted adversarial attacks, shows that adversarial robustness actually decreases slightly as the bottleneck is tightened.

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[2]We have explicitly observed this in unpublished work by one of our students which demonstrated that putting an explicit fixed Difference of Gaussian filter before the network could encourage shape bias.

References

Nicholas Carlini, Anish Athalye, Nicolas Papernot, Wieland Brendel, Jonas Rauber, Dimitris Tsipras, Ian Goodfellow, Aleksander Madry, and Alexey Kurakin. On evaluating adversarial robustness. *arXiv preprint arXiv:1902.06705*, 2019.

Joel Dapello, Tiago Marques, Martin Schrimpf, Franziska Geiger, David D Cox, and James J DiCarlo. Simulating a Primary Visual Cortex at the Front of CNNs Improves Robustness to Image Perturbations. *BioRxiv preprint*, 2020.

Robert Geirhos, Patricia Rubisch, Claudio Michaelis, Matthias Bethge, Felix A. Wichmann, and Wieland Brendel. Imagenet-trained CNNs are biased towards texture; increasing shape bias improves accuracy and robustness. In *International Conference on Learning Representations*, 2019. URL https://openreview.net/forum?id=Bygh9j09KX.

Ian Goodfellow, Jonathon Shlens, and Christian Szegedy. Explaining and harnessing adversarial examples. In *International Conference on Learning Representations*, 2015. URL http://arxiv.org/abs/1412.6572.

Ethan Harris, Daniela Mihai, and Jonathon Hare. Spatial and Colour Opponency in Anatomically Constrained Deep Networks. In *Workshop on Shared Visual Representations in Humans and Machines (SVRHM)*, 2019. URL https://arxiv.org/abs/1910.11086.

Ethan Harris, Daniela Mihai, and Jonathon Hare. How Convolutional Neural Network Architecture Biases Learned Opponency and Colour Tuning. *Neural Computation*, 2020. URL https://arxiv.org/abs/2010.02634.

Dan Hendrycks, Kevin Zhao, Steven Basart, Jacob Steinhardt, and Dawn Song. Natural adversarial examples. *arXiv preprint arXiv:1907.07174*, 2019.

David H Hubel and Torsten N Wiesel. Receptive fields and functional architecture of monkey striate cortex. *The Journal of physiology*, 195(1):215–243, 1968.

David H. Hubel and Torsten N. Wiesel. *Brain and visual perception: the story of a 25-year collaboration*. Oxford University Press, 2004.

Elizabeth N. Johnson, Michael J. Hawken, and Robert Shapley. The spatial transformation of color in the primary visual cortex of the macaque monkey. *Nature neuroscience*, 4(4):409, 2001.

Alex Krizhevsky, Geoffrey Hinton, et al. Learning multiple layers of features from tiny images. 2009.

Alexey Kurakin, Ian Goodfellow, and Samy Bengio. Adversarial examples in the physical world. *ICLR Workshop*, 2017. URL https://arxiv.org/abs/1607.02533.

Peter Lennie, John Krauskopf, and Gary Sclar. Chromatic mechanisms in striate cortex of macaque. *Journal of Neuroscience*, 10(2):649–669, 1990.

Jack Lindsey, Samuel A Ocko, Surya Ganguli, and Stephane Deny. A Unified Theory of Early Visual Representations from Retina to Cortex through Anatomically Constrained Deep CNNs. In *International Conference on Learning Representations*, 2019. URL https://openreview.net/forum?id=S1xq3oR5tQ.

Aleksander Madry, Aleksandar Makelov, Ludwig Schmidt, Dimitris Tsipras, and Adrian Vladu. Towards deep learning models resistant to adversarial attacks. In *International Conference on Learning Representations*, 2018. URL https://openreview.net/forum?id=rJzIBfZAb.

S. Moosavi-Dezfooli, A. Fawzi, and P. Frossard. Deepfool: A simple and accurate method to fool deep neural networks. In *2016 IEEE Conference on Computer Vision and Pattern Recognition (CVPR)*, pages 2574–2582, 2016.

Chris Olah, Alexander Mordvintsev, and Ludwig Schubert. Feature visualization. *Distill*, 2(11):e7, 2017.

Jonas Rauber, Wieland Brendel, and Matthias Bethge. Foolbox: A python toolbox to benchmark the robustness of machine learning models. In *Reliable Machine Learning in the Wild Workshop, 34th International Conference on Machine Learning*, 2017. URL `http://arxiv.org/abs/1707.04131`.

Jonas Rauber, Roland Zimmermann, Matthias Bethge, and Wieland Brendel. Foolbox native: Fast adversarial attacks to benchmark the robustness of machine learning models in pytorch, tensorflow, and jax. *Journal of Open Source Software*, 5(53):2607, 2020. doi: 10.21105/joss.02607. URL `https://doi.org/10.21105/joss.02607`.

Olga Russakovsky, Jia Deng, Hao Su, Jonathan Krause, Sanjeev Satheesh, Sean Ma, Zhiheng Huang, Andrej Karpathy, Aditya Khosla, Michael Bernstein, et al. ImageNet Large Scale Visual Recognition Challenge. *International journal of computer vision*, 115(3):211–252, 2015.

Denis Schluppeck and Stephen A Engel. Color opponent neurons in v1: a review and model reconciling results from imaging and single-unit recording. *Journal of vision*, 2(6):5–5, 2002.

Robert Shapley and Michael J. Hawken. Color in the Cortex: single- and double-opponent cells. *Vision Research*, 51(7):701 – 717, 2011. ISSN 0042-6989. doi: https://doi.org/10.1016/j.visres.2011.02.012. URL `http://www.sciencedirect.com/science/article/pii/S0042698911000526`. Vision Research 50th Anniversary Issue: Part 1.

Figure 8: Grey-scale and colour receptive field visualisations for networks with $N_{BN} = 1$ and $N_{BN} = 32$.

Figure 9: Cell responses to an example input for the same cells used in Figure 8 from networks with $N_{BN} = 1$ and $N_{BN} = 32$.

A   Receptive Field Visualisations

Figure 8 shows the receptive field visualisations in grey-scale and colour for cells in the Retina-2 layer of networks with $N_{BN} = 1$ and $N_{BN} = 32$. These are computed by taking the gradient of the response of the centre neuron to a blank stimulus (constant value of zero) and then applying min / max normalisation to obtain an image. The results show that the cells in networks with $N_{BN} = 1$ are somewhat centre-surround, but with quite a small extent. In contrast, the cells in networks with $N_{BN} = 32$ have much smoother receptive fields that are sensitive to changes over a much larger region of the input. Although it is true that we find exclusively centre-surround cells in the networks with $N_{BN} = 1$, we also find some centre-surround cells in the networks with $N_{BN} = 32$.

To try to better understand the functions performed by these cells, we plot the response of each cell to an example input in Figure 9. The results show that when $N_{BN} = 1$, the Retina-2 cells preserve most of the edge information in the image (with perhaps a mild edge enhancement effect) and respond in accordance with the brightness of the input (albeit inverted for two of the cells). In contrast, the Retina-2 cells in networks with $N_{BN} = 32$ distort the input heavily, performing specific colour opponent edge detection functions.

Figure 10: Robust accuracy of the CIFAR-10 trained models from Harris et al. [2019, 2020] with different bottleneck widths (line styles) against L_∞ adversarial attack budgets.

Figure 11: Model calibration, Area Under the Response Rate Accuracy curve (AURRA), on ImageNet-A [Hendrycks et al., 2019] as a function of bottleneck width. Networks with tighter bottlenecks exhibit improved calibration.

B CIFAR-10 Adversarial Robustness

To determine the adversarial robustness of the models trained on CIFAR-10 from Harris et al. [2019, 2020], we perform the same suite of attacks as in our ImageNet experiments (outlined in Appendix C) but over the whole CIFAR-10 test set. Figure 10 shows the worst case accuracy for these models following all attacks. The results again show that no significant difference is obtained through the introduction of a bottleneck.

C Artificial Adversarial Attacks

We use the following adversarial attacks with L_∞ perturbation budgets between 0.0 and 1.1. Each attack uses the FoolBox [Rauber et al., 2020] implementation with default parameters.

- Fast Gradient Sign Method [Goodfellow et al., 2015]
- Projected Gradient Descent [Madry et al., 2018]
- Basic Iterative Method [Kurakin et al., 2017]
- Additive Uniform Noise
- DeepFool [Moosavi-Dezfooli et al., 2016]

D Model Training

Models were trained on the ImageNet 2012 training set on nodes equipped with four NVidia Quadro
RTX 8000 GPUs. The following setup was used:

Batch size:	1024	
Optimiser:	SGD with the following:	
	initial lr:	0.1
	momentum:	0.9
	weight decay:	1e-4
	schedule:	lr drops by factor of 10 at 30 and 60 epochs
	epochs:	90
Data Augmentation:	Random resized cropping to 224x224	
	random horizontal flipping	
	normalisation using standard ImageNet mean and s.d.	

Code and all trained models will be made available following de-anonymisation of the manuscript.