# TITLE PAGE

**AN ENSEMBLE OF CONVOLUTIONAL NEURAL NETWORKS FOR DERMOSCOPIC IMAGES CLASSIFICATION**

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by

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# RECOMMENDATION SHEET

This is to certify that I have supervised the preparation of and read the thesis proposal prepared by **Asaph F. Vega, Alexander Reyes, and Dexter B. Camila** entitled **AN ENSEMBLE OF CONVOLUTIONAL NEURAL NETWORKS FOR DERMOSCOPIC IMAGES CLASSIFICATION** and that the said paper has been submitted for final examination by the Oral Examination Committee.

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| --- |
| **Prof. Joel C. De Goma MSc.** |
| Thesis Adviser |

Note:

This page will be replaced with the Acceptance Form, Revisions List, and Conformity of Revisions once Accepted.

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# Chapter 1

**IN.TRO.DUCTION**

The American Cancer Society states that Melanoma is widely considered the deadliest of skin cancers and the incidence rates continues to rise worldwide. While Merkel cell carcinoma has a higher chance to lead to a fatality, melanoma is the cause of mo.re dea.ths than a.ny other ty.pe of ski.n ca.n.ce.r overall [1]. This cancer is po.ten.tia.lly de.te.cta.ble using dermoscopy at a ve.ry ea.rly stage while it is still mana.geable to cure. Dermoscopy is defined as the method of capturing an illuminated, magnified image of skin lesions [2]. The advancement of machine learning and image analysis have presented the ability to identify distinct malignant melanoma from benign incidents allowing the possibility for early detection.

## Context of the Study

Mo.st recent methods in the field of skin lesion classification re.ly on ha.nd-crafted

Feat.ures, su.ch as A.B.C.D.E rule (the acronym stands for Asymmetry, Border, Color, Dermoscopic structure a.nd Evolving) [43], 3-point checklist [44], 7-.poi.nt chec.klist [45], Men.zies met.hod [46], and C.A.S.H (Col.or, Arch.itec.ture, Sym.met.ry, and Homo.geneity) [47]. Physicians often rely on person.al experience and evaluate ea.ch patient’s lesions on a case-by-case basis by considering the patient’s local lesion patterns [48]. The accuracy of diagnosis for melanoma detection without computer-based assistance is reported to be between 65% and 80% [49].

According to the study of Mishra and Celebi, automating the dermoscopy method allows for faster and more frequent analy.sis of sk.in lesi.ons increasing the possibility of early detection [3]. While the efficiency of dermoscopy methods when evaluated by a dermatologist or licensed operator is relatively free of discrepancy with regards to skin tone, current deep learning image analysis techniques do not share the same results [4][5]. This can be addressed by improving the skin lesion segmentation process. However, there are several reasons skin lesion segmentation is a challenging task. Distinguishing between normal skin and lesional skin when contrast is poor is the primary reason. Mishra and Celebi proposed pre- and post-processing steps to maximize skin lesion segmentation accuracy not yet implemented by the related studies described in this paper.

Bisla et al. addressed the problem of images with less than optimal conditions and contaminants that affect lesion segmentation accuracy [51]. To resolve this data purification problem, Bisla et al. utilized traditional data processing methods to find and remove hair and ruler on the images. They extended the hair-removal algorithm [6] by overlaying the processed image with the segmented lesion obtained from their segmentation network. They utilized a U-Net architecture [7] that had been successfully applied to the problem of medical image segmentation and won the ISBI Cell Tracking Challenge [8] in 2015. Applying the techniques described here could potentially increase lesion segmentation accuracy and will be further discussed in chapter 3.

Codella et al. proposed the Deep Residual Network (DRN). They report state-of-the-art performance results using ConvNets to extract image descriptors by using a pre-trained model from the ImageNet Large Scale Visual Recognition Challenge (ILSVRC) 2012 [48]. They also investigate a more recent network structure called Deep Residual Network (DRN) [8].

## The Opportunity

The original authors of the U-Net architecture conducted experiments showing effective results and ranked first when applied to its original purpose single cell segmentation. Since its inception, the U-Net architecture has been successfully applied to various applications in the biomedical field. By incorporating the pre- and post-processing lesion segmentation methods proposed by Mishra and Celebi [3], the implementation of the U-Net architecture to skin.. lesion segment..ation by Code.lla et al. [7], and the techniques known as Transfer learning put forth by the studies of Kawahara et al. [6], an opportunity for a comparative evaluation of classification without delegated segmentation (entire image as input) with the two approaches of segmentation followed by classification in order to assess which approach achieves better classification results presents itself.

## Research Objectives

The emergence of the deep learning paradigm and the advance in computational power have enabled the development of the automated medical image analysis that present remarkable performance and promising results. The primary objective of this study is to apply the most cutting-edge, state-of-the-art machine learning models to skin lesion segmentation and classification through transfer learning from existing models. This can be accomplished by fulfilling the following objectives:

* **Analyze** the aforementioned algorithms for prediction of skin lesion segmentations from dermoscopic images in the form of binary masks;
* **Modify** current methods for skin lesion disease detection by delegating segmentation and classification into separate CNN architectures;
* **Compare** the skin lesion segmentation and classification delegation effects on the accuracy of the overall architecture with results of classification alone.

## Research Questions

The following research questions were derived from the aforementioned objectives of this study:

* What affects would applying the Transfer Learning paradigm to the skin lesion segmentation model have on the training time of the skin lesion classification model?
* What impact will delegating the segmentation task to a separate, dedicated CNNs have on the performance of the overall architecture?
* What affects do segmented images of skin lesions have compared to the current techniques in which classification is conducted on the entire image?

## Significance of the Study

The accuracy of diagnosis for melanoma detection without computer-based assistance

is reported to be between 65% and 80% [49]. The focus of the study is on the problem of skin lesion classification and present a machine learning, deep neural network based solution for the problem of dermoscopic image classification containing a skin lesion as malignant or benign. The proposed solution is built around the VGG16 and U-Net convolutional neural network architectures and applies the transfer learning paradigm from top performing experiments.

With this study, we can improve the methodology for classifying skin lesions by implementing the state-of-the-art techniques of various studies discussed in this paper.

## Scope and Limitations (or Delimitations)

The overall objective of this study is to improve existing methods of skin lesion classification by applying Transfer Learning of top performing architectures and related state-of-the-art techniques discussed in this paper. The inclusion, but not modification, of skin lesion segmentation and classification methods discussed in this paper will be implemented. Furthermore, the attempt to improve skin lesion detection accuracy will be primarily focused whether the process of segmentation yields any comparative effects.

# Chapter 2

**REVIEW OF RELATED LITERATURE**

## Introduction

Dermatologists usually rely on using skin lesion-specific image cues to differentiate harmful lesions from benign lesions and to support their diagnosis. Dermoscopy (commonly known terms are epiluminescence micro.scopy or dermatoscopy) is a procedure that magnifies the inspected regions and removes the surface reflection of the region as well to get a much clearer image of the skin that improves the diagnostic accuracy [2]. The improvement of machine learning and image analysis have bestowed the power of detecting cancerous skin lesions from benign lesions permitting the likelihood of early detection.

## Skin Lesion Segmentation

As stated in the study of Mishra and Celebi, automating the dermoscopy technique by applying image processing and machine learning algorithms permits for faster and more of frequent analysis of skin lesions increasing the chance of early detection [4]. While the efficiency of dermoscopy methods when evaluated by a dermatologist or licensed operator is free of discrepancy with regards to skin tone, current deep learning image analysis techniques do not share the same results [4]. An operator has the ability to adjust the lighting, focus, and other parameters to produce the best possible image before analyzing whereas neural networks must be trained with predetermined images and thus cannot adjust any parameters when processing different skin types. Mishra and Celebi proposed pre- and post-processing steps for skin lesion segmentation, feature segmentation, and classification to maximize classification accuracy not yet implemented by the related studies described in this paper. The proposed model starts with a combination of pre-processing steps for lesion segmentation that plays an important role segmenting skin lesion. The proposed pre-processing steps includes elimination of variable lighting effects [7] [8] [9] [10], changing the data into a precise color region. [11], choosing correct channel of color. [12], enhancement of the chosen color channel. [13], increasing contrast. [14] [15], data purification (removing hair), normalization of color variation, image smoothing. [16] [17] [18] [19] [20], removing the vignette effect. [21], and localizing the lesion. [22]. The model has feature segmentation which is different from lesion segmentation. This method segments the feature of the lesions according to its pattern. In every feature segmentation can have multiple segments around the lesion region. Feature segmentation also includes pre-processing steps which are feature dependent, segmentation and post-processing steps. This feature segmentation procedure uses a completely distinctive approach that is obtain to manage artifacts that oftentimes obstructs the region of interest. The model used feature generation and classification to dictate if the lesion is malignant or harmless. The attributes used in feature segmentation was used to generate clinical feature. The attributes were used in the classifier to be able to differentiate melanomas from harmless lesions. It was concluded that automating dermoscopy by implementing image processing techniques and machine learning algorithms makes it faster and have more frequent analysis of skin lesions increasing the chance of its early detection and treatment [4].

Codella et al. proposed a system that employs hand-crafted feature extraction techniques including local binary patterns (LBP) [29]. These features have been used in previous research that have achieved best performance in dermoscopic images and other medical image datasets [32]. Image descriptors were extracted using a deep learning framework by using pre-trained models from Image Large Scale Visual Recognition Challenge (ILSVRC). Much more recent network structure to win the ImageNet 2015 which eases the optimization and convergence of extremely deep networks was also investigated [33]. They have discovered that object's important characteristics contribute complimentary data to the classifier which improved performance [32]. This new state of the art performance showed improvements in the area under receiver by a factor of 2.9 [29]. They also investigate a more recent network structure called Deep Residual Network (DRN) [29].

Kawahara et al. delve into the concept of engaging a pre-trained ConvNet as a feature extractor to differentiate amongst ten categories of non-dermoscopic skin images [42]. The paper showed how filt.ers from a C.N.N pre-trained on natural images generalize to classifying 10 different class.es of non-dermoscopic skin images, perform.ing better than prior researches’ results. They hypothes.ized that subtra.cting the mean RGB pixel values computed over every individual image (per-image-mean) can improve the discriminant values within the ensuing feature vector. The per-image-mean gave improvements to classification’s accuracy [42].

## Skin Lesion Classification

Li et al. illustrated the effectiveness of deep convolutional net.works and lower level image feature descrip..tors for skin lesion analysis even with limited and unbalanced training data. An image classifier and an interpretation method were used in this study. Convolutional Neural Networks was used as the image classifier since it has hierarchical feature learning ability that is broadly used in image classification and recognition [26]. The Convolutional Neural Networks based techniques performed better than traditional techniques particularly in the recent ImageNet challenges [27]. Two CCN image classification architecture was modified and fine-tuned: ResNet50 and VGG for encoding image features. LightGBM that boosts tree-based algorithm was also used to combine different CNN model features. The interpretation method used in the study was based on Zingraf et al.’s study [28] which visualizes how the CNNs respond to a specific corrupted input image in order to justify a specific classification created by the network. The method corrupts the pixels in a sliding window that scans the focused region, consequently, analyzes the distinction of prediction outcome. The results showed the effectiveness of deep convolutional networks and low-level image feature descriptors. As this study was conducted for the Open Challenge of Skin Lesion Analysis Towards Melanoma Detection (ISIC 2018), the dataset was limited and contained samples of predominantly lighter skin tones.

Haofu Liao proposed to create a universal skin disease classification using deep convolutional neural network. Although, the human engineered feature extraction is not advisable when pertaining to the universal skin disease classification system as well as the hand-crafted features in which it is purposed for a limited number of skin disease. To make the study feasible he used feature learning so that the machine will be trained to decide on which feature is best to be use [34]. Convolutional neural network became well-known in object classification and feature learning. High performance GPU allows the training of large-scale datasets, according to various researchers from Image..Net Lar.ge Scale Visual Recognition Challenge (ILSVRC) portrays that advanced Convolutional Neural Network Architectures are able to exceed humans in object classification [35]. It has been shown that in several cases transfer learning may be accustomed efficiently train a deep CNN [37, 38]. In transfer learning, rather than coaching the network from indiscriminately initialized parameters, individuals takes a pre-trained network and fine-tunes its weights by continued the backpropagation. In their approach, they did transfer learning by fine-tuning ImageNet [35] pre-trained models with Caffe [36], a deep learning framework that supports communicative and efficient deep CNN training. As they concluded that building a universal skin disease classification system using deep CNN is feasible by tackling the problem by fine-tuning ImageNet pre-trained models.

The aforementioned studies produced promising results when classifying skin lesions. The work of Kawahara et al. explores the idea of using a pretrained CNN as a feature extractor to distinguish among 10 classes of non-dermoscopic skin images. Liao describes an attempt to construct a universal skin disease classification by applying transfer learning on a deep CNN and fine-tuned its weights. Codella et al. report state-of-the-art performance using CNNs to extract image descriptors by using a pre-trained model from the ImageNet Large Scale Visual Recognition Challenge (ILSVRC) in 2012. They also investigate a more recent structure called Deep Residual Network (DRN).

## Survey for Feature Segmentation

Table 2.1

|  |  |  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- | --- | --- |
| STUDY | DATASET | DISEASES | METHODS | ACCURACY | | SENSITIVITY | SPECIFICITY |
| Li et al. [26] | 10,015 images from ISIC 2018 | 327 images of Actine Keratosis, 514 images of basal cell carcinoma, 115 images of dermatofibroma, 1,113 images of melanoma, 6,705 images of nevus, 1,099 images of pigmented benign keratosis, 142 images of vascular lesion | VGG + ResNet50 | 0.85 | | N/A | N/A |
| Haofu Liao [34] | 23,000 images from Dermnet | Acne, Rosacea, Malignant Lesions, Atopic Dermatitis, Bullous Disease, Bacterial Infections, Eczema, Exanthems, Drug Eruptions, Hair Diseases, STDs, Pigmentation Disorders, Connective Tissue Diseases, Melanoma, Nevi & Moles, Nail Diseases, Contact Dermatitis, Psoriasis & Lichen Planus, Infestations & Bites, Benign Tumors, Systemic Disease, Fungal Infections, Urticaria, Vascular Tumors, Vasculitis, and Viral Infections | VGG16 | Top-1 | Top-5 | N/A | N/A |
| 72.7% | 91.0% |
| VGG19 | 73.1% | 90.9% | N/A | N/A |
| GoogleNet | 71.8% | 90.7% | N/A | N/A |
| 1,300 images from OLE | Rosacea, Actinic Keratosissis, Basal Cell Carcinoma, Squamous Cell Carcinoma, Atopic Dermatitis, Verruca, Nummular Eczema, Lupus Erythematosus, Melanoma, Melanocytic Nevus, Contact Dermatitis, Lichen Planus, Pityriasis Rosea, Psoriasis, Seborrheic Keratosis, Tinea Corposis, Tinea Versicolor, Urticaria, and Herpes | VGG19 | 24.8% | 61.7% | N/A | N/A |
| VGG19 Improved | 31.1% | 69.5% | N/A | N/A |
| Barata et al. [39] | 176 images from Hospital Pedro Hispano – Matosinhos | Not Specified | Global Features | Not Specified | | 96% | 80% |
| Local Features | 100% | 75% |

## Survey for Image Classification

Table 2.2

|  |  |  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- | --- | --- |
| STUDY | DATASET | DISEASES | METHODS | ACCURACY | | SENSITIVITY | SPECIFICITY |
| Li et al. [26] | 10,015 images from ISIC 2018 | 327 images of Actine Keratosis, 514 images of basal cell carcinoma, 115 images of dermatofibroma, 1,113 images of melanoma, 6,705 images of nevus, 1,099 images of pigmented benign keratosis, 142 images of vascular lesion | VGG + ResNet50 | 0.85 | | N/A | N/A |
| Haofu Liao [34] | 23,000 images from Dermnet | Acne, Rosacea, Malignant Lesions, Atopic Dermatitis, Bullous Disease, Bacterial Infections, Eczema, Exanthems, Drug Eruptions, Hair Diseases, STDs, Pigmentation Disorders, Connective Tissue Diseases, Melanoma, Nevi & Moles, Nail Diseases, Contact Dermatitis, Psoriasis & Lichen Planus, Infestations & Bites, Benign Tumors, Systemic Disease, Fungal Infections, Urticaria, Vascular Tumors, Vasculitis, and Viral Infections | VGG16 | Top-1 | Top-5 | N/A | N/A |
| 72.7% | 91.0% |
| VGG19 | 73.1% | 90.9% | N/A | N/A |
| GoogleNet | 71.8% | 90.7% | N/A | N/A |
| 1,300 images from OLE | Rosacea, Actinic Keratosissis, Basal Cell Carcinoma, Squamous Cell Carcinoma, Atopic Dermatitis, Verruca, Nummular Eczema, Lupus Erythematosus, Melanoma, Melanocytic Nevus, Contact Dermatitis, Lichen Planus, Pityriasis Rosea, Psoriasis, Seborrheic Keratosis, Tinea Corposis, Tinea Versicolor, Urticaria, and Herpes | VGG19 | 24.8% | 61.7% | N/A | N/A |
| VGG19 Improved | 31.1% | 69.5% | N/A | N/A |
| Barata et al. [39] | 176 images from Hospital Pedro Hispano – Matosinhos | Not Specified | Global Features | Not Specified | | 96% | 80% |
| Local Features | 100% | 75% |

## Summary

The techniques discussed here have potential to improve both convolutional neural networks for skin lesion segmentation and classification. The pre-processing methods proposed by Mishra and Celebi include removing variable lighting effects, normalizing color variation, and deleting contaminants affecting the subject (i.e. data purification). These processes are to be applied to the data before entering the lesion segmentation stage. Post-processing steps were then applied to the output of the lesion segmentation stage before applying a binary mask of the lesion border. The process was then reiterated replacing the lesion segmentation stage with Clinical Feature segmentation in the next iteration and finishing with binary masks of clinical feature segments. Clinical features are specific structures in the lesion which separates them from benign lesions. A final iteration was then conducted involving feature generation, feature selection, and finally, model optimization before the final prediction.

The aforementioned studies produced promising results when classifying skin lesions. The work of Kawahara et al. explores the idea of using a pretrained CNN as a feature extractor to distinguish among 10 classes of non-dermoscopic skin images. Liao describes an attempt to construct a universal skin disease classification by applying transfer learning on a deep CNN and fine-tuned its weights. Codella et al. report state-of-the-art performance using CNNs to extract image descriptors by using a pre-trained model from the ImageNet Large Scale Visual Recognition Challenge (ILSVRC) in 2012. They also investigate a more recent structure called Deep Residual Network (DRN).

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# Chapter 3

**THEORETICAL FRAMEWORK**

There are two main components of this study that will discussed here. During the past decades many approaches have been proposed to automatically generate image representations that can provide support to tasks like image classification, object detection, recognition or semantic segmentation. Most of them have relied on hand-engineered low-level descriptors. But since the publication of AlexNet in 2012, state-of-the-art methods in computer vision mostly rely on learning representations using deep convolutional neural networks.

Most current methods in the field of melanoma classification still rely on hand-crafted features. Typically, after the feature extraction based on these descriptions, machine learning methods such as k-nearest neighbors (kNN), Artificial Neural Networks (ANNs), logistic regression, decision trees and support vector machines (SVMs) have been explored to solve the classification task with moderate success [10].

Examples of related work using hand-crafted features and popular classifiers include:

• Codella et al. [11] use hand-crafted feature extraction techniques including color histogram, edge histogram, and a multi-scale variant of color local binary patterns (LBP).

• Barata et al. [12] explore two different methods for the detection of melanoma in dermoscopy images based on global and local features. They conclude that color features perform much better than texture features alone.

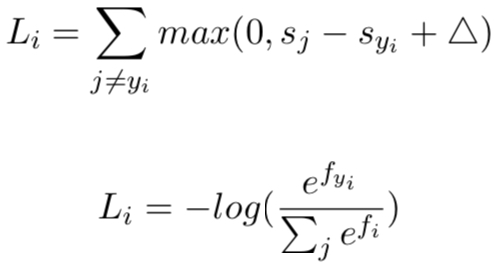
More recently, the emergence of a machine learning paradigm known as deep learning has enabled the development of medical image analysis systems that can display remarkable accuracy, to the point of raising concerns about the future of the human radiologist [13][14].

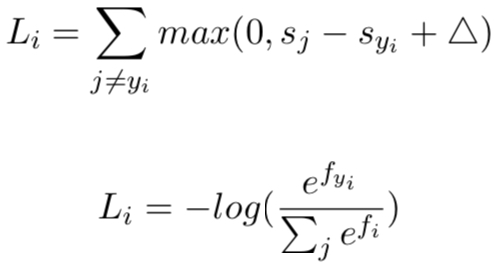
The next sections provide an overview of the deep learning techniques for image segmentation and image classification used in this study.

# Convolutional Neural Networks

Convolutional Neural Networks (also known as CNNs or ConvNets) are inspired in the behavior of biological systems through artificial neurons with learnable weights and biases. The layered architecture that Neural Networks performs based on matrix multiplications enables its application for image classification tasks. For this reason, ConvNets architectures assume that the input are images that have to be transformed into an output holding the class score predicted. The loss function is used to measure how well the predicted scores agrees with the ground truth labels in the input data. Most common loss functions are the Multiclass Support Vector Machine (SVM) (Equation 2.2) and the Softmax (Equation 2.3).

Equation 2.1





Equation 2.2

# Image Segmentation

Typically, segmentation is used as a preprocessing method in the classification process to remove potentially non-relevant information from the classification process [21]. ConvNets are typically used on image classification tasks: in the case of supervised learning, the input are images sorted by class, and the output to a certain image is a single class label. However, in the biomedical image processing field, a localization is often also required in addition to a global scale label, i.e. the network assigns a class label to each pixel. This is the main idea of a semantic segmentation model [15] using ConvNets.

Novel proposals for semantic segmentation [16] using convolutional neural networks introduce the idea of deconvolutional networks on the top of common ConvNets. This backward strided convolution is convenient to generate a segmentation map of the input through a sequence of deconvolution operations. These layers (shown in Figure 2.1) compute a dense pixel-wise class probability map by consecutive operations of unpooling, deconvolution, and rectification.

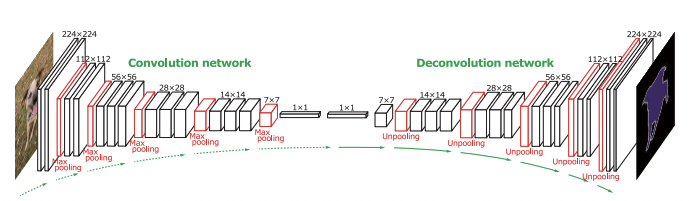


Figure 3.2 Semantic segmentation networks [3] using deep deconvolutional layers

1. **Transfer Learning**

Training an entire convolutional neural network in the medical imaging field is not always possible, due to the fact that datasets are often not large enough. Alternatively, random initialization of weights is replaced by a pretrained network on large datasets, i.e. ImageNet [18], that contains 1.2 million images labeled with 1,000 classes. This technique is known as Transfer Learning [19] and it is very common in machine learning scenarios. The main objective is the improvement of learning in a new task by transferring knowledge from a related task that has already been learned.

In practice, Transfer Learning from a pretrained ConvNet is typically used in these two different ways [20]:

• **Fixed Feature Extraction**. Use the output of one intermediate layer to train a machine learning classifier.

• **Fine-tuning**. Replace the last fully connected layer of the network with a new classification task and use the training data from the new domain to update the weights of some layers. In general, the earlier layers of the ConvNets are not updated and only the deeper ones are updated to the new task.

# Chapter 4

**METHODOLOGY**

This study in the Deep Learning field about the impact and effects of removing skin image background by applying image segmentation methods for a subsequent classification of melanoma. The goal of this study is to better understand the effects on performance results when segmenting a dermoscopic image by isolating and comparing results from both unaltered, bit-wise segmented, and U-Net segmented skin lesion images. Specifically, it hopes to better understand whether the values outside the lesion are detrimental to lesion classification or are instead beneficial to lesion classification by providing contextual information relevant to each lesion.

In order to achieve the project goal, two successful and well-known Convolutional Neural Networks architectures in the image semantic segmentation and image classification tasks have been adopted. The order of execution will be:

1. **Skin lesion segmentation**. The first task will perform an automatic prediction of lesion segmentation from dermoscopic images taking the form of binary masks using the U-Net architecture.

2. **Skin lesion classification**. The second task will perform the automatic classification as either melanoma or non-melanoma. In order to find the best classification model, this task will be divided into three subtasks according to different type of input skin images:

(a) **Unaltered lesion classification**. The basic model will perform the classification over the original skin RGB images contained in the ISIC dataset.

(b) **Manually segmented lesion classification**. The manually segmented images will be generated by performing a bit-wise operation on the original images and its corresponding original binary mask contained in the ISIC dataset.

(c) **U-Net Segmented lesion classification**. The third, and most complex model will perform the classification over the automatically segmented images from the U-Net segmentation model.

## Data Gathering

Datasets from a cross-sectional study of the clinical, dermoscopic and histopathological features of pigmented skin lesions on patients seen at the Out-Patient Departments of Quirino Memorial Medical Center and Ospital ng Makati will be accessed to conduct this research. Additional. data from the Research Institute for Tropical Medicine of the Philippines will also be requested for supplementation. However, in order to gather the aforementioned datasets, a letter requesting permission to gather and conduct the Mapùa University affiliated research will be employed.

Once permission and access has been granted, retrieval of the image datasets will be conducted from the requested sources*.* The size of the datasets is expected to be large and will require sufficient storage space and time to complete the transfer to more accessible mediums. The data will then be stored in a remote server to provide cloud services for processing access to the data.

## Data Pre-processing

This project takes advantage of ConvNets properties regarding input preprocessing: few previous processing techniques are needed. Although some basic preprocessing forms are performed:

• **Mean subtraction**. In order to center the cloud of RGB values from input data around zero along every dimension of the image, a mean subtraction is applied across the image features.

• **Image normalization**. By dividing each RGB dimension of input images by its standard deviation, a normalization is obtained from its original 0- and 255-pixel values to 1 and 0 normalized values. This preprocessing technique will avoid further issues caused by poor contrast images.

• **Image cropping and resizing**. Input images are preprocessed to be accepted by the architecture though cropping the image to the same aspect ratio as needed and resizing the original image to 64x80 pixels for the U-Net and 224x224 pixels for the VGG-16.

## Skin Lesion Segmentation

The overview of the feature extraction task follows the main structure described in section 3.1. Training and Validation data from the Segmentation dataset is processed with a Python script in order to load the images and masks and convert them into NumPy binary format files (.npy) that will allow a faster loading during the learning and prediction algorithm. Data is previously organized into train, train masks and test folders.

The proposed model is the U-Net network trained from scratch, which means that weights are randomly initialized and optimized with backpropagation. The network output is an array containing the 379 test binary masks, which can be converted into JPG images of 64 to 80 pixels dimension. This means a posterior post-processing technique will be needed to enlarge the images.

# Applying Transfer Learning

The same training methodology is proposed to face the two tasks of this project. The only difference between the segmentation and classification tasks are the models used. As previously commented in Section 3.3, the U-Net architecture is the architecture proposed for the segmentation task, while the VGG-16 is associated with a classification model. Following the diagram of Figure 4.1, the training data is trained through the learning algorithm defined by each model, which applies the stochastic gradient descent (SGD).

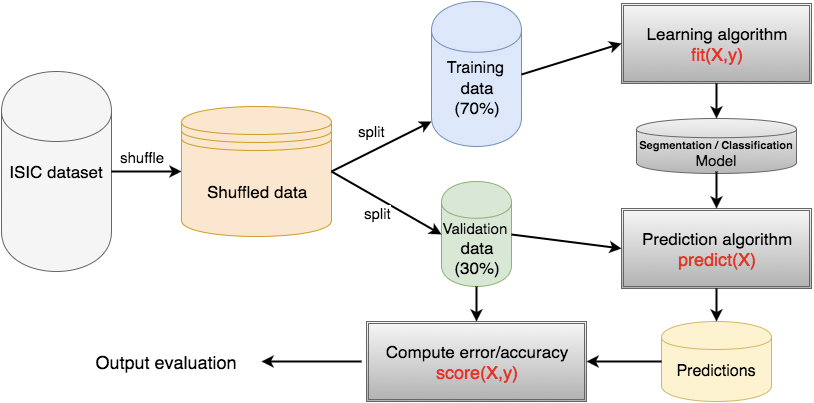


Figure 4.1 Training methodology

During training, some parameters must be considered to be altered in order to get the best performance of the network proposed regarding the problem to be solved. Typical ConvNets parameters are the following:

• **Batch size**. The batch size is attributed to the number of training images in one forward or backward pass. It is important to highlight that the higher the batch size, the more memory will be needed.

• **Iterations**. The number of iterations is the number forward or backward of passes: each pass using a batch size number of images.

• **Epoch**. The number of epochs measures how many times every image has been seen during training (i.e. one epoch means that every image has been seen once). It can be also understood as a one forward pass and one backward pass of all the training examples. It is numerically computed as:



Equation 4.1

• **Loss function**. Loss function (also called cost function) evaluates the penalty between the prediction and the ground truth label in every batch.

• **Learning rate**. The learning rate parameter defines the step size for which the weights of a model are updated regarding the stochastic gradient descent.

• **Decay**. The weight decay is an additional weight update parameter that induces the weights to exponentially decay to zero once the update process is over.

• **Optimizer**. Keras framework provides optimizers [58] in order to find the most optimal set of hyperparameters for the model.

# Classification

The main issue of the classification task is to avoiding overfitting caused by the small number of images of skin lesion in most dermatology datasets. In order to solve this problem, the objective of the proposed model is to firstly extract features from images with the original reduced network VGG-16 (with 138m parameters) over the VGG-19 (with 144m parameters) and secondly load those extracted representations on a fine-tuned VGG-16 architecture.

# Validation

In order to validate the model, the dataset will be split between training data and validation data (70% and 30% respectively). Once the model has learned the weights, a prediction algorithm classifies the validation data according to the training. A final model evaluation is performed by comparing the predictions with the ground truth data.

# APPENDICES

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