The face module emerges from domain-general visual experience: a deprivation study on deep convolution neural network

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Abstract

Can faces be accurately recognized with zero experience on faces? The answer to this question is critical because it examines the role of experiences in the formation of domain-specific modules in the brain. However, thorough investigation with human and non-human animals on this issue cannot easily dissociate the effect of the visual experience from that of genetic inheritance, i.e., the hardwired domain-specificity. The present study addressed this problem by building a model of selective deprivation of the experience on faces with a representative deep convolutional neural network (DCNN), AlexNet. We trained a new AlexNet with the same image dataset, except that all images containing faces of human and nonhuman primates were removed. We found that the experience-deprived AlexNet (d-AlexNet) did not show significant deficits in face categorization and discrimination, and face-selective modules also automatically emerged. However, the deprivation made the d-AlexNet to process faces in a more parts-based fashion, similar to the way of processing objects. In addition, the face representation of the face-selective module in the d-AlexNet was more distributed and the empirical receptive field was larger, resulting in less degree of selectivity of the module. In sum, our study provides undisputable evidence on the role of nature versus nurture in developing the domain-specific modules that domain-specificity may evolve from non-specific stimuli and processes without genetic predisposition, which is further fine-tuned by domain-specific experience.

Keywords: face perception, face domain, deep convolutional neural network, visual deprivation, experience

1   Introduction

A fundamental question in cognitive neuroscience is how nature and nurture form our cognitive modules. In the center of the debate is the origin of face recognition ability. Numerous studies have revealed both behavioral and neural signatures of face-specific processing, indicating a face module in the brain (for reviews, see Freiwald, Duchaine, & Yovel, 2016; Kanwisher & Yovel, 2006). Further studies from behavioral genetics revealed the contribution of genetics on the development of the face-specific recognition ability in humans (Wilmer et al., 2010; Zhu et al., 2010). Collectively, these studies suggest an innate domain-specific module for face cognition. However, it is unclear whether the visual experience is also necessary for the development of the face module.

A direct approach to address this question is visual deprivation. Two studies on monkeys selectively deprived the visual experience of faces since birth, while leaving the rest of experiences untouched (Arcaro, Schade, Vincent, Ponce, & Livingstone, 2017; Sugita, 2008). They report that face-deprived monkeys are still capable of categorizing and discriminating faces (Sugita, 2008), though less prominent in selectively looking preference to faces over non-face objects (Arcaro et al., 2017). Further examination of the brain of the experience-deprived monkeys fails to localize typical face-selective cortical regions with the standard criterion; however, in the inferior temporal cortex where face-selective regions are normally localized, face-selective activation (i.e., neural responses to faces larger than nonface objects) is observed (Arcaro et al., 2017). Taken together, without visual experiences of faces, rudimental functions to process faces may still evolve to some extent.

Two related but independent hypotheses may explain the emergence of the face module without face experiences. An intuitive answer is that the rudimental functions are hardwired in the brain by genetic predisposition (McKone, Crookes, Jeffery, & Dilks, 2012; Wilmer et al., 2010). Alternatively, we argue that the face module may emerge from experiences on nonface objects and related general-purpose processes, because representations for faces may be constructed by abundant features derived from nonface objects. Unfortunately, studies on humans and monkeys are unable to thoroughly decouple the effect of nature and nurture to test these two hypotheses.

Recent advances in deep convolutional neural network (DCNN) provide an ideal test platform to examine the role of visual experiences alone on face modules without genetic predisposition. Previous studies have shown that DCNNs are similar to human visual cortex both structurally and functionally (Kriegeskorte, 2015), but free of any predisposition on functional modules. Therefore,

with DCNNs we can manipulate experiences without considering interactions from genetic

predisposition. In this study, we asked whether DCNNs can achieve face-specific recognition ability

when visual experiences on faces were selectively deprived.

To do this, we trained a representative DCNN, AlexNet (Krizhevsky, 2014; Krizhevsky,

Sutskever, & Hinton, 2012), to categorize nonface objects with face images carefully removed from

the training dataset. Once this face-deprived DCNN (d-AlexNet) was trained, we compared its

behavioral performance to that of a normal AlexNet of the same architecture but with faces present

during training in both face categorization (i.e., differentiating faces from nonface objects) and

discrimination (i.e., discriminating faces among different individuals) tasks. We predicted that the d-

AlexNet, though without predisposition and experiences of faces, may still develop face selectivity

through its visual experiences of nonface objects.

2   Materials and methods

2.1   Stimuli

Deprivation dataset The deprivation dataset was constructed to train the d-AlexNet. It was based on

the ImageNet Large-Scale Visual Recognition Challenge (ILSVRC) 2012 dataset (Deng et al., 2009),

which contains 1,281,167 images for training and 50,000 images for validation, in 1000 categories.

These images were first subjected to automated screening with an in-house face-detection toolbox

based on VGG-Face (Parkhi, Vedaldi, & Zisserman, 2015), and then further screened by two human

raters, who separately judged whether a given image contains faces of humans or non-human

primates regardless of the orientation and intactness of the face, or anthropopathic artwork, cartoons,

and artifacts. We removed images judged by either rater as containing any above-mentioned contents.

Finally, we removed categories whose remaining images were less than 640 images (approximately

half of the original number of images in a category). The resultant dataset consists of 736 categories,

with 662,619 images for training and 33,897 for testing the performance.

Classification dataset To train a classifier that can classify faces, we constructed a classification

dataset consisting of 204 categories of non-face objects and one face category, each of 80 exemplars.

For the non-face categories, we manually screened Caltech-256 (Griffin, Holub, & Perona, 2007) to

remove images containing human, primate, or cartoon faces, and then removed categories whose

remaining images were less than 80. In each of the 204 remaining non-face categories, we randomly

chose 70 images for training and another 10 for calculating classification accuracy. The face category

was constructed by randomly selecting 1000 faces images from Faces in the Wild (FITW) dataset

(Berg, Berg, Edwards, & Forsyth, 2005). Among them, 70 was used as training data and another 10

for classification accuracy. In addition, to characterize DCNN's ability in differentiating faces from

object categories, we compiled a second dataset consisting of all images in the face category except

those used in training.

Discrimination dataset To train a classifier that can discriminate faces at individual level, we

constructed a discrimination dataset consisting of face images of 133 individuals, 300 images each,

selected from the Casia-WebFace database (Yi, Lei, Liao, & Li, 2014). For each individual in the

dataset, 250 were randomly chosen for training and another 50 for calculating discrimination

accuracy.

Representation dataset To examine representational similarity of faces and non-face images

between the d-AlexNet and the normal one, we constructed a representation dataset with two

categories, faces and bowling pins as an 'unseen' non-face object category that was not presented to

the DCNNs during training. Each category consisted of 80 images. The face images were a random

subset of FITW, and images of bowling pins were randomly chosen from the corresponding category

in Caltech-256.

Movies clips for DCNN-brain correspondence analysis We examined the correspondence between

the face-selective response of the DCNNs and brain activity using a set of 18 clips of 8-min natural

color videos from the Internet that are diverse yet representative of real-life visual experiences (Wen

et al., 2017).

2.2   The deep convolutional neural network

Our model of selective deprivation, the d-AlexNet, was built with the architecture of the well-known

DCNN 'AlexNet' (Krizhevsky et al., 2012, see Figure 1a for illustration). AlexNet is a feed-forward

hierarchical convolutional neural network consisting of five convolutional layers (denoted as Conv1

– Conv5, respectively) and three fully connected layers denoted as FC1 – FC3. Each convolutional

layer consists of a convolutional sublayer, followed by a ReLU sublayer, and Conv1, 2, and 5 are

further followed by a pooling sublayer. Each convolutional sublayer consists of a set of distinct

channels. Each channel convolves the input with a distinct linear filter (or kernel) which extracts

filtered outputs from all locations within the input with a particular stride size. FC1 to FC3 are fully
connected layers. FC3 is followed by a sublayer using a softmax function to output a vector of
probabilities that represents the probability of the visual input containing the corresponding object
category (Krizhevsky et al., 2012).

The d-AlexNet used the architecture of AlexNet but changed the number of units in FC3 to 736,
so was the following softmax function, to match the number of categories in the deprivation dataset.
Same to the pre-training AlexNet in pytorch 1.2.0 (https://pytorch.org/, Paszke et al., 2017), the d-
AlexNet was initialized with values drawn from a uniform distribution, and was then trained on the
deprivation dataset following the approach specified in Krizhevsky et al., (2014). We used the pre-
trained AlexNet from pytorch 1.2.0 as the normal DCNN, referred to as the AlexNet in this paper for
brevity.

The present study referred to channels in the convolutional sublayers by the layer they belong to
and a channel index, following the convention of pytorch 1.2.0. For instance, Layer 5-Ch256 refers to
the 256th convolutional channel of Layer 5.

2.3   Transfer learning for classification and discrimination

To examine to what extent our manipulation of the visual experience affected the categorical
processing of faces, we replaced the fully-connected layers of each DCNN with a two-layer face-
classification classifier. The first layer was a fully connected layer with 43,264 units as inputs and
4,096 units as outputs with sigmoid activation function, and the second was a fully connected layer
with 4,096 units as inputs and 205 units as outputs, each of which corresponded to one category of
the classification dataset. This classifier, therefore, classified each image into one category of the
classification dataset. The face-classification classifier was trained for each DCNN with the training
images in the classification dataset for 90 epochs.

To examine to what extent our manipulation of the visual experience affected face
discrimination, we similarly replaced the fully connected layers of each DCNN with a discrimination
classifier. The discrimination classifier differed from the classification classifier only in its second
layer, which had 133 units instead as outputs, each corresponding to one individual in the
discrimination dataset. The face-discrimination classifier was trained for each DCNN with the
training images in the discrimination dataset for 90 epochs.

2.4 The face selective channels in DCNNs

To identify the channels selectively responsive to faces, we submitted images in the classification
dataset to each DCNN, recorded the average activation in each channel of Conv5 after ReLU in
response to each image, and then averaged the channel-wise activation within each category. We
selected channels where the face category evoked the highest activation, and used the Mann-Whitney
U test to examine the activation difference between faces and objects that had the second-highest
activation in these channels ($p < .05$, Bonferroni corrected). The selectivity of each face channel thus
identified was indexed by the selective ratio. The selective ratio was calculated by dividing the face
activation by the second-highest activation. In addition, we measured the lifetime sparseness of each
face-selective channel as an index for selectivity of faces among all non-face objects. We first
normalized the mean activations of a face channel in Layer5 to all the categories to the range of 0-1,
and then calculated lifetime sparseness with the formula:

$$S = (\sum_{i=1,n} r_i / n)^2 / \sum_{i=1,n} (r_i^2 / n)$$

where r_i is the normalized activations to the ith object category. The smaller this value is, the higher
the selectivity is.

2.5 DCNN-Brain Correspondence

We submitted the movie clips to the DCNNs. Following Wen (2017)'s approach, we extracted and
log-transformed the channel-wise output (the average activation after ReLU) of each face-selective
channel using DNNBrain, an in-house toolbox (Chen et al., submitted), and then convolved it with a
canonical hemodynamic response function (HRF) with a positive peak at 4s. The HRF convolved
channel-wise activity was then down-sampled to match the sampling rate of functional magnetic
resonance imaging (fMRI) and the resultant timeseries was standardized before further analysis.

Neural activation in the brain was derived from the preprocessed data in Wen (2017). The
fMRI data were recorded while human participants viewed each movie clips twice. We averaged the
standardized time series across repetition and across subjects for each clip. Then, for each DCNN, we
conducted multiple regression for each clip, with the activation time series of each brain vertex as the
dependent variable and that of face-selective channels in this network as independent variables. For
the d-AlexNet, all face-selective channels were included. For the AlexNet, we included the same
number of face-selective channels with the highest face selectivity to match the complexity of the

regression model. We used the R^2 of each vertex as the index of the overall Goodness of fit of the

regression in that vertex. The R^2 values were then averaged across clips. The larger the R^2 value, the

higher correspondence between the DCNN and the brain in response to movie clips.

To determine whether cortical regions with large R^2 values were traditional face-selective

regions, we delineated the bilateral fusiform face areas (FFA) and the occipital face area (OFA) with

the maximum-probability atlas of face-selective regions (Zhen et al., 2015). The map was based on

fMRI data of 202 healthy adult subjects and denoted the probability across the participants of each

brain vertex to be delineated into specific face-selective regions based on the task-state activation of

faces versus objects. Two hundred of vertexes of the highest probability of the left FFA and 200 of

the right FFA were included in the ROI of FFA, and the ROI of OFA was delineated in the same

way. The correspondence with brain activation in each ROI and the impact of the visual experience

was examined by submitting the vertex-wise R^2 into a two-way ANOVA with visual experience (d-

AlexNet vs. AlexNet) as within-subject factor and regional correspondence (OFA and FFA) as

between-subject factor.

2.6 Face inversion effect in DCNNs

The average activation amplitude of the top 2 face-selective channels of each DCNN in response to

upright and inverted version of 20 faces from the Reconstructing Faces dataset (VanRullen & Reddy,

2019) was measured. The inverted faces were generated by vertically flipping the upright ones. The

face inversion effect in the d-AlexNet was measured with paired sample t-tests (two-tailed) and the

impact of the experience on the face inversion effect was examined by two-way ANOVAs with

visual experience (d-AlexNet vs. AlexNet) and inversion (upright vs inverted) as within-subject

factors.

2.7 Representational similarity analysis

To examine whether faces in the d-AlexNet were processed in an object-like fashion, we compared

the within-category representational similarity of faces to that of bowling pins, an 'unseen' non-face

object category never exposed to either DCNN. Specifically, for each image in the representation

dataset, we arranged the average activations of each channel of Conv5 after ReLU into vectors, and

then for each pair of images we calculated and then Fisher-z transformed the correlation between

their vectors, which served as an index of pairwise representational similarity. Within-category

similarity between pairs of face images and that between pairs of object images were calculated

separately. A 2 × 2 ANOVA was conducted with visual experience (d-AlexNet vs AlexNet) and

category (face vs object) as independent factors. In addition, cross-category similarity between faces

and bowling pins was also calculated for each DCNN, and a paired sample t-test (two-tailed) on two

DCNNs was conducted.

2.8   Sparse coding and empirical receptive field

To quantify the degree of sparseness of the face-selective channels in representing faces, we

submitted the same set of 20 natural images containing faces from FITW to each DCNN, and

measured the number of activated units (i.e., the units showing above-zero activation) in the face-

selective channels. The more non-zero units of the face-selective channels, the less sparse of the

representation for faces. The coding sparseness of the two DCNNs was compared with a paired-

sample t-test.

We also calculated the size of the empirical receptive field of the face-selective channels.

Specifically, we obtained activation maps of 1000 images randomly chosen from FITW. Using an in-

house toolbox DNNBrain (Chen et al., submitted), we up-sampled each activation maps to the same

size of the input. For each image, we averaged the up-sampled activation within the theoretical

receptive field of each unit (the part of the image covered by the convolution of this unit and the

preceding computation, decided by the network architecture), and selected the unit with the highest

average activation. We then cropped the up-sampled activation map by the theoretical receptive field

of this unit, to locate the image part that activated this channel most across all the units. Then, we

averaged corresponding cropped activation maps across all the face images, and the resultant map

denotes the empirical receptive field of this channel, delineating the part of the theoretical receptive

field that causes this channel to respond strongly in viewing its preferred stimuli.

3   Results

The d-AlexNet was trained with a dataset of 662,619 non-face images consisting of 736 non-face

categories, generated by removing images containing faces from the ILSVRC 2012 dataset (Figure

1a). The d-AlexNet was initialized and trained in the same way as the AlexNet, and the resultant top-

1 accuracy (57.29%) and the top-5 accuracy (80.11%) were comparable with the pre-trained

AlexNet.

We first examined the performance of the d-AlexNet in two representative tasks of face

processing, face categorization (i.e., differentiating faces from non-face objects) and face

discrimination (i.e., identifying different individuals). The output of Conv5 after ReLU of the d-

AlexNet was used to classify objects in the classification dataset. The averaged categorization

accuracy of the d-AlexNet (67.40%) was well above the chance level (0.49%), and comparable to

that in the AlexNet (68.60%, $t(204) = 1.26$, $p = 0.209$, Cohen's d = 0.007, Figure 1b). Critically, the

d-AlexNet, although with no experience on faces, succeeded in the face categorization task, with an

accuracy of 86.50% in categorizing faces from non-face objects. Note that the accuracy was

numerically smaller than the AlexNet's accuracy in categorizing faces (93.90%) though (Figure 1c).

A similar pattern was observed in the face discrimination task. In this task, the output of Conv5

after ReLU of each DCNN was used to identify 33,250 face images into 133 identities in the

discrimination dataset. As expected, the AlexNet was capable of face discrimination (65.9%), well

above the chance level (0.75%), consistent with previous studies (AbdAlmageed et al., 2016;

Grundstrom, Chen, Ljungqvist, & Astrom, 2016). Critically, the d-AlexNet also showed the

capability of discriminating faces, with an accuracy of 69.30% that was even significantly higher

than that of the AlexNet, $t(132) = 3.16$, $p = .002$, Cohen's d = 0.20, (Figure 1d). Taken together,

visual experiences on faces seemed not necessary for developing basic functions of processing faces.

Figure 1. (a) An illustration of the screening to remove images containing faces for the d-AlexNet.
The 'faces' shown in the figure were AI-generated for illustration purpose only, and therefore have
no relation to real person. In the experiment, face images were from the ImageNet, with real persons'
faces. (b) The classification performance across categories of the two DCNNs was comparable. (c)
Both DCNNs achieved high accuracy in categorizing faces from other images. (d) Both DCNNs'
performance in discriminating faces was above the chance level, and the d-AlexNet's accuracy was
significantly higher than that of the AlexNet. The error bar denotes standard error. The asterisk
denotes statistical significance (α = .05). n.s. denotes no significance.

Was a face module formed in the d-AlexNet to support these functions? To answer this

question, we searched all the channels in Conv5 of the d-AlexNet, where face-selective channels

have been previously identified in the AlexNet (Baek, Song, Jang, Kim, & Paik, 2019). To do this,

we calculated the activation of each channel in Conv5 after ReLU in response to each category of the

classification dataset, and then identified channels that showed significantly higher response to faces

than non-face images with Mann-Whitney U test (ps < .05, Bonferroni corrected). Two face-selective

channels (Ch29 and Ch50) met this criterion in the d-AlexNet (for an example channel, see Figure

2a, right), whereas four face-selective channels (Ch195, Ch125, Ch60, and Ch187) were identified in

the AlexNet (for an example channel, see Figure 2a, left). The face-selective channels in two DCNNs

differed in selectivity. The averaged selective ratio, the ratio of the activation magnitude to faces by

that to non-face objects, was 1.29 (range: 1.22 - 1.36) in the d-AlexNet, much lower than that in the

AlexNet (average ratio: 3.63, range: 1.43 - 6.66). The lifetime sparseness, which measures

the breadth of tuning of a channel in response to a set of categories, also showed a similar result. The

average lifetime sparseness index of the face channels in the AlexNet (mean = 0.25, range: 0.11 –

0.51) was smaller than that in the d-AlexNet (mean = 0.71, range: 0.70 – 0.71), indicating higher face

selectivity in the AlexNet than that in the d-AlexNet. Taken together, this finding suggested that the

face-selective channels already emerged in the d-AlexNet, though the face selectivity was weaker.

How did the face-selective channels correspond to face-selective cortical regions in humans,

such as the FFA and OFA? To answer this question, we calculated the coefficient of determination

(R^2) of the multiple regression with the output of the face-selective channels as regressors and the

fMRI signals from human visual cortex in response to movies on natural vision as the regressand. As

shown in Figure 2b (right), the face-selective channels identified in the d-AlexNet corresponded to

the bilateral FFA, OFA, and the posterior superior temporal sulcus face area (pSTS-FA). Similar

correspondence was also found with the top two face-selective channels in the AlexNet (Figure 2b,

left). Direct visual inspection revealed that the deprivation weakened the correspondence between the

face-selective channels and face-selective regions in human. This observation was confirmed by the

main effect of visual experiences (F (1,798) = 161.97, p < .001, partial η^2 = 0.17) in a two-way

ANOVA of visual experiences (d-AlexNet vs. AlexNet) by regional correspondence (the OFA versus

the FFA). In addition, the main effect of the regional correspondence showed that the response

profile of the face-selective channels in the DCNNs fitted better with the activation of the FFA than

that of the OFA (F (1,798) = 98.69, p = .001, partial η^2 = 0.11), suggesting that the face-selective

channels in DCNNs may in general prefer to process faces as a whole than face parts. Critically, the

two-way interaction was significant (F (1,798) = 84.9, $p <$ .001, partial η^2 = 0.10), indicating that the

experience affected the correspondence to the FFA and OFA disproportionally. A simple effect

analysis revealed that the correspondence to the FFA (MD = 0.023, $p <$ .001) was increased by face-

specific experiences to a significantly larger extent than that to the OFA (MD = 0.004, $p =$ .013,

Figure 2c). Since the FFA is more involved in holistic processing of faces and the OFA is more

dedicated to the part-based analysis, the disproportional decrease in correspondence between the

face-selective channels in the d-AlexNet and the FFA implied that the role of the experience on faces

was to facilitate the processing of faces as a whole.

To test this conjecture, we examined how the d-AlexNet responded to inverted faces, a

behavioral signature of face-specific processing. As expected, there was a face inversion effect in the

AlexNet's face-selective channels, with the magnitude of the activation to upright faces significantly

larger than that to inverted faces (t (19) = 6.45, $p <$ .001, Cohen's d = 1.44) (Figure 2d). However, no

inversion effect was observed in the d-AlexNet, as the magnitude of the activation to upright faces

was not significantly larger than that to inverted faces (t (19) = 0.86, $p =$ .40). The lack of the

inversion effect in the d-AlexNet was further supported by a two-way interaction of visual experience

by orientation of faces, F (1, 19) = 7.79, $p =$ .012, partial η^2 = 0.29. That is, unlike the AlexNet, the

d-AlexNet processed upright faces in the same fashion as inverted faces.

Figure 2. (a) The category-wise activation profiles of example face-selective channels of the AlexNet
(left) and the d-AlexNet (right). The 'faces' shown here were AI-generated for illustration purpose
only. (b) The R^2 maps of the regression with the activation of the d-AlexNet's (right) or the
AlexNet's face-selective channels (left) as the independent variables. The higher R^2 in multiple
regression, the better in correspondence between the face channels in the DCNNs and the face-
selective regions in the human brain. The crimson lines delineate the ROIs of the OFA and the FFA.
(c) The face-channels of both DCNNs corresponded better with the FFA than the OFA, and the
difference between the AlexNet and the d-AlexNet was larger in the FFA. (d) Face inversion effect.
The average activation amplitude of the top two face-selective channels differed in response to
upright and inverted faces in the AlexNet but not the d-AlexNet. The error bar denotes standard error.
The asterisk denotes statistical significance ($\alpha = .05$). n.s. denotes no significance.

Previous studies on human suggested that inverted faces are processed in an object-like fashion. That is, it relies more on the parts-based analysis than the holistic processing. Therefore, we speculated that in the d-AlexNet faces were also represented more like non-face objects. To test this speculation, we first compared the representational similarity among responses in Conv5 to faces and bowling-pins, a novel object category that was not exposed to either DCNNs. As expected, the two-way interaction of experience (AlexNet versus d-AlexNet) by category (faces versus bowling-pins) was significant (F (1, 6,318) = 4,110.88, p <.001, partial η^2 = 0.39), and the simple effect analysis suggested that the representation for faces in the AlexNet was more similar between each other than in the d-AlexNet (MD = 0.16, p <.001), whereas the within-category representation similarity for bowling-pins showed the same but numerically smaller between-DCNN difference (MD = 0.005, p = .002) (Figure 3a).

A more critical test was to examine how face-specific experiences made faces being processed differently from objects. Here we calculated between-category similarities between faces and bowling-pins. We found that the between-category similarity between faces and bowling-pins was significantly higher in the d-AlexNet than that in the AlexNet (t (3,159) = 42.42, MD = 0.07, p < .001, Cohen's d = 0.76) (Figure 3b), suggesting that faces in the d-AlexNet were indeed represented more like objects. In short, although d-AlexNet was able to perform face tasks similar to the one with face-specific experiences, it represented faces in an object-like fashion.

Finally, we asked how faceness was achieved in DCNNs with face-specific experiences. Neurophysiological studies on monkeys demonstrate experience-associated sharpening of neural response, with fewer neurons activated after learning. Here we performed a similar analysis by measuring the number of non-zero units (i.e., units with above-zero activation) of the face-selective channels activated by natural images containing faces. As shown in the activation map (Figure 3c), a smaller number of units were activated by faces in the AlexNet than that in the d-AlexNet (t (19) = 3.317, MD =15.78, Cohen's d = 0.74) (Figure 3d), suggesting that the experience on faces made the representation to faces sparser, and thus more effective. Another effect of visual experiences observed in neurophysiological studies is that experiences reduce the size of neurons' receptive field. Here we also mapped the empirical receptive field of the face-selective channels. Similarly, we found that the empirical receptive field of the AlexNet was smaller than that of the d-AlexNet. That is,

within the theoretical receptive field, the empirical receptive field of the face-selective channels in
the AlexNet was tuned to focus on a smaller region by face-specific experiences (Figure 3e).

Figure 3. (a) The within-category similarity in the face category and an unseen non-face category
(bowling pins) in the DCNNs. (b) The between-category similarity between faces and bowling pins.
(c) The activation maps of a typical face-selective channel of each DCNN in responses to natural
images containing faces. Each pixel denotes activation in one unit. The images shown here were
historical portrait paintings downloaded from the Internet for illustration purpose only, and are
different from the images used in this study. (d) The extent of activation of the face-selective
channels of each DCNN in responses to natural images containing faces. (e) The empirical receptive
fields of the face-selective channels of each DCNN. The error bar denotes standard error. The
asterisk denotes statistical significance ($\alpha = .05$).

4   Discussion

This study presented a DCNN model of selective visual deprivation of faces. We found that without genetic predisposition and face-specific visual experiences, DCNNs were still capable of face perception. In addition, face-selective channels were also present in the d-AlexNet, which corresponded to human face-selective regions. That is, the visual experience of faces was not necessary for an intelligent system to develop a face-selective module. On the other hand, besides the slightly compromised selectivity of the module, the deprivation led the d-AlexNet to process faces in a more parts-based fashion, similar to the way of processing objects. Indeed, face-inversion effect was absent in the d-AlexNet, and the representation of faces was more similar to objects as compared to the AlexNet. Finally, the functionality of face-specific experiences that led the AlexNet to process faces as a whole might be achieved by fine-tuning the sparse coding and the size of the receptive field of the face-selective channels. In sum, our study addressed a long-standing debate on nature versus nurture in developing the face-specific module, and illuminated the role of visual experiences in shaping the module.

The observation that without domain-specific visual experience, the face-selective processing and module still emerged in the d-AlexNet seems surprising; yet this finding is consistent with previous studies on non-human primates and new-born human infants (Bushneil, Sai, & Mullin, 1989; Goren, Sarty, & Wu, 1975; Morton & Johnson, 1991; Sugita, 2008; Valenza, Simion, Cassia, & Umiltà, 1996), where the face-specific experience is found not necessary for face detection and recognition. However, the experience-independent face processing is largely attributed to either innate face-specific mechanisms (McKone et al., 2012; Morton & Johnson, 1991) or domain-general processing with predisposed biases (Cassia, Turati, & Simion, 2004; Simion & Di Giorgio, 2015; Simion, Macchi Cassia, Turati, & Valenza, 2001). Our study argued against this conjecture, because unlike any biological system, DCNNs have no predefined genetic inheritance or processing biases. Therefore, the face-specific processing observed in DCNNs had to derive from domain-specific factors.

We speculated that the face module in the d-AlexNet may result from a tremendous amount of features represented in the multiple layers of the network, with which face-like features were selected to construct face-specific module. In fact, previous studies on DCNNs have shown that DCNN's lower layers showed sensitivity to myriad visual features similar to primates' primary visual cortex (Krizhevsky et al., 2012), while the higher layers are tuned to complex features resembling those

represented in the ventral visual pathway (Güçlü & van Gerven, 2015; Khaligh-Razavi &

Kriegeskorte, 2014; Pospisil, Pasupathy, & Bair, 2018; Yamins et al., 2014). With such repertoire of

rich features, a representational space for faces, or for any natural object, may be constructed by

selecting face-like features and features that are potentially useful in a variety of face tasks.

Supporting evidence for this conjecture came from the observation that the d-AlexNet processed

faces in an object-like fashion. For example, the face inversion effect, a behavioral signature of face-

specific processing in human (Kanwisher, Tong, & Nakayama, 1998; Rossion & Gauthier, 2002;

Yin, 1969) was absent in the d-AlexNet. That is, similar to inverted faces, upright faces may also be

processed like objects in the d-AlexNet. A more direct illustration of the object-like representation of

faces came from the analysis on the representational similarity between faces and objects. As

compared to the AlexNet, faces in the representational space of the d-AlexNet were less congregated

among each other; instead they were more intermingled with non-face object categories. The finding

that face representation was no longer qualitatively different from object representation may help

explaining the performance of the d-AlexNet. Because faces were less segregated from objects in the

representational space, the d-AlexNet's accuracy of face categorization was worse than that of the

AlexNet. In contrast, within the face category, individual faces were less congregated in the

representational space; therefore, the discrimination of individual faces became easier instead,

suggested by the slightly higher face discrimination accuracy in the d-AlexNet than the AlexNet. In

short, when the representational space of the d-AlexNet was formed exclusively based on features

from non-face stimuli, faces were represented no longer qualitatively different from non-face objects,

which inevitably led to 'object-like' face processing.

The face-specific processing is likely achieved through prior exposure to faces. At first glance,

the effect of face-specific experiences seemed quantitative, as in the AlexNet, both the selectivity to

faces and the number of the face-selective channels were increased, and the correspondence between

the face-selective channels and the face-selective regions in human brain was tighter. However,

careful scrutiny of the difference between the two DCNNs revealed that the changes led by the

experience may be qualitative. For example, the deprivation of visual experiences disproportionally

weakened the DCNN-brain correspondence in the FFA as comparing to the OFA, given that the FFA

is engaged more in the configural processing and the OFA in parts-based analysis (Liu, Harris, &

Kanwisher, 2010; Nichols, Betts, & Wilson, 2010; Zhao et al., 2014). Therefore, the 'face-like' face

processing may come from the fact that face-specific experiences led the representation of faces more

congregated within face category and more separable from the representation of non-face objects

stimuli (see also Gomez, Barnett, & Grill-Spector, 2019). In this way, a relative encapsulated

representation may help developing a unique way of processing faces, qualitatively different from

non-face objects.

The advantage of the computational transparency of DCNNs may shed light on the

development of domain specificity of the face module. First, we found that face-specific experiences

increased the sparseness of face representation, as fewer units of the face channels were activated by

faces in the AlexNet. The experience-dependent sparse coding has been widely discovered in the

visual cortex, such as the V4, MT, and IT (for reviews, see Desimone, 1996; Grill-Spector, Henson,

& Martin, 2006; Wiggs & Martin, 1998). The experience-induced increase of sparseness is thought to

reflect a preference-narrowing process that tunes neurons to a smaller range of stimuli (Kohn &

Movshon, 2004); therefore, with sparse coding faces are less likely to be intermingled with non-face

objects, which may lead to more congregated representations in the representational space in the

AlexNet, as compared to the d-AlexNet. Second, we found that the empirical receptive field of the

face channel in the AlexNet was smaller than that in the d-AlexNet, suggesting that the visual

experience on faces decreased the size of the receptive field of the face channels. This finding fits

perfectly with neurophysiological studies that the size of receptive fields of visual neurons is reduced

after eye-opening (Braastad & Heggelund, 1985; Cantrell, Cang, Troy, & Liu, 2010; Koehler,

Akimov, & Renteria, 2011; Tavazoie & Reid, 2000). Importantly, along with the refined receptive

fields, the selectivity of neurons increases (Spilmann, 2014), possibly because neurons can avoid

distracting information by focusing on a more restricted part of stimuli, which may further allowed

finer representation of the selected regions. This is especially important for processing faces because

faces are highly homogeneous, and some information is identical across faces, such as parts

composition (eyes, noses, and mouth) and their configural arrangements. Therefore, the reduced

receptive field of the face channels may facilitate selective analyses of discriminative face features

while avoiding irrelevant information. Further, the sharpening of the receptive field and the fine-

tuned selectivity may result in superior discrimination ability on faces, and allow faces to be

processed at the sub-ordinate level (i.e., identification), whereas the rest of objects are largely

processed at the basic level (i.e., categorization).

It has long been assumed that domain-specific visual experiences and inheritance are the pre-

requisites in the development of the face module. In our study with DCNNs as a model, we

completely decoupled the genetic predisposition and face-specific visual experiences, and found that

the representation for faces can be constructed with features from non-face objects to realize basic

functions for face recognition. Therefore, in many situations, the difference between faces and

objects is 'quantitative' rather than 'qualitative', as they are represented in a continuum of the

representational space. In addition, we also found that face-specific experiences likely fine-tuned the

face representation, and thus transformed the 'object-like' face processing into 'face-specific'

processing. However, we shall be cautious that our finding may not be applicable for the

development of face module in human, as in the biological brain experience-induced changes are

partly attributed to the inhibition from lateral connections (Grill-Spector et al., 2006; Norman &

O'Reilly, 2003), whereas there is no lateral or feedback connection in DCNNs. Recent studies have

shown similar representation for faces between DCNNs and humans (Song, Qu, Xu, & Liu, 2020),

suggesting that a common mechanism may be shared by both artificial and biological intelligent

systems. Future studies are needed to examine the applicability of our finding to humans. On the

other hand, our study also illustrated the advantages of using DCNNs as a model to understand

human mind because of its computational transparency and its dissociation of factors in nature and

nurture. Thus, our study invites future studies with DCNNs to understand the development of domain

specificity in particular and a broad range of cognitive modules in general.

5   Conflict of Interest

The authors declare no competing interests.

6   Author Contributions

J. L. conceived and designed the study. Y.Z. analyzed the data with input from all authors. S.X. wrote the

manuscript with input from J. L., Y.Z. and Z. Z.

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9   Reference

AbdAlmageed, W., Wu, Y., Rawls, S., Harel, S., Hassner, T., Masi, I., et al. (2016). Face Recognition Using Deep Multi-Pose Representations. In *2016 Ieee Winter Conference on Applications of Computer Vision*.

Arcaro, M. J., Schade, P. F., Vincent, J. L., Ponce, C. R., & Livingstone, M. S. (2017). Seeing faces is necessary for face-domain formation. *Nature Neuroscience, 20*(10), 1404-+.

Baek, S., Song, M., Jang, J., Kim, G., & Paik, S.-B. (2019). Spontaneous generation of face recognition in untrained deep neural networks. *bioRxiv*, 857466.

Berg, T. L., Berg, A. C., Edwards, J., & Forsyth, D. A. (2005). *Who's in the picture.* Paper presented at the Advances in neural information processing systems.

Braastad, B. O., & Heggelund, P. (1985). Development of spatial receptive-field organization and orientation selectivity in kitten striate cortex. *Journal of Neurophysiology, 53*(5), 1158-1178.

Brock, A., Donahue, J., & Simonyan, K. (2018). Large scale gan training for high fidelity natural image synthesis. *arXiv preprint arXiv:1809.11096*.

Bushneil, I., Sai, F., & Mullin, J. (1989). Neonatal recognition of the mother's face. *British Journal of Developmental Psychology, 7*(1), 3-15.

Cantrell, D. R., Cang, J., Troy, J. B., & Liu, X. (2010). Non-Centered Spike-Triggered Covariance Analysis Reveals Neurotrophin-3 as a Developmental Regulator of Receptive Field Properties of ON-OFF Retinal Ganglion Cells. *Plos Computational Biology, 6*(10).

Cassia, V. M., Turati, C., & Simion, F. (2004). Can a nonspecific bias toward top-heavy patterns explain newborns' face preference? *Psychological Science, 15*(6), 379-383.

Deng, J., Dong, W., Socher, R., Li, L.-J., Li, K., & Fei-Fei, L. (2009). *Imagenet: A large-scale hierarchical image database.* Paper presented at the 2009 IEEE conference on computer vision and pattern recognition.

Desimone, R. (1996). Neural mechanisms for visual memory and their role in attention. *Proceedings of the National Academy of Sciences of the United States of America, 93*(24), 13494-13499.

Freiwald, W., Duchaine, B., & Yovel, G. (2016). Face Processing Systems: From Neurons to Real-World Social Perception. In S. E. Hyman (Ed.), *Annual Review of Neuroscience, Vol 39* (Vol. 39, pp. 325-346).

Gomez, J., Barnett, M., & Grill-Spector, K. (2019). Extensive childhood experience with Pokemon suggests eccentricity drives organization of visual cortex. *Nature Human Behaviour, 3*(6), 611-624.

Goren, C. C., Sarty, M., & Wu, P. Y. (1975). Visual following and pattern discrimination of face-like stimuli by newborn infants. *Pediatrics, 56*(4), 544-549.

Griffin, G., Holub, A., & Perona, P. (2007). Caltech-256 object category dataset.

Grill-Spector, K., Henson, R., & Martin, A. (2006). Repetition and the brain: neural models of
stimulus-specific effects. *Trends in Cognitive Sciences, 10*(1), 14-23.

Grundstrom, J., Chen, J., Ljungqvist, M. G., & Astrom, K. (2016). Transferring and Compressing
Convolutional Neural Networks for Face Representations. In A. Campilho & F. Karray
(Eds.), *Image Analysis and Recognition* (Vol. 9730, pp. 20-29).

Güçlü, U., & van Gerven, M. A. (2015). Deep neural networks reveal a gradient in the complexity of
neural representations across the ventral stream. *Journal of Neuroscience, 35*(27), 10005-
10014.

Kanwisher, N., Tong, F., & Nakayama, K. (1998). The effect of face inversion on the human
fusiform face area. *Cognition, 68*(1), B1-B11.

Kanwisher, N., & Yovel, G. (2006). The fusiform face area: a cortical region specialized for the
perception of faces. *Philosophical Transactions of the Royal Society B-Biological Sciences,*
361(1476), 2109-2128.

Karras, T., Laine, S., & Aila, T. (2019). *A style-based generator architecture for generative*
adversarial networks. Paper presented at the Proceedings of the IEEE conference on
computer vision and pattern recognition.

Khaligh-Razavi, S.-M., & Kriegeskorte, N. (2014). Deep supervised, but not unsupervised, models
may explain IT cortical representation. *PLoS computational biology, 10*(11), e1003915.

Koehler, C. L., Akimov, N. P., & Renteria, R. C. (2011). Receptive field center size decreases and
firing properties mature in ON and OFF retinal ganglion cells after eye opening in the mouse.
Journal of Neurophysiology, 106(2), 895-904.

Kohn, A., & Movshon, J. A. (2004). Adaptation changes the direction tuning of macaque MT
neurons. *Nature Neuroscience, 7*(7), 764-772.

Kriegeskorte, N. (2015). Deep neural networks: a new framework for modeling biological vision and
brain information processing. *Annual review of vision science, 1*, 417-446.

Krizhevsky, A. (2014). One weird trick for parallelizing convolutional neural networks. *arXiv*
preprint arXiv:1404.5997.

Krizhevsky, A., Sutskever, I., & Hinton, G. E. (2012). *Imagenet classification with deep*
convolutional neural networks. Paper presented at the Advances in neural information
processing systems.

Liu, J., Harris, A., & Kanwisher, N. (2010). Perception of Face Parts and Face Configurations: An
fMRI Study. *Journal of Cognitive Neuroscience, 22*(1), 203-211.

McKone, E., Crookes, K., Jeffery, L., & Dilks, D. D. (2012). A critical review of the development of
face recognition: Experience is less important than previously believed. *Cognitive*
Neuropsychology, 29(1-2), 174-212.

Morton, J., & Johnson, M. H. (1991). CONSPEC and CONLERN: a two-process theory of infant
face recognition. *Psychological Review, 98*(2), 164.

Nichols, D. F., Betts, L. R., & Wilson, H. R. (2010). Decoding of faces and face components in face-
sensitive human visual cortex. *Frontiers in Psychology, 1*.

Norman, K. A., & O'Reilly, R. C. (2003). Modeling hippocampal and neocortical contributions to
566   recognition memory: A complementary-learning-systems approach. *Psychological Review,*
567   *110*(4), 611-646.

568   Parkhi, O. M., Vedaldi, A., & Zisserman, A. (2015). *Deep face recognition.* Paper presented at the
569   bmvc.

570   Paszke, A., Gross, S., Chintala, S., Chanan, G., Yang, E., DeVito, Z., et al. (2017). Automatic
571   differentiation in pytorch.

572   Pospisil, D. A., Pasupathy, A., & Bair, W. (2018). 'Artiphysiology'reveals V4-like shape tuning in a
573   deep network trained for image classification. *Elife, 7*, e38242.

574   Rossion, B., & Gauthier, I. (2002). How does the brain process upright and inverted faces?
575   *Behavioral and cognitive neuroscience reviews, 1*(1), 63-75.

576   Simion, F., & Di Giorgio, E. (2015). Face perception and processing in early infancy: inborn
577   predispositions and developmental changes. *Frontiers in Psychology, 6*.

578   Simion, F., Macchi Cassia, V., Turati, C., & Valenza, E. (2001). The origins of face perception:
579   specific versus non‐specific mechanisms. *Infant and Child Development: An International*
580   *Journal of Research and Practice, 10*(1‐2), 59-65.

581   Song, Y., Qu, Y., Xu, S., & Liu, J. (2020). Implementation-independent representation for deep
582   convolutional neural networks and humans in processing faces. *bioRxiv*.

583   Spilmann, L. (2014). Receptive fields of visual neurons: The early years. *Perception, 43*(11), 1145-
584   1176.

585   Sugita, Y. (2008). Face perception in monkeys reared with no exposure to faces. *Proceedings of the*
586   *National Academy of Sciences of the United States of America, 105*(1), 394-398.

587   Tavazoie, S. F., & Reid, R. C. (2000). Diverse receptive fields in the lateral geniculate nucleus during
588   thalamocortical development. *Nature Neuroscience, 3*(6), 608-616.

589   Valenza, E., Simion, F., Cassia, V. M., & Umiltà, C. (1996). Face preference at birth. *Journal of*
590   *experimental psychology: Human Perception and Performance, 22*(4), 892.

591   VanRullen, R., & Reddy, L. (2019). Reconstructing faces from fMRI patterns using deep generative
592   neuralnetworks. *Communications biology, 2*(1), 193-193.

593   Wen, H., Shi, J., Zhang, Y., Lu, K.-H., Cao, J., & Liu, Z. (2017). Neural encoding and decoding with
594   deep learning for dynamic natural vision. *Cerebral Cortex, 28*(12), 4136-4160.

595   Wiggs, C. L., & Martin, A. (1998). Properties and mechanisms of perceptual priming. *Current*
596   *Opinion in Neurobiology, 8*(2), 227-233.

597   Wilmer, J. B., Germine, L., Chabris, C. F., Chatterjee, G., Williams, M., Loken, E., et al. (2010).
598   Human face recognition ability is specific and highly heritable. *Proceedings of the National*
599   *Academy of Sciences of the United States of America, 107*(11), 5238-5241.

600   Yamins, D. L., Hong, H., Cadieu, C. F., Solomon, E. A., Seibert, D., & DiCarlo, J. J. (2014).
601   Performance-optimized hierarchical models predict neural responses in higher visual cortex.
602   *Proceedings of the National Academy of Sciences, 111*(23), 8619-8624.

603   Yi, D., Lei, Z., Liao, S., & Li, S. Z. (2014). Learning face representation from scratch. *arXiv preprint*
604   *arXiv:1411.7923*.

Yin, R. K. (1969). Looking at upside-down faces. *Journal of Experimental Psychology, 81*(1), 141-
&.

Zhao, M., Cheung, S.-h., Wong, A. C. N., Rhodes, G., Chan, E. K. S., Chan, W. W. L., et al. (2014).
Processing of configural and componential information in face-selective cortical areas.
Cognitive Neuroscience, 5(3-4), 160-167.

Zhen, Z., Yang, Z., Huang, L., Kong, X.-z., Wang, X., Dang, X., et al. (2015). Quantifying
interindividual variability and asymmetry of face-selective regions: a probabilistic functional
atlas. *Neuroimage, 113*, 13-25.

Zhu, Q., Song, Y., Hu, S., Li, X., Tian, M., Zhen, Z., et al. (2010). Heritability of the Specific
Cognitive Ability of Face Perception. *Current Biology, 20*(2), 137-142.