




Conditional adversarial segmentation and deep learning approach for skin lesion sub-typing from dermoscopic images

P. Mirunalini¹ · Karthik Desingu¹ · S. Aswatha¹ · R. Deepika¹ · V. Deepika¹ · S. M. Jaisakthi² 

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Abstract

Automatic skin lesion subtyping is a crucial step for diagnosing and treating skin cancer and acts as a first level diagnostic aid for medical experts. Although, in general, deep learning is very effective in image processing tasks, there are notable areas of the processing pipeline in the dermoscopic image regime that can benefit from refinement. Our work identifies two such areas for improvement. First, most benchmark dermoscopic datasets for skin cancers and lesions are highly imbalanced due to the relative rarity and commonality in the occurrence of specific lesion types. Deep learning methods tend to exhibit biased performance in favor of the majority classes with such datasets, leading to poor generalization. Second, dermoscopic images can be associated with irrelevant information in the form of skin color, hair, veins, etc.; hence, limiting the information available to a neural network by retaining only relevant portions of an input image has been successful in prompting the network towards learning task-relevant features and thereby improving its performance. Hence, this research work augments the skin lesion characterization pipeline in the following ways. First, it balances the dataset to overcome sample size biases. Two balancing methods, synthetic minority oversampling TEchnique (SMOTE) and Reweighting, are applied, compared, and analyzed. Second, a lesion segmentation stage is introduced before classification, in addition to a preprocessing stage, to retain only the region of interest. A baseline segmentation approach based on Bi-Directional ConvLSTM U-Net is improved using conditional adversarial training for enhanced segmentation performance. Finally, the classification stage is implemented using EfficientNets, where the B2 variant is used to benchmark and choose between the balancing and segmentation techniques, and the architecture is then scaled through to B7 to analyze the performance boost in lesion classification. From these experiments, we find that the pipeline that balances using SMOTE and uses the adversarially trained segmentation network achieves the best baseline performance of 91% classification accuracy with EfficientNet B2. Based on the scaling experiments, we find that optimal performance is reached with the B6 architecture that classifies with a 97% accuracy. Furthermore, the proposed pipeline for lesion characterization outperforms the state of the art performance on the ISIC dataset.

Keywords Adversarial learning · Skin lesion · Dermoscopic images · EfficientNet · SMOTE · Reweighting · BCDU-Net · Deep learning · Adversarial segmentation

1 Introduction

The number of people diagnosed with skin cancer is increasing annually. Skin lesions are caused by various dermatological disorders and can either be benign or malignant. Therefore, not only it is crucial to discriminate benign and malignant lesions for proper treatment, but also the lesion types, since early detection of malignant lesions

can significantly reduce the risk of fatality. However, the manual inspection of skin lesions is prone to failure, time consuming, and subjective, since most skin lesions have high visual similarity that is often hard to discern for the human eye. Routine clinical settings based on manual inspection have achieved less than 80% accuracy in detecting melanomas from dermoscopic images [53]. Hence, the clinical workflow can benefit from an automatic method that can classify skin lesion types as a preliminary diagnosing tool.

The occurrence of different types of skin lesions is highly disproportionate, some being extremely rare and

P. Mirunalini, Karthik Desingu and S M Jaisakthi have contributed equally to this work.

Extended author information available on the last page of the article

others common. As a result, dermoscopic image datasets available for skin lesions also exhibit such imbalance. Diagnostic systems, especially those based on machine learning techniques, are prone to sample size bias when trained using such unbalanced data sets [11–13, 40]. Hence, it is imperative to eliminate such biases when developing diagnostic algorithms using unbalanced datasets.

Dermoscopic images are frequently characterized by artifacts such as hair, vein, marker annotations, that are irrelevant to skin lesion identification. The presence of artifacts has a negative impact on the ability of machine learning-based classifiers to learn representative features that can distinguish types of lesion. Therefore, segmenting to retain only the relevant portions of the lesion from the dermoscopic images before feeding them to the classifier will improve performance. However, since lesions typically occur in irregular shapes, the segmentation process is complex and challenging. In such scenarios, deep learning-based segmentation methods are known to outperform traditional heuristic-based and algorithmic image processing methods. Deep learning methods can learn complex features that traditional hand-crafted features used in traditional and pixel-statistics based image processing methods cannot represent.

Hence, in this paper we sought to employ a deep learning-driven skin lesion diagnosis pipeline to leverage their proven ability to learn discriminative features from their input incrementally through their hidden-layer architecture. We examine and prove the effectiveness of an automatic two-level diagnostic system that first segments skin lesions regions from dermoscopic images using an adversarially trained segmentation network, and then classifies the lesion type from these images using a class of convolutional neural network (CNN) architectures called EfficientNets [50]. To address sample-size imbalance in skin lesion datasets, we analyze the effectiveness of two methods: class reweighted training and synthetic data sampling. Finally, we propose a pipeline that demonstrates best classification performance on the study dataset along with discussions about the effectiveness and strengths of different stages of the pipeline.

The rest of this article is organized as follows: Sect. 2 presents a brief review of the related literature, summarizing the key research gaps and the contributions of this paper. The study methodology and considerations are shown in Sect. 3. Section 4 details the conducted experiments and their outcomes, while Sect. 5 discusses the comparison of our proposed approach with methods introduced in existing literature. Section 6 presents a discussion of the results along with strengths and limitations of the proposed pipeline. The implementation of this

research work will be made available at: <https://github.com/karthik-d/lesion-characterization-using-cgan>.

2 Related work

The accurate classification of skin lesions in their early stages enables dermatologists to treat patients well in time. The time-consuming nature and possibility of inter and intra evaluator bias in the manual diagnosis of skin cancer can be overcome by employing a computerized analysis system. Several proposals were made to overcome the challenges faced in classifying dermoscopic images. Kassem et al. [27] proposed a model utilizing transfer learning and a pre-trained GoogleNet model by adding more filters to each layer to enhance features and reduce noise. Gessert et al. [15] proposed a classification model for dermoscopic images using ensemble of deep learning methods which also predicts an unknown class using a data-driven approach. In addition, it addresses the class-imbalance problem using a loss balancing approach. And in [19], the authors compared the diagnosis performance between dermatologists and an Inception-v4 model at two levels: one using just dermoscopic images and the other that integrated clinical information into model prediction, finding the latter approach to perform better. The effect of using multiple image sizes with same network was studied using multi-scale multi-CNN fusion approach; this boosted the classification performance compared to a single network or a single image scale [34]. A hyper-connected CNN (HcCNN) with a multi-scale attention (MsA) block was used to improve classification performance of skin lesions by prioritizing semantically important subtle regions [6]. Techniques based on ensemble network and majority voting from pre-trained models such as ResNet50, MobileNetV3, and EfficientNet have also been experimented with skin classification process [51]. Zhou et al. [56], proposed a model which includes several deep CNN models such as DenseNet121, SeResNeXt50, SeResNeXt101, and EfficientNets, and experimented by taking different combinations of these base models to tune the precision and recall performance.

A range of heuristic-based and hybrid radiomic-deep learning methods have also been proposed in the past. In [33], self-ensemble and stacking-based framework was used to classify skin acne types. This approach stacks the best subset of all features obtained during training epochs for classification. Muthulakshmi and Hemapriya [38] proposed radiomics feature extraction model for the detection of Kaposi sarcoma and proposed medical image fusion on deep feature learning using pre-built ResNet50 and VGG16 and also applied an ensemble learning approach. Intelligent multi-level thresholding and deep learning (IMLT-DL)

model was proposed in [29]. In this work the authors used top hat filtering and inpainting methods to detect the lesion regions and then applied Mayfly optimization. Multilayer Kapur's thresholding was used to locate the trouble spots. Finally, a neural network was used to extract features and was classified with gradient boosting trees.

It has been shown that the accuracy of deep learning models can be improved by resolving the class imbalance problem in the training dataset [37]. To address imbalance, various approaches have been proposed in the past. Venkataraman et al. [52] studied the effect of using data augmentation and SMOTE for balancing the data set for classifying tuberculosis, pneumonia, and COVID-19. Tahir et al. [49] proposed a CNN based model, namely deep learning-based skin cancer classification network (DSCC_Net) and evaluated it on ISIC 2020, HAM10000, and DermIS datasets. To deal with class imbalance, the authors used the SMOTE Tomek method [5] to balance the minority classes and obtained 92.51% using a MobileNet classifier.

The presence of artifacts such as hair and air bubbles, intrinsic factors such as size, texture, contrast, shape, color, and variations in image acquisition conditions make lesion image processing a challenging task [27, 36]. Although deep learning has been widely used to deal with such visual complexity for medical image analysis, challenges such as unavailability of quality data, unavailability of large balanced datasets, high inter-class and intra-class variations [46], the presence of noise and unwanted artifacts in dermoscopic images, among others make them all the more challenging and impede feature extraction using a neural network [4, 24].

In addition to using image preprocessing techniques to remove specific artifacts, segmenting dermoscopic images to retain only lesion regions can be crucial to improve lesion classification performance. A range of image segmentation algorithms have been proposed for lesion segmentation in the past. The ABCD algorithm (Asymmetry, Border, Color, Diameter of lesions), a rule-based skin lesion segmentation system [39], and its derivatives such as ABCDE (Asymmetry, Border, Color, Diameter and Evolving) [1] and ABCDEF (Asymmetry, Border, Color, Diameter, Evolving and Firm) [26] are popular algorithmic image processing approaches. Other image processing and machine learning approaches used in the past include thresholding [8], active contours and region growing [7, 14], and clustering techniques [10]. Jaisakthi et al. [25], proposed an automatic skin lesion segmentation model using Grabcut and the K-means algorithm, where the K-means algorithm helps improving the boundaries of the segments by utilizing the color features learnt from the training images. However, these approaches rely on hand-crafted feature extraction from dermoscopic images; this

requires extensive manual effort to pick and experiment with feature extraction algorithms and limits discriminatory capability of the extracted features owing to very high visual complexity in lesion images. Hence, these traditional methods often do not yield good results for large and complex datasets; even when they do, they seldom are readily transferable to other datasets, and need to be adapted to suit the new dataset.

Deep learning based techniques alleviate the issue of feature extraction by elegantly blending the feature extraction process with learning. Various studies have proposed or applied different deep learning architectures for ROI segmentation. Manu et al. [18] designed an end-to-end ensemble segmentation methods that combines Mask R-CNN and DeeplabV3+ with preprocessing and post processing methods to produce accurate lesion segmentation for the ISIC 2017 dataset. While this approach generates bounding boxes for objects accurately, its performance on generating segmentation masks is quite low the obtained boundaries are not complete, often excluding out lesion regions in the segmented image. Liu et al. [32] proposed a new CNN-based method for segmentation of skin lesions with auxiliary task learning to identify area of infection. Jafari et al. [23] proposed an automated skin lesion segmentation model using deeper networks and a residual learning technique. Sohaib et al. [20] proposed a modified U-Net architecture with a 46-layered structure to obtain a good segmentation rate on the ISIC 2018 dataset after preprocessing the images to remove artifacts.

CNN as well as other deep learning-based methods requires a large number of training samples, especially for tasks such as segmentation, since every pixel of the image has to be assigned a class as opposed to say, a simpler image classification task. Studies in the past have addressed this issue using data augmentation methods and generative adversarial networks (GANs). Pollastri et al. [42] proposed a modified Deep Conventional generative adversarial network (DCGAN) model to generate both skin lesion images and their segmentation masks for dermoscopic images. Lazo et al. [30] proposed a GAN model built using the pix2pix network architecture called PatchGAN. Sivanesan et al. [45] proposed a conventional GAN model with pix2pixHD and U-Net+ architecture. Shubham et al. [21] proposed a unique adversarial learning-based approach namely Efficient-GAN (EGAN) that creates precise lesion masks using an unsupervised generative network. These studies suggest that data scarcity can be dealt with using generative models and data augmentation.

Very limited studies, however, have conducted experiments to evaluate the impact of skin lesion segmentation on the performance of the lesion classifiers. A fruit fly optimization algorithm and machine learning based technique

to segment and classify skin cancer was proposed in [47]. Yu et al. [54] proposed a two staged lesion classification model with the help of deep residual network. This study showed that filtering out the useless background considerably increased the performance of the classifier, in strong agreement with our hypothesis that segmentation enhances classification performance. Zhang et al. [55] proposed an attention residual learning convolutional neural network that includes an attention method to learn representative image features without explicit segmentation, and the proposed model was able to adaptively focus on discriminative parts and achieved state-of-the-art performance. Almasni et al. [2] proposed an integrated diagnostic framework that combines skin lesion segmentation using deep segmentator called full-resolution convolutional network (FrCN) and classification accuracy were compared with different deep learning models and best performance was achieved using Inception-ResNet-v2. Diaz et al. [17] presented DermaKNet, a CAD system based on CNNs, which first segments the dermoscopic image by passing it to a lesion segmentation network (LSN) for generating the binary mask which then arguments and passes it to the diagnosis network (DN) that provides final diagnosis. Mahbod et al. [35] investigated the impact of the skin lesion segmentation on the performance of dermoscopic image classification using ISIC 2017 image dataset. In [31], the authors proposed two deep learning frameworks, the Lesion Indexing Network (LIN) by adopting FCRN-88 and the lesion feature network, which is a CNN-based framework to address lesion segmentation, classification, and dermoscopic feature extraction. Despite sparse exploration, lesion segmentation before classification is a promising approach to improve the performance of skin lesion diagnostic systems as justified by notable performance boosts in these prior studies.

2.1 Adaptations, gaps, and key contributions

Visual inspection of skin lesions with the naked eye cannot easily distinguish between different types of skin cancer and may result in an incorrect diagnosis. The use of deep learning with dermoscopy can help automate the diagnosis of skin lesions. The key challenges to skin lesion characterization as identified above include limited exploration of the segmentation prior to classification approach, sample size bias due to imbalanced datasets, and artifacts on dermoscopic images among others. We attempt to address these challenges by selecting some of the most promising approaches in literature; adapting, modifying, and/or proposing new approaches; and performing experiments to optimize the best performing lesion subtyping pipeline. The rest of this section highlights these contributions and their rationale.

EfficientNet to classify lesion segments Tan et al. [50] first demonstrated that a mobile size EfficientNet model can be scaled up very effectively, surpassing state-of-the-art accuracy with an order of magnitude fewer parameters and FLOPS, on both ImageNet and five commonly used transfer learning datasets. Furthermore, prior studies have shown that the EfficientNet backbone is effective for dermoscopic image classification. Hence, in this study, we adopt this backbone and systematically scale its complexity to identify the optimal lesion processing pipeline. Furthermore, the regime of segmentation before classification has sufficiently explored; hence, through the analysis of different stages of the classification pipeline carried out in this work, we have attempted to further explore the effectiveness of this approach.

Adversarial training for segmentation The classification of the entire dermoscopic images may not be effective, as they may contain some background features that affect the performance of the identification process. Most of the papers [15, 27] have performed classification directly on entire dermoscopic images and have performed inferiorly. Further, studies that did perform segmentation before classification [2, 17, 20, and 55] have mostly used traditional training approaches with U-Net and CNN based segmentation architectures. However, as noted earlier, complex tasks such as segmentation using deep learning require a large number of training examples when trained using traditional approaches. Hence, in this study, we employ an adversarial training approach on a BCDU-Net architecture to leverage the strength of GANs in dealing with data scarcity; with a discriminator feedback to enhance and speed-up segmentation mask learning, the segmentation network is able to produce more accurate segmentation masks with limited training data.

Preprocessing lesion segment regions to remove artifacts From the workflow proposed in Yu et al. [54], without applying preprocessing methods could leave artifacts in the segmented lesion regions that negatively impact classification performance. Hence, in addition to lesion segmentation, we apply preprocessing methods on the images prior to classification to remove unwanted artifacts using contrast limited adaptive histogram equalization (CLAHE) and image inpainting algorithms to remove artifacts such as veins, hair, etc. and for illumination correction.

Handling imbalanced datasets The dermoscopic image dataset used in this work is heavily imbalanced, with class sizes varying from 200 to 12000. The resulting sample size bias problem was not handled by majority of the previous studies. The authors of [15, 17, 27, 56] have handled this problem by randomly adding images to the minority classes and by using simple image augmentation methods.

These methods increase the risk of overfitting as they produce images similar to the existing ones. Hence, to overcome the possibility of overfitting, experiments were conducted using SMOTE oversampling technique [44, 52] as well as class reweighted training to identify an effective method for dealing with imbalance. Moreover, SMOTE is a relatively superior synthetic image generation approach as it generates new samples from the existing images, thereby increasing the generalization capabilities of the classifier. Finally, the observed strengths of each data balancing method are analyzed and discussed.

Generated ground truth masks for ISIC 2019 dataset The ISIC 2019 dataset does not contain ground truth for lesion segments. As part of this work, lesion segmented masks were generated for the ISIC 2019 dataset using the ISIC 2018 dataset to train the segmentation network. Further, the generated masks were validated with the help of clinicians. By doing so, the sample size available was increased to train the final lesion classification network that uses segmented lesion regions as inputs.

3 Methodology

In this work, two different deep learning based automatic diagnostic systems were proposed. The first system performs classification on the entire dermoscopic images to identify the lesion types, while the second system involves a two-level process, where the region of interest (ROI) was obtained through deep learning-based segmentation followed by the classification of the segmented region. In both approaches, the problem of unbalanced dataset was overcome using either the SMOTE or reweighting technique. The results were compared and analyzed using an exhaustive experimental setup. The proposed architecture is shown in Fig. 1.

3.1 Preprocessing

Dermoscopic images are characterized by a range of artifacts, and eliminating them before the training phase can significantly boost the performance of the system. To this end, the following methods were applied on the dermoscopic images as preprocessing steps to reduce the artifacts present and to enhance the image quality for further processing.

3.1.1 Hair and vein removal

The dermoscopic images in the dataset have different artifacts that include hair and veins. Since, these artifacts obscure the region of interest of the lesion images, learning

discriminating features becomes difficult for the neural network. Hence, the images were preprocessed by detecting and deleting the hair nodules using the Frangi vesselness algorithm, and the removed portions were refilled using the inpainting method based on the fast march method (FMM) [25].

3.1.2 Uneven illumination correction

Dermoscopic images obtained using noninvasive microscopic technique usually suffers from uneven illumination during image acquisition [44]. This influences the automatic analysis system to suffer from inaccurate segmentation which results in decreased classification accuracy. So, the CLAHE algorithm [41, 57] was used as an illumination correction technique to overcome this problem. The CLAHE algorithm is applied to the L channel of the LAB color space converted image, which splits each image into smaller blocks and applies contrast limiting followed by histogram equalization for each block. Finally, the contrast-enhanced L channel is merged to form a LAB image and subsequently converted to an RGB image [22]. Thus, all the images were preprocessed using the aforementioned steps and passed on for subsequent analysis.

3.2 Classification of dermoscopic images

The preprocessed images were classified into various categories of cancer types using different deep architectural models. In this proposed work, experiments were conducted to study the performance of the classifiers by training the architecture using, with and without segmenting the lesion region from the dermoscopic images. For segmenting the lesion region Bi-Directional ConvLSTM U-Net (BCDU-Net) [3] was applied and for classification EfficientNet was used. Additionally, the performance of the classifiers were analyzed, by training the classifiers using balanced and unbalanced dataset. For balancing the dataset SMOTE and Reweighting techniques were utilized. The performance of the above mentioned methods were investigated and the results were compared.

Several experiments were conducted to analyze the performance of the proposed methodology, by training the architecture with different strategies as listed:

- Classification of dermoscopic images with and without segmentation.
- Classification of dermoscopic images with balanced and unbalanced training sets.

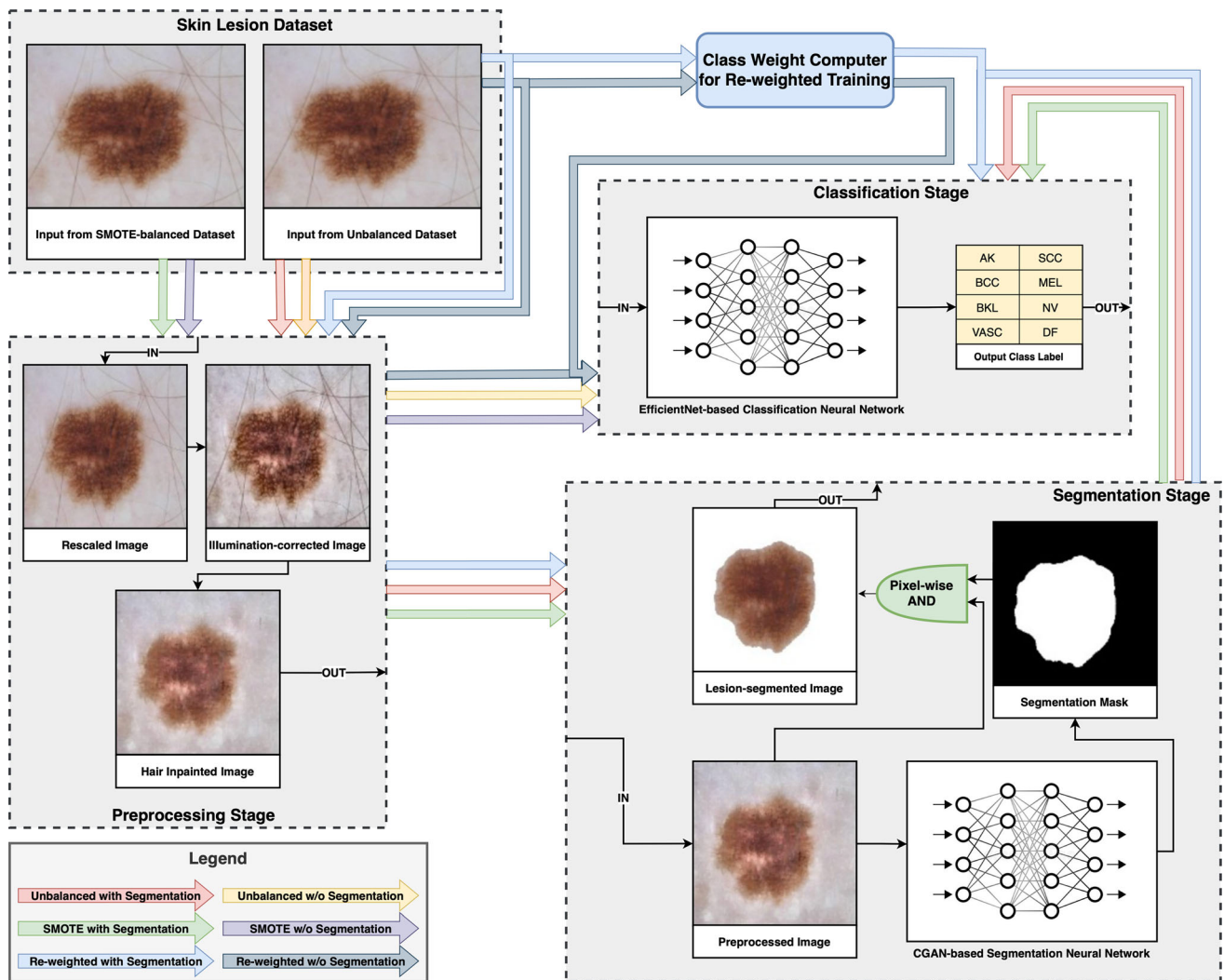


Fig. 1 Overall pipeline of the study, color-coded by the different experiments designed to analyze the effect of dataset balancing techniques and segmentation on lesion characterization

3.2.1 Classification of dermoscopic images with and without segmentation

Two different methods were proposed to classify dermoscopic images into eight different categories of cancer types that are shown in Table 1. The first method uses entire dermoscopic images while the latter uses the segmented lesion region to train the classification architecture. The BCDU-Net architecture was used for producing segmentation masks of lesions and was trained under two different settings, namely the baseline (simple supervised learning) and the generator-discriminator (CGAN method) settings. The EfficientNet architecture was used for classification, chosen for its success with skin lesion image analysis based on previous studies. EfficientNets are a family of CNNs with increasing network complexities that uses a technique called compound scaling method. Rather

Table 1 Skin lesion classes, their abbreviations, sample sizes, and weights computed for reweighted training in the ISIC 2019 dataset

Lesion type (abbreviation)	Samples	Weights
Melanoma (MEL)	4522	0.6
Melanocytic nevus (NV)	12,875	0.2
Basal cell carcinoma (BCC)	3323	0.9
Actinic keratosis (AK)	867	3.5
Benign keratosis (BKL)	2624	1.2
Deramatofibroma (DF)	239	12.9
Vascular lesion (VC)	253	12.8
Squamous cell (SCC)	628	4.9

than scaling randomly width, depth, or resolution, compound scaling method uniformly scales in all dimensions.

This approach paves the way to achieve improved accuracy for fewer parameters and less computations than previous models. Specifically, the basic architecture version B0 is scaled up to obtain different variants of EfficientNets, named B1 through B7. In our work we have used B2 variant as a baseline to analyze the classification performance.

Segmentation using Bi-Directional ConvLSTM U-Net - Baseline method An extended version of the U-Net with densely connected convolutions called BCDU-Net uses BConvLSTM layers for skip connections and reuses feature maps with densely connected convolutions. The encoding path of the architecture has four steps. It has two convolutional 3×3 filters followed by a 2×2 max-pooling function and ReLU. The feature maps are doubled at each step by collecting all the image representations, effecting an increase in dimension of these representations layer-by-layer. The high dimensional image representation of the final layer of the encoding path leads to rich feature representations.

Following the encoding path, there are three densely connected convolution layers. These densely connected convolution layers improve its performance through collective knowledge, and these feature maps are reused throughout the network. There are N consecutive blocks, and each block comprises two consecutive convolution layers. The decoding path performs up-sampling to the outputs of the previous layers. The feature maps from the contracting and expanding path are then processed using BConvLSTM. The output of the BConvLSTM is computed as shown in Eq. 1.

$$Y_t = \tanh\left(W_y^{\vec{H}} * \vec{H}_t + W_y^{\overleftarrow{H}} * \overleftarrow{H}_t + b\right), \quad (1)$$

where \vec{H}_t and \overleftarrow{H}_t denote the hidden state tensors for the forward and backward states, respectively, b is the bias term and Y_t indicates the final result. The upsampling function in each layer increases the size of each feature map and decreases the feature channels. At the end of the expanding path, the discriminative features learned by the encoder are projected onto the pixel space. Thus, the corresponding mask for the input image was generated using the baseline setup.

Segmentation using CGAN method - Discriminator-Generator model

To achieve more precise segmentation results than the BCDU-Net, CGAN was used. The CGAN model has been trained with the aforementioned Bi-Directional ConvLSTM U-Net (BCDU-Net) network, coupled with a pixel discriminator network under a conditional generative adversarial network (CGAN) setting, to perform skin lesion segmentation. In this setting, the generator was trained to

generate masks of lesions, while simultaneously, a pixel discriminator was trained to identify ground truth masks as “real” and predicted masks as “fake”. In effect, the generator eventually learns to produce masks that are close to the “real” image (the segmentation mask). The two simultaneous losses drive the generator to produce precise segmentation masks of lesions. The trained segmentation model was then applied to the entire classification data set to generate segmentation masks and validated the segmentation regions obtained with the help of medical experts. The overall architecture of the proposed segmentation network and adversarial training approach is shown in Fig. 2. The baseline BCDU-Net, an encoder-decoder neural network architecture, was used as the generator of the CGAN model. This generator produces output images, which are conditional on the input image and the produced images were classified as real or fake by the pixel discriminator network. The discriminator is a fully convolutional neural network that evaluates the generated images at the pixel level. This fine-grained discrimination helps in capturing finer details in the generated output by comparing the generated target images and the ground truth images, pixel by pixel. Since, the discriminator leverages both conditional information and pixel-wise evaluation, it is more appropriate for pixel-wise segmentation and generates accurate, well-defined lesion segmentation. The discriminator feedback is used to improve the efficiency of the generator in generating high quality masks (shown in Fig. 2).

The proposed CGAN approach can significantly increase the segmentation performance of a baseline segmentation network leveraging the conditioning imposed by the generator-discriminator learning setup. The proposed generator-discriminator model generates excellent quality segmentation masks for images with low contrast, unclear, and irregular boundaries, in comparison to the baseline segmentation network in this work. This performance improvement is analyzed in subsequent sections. EfficientNet models were then used to classify the segmented lesion images into different cancer subtypes.

Post-processing Post-processing involves the generation of the segmented region of the image by performing a logical AND operation of the binary mask with the original skin lesion image. The segmented skin lesion images obtained are then classified using the same EfficientNet variants, and the performance of the two approaches were compared.

3.2.2 Classification of dermoscopic images with balanced and unbalanced training set

The classification of the dermoscopic images was performed in two ways. Since the dataset is highly unbalanced

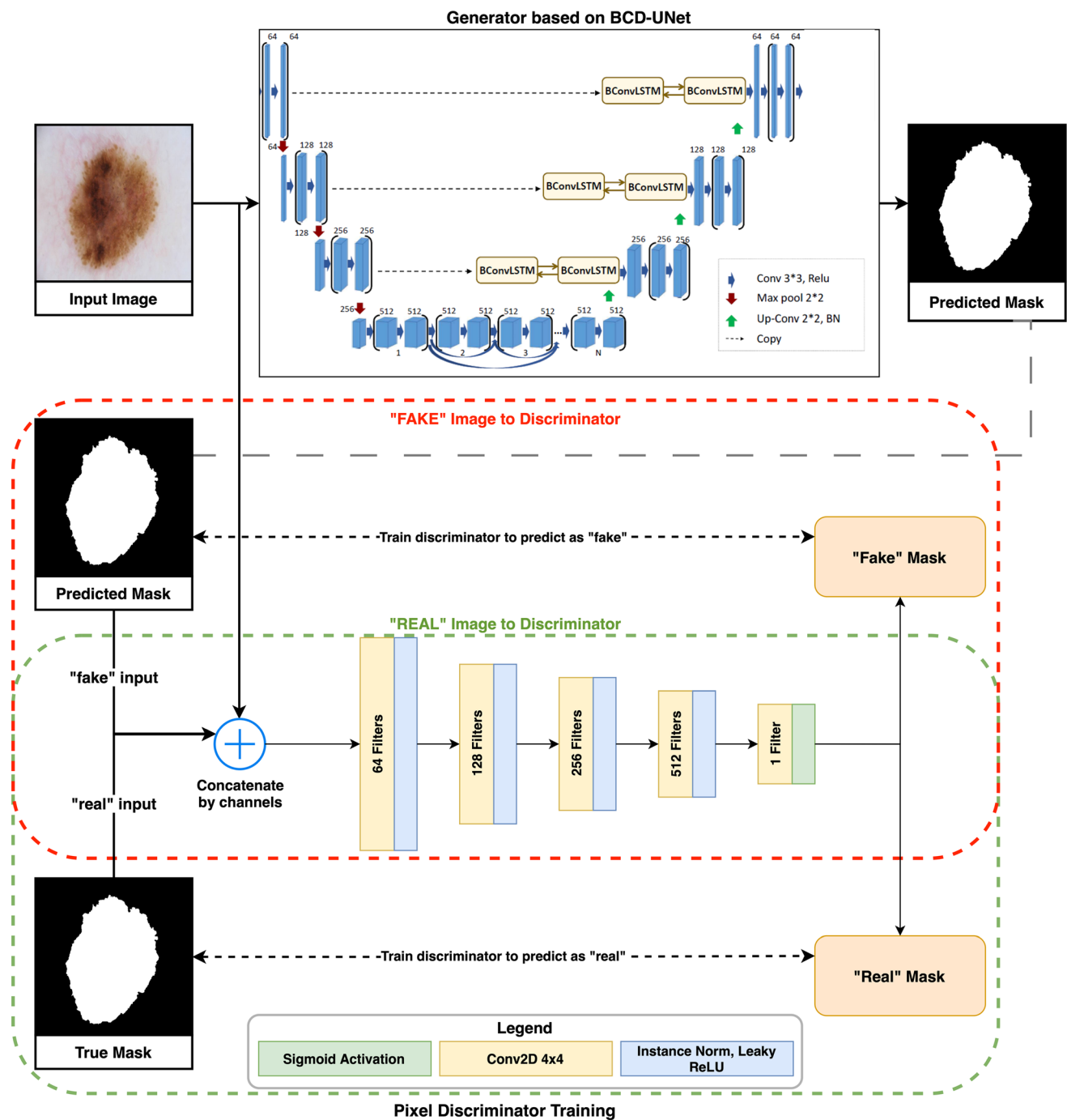


Fig. 2 Conditional adversarial training setup with BCD-UNet -based generator network and convolutional pixel discriminator to apply CGAN strategy for lesion segmentation training. *BCD-UNet architecture diagram adapted from [3]*

with an uneven number of images in each class, there is a possibility of bias toward the class having a larger number of samples. Hence, to overcome the problem of imbalance, SMOTE and reweighting techniques were applied to balance the dataset. Then, the EfficientNet architecture was used to classify both the entire dermoscopic images and segmented lesion regions. The performance of both classifiers was compared before and after balancing the dataset.

Dataset Balancing using SMOTE SMOTE [9] generates synthetic samples from the minority class to balance the dataset. The synthetic instance is created by interpolating the positive class instance with one of the k-nearest neighbors at random in the feature space. The generated synthetic data point is neither the exact copy of the positive sample nor too different from the minority class. This

algorithm helps to overcome the over fitting problem posed by random oversampling method.

Dataset balancing using reweighting In reweighting, higher weights are assigned to the minority classes and lower weights are assigned to the majority classes to influence the loss function. This method can be an alternative technique for balancing the imbalanced image dataset without the need for image resampling. The estimated class weights for unbalanced dataset were determined using the following Eq. 2.

$$\text{weight}_i = \frac{\text{total no. of samples}}{\text{no. of classes} \times \text{no. of samples in } i}, \quad (2)$$

where i denotes the class for which the weight is to be determined. The proposed system was assessed using different experiments by training the entire dermoscopic images and segmented lesions, before and after balancing the imbalanced data set.

4 Experiments and results

4.1 Skin lesion dataset

Two datasets from the ISIC Skin Lesions cohort were used as benchmark to evaluate the methods proposed in this work. The following sections describe the composition and use of these datasets in this study.

4.1.1 ISIC 2018 segmentation dataset

The 2018 segmentation dataset consists of 3694 dermoscopic images with 2594 training images, 100 validation images and 1000 testing images in the JPEG format. The binary mask ground truth for the training images are available as PNG images. A sample skin lesion image and its corresponding binary mask is shown in Fig. 3.

4.1.2 ISIC 2019 classification dataset

The dataset for ISIC 2019 challenge contains 33,569 dermoscopic images with 25,331 images in 8 categories for training as shown in Table 1. The test set contains 8238 test images given in JPEG format. The classes melanoma, vascular lesion and squamous cell carcinoma are malignant type of skin cancer and the remaining classes fall under benign lesion types. In this dataset, the ground truth contains image ID and the class label of skin lesion for each images.

In this research work, we propose to classify the images by segmenting the lesion region, but the dataset doesn't contain the segmentation mask, so we created the

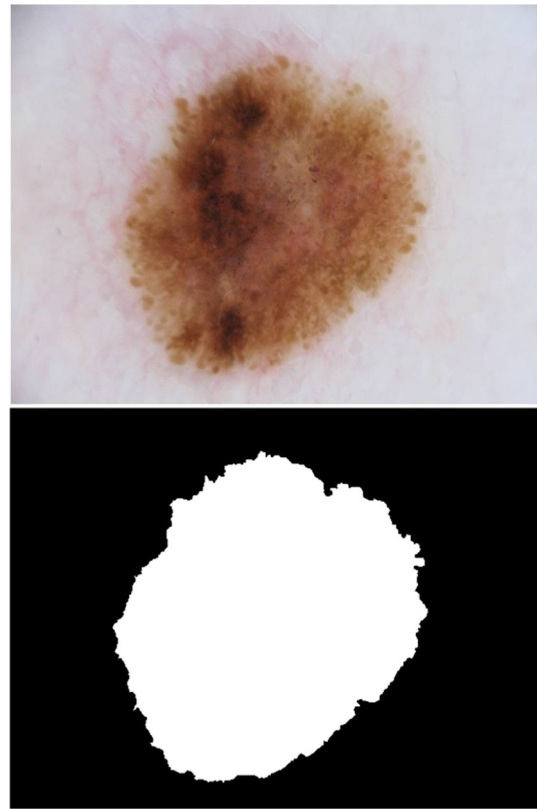


Fig. 3 ISIC 2018 segmentation dataset depicting a skin lesion image (top) and its segmentation ground truth as a binary mask (bottom)

segmentation mask using CGAN and validated with the medical expert.

4.2 Dataset partitioning

Two datasets were used as source data for this research work, the training subset is provided with corresponding ground truth. However, no validation subset is demarcated for the classification dataset, and the segmentation ground truth is not publicly available for the designated testing subset. Hence, the training subsets of both the datasets were first divided into 80% and 20% splits, randomly and universally for all experiments. During classifier training, 20% of the first split of the classifier data was used for epoch-wise validation. Whereas for training the segmentation network, the ISIC-designated validation set of 100 samples in the segmentation dataset was used for epoch-wise validation.

4.2.1 Unbalanced classification dataset

After the aforementioned split, the 25,331 training images of the ISIC 2019 classification dataset were divided into 20,265 training images and 5066 testing images. The split was performed class-wise, so that the original proportion of

samples per class is conserved in each subset of the split. The long-tailed distribution of samples in the dataset is evident in Table 1. The number of samples present in each class is not equally distributed, so it is termed as an *imbalanced dataset*; we use this classification dataset for some experiments and these experiments are labeled with the term *unbalanced* to denote that no technique was applied to balance them. In this work, reweighting and SMOTE techniques were used to balance the dataset.

4.2.2 Classification dataset balanced by reweighting

The reweighting technique for balancing assigns weights to the classes during training based on their relative proportions in the dataset, according to Eq. 2. The class weights computed are shown in Table 1.

Classification dataset balanced using SMOTE To obtain this version of the dataset, the SMOTE technique was directly applied on the original training set of 25,331 images in the ISIC 2019 dataset. SMOTE oversamples all minority classes to a count of 7000 images, resulting in a total of 61,875 images. These images were split into a training set comprising 49,500 images, and a testing set with 12,375 images. The split was performed class-wise to conserve the relative proportions of samples per class.

Segmentation Dataset The 2594 training samples in the ISIC 2018 segmentation dataset were split in 80–20 ratio to obtain 2075 training and 519 testing images. The ISIC-designated validation set of 100 samples were used for epoch-wise validation during training experiments on the segmentation networks.

4.3 Experiments for lesion segmentation

This research work compares the classification performance on lesion-segmented and entire dermoscopic images. However, the classification datasets (unbalanced and balanced, both) do not contain segmentation masks to obtain the lesion regions. Hence, the segmentation dataset, that contains 2594 dermoscopic images with lesion segmentation masks, was used to train a segmentation model. This trained segmentation model was later used to predict lesion masks for all the images in the classification datasets.

To validate the segmentation masks predicted by the trained model, 50 representative images were chosen from each class of the ISIC 2019 classification dataset. Segmentation masks for these 400 samples were then prepared by three medical experts independently, to obtain three *annotated segmentation masks* for each sample. An inter-rater agreement was then computed based on regional

overlap using the Generalized Conformity Index (GCI) metric [28] as shown in equation 3,

$$GCI = \frac{\sum_{\text{pairs}(i > j)} X_i \cap X_j}{\sum_{\text{pairs}(i > j)} X_i \cup X_j}, \quad (3)$$

where X_i represents the set of nuclear region pixels of the segmented ground truth obtained from rater i and $\sum_{\text{pairs}(i > j)} \dots$ represents the summation over every unique pair of raters.

A GCI score of 0.93 was obtained across the three raters. A bitwise OR operation was performed, thereupon, across the three *annotated segmentation masks* per sample to obtain a single *ground truth mask*. The OR operation ensures that any pixel that is marked as a lesion by at least one expert is treated as a lesion in the mask of *ground truth*. These 400 sets, each containing a dermoscopic image and its ground truth mask, was used as the testing set for the segmentation model.

After mask generation, the BCDU-Net architecture was used for segmentation and trained under two different settings, namely the baseline (simple supervised learning) and the generator-discriminator (CGAN method) settings. In the baseline setting, the model converged after 120 epochs with a validation F1-Score of 0.87 for segmentation. Whereas in the CGAN setting, that uses a pixel discriminator, the model converged in 100 epochs and obtained a validation F1-Score of 0.91. The segmentation performances, evaluated on the expert annotated test set of 400 annotated samples, were shown in Table 2. A qualitative comparison of the performance of the two models is shown in Fig. 4. The CGAN setting clearly outperforms the baseline due to the discriminator's feedback to the generator during each cycle of model training propagation. Hence, for all classification experiments, the CGAN generator model was used to predict the lesion masks because the Conditional Adversarial Segmentation approach combines the power of deep learning for image segmentation with the contextual awareness provided by conditional adversarial networks. This integration aims to enhance the precision and context-awareness of segmentation models, making them more suitable for complex segmentation tasks.

4.4 Experiments for lesion classification

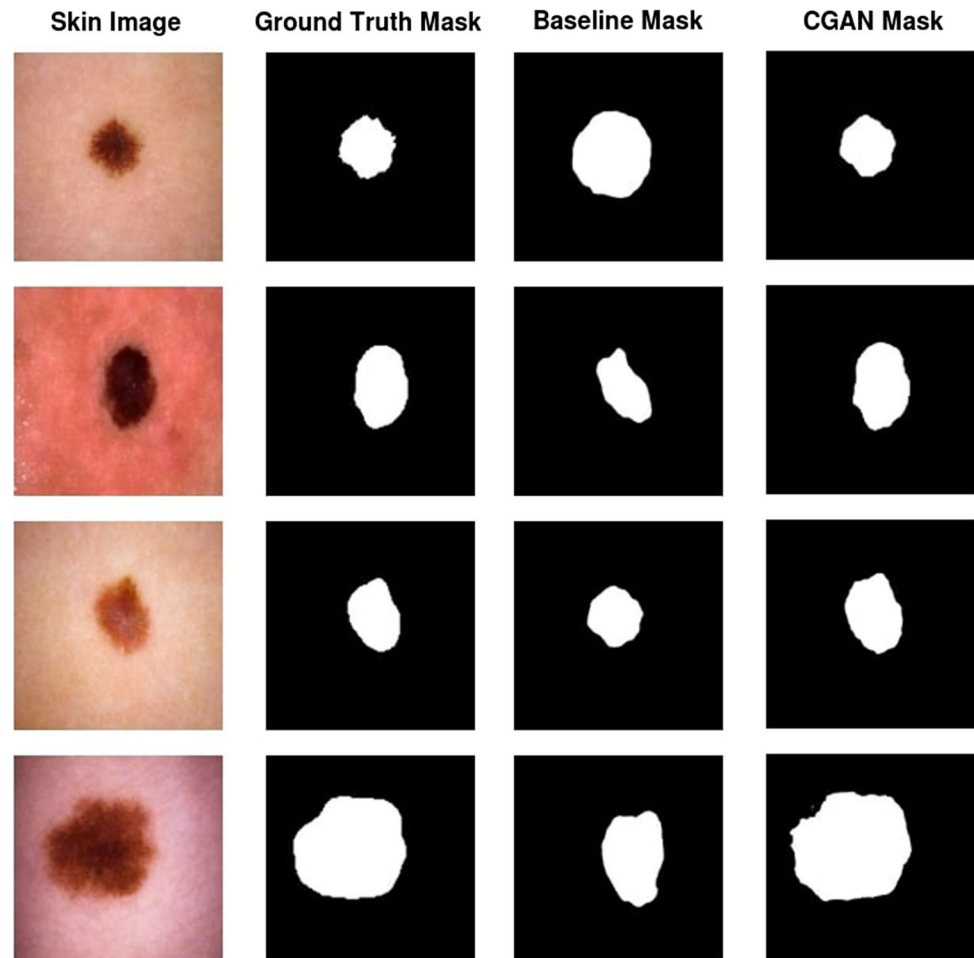
The performance of the classification architecture trained using entire dermoscopic images (EDI), and lesion segmented dermoscopic images (LSDI) were evaluated using different experimental setup to achieve the following objectives:

- To ascertain the classifier performance trained on the unbalanced dataset

Table 2 Quantitative comparison of segmentation performance trained under the baseline and CGAN settings

Method	Accuracy	Specificity	Sensitivity	Precision	F1-score
BCDU-Net: Baseline Setting	0.92	0.95	0.85	0.86	0.86
BCDU-Net + Pixel Discriminator: CGAN Setting	0.95	0.96	0.89	0.89	0.89

Evaluated on expert-annotated testing samples

Fig. 4 Qualitative comparison of segmentation performance trained under the baseline and CGAN settings

- To ascertain the classifier performance trained on the dataset balanced by reweighting
- To ascertain the classifier performance trained on the dataset balanced using SMOTE

4.4.1 Classification of entire dermoscopic images (EDI)

Experiments were performed to ascertain the classifier's performance in classifying entire dermoscopic images. Since, some of the classes in the dataset have a disproportionate sample size, the performances of the model before balancing the dataset, after balancing using SMOTE and reweighting techniques were compared.

Classifier trained on unbalanced data An EfficientNetB2 classifier model was trained for 50 epochs using dataset. A learning rate of $1e-03$ was used and a validation accuracy of 98.4%, and testing accuracy of 80.3% was obtained.

Classifier trained on balanced data using SMOTE Class imbalanced dataset is a major hurdle which leads to biased model performance, incorrect classification of minority classes and this challenges can be addressed with SMOTE method that synthetically generates samples for minority class. In this work, the dataset was first balanced using SMOTE and then classified using EfficientNetB2 classifier model. The model was trained for 50 epochs at a learning rate of $1e-03$, using the balanced dataset. A validation

accuracy of 97.4%, and testing accuracy of 89.1% were obtained and the results were compared.

Classifier trained on balanced data using reweighting Reweighting is a popular method used for class balancing which ensures better accuracy. This method effectively rebalances the loss by assigning weights for different classes. After balancing, an EfficientNetB2 model was trained with the class weights shown in Sect. 4.2 for 50 epochs at a learning rate of $1e-03$, to obtain a validation accuracy of 95.4%, and a testing accuracy of 79.9%.

The class-wise comparison between unbalanced and balanced using the SMOTE and reweighting techniques is shown in Table 3. It is tacit that the NV class has the highest number of samples, whereas VASC has the lowest number of samples. Therefore, NV attains the highest accuracy of 0.91% without balancing and VASC obtained a lesser accuracy comparatively. When comparing the SMOTE and reweighting methods, SMOTE accomplished better performance than reweighting.

4.4.2 Classification of lesion segmented dermoscopic images

Experiments were performed to ascertain the classifier's performance in classifying lesion segmented dermoscopic images. Segmented skin lesions were obtained using segmentation masks predicted by a previously trained BCDU-Net based segmentation model and the CGAN setting. Since the segmentation model achieved better results when trained under the CGAN setting, this version of the model was used to produce the segmentation masks for all classification experiments. The performance of the model before balancing the dataset and after balancing using SMOTE and reweighting technique was compared and presented. Figures 5 and 6 show the sample predictions from the ISIC 2019 dataset.

Classifier trained on unbalanced data An EfficientNetB2 classifier model was trained using dataset. It converged

after 50 epochs at a learning rate of $1e-03$, and attained a validation accuracy of 99.3%, and a testing accuracy of 79.6%.

Classifier trained on balanced data using reweighting An EfficientNetB2 model was trained with the class weights shown in Sect. 4.2 for 50 epochs at a learning rate of $1e-03$, to obtain a validation accuracy of 88.7%, and a testing accuracy of 74.3%.

Classifier trained on balanced data using SMOTE An EfficientNetB2 model was trained for 50 epochs at a learning rate of $1e-03$, using dataset. A validation accuracy of 98.6% and a testing accuracy of 91.1% were obtained. Tables 4 and 5 present class based results of the classifier model trained with segmented and unsegmented images before and after balancing the dataset. Table 4 shows that VASC, SCC, MEL and DF produced accuracy of 0.71, 0.58, 0.67 and 0.59, respectively which is an improved accuracy for segmented classification over unsegmented classification. This improvement in accuracy is due to the fact that the lesion and the background region are highly similar. So segmenting the lesion region and classifying based on the ROI showed good accuracy for the corresponding classes.

From Table 5 it is evident that segmenting the lesion region and balancing the dataset using SMOTE, produced very high accuracy when compared to unbalanced dataset shown in Table 4. For the classes AK, BKL, SCC, MEL and DF showed a very high improvement in terms of accuracy from 0.54, 0.60, 0.58, 0.67 and 0.59 to 0.91, 0.83, 0.91, 0.84 and 0.99, respectively, because segmenting and balancing the dataset. The comparison between segmented, unsegmented and balancing using SMOTE and reweighting technique is shown in Table 6. From the results it is visible that segmenting the lesion from the images and balancing the data set using SMOTE technique outperforms for all the classes.

Table 3 Comparison between unbalanced and balanced dataset using SMOTE and reweighted training techniques

Classes	Unbalanced classification				SMOTE				Reweighted training			
	Acc	Pre	Rec	F1-S	Acc	Pre	Rec	F1-S	Acc	Pre	Rec	F1-S
AK	0.57	0.61	0.57	0.59	0.90	0.91	0.90	0.90	0.51	0.50	0.51	0.51
BCC	0.86	0.73	0.86	0.79	0.83	0.85	0.83	0.84	0.70	0.84	0.70	0.71
BKL	0.68	0.62	0.68	0.65	0.81	0.85	0.81	0.83	0.68	0.69	0.68	0.69
VASC	0.68	0.88	0.68	0.77	1.00	0.98	1.00	0.99	0.82	0.88	0.82	0.85
SCC	0.56	0.75	0.56	0.64	0.96	0.89	0.96	0.92	0.61	0.75	0.61	0.67
MEL	0.60	0.79	0.60	0.68	0.70	0.84	0.70	0.76	0.71	0.71	0.71	0.71
NV	0.91	0.87	0.91	0.89	0.93	0.87	0.93	0.90	0.91	0.86	0.91	0.89
DF	0.57	0.86	0.57	0.68	0.99	0.95	0.99	0.97	0.64	0.78	0.64	0.70

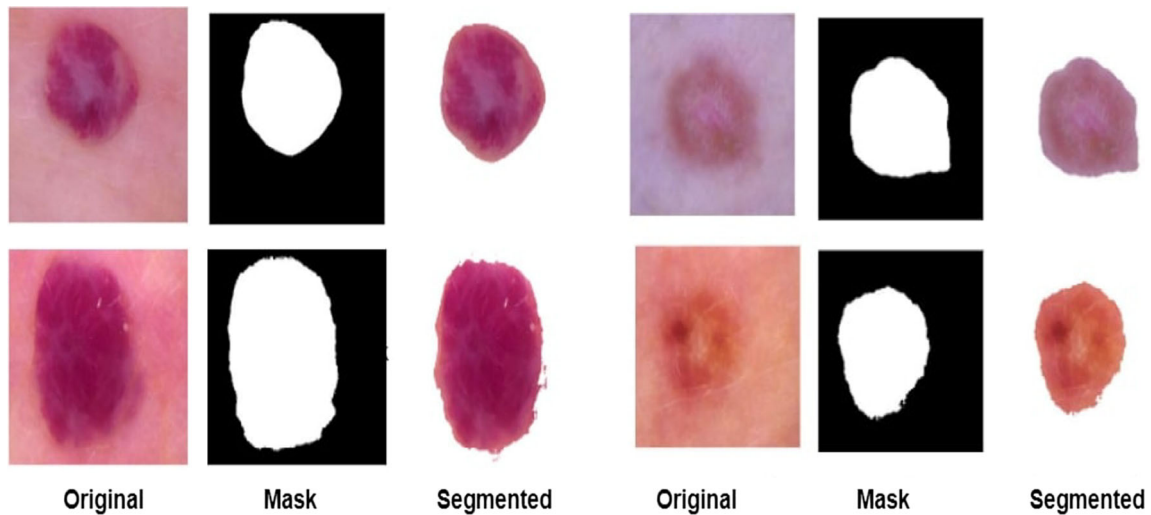


Fig. 5 Comparison of predictions from the BCDU-Net for (left) VASC and (right) DF

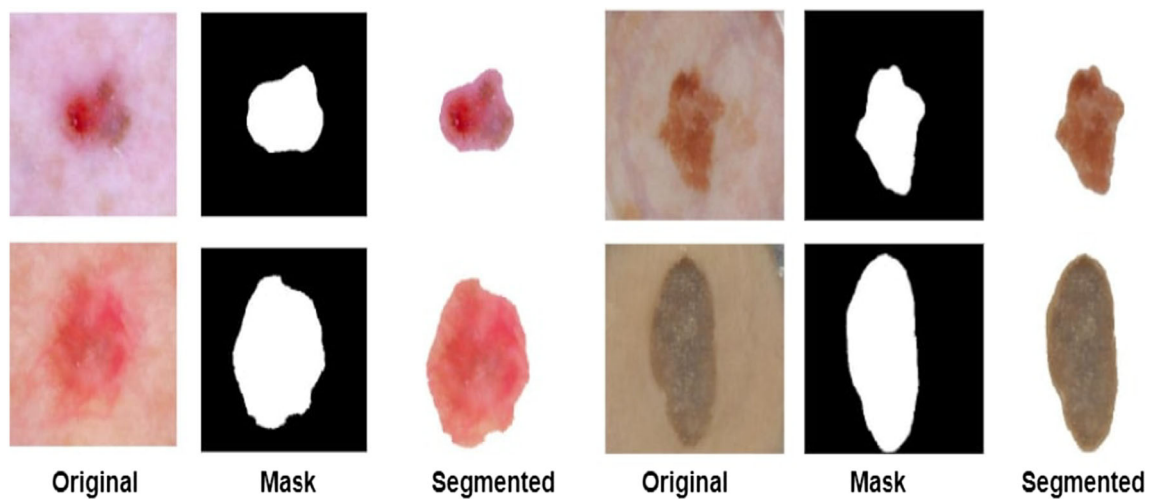


Fig. 6 Comparison of predictions from the BCDU-Net for (left) BCC and (right) BKL

Table 4 Comparison between unsegmented and segmented classification (unbalanced)

Classes	Unsegmented classification				Segmented classification			
	Acc	Precision	Recall	F1-score	Acc	Precision	Recall	F1-score
AK	0.57	0.61	0.57	0.59	0.54	0.62	0.54	0.58
BCC	0.86	0.73	0.86	0.79	0.76	0.76	0.76	0.76
BKL	0.68	0.62	0.68	0.65	0.60	0.66	60	0.63
VASC	0.68	0.88	0.68	0.77	0.71	0.82	0.71	0.76
SCC	0.56	0.75	0.56	0.64	0.58	0.68	0.58	0.63
MEL	0.60	0.79	0.60	0.68	0.67	0.73	0.67	0.70
NV	0.91	0.87	0.91	0.89	0.91	0.86	0.91	0.88
DF	0.57	0.86	0.57	0.68	0.59	0.87	0.59	0.70

The highest accuracies obtained for that class are highlighted in bold

Table 5 Comparison between unsegmented and segmented classification (balanced)

Classes	Unsegmented classification				Segmented classification			
	Acc	Precision	Recall	F1-score	Acc	Precision	Recall	F1-score
AK	0.90	0.91	0.90	0.90	0.91	0.86	0.91	0.89
BCC	0.83	0.85	0.83	0.84	0.76	0.87	0.76	0.81
BKL	0.81	0.85	0.81	0.83	0.83	0.77	0.83	0.80
VASC	1.00	0.99	1.00	0.99	0.99	0.98	0.99	0.99
SCC	0.96	0.89	0.96	0.92	0.91	0.95	0.91	0.93
MEL	0.70	0.84	0.70	0.76	0.84	0.83	0.84	0.83
NV	0.93	0.87	0.93	0.90	1.00	1.00	1.00	1.00
DF	0.99	0.95	0.99	0.97	0.99	0.97	0.99	0.98

The highest accuracies obtained for that class are highlighted in bold

Table 6 Comparison between SMOTE and reweighting for unsegmented and segmented classification

Classes	Unsegmented		Segmented	
	SMOTE	Weights	SMOTE	Weights
AK	0.9	0.51	0.91	0.51
BCC	0.83	0.70	0.76	0.70
BKL	0.81	0.68	0.83	0.58
VASC	1.00	0.82	0.99	0.82
SCC	0.96	0.61	0.91	0.62
MEL	0.7	0.71	0.84	0.69
NV	0.93	0.91	1.00	0.81
DF	0.99	0.64	0.99	0.57

The highest accuracies obtained for that class are highlighted in bold

5 Comparison with existing work

We have compared our results in terms of precision, recall, F1-score and class average and contrasted them with the results of previous investigations in the literature. The comparison results are shown in the following Tables 7, 8, 9 and 10.

Table 7 Comparison of methods—precision

Method	Precision							
	AK	BCC	BKL	DF	MEL	NV	SCC	VASC
Ensemble of multi-Res EfficientNets + SEN154 2 [16]	0.43	0.65	0.71	0.47	0.76	0.93	0.42	0.50
Ensemble of EfficientNetB3-B4-SeResNeXt101 [56]	0.33	0.55	0.58	0.37	0.70	0.87	0.29	0.44
Ensemble, ood threshold 100% [43]	0.38	0.53	0.57	0.50	0.76	0.83	0.42	0.61
<i>Proposed methods</i>								
Unsegmented, unbalanced	0.61	0.73	0.62	0.86	0.79	0.87	0.75	0.88
Unsegmented, balanced using SMOTE	0.91	0.85	0.85	0.95	0.84	0.87	0.89	0.98
Segmented, unbalanced	0.62	0.76	0.66	0.87	0.73	0.86	0.68	0.82
Segmented, balanced using SMOTE	0.86	0.87	0.77	0.97	0.83	1.00	0.95	0.98

The highest accuracies obtained for that class are highlighted in bold

Precision estimates the reliability of correct predictions. From our experiments it is proved that our classifiers produced improved precision for most of the classes when compared with [15, 43, 56]. This proves our system produces high diagnostic accuracy which helps in critical decision making. A high recall rate helps in minimizing the number of false negative which could avoid in delayed treatment or an incorrect prognosis. And our experiments have proved to produce good results in real time systems with high recall rate. From all the experiments conducted in this research work it is demonstrated that segmenting and balancing the dataset produced improved precision and recall which results in high F1-score. From Table 10 it is apparent that after taking the average of the performance metrics the proposed system produces 19% increase in average F1-score.

Table 8 Comparison of methods—recall

Method	Recall							
	AK	BCC	BKL	DF	MEL	NV	SCC	VASC
Ensemble of multi-Res EfficientNets + SEN154 2 [16]	0.48	0.72	0.39	0.58	0.59	0.71	0.44	0.64
Ensemble of EfficientNetB3-B4-Seresnext101 [56]	0.60	0.84	0.56	0.68	0.68	0.81	0.52	0.60
Ensemble, ood threshold 100% [43]	0.32	0.85	0.55	0.57	0.68	0.87	0.45	0.56
<i>Proposed methods</i>								
Unsegmented, unbalanced	0.57	0.86	0.68	0.57	0.60	0.91	0.56	0.68
Unsegmented, balanced using SMOTE	0.90	0.83	0.81	0.99	0.70	0.93	0.96	1.00
Segmented, unbalanced	0.54	0.76	0.60	0.59	0.67	0.91	0.58	0.71
Segmented, balanced using SMOTE	0.91	0.76	0.83	0.99	0.84	1.00	0.91	1.00

The highest accuracies obtained for that class are highlighted in bold

Table 9 Comparison of methods—F1-score

Method	F1-score							
	AK	BCC	BKL	DF	MEL	NV	SCC	VASC
Ensemble of multi-Res EfficientNets + SEN154 2 [16]	0.45	0.68	0.51	0.52	0.67	0.86	0.43	0.56
Ensemble of EfficientNetB3-B4-SeResNeXt101 [56]	0.42	0.66	0.57	0.48	0.69	0.84	0.37	0.51
Ensemble, ood threshold 100% [43]	0.35	0.66	0.56	0.53	0.72	0.85	0.43	0.59
<i>Proposed methods</i>								
Unsegmented, unbalanced	0.59	0.79	0.65	0.68	0.68	0.89	0.64	0.77
Unsegmented, balanced using SMOTE	0.90	0.84	0.83	0.97	0.76	0.90	0.92	0.99
Segmented, unbalanced	0.58	0.76	0.63	0.70	0.70	0.88	0.63	0.76
Segmented, balanced using SMOTE	0.89	0.81	0.80	0.98	0.83	1.00	0.93	0.99

The highest accuracies obtained for that class are highlighted in bold

Table 10 Comparison of methods—evaluation metrics (class average)

Methods	Accuracy	Precision	Recall	F1-score
Ensemble of multi-Res EfficientNets + SEN154 2 [16]	0.92	0.60	0.51	0.52
Ensemble of EfficientNetB3-B4-SeResNeXt101 [56]	0.91	0.51	0.61	0.53
Ensemble, ood threshold 100% [43]	0.92	0.58	0.54	0.52
Unsegmented, unbalanced	0.80	0.76	0.68	0.71
Unsegmented, balanced using SMOTE	0.89	0.89	0.89	0.89
Segmented, unbalanced	0.80	0.75	0.67	0.71
Segmented, balanced using SMOTE	0.91	0.90	0.90	0.90

The highest accuracies obtained for that class are highlighted in bold

6 Discussion

6.1 Effects of balancing the dataset

This section examines the effects of balancing the dataset before training the model for classification. From Table 1, it can be seen that the distribution of samples is highly disproportionate, producing a long-tailed distribution. Since, deep learning methods cannot handle such heavily imbalanced datasets due to consequent bias towards classes with more samples, there is a need to balance the dataset.

This is supported by the empirical results obtained in this work, shown in Table 3. Classifiers were trained on both unbalanced and balanced datasets, and the accuracy of classification is significantly better when using the balanced datasets.

Furthermore, the method used to balance the dataset plays an important role in determining the model performance. It is clear from Table 6 that the reweighting technique does little to improve the performance. In fact, the classifier performance is worse than that obtained using the unbalanced dataset for some of the classes. On the

contrary, the SMOTE technique has been very effective in improving classifier performance. This is because the reweighting technique only changes the priority towards the classes by assigning balanced class weights for each class, following Eq. 2. Clearly, the bias towards high sample size classes is not eliminated by such a simplistic relation to repeatedly propagate samples from the under-represented classes. The SMOTE technique that generates new synthetic samples by oversampling minority classes, provides a complex enough balancing strategy to notably diminish the inter-class bias.

SMOTE stands out as a computationally less intensive alternative compared to reweighting methods. Reweighting methods involve adjusting the weights assigned to instances within the dataset to balance the influence of the minority and majority classes during model training resulting in high computational cost. Recalculating probabilities and updating model parameters for each instance can be resource-intensive, especially when dealing with large datasets. The efficiency of SMOTE lies in its ability to augment the dataset with synthetic samples, allowing the model to learn the underlying patterns of the minority class without a substantial increase in computational demands. SMOTE offers a practical and computationally efficient solution for handling imbalanced datasets and considered to be a valuable tool for addressing class imbalance without compromising efficiency.

The other most popular methods for balancing datasets are random under-sampling and random over-sampling. Random under-sampling discards instances from the majority class, potentially leading to loss of valuable information, while random over-sampling duplicates instances, which may not contribute new insights and can lead to overfitting. Hence, these methods may not contribute to improved generalization because random under-sampling may removes instances without considering their importance, and random over-sampling duplicates existing information. Also these methods may not be as computationally efficient, especially in the case of random over-sampling, which increases the dataset size.

6.2 Effects of segmenting the skin lesions

The original unsegmented dataset comprises irrelevant information that does not help in discerning between lesion types. The non-lesion portions are predominantly just regions of clear skin, but on occasion, they also include other objects such as reference stickers and marker annotations. When comparing the classification results using entire lesion segmented dermoscopic images (Table 5), it is evident that classifiers trained using entire lesion segmented images consistently outperform their entire image counterparts. And this observation holds for both balanced,

as well as unbalanced datasets. Hence, the nonlesion regions act as artifacts that negatively impact classification accuracy. The classifier performance significantly improves when only relevant portions of the image are supplied, since the classifier trains to learn representative features exclusively from the portions relevant for classification.

6.3 Scaling the classification model

Compound scaling was performed using EfficientNet variants to analyze the improvement in classifier performance at the expense of increased usage of compute resources and training time. The scaling was performed on the best experimental setting for classification with SMOTE balanced dataset and lesion segmentation. It is evident from Table 11 that the B6 and B7 configurations produce significantly better classifiers than their lower versions, suggesting that the proposed skin lesion classification pipeline scales well to improve performance. Furthermore, since B6 performs almost as well as B7, lesser resource usage is required to tune to best performance.

7 Conclusion

Accurately characterizing skin lesion type from dermoscopic images is essential to provide timely treatment, as manual identification of lesions is time and resource intensive and prone to human error. Performing lesion segmentation before classification improves performance, as it considers only the exact regions of the lesion, ignoring unwanted portions of images. Specific preprocessing methods further help alleviate artifacts in the lesion regions. We finally propose, after comparing different components of the classification pipeline through experiments, a novel pipelined approach which accurately delineates the lesion region present in the dermoscopic images and uses it to identify the lesion type. In this pipeline, the lesion regions are extracted using a BCDU-

Table 11 Comparison of performances of different EfficientNet variants to compound scale the model for best classification performance

EfficientNet variant	Accuracy	Precision	Recall	F1-score
B2	91.03	0.90	0.90	0.90
B3	91.44	0.89	0.93	0.90
B4	92.19	0.91	0.92	0.87
B5	94.88	0.95	0.94	0.94
B6	96.87	0.96	0.97	0.96
B7	96.68	0.96	0.97	0.95

The highest accuracies obtained for that class are highlighted in bold

Net model trained adversarially using a conditional GAN approach; this segmentation network achieves improved lesion segmentation accuracy than the baseline setting that uses traditional training. The dataset used in this work is highly imbalanced and is known to lead to sample size bias, resulting in reduced classification accuracy. To improve classification, we experimented with two balancing techniques: SMOTE and reweighting; we found that the classification performance was better with SMOTE balancing method and avoided overfitting.

The proposed pipeline, however, is a sequential workflow that first performs segmentation followed by classification using two independent networks. Hence, the inference time overhead can be substantial, depending on the compute power available and complexity of the network used. Future work can explore using a common backbone for both segmentation and classification; in doing so, a single network can capture lesion regions and subtype the lesion simultaneously, saving inference time. Moreover, although deep learning approaches have demonstrated remarkable performance in medical image analysis, its applicability and generalizability to unseen, out-of-distribution data are limited by its dependence on high-quality annotations [48]. Few-shot learning is a promising approach that future studies could explore to better generalize trained models to new datasets, to perform reasonably with very few examples from a given data cohort, as well as to minimize inference time overheads.

Author contributions Study design and conceptualization were done by KD, MP, JSM; Experiments and implementation were done by KD, AS, DR, and DV; Validation were done by MP, JSM, KD; Manuscript preparation, review, and editing were done by JSM, MP, KD. All authors have read and agreed to the published version of the manuscript.

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Data availability Data supporting the findings of this study are available from the ISIC challenge websites (2018 and 2019) at <https://challenge.isic-archive.com/data/#2018> and <https://challenge.isic-archive.com/data/#2019>. The generated and validated skin lesion masks will be made available by the corresponding author upon reasonable request.

Code availability The implementation of this research work will be made available at <https://github.com/karthik-d/lesion-characterization-using-cgan>.

Declarations

Conflict of interest The authors declare that they have no conflict of interest.

Compute resources used The implementation of these experiments was carried out using an Intel Xeon 2.20GHz CPU, CUDA-enabled Tesla T4 and Tesla P100 GPUs with RAM sizes of 25 GB each, and a NVIDIA-SMI 460.32.03 TPU with a RAM size of 35 GB.

Ethical approval The data used in this study is publicly available at: <https://challenge.isic-archive.com/data/#2018> and <https://challenge.isic-archive.com/data/#2019>.

References

1. Abbasi NR, Shaw HM, Rigel DS et al (2004) Early diagnosis of cutaneous melanoma: revisiting the ABCD criteria. *JAMA* 292(22):2771–2776
2. Al-Masni MA, Kim DH, Kim TS (2020) Multiple skin lesions diagnostics via integrated deep convolutional networks for segmentation and classification. *Comput Methods Programs Biomed* 190:105351
3. Azad R, Asadi-Aghbolaghi M, Fathy M, et al (2019) Bi-directional convlstm u-net with Densley connected convolutions. In: *Proceedings of the IEEE/CVF international conference on computer vision workshops*, pp 0–0
4. Barata C, Celebi ME, Marques JS (2019) A survey of feature extraction in dermoscopy image analysis of skin cancer. *IEEE J Biomed Health Inform* 23(3):1096–1109. <https://doi.org/10.1109/JBHI.2018.2845939>
5. Batista GE, Bazzan AL, Monard MC et al (2003) Balancing training data for automated annotation of keywords: a case study. *Wob* 3:10–8
6. Bi L, Feng DD, Fulham M et al (2020) Multi-label classification of multi-modality skin lesion via hyper-connected convolutional neural network. *Pattern Recogn* 107:107502
7. Celebi ME, Aslandogan YA, Stoecker WV et al (2007) Unsupervised border detection in dermoscopy images. *Skin Res Technol* 13:454–462
8. Celebi ME, Wen Q, Hwang S et al (2013) Lesion border detection in dermoscopy images using ensembles of thresholding methods. *Skin Res Technol* 19:e252–e258
9. Chawla NV, Bowyer KW, Hall LO et al (2002) Smote: synthetic minority over-sampling technique. *J Artif Intell Res* 16:321–357
10. Delgado-Gómez D, Butakoff C, Ersbøll BK et al (2008) Independent histogram pursuit for segmentation of skin lesions. *IEEE Tran Biomed Eng* 55:157–161
11. Desingu K, Palaniappan M, Kumar J (2021) Snake species classification using transfer learning. In: *Conference and labs of the evaluation forum*, vol 2936, pp 1571–1578
12. Desingu K, Bhaskar A, Palaniappan M, et al (2022) Classification of fungi species: a deep learning based image feature extraction and gradient boosting ensemble approach. In: *Conference and labs of the evaluation forum*. <https://ceur-ws.org/Vol-3180/paper-162.pdf>
13. Desingu K, Chandrabose A (2022) Few-shot classification of skin lesions from dermoscopic images by meta-learning representative embeddings. <https://doi.org/10.48550/ARXIV.2210.16954>
14. Erkol B, Moss RH, Stanley RJ et al (2005) Automatic lesion boundary detection in dermoscopy images using gradient vector flow snakes. *Skin Re Technol* 11:17–26
15. Gessert N, Nielsen M, Shaikh M et al (2020) Skin lesion classification using ensembles of multi-resolution EfficientNets with meta data. *MethodsX* 7:100864. <https://doi.org/10.1016/j.mex.2020.100864> (<https://www.sciencedirect.com/science/article/pii/S2215016120300832>)


16. Gessert N, Nielsen M, Shaikh M et al (2020) Skin lesion classification using ensembles of multi-resolution EfficientNets with meta data. *MethodsX* 7:100864
17. Gonzalez-Diaz I (2018) Dermaknet: incorporating the knowledge of dermatologists to convolutional neural networks for skin lesion diagnosis. *IEEE J Biomed Health Inform* 23(2):547–559
18. Goyal M, Oakley A, Bansal P et al (2020) Skin lesion segmentation in dermoscopic images with ensemble deep learning methods. *IEEE Access* 8:4171–4181. <https://doi.org/10.1109/ACCESS.2019.2960504>
19. Haenssle H, Fink C, Schneiderbauer R et al (2018) Man against machine: diagnostic performance of a deep learning convolutional neural network for dermoscopic melanoma recognition in comparison to 58 dermatologists. *Ann Oncol: Off J Eur Soc Med Oncol* 29:1836–1842
20. Hasan SN, Gezer M, Azeez RA, et al (2019) Skin lesion segmentation by using deep learning techniques. In: 2019 Medical technologies congress (TIPTKNO). IEEE, pp 1–4
21. Innani S, Dutande P, Baid U et al (2023) Generative adversarial networks based skin lesion segmentation. *Sci Rep* 13(1):13467
22. Izadi S, Mirikharaji Z, Kawahara J, et al (2018) Generative adversarial networks to segment skin lesions. In: 2018 IEEE 15th international symposium on biomedical imaging (ISBI 2018), pp 881–884. <https://doi.org/10.1109/ISBI.2018.8363712>
23. Jafari MH, Karimi N, Nasr-Esfahani E, et al (2016) Skin lesion segmentation in clinical images using deep learning. In: 2016 23rd international conference on pattern recognition (ICPR). IEEE, pp 337–342
24. Jaisakthi S, Desingu K, Mirunalini P et al (2023) A deep learning approach for nucleus segmentation and tumor classification from lung histopathological images. *Netw Model Anal Health Inform Bioinform* 12(1):22
25. Jaisakthi SM, Mirunalini P, Aravindan C (2018) Automated skin lesion segmentation of dermoscopic images using grabcut and k-means algorithms. *IET Comput Vision* 12(8):1088–1095
26. Jensen JD, Elewski B (2015) The ABCDEF rule: combining the “ABCDE rule” and the “ugly duckling sign” in an effort to improve patient self-screening examinations. *J Clin Aesth Dermatol* 8(2):15
27. Kassem MA, Hosny KM, Fouad MM (2020) Skin lesions classification into eight classes for ISIC 2019 using deep convolutional neural network and transfer learning. *IEEE Access* 8:114822–114832
28. Kouwenhoven E, Giezen M, Struikmans H (2009) Measuring the similarity of target volume delineations independent of the number of observers. *Phys Med Biol* 54(9):2863. <https://doi.org/10.1088/0031-9155/54/9/018> (<https://dx.doi.org/10.1088/0031-9155/54/9/018>)
29. Krishna SR, Gudur A, Jain S, et al (2024) Deep learning for automatic diagnosis of skin cancer using dermoscopic images. *J Artif Intell Technol*
30. Lazo C (2021) Segmentation of skin lesions and their attributes using generative adversarial networks. [arXiv:2102.00169](https://arxiv.org/abs/2102.00169)
31. Li Y, Shen L (2018) Skin lesion analysis towards melanoma detection using deep learning network. *Sensors* 18(2):556
32. Liu L, Tsui YY, Mandal M (2021) Skin lesion segmentation using deep learning with auxiliary task. *J Imaging* 7(4):67
33. Liu S, Chen R, Gu Y et al (2023) AcneTyper: an automatic diagnosis method of dermoscopic acne image via self-ensemble and stacking. *Technol Health Care* 31(4):1171–1187
34. Mahbod A, Schaefer G, Wang C et al (2020) Transfer learning using a multi-scale and multi-network ensemble for skin lesion classification. *Comput Methods Programs Biomed* 193:105475
35. Mahbod A, Tschandl P, Langs G et al (2020) The effects of skin lesion segmentation on the performance of dermoscopic image classification. *Comput Methods Programs Biomed* 197:105725
36. Mirikharaji Z, Abhishek K, Bissoto A et al (2023) A survey on deep learning for skin lesion segmentation. *Med Image Anal* 88:102863. <https://doi.org/10.1016/j.media.2023.102863>
37. Mohamed EH, El-Behaidy WH (2019) Enhanced skin lesions classification using deep convolutional networks. In: 2019 Ninth international conference on intelligent computing and information systems (ICICIS), pp 180–188. <https://doi.org/10.1109/ICICIS46948.2019.9014823>
38. Muthulakshmi V, Hemapriya N (2023) CNN-based dermoscopic analysis of vascular skin lesions in the prognosis of skin lesion sarcoma based on ensemble learning. *J Intell Fuzzy Syst* 1–20
39. Nachbar F, Stolz W, Merkle T et al (1994) The ABCD rule of dermatoscopy: high prospective value in the diagnosis of doubtful melanocytic skin lesions. *J Am Acad Dermatol* 30(4):551–559
40. Palaniappan M, Desingu K, Bharathi H, et al (2022) Deep learning and gradient boosting ensembles for classification of snake species. In: Conference and labs of the evaluation forum. <https://ceur-ws.org/Vol-3180/paper-173.pdf>
41. Pizer SM, Amburn EP, Austin JD et al (1987) Adaptive histogram equalization and its variations. *Comput Vision Graph Image Process* 39(3):355–368. [https://doi.org/10.1016/S0734-189X\(87\)80186-X](https://doi.org/10.1016/S0734-189X(87)80186-X) (<https://www.sciencedirect.com/science/article/pii/S0734189X8780186X>)
42. Pollastri F, Bolelli F, Palacios RP, et al (2018) Improving skin lesion segmentation with generative adversarial networks. In: 2018 IEEE 31st international symposium on computer-based medical systems (CBMS). IEEE, pp 442–443
43. Pollastri F, Parreño M, Maroñas J et al (2021) A deep analysis on high-resolution dermoscopic image classification. *IET Comput Vision* 15(7):514–526
44. Shah V, Autee P, Sonawane P (2020) Detection of melanoma from skin lesion images using deep learning techniques. In: 2020 International conference on data science and engineering (ICDSE). IEEE, pp 1–8
45. Sivanesan U, Braga LH, Sonnadara RR, et al (2019) Unsupervised medical image segmentation with adversarial networks: from edge diagrams to segmentation maps. [arXiv:1911.05140](https://arxiv.org/abs/1911.05140)
46. Song Y, Cai W, Huang H et al (2015) Large margin local estimate with applications to medical image classification. *IEEE Trans Med Imaging* 34(6):1362–1377. <https://doi.org/10.1109/TMI.2015.2393954>
47. Sonia R, Joseph J, Kalaiyarasi D et al (2024) Segmenting and classifying skin lesions using a fruit fly optimization algorithm with a machine learning framework. *Automatika* 65(1):217–231
48. Suganyadevi S, Seethalakshmi V, Balasamy K (2022) A review on deep learning in medical image analysis. *Int J Multimed Inf Retr* 11(1):19–38
49. Tahir M, Naeem A, Malik H et al (2023) Dscnet: multi-classification deep learning models for diagnosing of skin cancer using dermoscopic images. *Cancers* 15(7):2179
50. Tan M, Le Q (2019) Efficientnet: Rethinking model scaling for convolutional neural networks. In: International conference on machine learning, PMLR, pp 6105–6114
51. Trang K, Nguyen HA, TonThat L, et al (2022) An ensemble voting method of pre-trained deep learning models for skin disease identification. In: 2022 IEEE international conference on cybernetics and computational intelligence (CyberneticsCom), pp 445–450. <https://doi.org/10.1109/CyberneticsCom55287.2022.9865634>
52. Venkataramana L, Prasad DVV, Saraswathi S et al (2022) Classification of Covid-19 from tuberculosis and pneumonia using deep learning techniques. *Med Biol Eng Comput* 60(9):2681–2691

53. Vestergaard M, Macaskill P, Holt P et al (2008) Dermoscopy compared with naked eye examination for the diagnosis of primary melanoma: a meta-analysis of studies performed in a clinical setting. *Br J Dermatol* 159(3):669–676
54. Yu L, Chen H, Dou Q et al (2016) Automated melanoma recognition in dermoscopy images via very deep residual networks. *IEEE Trans Med Imaging* 36(4):994–1004
55. Zhang J, Xie Y, Xia Y et al (2019) Attention residual learning for skin lesion classification. *IEEE Trans Med Imaging* 38(9):2092–2103
56. Zhou S, Zhuang Y, Meng R (2019) Multi-category skin lesion diagnosis using dermoscopy images and deep cnn ensembles. *linea*, ISIC Challenge
57. Zuiderveld K (1994) Contrast Limited Adaptive Histogram Equalization. Academic Press Professional Inc, Williston, pp 474–485

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Authors and Affiliations

P. Mirunalini¹ · Karthik Desingu¹ · S. Aswatha¹ · R. Deepika¹ · V. Deepika¹ · S. M. Jaisakthi² 

✉ S. M. Jaisakthi
jaisakthi.murugaiyan@vit.ac.in

P. Mirunalini
miruna@ssn.edu.in

Karthik Desingu
karthik19047@cse.ssn.edu.in

S. Aswatha
aswatha18026@cse.ssn.edu.in

R. Deepika
deepika18041@cse.ssn.edu.in

V. Deepika
deepika18042@cse.ssn.edu.in

¹ Department of Computer Science Engineering, Sri Sivasubramaniya Nadar College of Engineering, Chennai 603110, India

² School of Computer Science and Engineering, Vellore Institute of Technology, Chennai Campus, Chennai 600127, India