# Skin Lesion Segmentation and Classification for ISIC 2018 by Combining Deep CNN and Handcrafted Features

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#### Abstract

This short report describes our submission to the ISIC 2018 Challenge in Skin Lesion Analysis Towards Melanoma Detection [1] for Task1 and Task 3. This work has been accomplished by a team of researchers at the University of Dayton Signal and Image Processing Lab. Our proposed approach is computationally efficient are combines information from both deep learning and handcrafted features. For Task3, we form a new type of image features, called hybrid features, which has stronger discrimination ability than single method features. These features are utilized as inputs to a decision-making model that is based on a multiclass Support Vector Machine (SVM) classifier. The proposed technique is evaluated on online validation databases. Our score was 0.841 with SVM classifier on the validation dataset.

#### 1 Introduction

The ISIC 2018 Skin Lesion Analysis Towards Melanoma Detection [1] is broken into three separate tasks: 1) Lesion Segmentation, 2), Lesion Attribute Detection and 3) Disease Classification. This report addresses Task 1 and Task 3 which are Lesion Segmentation and Disease Classification. The goal of the Task 3 is to classify skin lesion images into 7 Possible disease categories – Melanoma (MEL), Melanocytic nevus (NV), Basal cell carcinoma (BCC), Actinic keratosis intraepithelial carcinoma (AKIEC), Benign keratosis (BKL), Dermatofibroma (DF), and Vascular lesion (VASC) [1]. Participants are being ranked using a normalized multiclass accuracy metric (balanced across categories) [1].

## 2 Task 1 Methodology

We proposed a hybrid Deep and Handcrafted system for lesion segmentation. Our hybrid technique makes use of the following two methods: Deep CNN and Gaussian mixture models (GMMs) [3]. Both of These are described below.

# **2.1** Deep and efficient UNet

We implemented an elegant architecture, the so-called "fully convolutional network" [2]. The network architecture illustrated in Fig. 1. The pattern in our segmentation networks requires the downsampling of an image between convolutional and ReLU layers, and then upsample the output to match the input size. The network starts with an image "Input Layer," which describes the image size that the network can process. All training and testing images resized to [224 224]

3]. The Down sampling network starts with the 3x3 convolution, Batch normalization layer and followed by a rectified linear unit ReLU layers. At the end of each downsample stage, there is 2x2 max pooling operation with stride 2. After stage 2 in downsampling, we double the number of feature channels. The upsampling is done using the transposed convolution layer commonly referred to as "deconv" or "deconvolution" layer. Every step in the upsampling of the feature map followed by a 3x3 convolution, Batch normalization and followed by a ReLU. The cropping is significant due to the loss of border pixels in every convolution. At the end of the network, we have a 1x1 convolution used to map the feature vector to the number of classes. In total the network has 109 layers.

Stage 5	Stage 4	Stage 3	Stage 2	Stage 1
conv5_1 bn_conv5_1 relu5_1 conv5_2 bn_conv5_2 bn_conv5_2 relu5_2 conv5_3 bn_conv5_3 relu5_3 conv5_4 bn_conv5_4 relu5_4 pool5 decoder5_unpool decoder5_conv4 decoder5_relu_4 decoder5_relu_4 decoder5_relu_4 decoder5_relu_3 decoder5_relu_3 decoder5_relu_3 decoder5_relu_2 decoder5_conv1 decoder5_relu_2 decoder5_relu_1 decoder5_relu_1 decoder5_relu_1	conv4_1 bn_conv4_1 relu4_1 conv4_2 bn_conv4_2 relu4_2 conv4_3 relu4_3 conv4_4 bn_conv4_4 relu4_4 pool4 decoder4_unpool decoder4_conv4 decoder4_relu_4 decoder4_relu_4 decoder4_relu_4 decoder4_relu_3 decoder4_relu_3 decoder4_relu_3 decoder4_relu_3 decoder4_relu_2 decoder4_relu_2 decoder4_relu_2 decoder4_relu_1 decoder4_relu_1 decoder4_relu_1	conv3_1 bn_conv3_1 relu3_1 conv3_2 bn_conv3_2 bn_conv3_3 relu3_2 conv3_3 bn_conv3_4 bn_conv3_4 bn_conv3_4 pool3 decoder3_unpool decoder3_conv4 decoder3_relu_4 decoder3_relu_4 decoder3_relu_3 decoder3_relu_3 decoder3_relu_3 decoder3_relu_3 decoder3_relu_3 decoder3_relu_2 decoder3_relu_2 decoder3_relu_2 decoder3_relu_2 decoder3_relu_2 decoder3_relu_2 decoder3_relu_1 decoder3_relu_1 decoder3_relu_1	conv2_1 bn_conv2_1 relu2_1 conv2_2 bn_conv2_2 pool2 decoder2_unpool decoder2_conv2 decoder2_bn_2 decoder2_relu_2 decoder2_conv1 decoder2_relu_1	inputImage conv1_1 reiu1_1 conv1_2 bn_conv1_2 bn_conv1_2 pool1  decoder1_unpool decoder1_conv2 decoder1_bn_2 decoder1_bn_2 decoder1_conv1 decoder1_conv1 decoder1_bn_1 softmax_reiu_1 pixelLabels

Fig. 1: U-net architecture where each red circle corresponds to a layer. The name of layers is denoted on the side of the circle.

### **2.2** Gaussian mixture models (GMMs)

This model uses the probability density functions of the tissue types to distinguish lesion tissue from normal skin tissue [3]. Our team has submitted this approach in separate submissions using traditional classifiers with hand-crafted features.

To fuse the two system together and obtain the final results. We come up with the best threshold that switches to select between a traditional segmentation approach [3] and deep learning method. This switch is based on estimated lesion area. Since the GMM seems better for smaller lesions and Unet better for larger ones. Therefore, If the lesion area of UNet is smaller than 4508 when the image size is 224 x224, we choose the GMMs mask. Otherwise, select the UNet model result as shown in the Fig. 2.

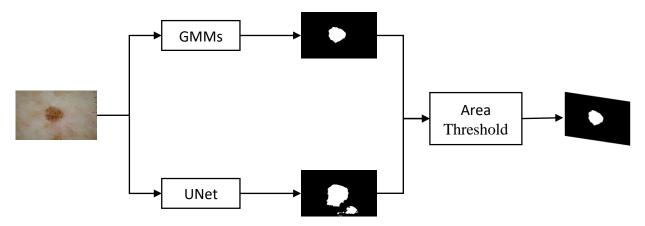


Fig. 2: Shows a hybrid Deep and Handcrafted system

## 3. Task 3 Methodology

Fig. 3 illustrates our proposed classification approach. First, we have trained two convolutional neural networks (CNN) on the available training data for Lesion Diagnosis. The training data consists of 10015 images. All training examples have been resampled to 244 x 224 x 3. We have trained out CNN's without any data augmentations or geometrically transformed such as rotation, translation, scaling and flipping. Second the 200 handcrafted feature [3]. The features are computed from the RGB image with respect to the lesion segmentation. Finally, the handcrafted features [3] are concatenated with CNN features to form the final feature vector. This final feature set is fed to a decision-making model based on multiclass support vector machine (SVM) classifier.

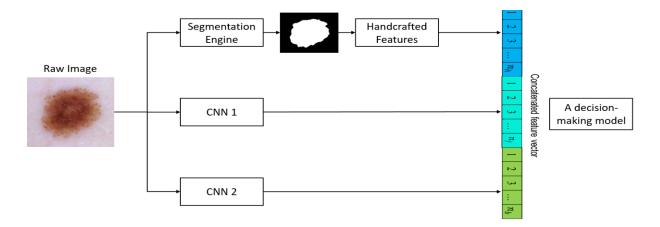


Fig. 3: Shows Proposed classification system.

# 4. Experimental Results

#### 4.1 Task 1

The results for Task 1 have been obtained using the provided validation dataset. The validation scores are for our information and are not proposed to be made public. We have tested our model on the provided validation data (193 samples in total). The mean overlap score on the testing data is 0.735.

#### 4.2 Task 3

In this Task, we employ the ISIC 2018 validation dataset [1] for system evaluation. The results show that our model obtains promising performance with class averaged recall of 0.841.

## 5. Conclusion

This research proposes a robust system for lesion segmentation and disease from dermoscopy images, which offers the vision of achieving an improved and more accurate classification of lesions from images. Our proposed method is based on the use of hybrid features, which are a combination of handcrafted features and deep learning features. Hybrid features provide richer information than that obtained using a single feature extraction method. Hybrid features significantly enhance the segmentation in Task 1 and classification Task 3 accuracy compared to the use of a single method.

### 6. References

- [1] The International Skin Imaging Collaboration (ISIC) Website, "ISIC 2018: Skin Lesion Analysis Towards Melanoma Detection," <a href="https://challenge2018.isic-archive.com/">https://challenge2018.isic-archive.com/</a>.
- [2] Long, J., Shelhamer, E., Darrell, T.: Fully convolutional networks for semantic segmentation (2014), arXiv:1411.4038 [cs.CV]
- [3] Hardie, Russell C., et al. "Skin Lesion Segmentation and Classification for ISIC 2018 Using Traditional Classifiers with Hand-Crafted Features." arXiv preprint arXiv:1807.07001(2018).