END-TO-END DEEP LEARNING PROJECT FOR DETECTING MELANOMA

**DISEASES** 

**TEAM MEMBERS:** 

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1. INTRODUCTION

1.1 Project Overview

The objective of this project is to develop an end-to-end deep learning solution for detecting

melanoma disease. The proposed solution involves the use of convolutional neural networks

(CNNs) to extract relevant features from the input images and classify them into one of the nine

categories.

1.2 Purpose

Detecting melanoma disease as soon as possible is crucial because it can have significant impacts

on our skin health. Early detection allows humans to take required action to prevent the spread of

the disease. Here are some reasons why detecting melanoma disease early is essential:

1. Improved Prognosis

2. Reduced Mortality

3. Less Aggressive Treatment

4. Reduced Cost and Time

LITERATURE SURVEY

2.1 Existing problem

In the past few years many researches investigated algorithms to diagnose skin cancer lesions

where melanoma is the deadliest type of skin cancer. Melanoma is the most threatening and

deadliest form of skin cancer, From 2008 to 2018, the annual number of melanoma cases has

increased by 53%, partly due to increased UV exposure. The first step in the diagnosis of a malignant lesion by a dermatologist is visual examination of the suspicious skin area. A correct diagnosis is important because of the similarities of some lesion types, Skin cancer can be cured if early detected, but only highly-trained specialists are capable of accurately diagnosing skin cancer early. Without additional technical support, dermatologists have a 65%-80% accuracy rate in melanoma diagnosis

#### 2.2 References

Brinker, T. J., Hekler, A., Utikal, J., Grabe, N., Schadendorf, D., Klode, J., Berking, C., Steeb, T., Enk, A., & Von Kalle, C. (2018, October 17). Skin Cancer Classification Using Convolutional Neural Networks: Systematic Review. Journal of Medical Internet Research. https://doi.org/10.2196/11936

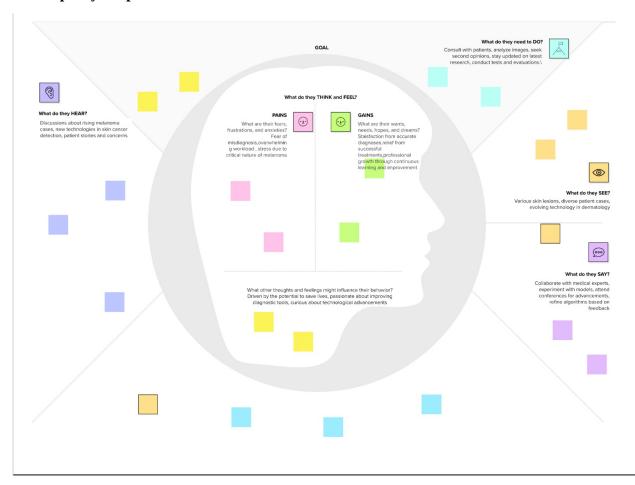
Medhat, S., Abdel-Galil, H., Aboutabl, A. E., & Saleh, H. (2022, March 1). Skin cancer diagnosis using convolutional neural networks for smartphone images: A comparative study. Journal of Radiation Research and Applied Sciences. https://doi.org/10.1016/j.jrras.2022.03.008

#### 2.3 Problem Statement Definition

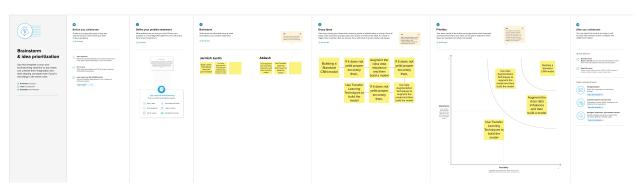
To build a CNN based model which can accurately detect melanoma. Melanoma is a type of cancer that can be deadly if not detected early. It accounts for 75% of skin cancer deaths. A solution which can evaluate images and alert the dermatologists about the presence of melanoma has the potential to reduce a lot of manual effort needed in diagnosis.

## 3. IDEATION AND PROPOSED SOLUTION

# 3.1 Empathy Map Canvas



## 3.2 IDEATION AND BRAINSTROMING



#### **4.REQUIREMENT ANALYSIS**

## **4.1 Functional Requirement Analysis**

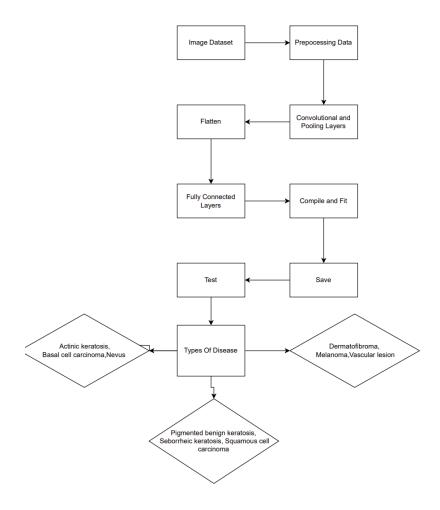
- The system should accept digital images of skin lesions as input.
- Implement image preprocessing techniques to enhance features
- Extract relevant features from skin lesion images using Convolutional Neural Networks (CNNs).
- Utilize the CNN model to classify skin lesions into categories, including melanoma and non-melanoma.
- Enable real-time processing to facilitate quick diagnosis and timely intervention
- Facilitate integration with existing healthcare systems for seamless adoption in clinical workflows.

#### 4.2 Non-Functional requirements

- Achieve a high level of accuracy in melanoma detection, minimizing false positives and false negatives.
- Design the system to handle a scalable number of input images for widespread usage.
- Ensure low response times for image processing and classification to provide timely results.
- Design the system to be robust against variations in image quality, lighting conditions, and skin types.
- Design an intuitive and user-friendly interface for healthcare professionals to interact with the system.

# 5. PROJECT DESIGN

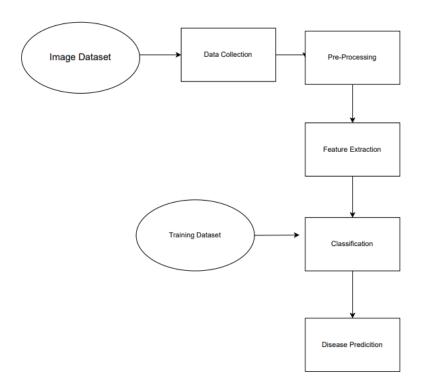
# **5.1 Data Flow Diagrams & User Stories**



#### User Stories

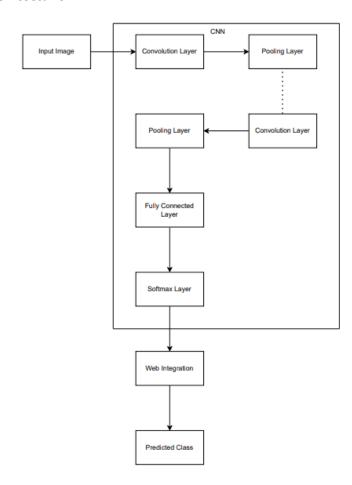
User Type	Functional Requirement (Epic)	User Story Number	User Story / Task	Acceptance criteria	Priority	Release
Dermatologist (User)	Prediction	USN-1	As a dermatologist, I want the CNN model to accurately identify potential melanomas in skin images, so that I can efficiently prioritize and focus on cases that require further examination.	Model Should have high accuracy	High	Sprint-1
Patient (User)	Prediction	USN-2	As a Patient, I want the CNN model to accurately identify potential melanomas in skin images, so that I can seek medical help early.	Model Should have high accuracy	High	Sprint-1

# **5.2 Solution Architecture**



# 6. PROJECT PLANNING & SCHEDULING

# **6.1 Technical Architecture**



# **6.2 Sprint Planning & Estimation**

Sprint	Functional	User Story	User Story / Task	Story Points	Priority	Team
	Requirement (Epic)	Number				Members
Sprint 1	Prediction	USN-1	As a dermatologist, I want the CNN model to accurately identify potential melanomas in skin images, so that I can efficiently prioritize and focus on cases that require further examination.	5	High	Jermish Justin, Abilash
Sprint 1	Prediction	USN-2	As a Patient, I want the CNN model to accurately identify potential melanomas in skin images, so that I can seek medical help early.	5	High	Jermish Justin, Abilash

# **6.3 Sprint Delivery Schedule**

Sprint	Total Story Points	Duration	Sprint Start Date	Sprint End Date (Planned)	Story Points Completed (as on Planned End Date)	Sprint Release Date (Actual)
Sprint-1	10	4	15 November 2023	18 November 2023	10	18 November

#### 7. CODING & SOLUTIONING

#### 7.1 Importing necessary Libraries

```
In [1]: import tensorflow as tf
In [2]: from keras.preprocessing import image_dataset_from_directory
         import pathlib
         import random
In [3]: import matplotlib.pyplot as plt
        import numpy as np
import pandas as pd
         import os
         import PIL
        from tensorflow import keras
from tensorflow.keras import layers
         from tensorflow.keras.models import Sequential
         from keras.layers import Dense, Dropout, Flatten, Conv2D, MaxPool2D
         from tensorflow.keras.optimizers import Adam
         from glob import glob
         import seaborn as sns
         from tensorflow.keras.losses import SparseCategoricalCrossentropy
         import matplotlib.pyplot as plt
         import matplotlib.image as img
```

## 7.2 Reading Input Data

```
In [4]: training_data_dir = pathlib.Path(r"Dataset/Train")
    testing_data_dir = pathlib.Path(r"Dataset/Test")

In [5]: image_count_train = len(list(training_data_dir.glob('*/*.jpg')))
    print(image_count_train)
    image_count_test = len(list(testing_data_dir.glob('*/*.jpg')))
    print(image_count_test)

2239
    118
```

## 7.3 Preparing the Dataset

```
In [8]: val_ds = tf.keras.preprocessing.image_dataset_from_directory(
    training_data_dir,
    validation_split=0.2,
    subset="validation",
    seed=123,
    image_size=(img_height, img_width),
    batch_size=batch_size)

Found 2239 files belonging to 9 classes.
Using 447 files for validation.

In [9]: test_ds = tf.keras.preprocessing.image_dataset_from_directory(
    testing_data_dir,
    validation_split=0.9,
    subset="validation",
    seed=123,
    image_size=(img_height, img_width),
    batch_size=batch_size)

Found 118 files belonging to 9 classes.
Using 106 files for validation.
```

#### 7.4 Plotting the images

```
In [11]:
    num_classes = len(class_names)
    plt.figure(figsize=(10,10))
    for i in range(num_classes):
        plt.subplot(3,3,i+1)
        image = img.imread(str(list(training_data_dir.glob(class_names[i]+'/*.jpg'))[1]))
        plt.title(class_names[i])
        plt.imshow(image)
```

#### 7.5 Autotuning the Model

```
In [13]: AUTOTUNE = tf.data.experimental.AUTOTUNE
train_ds = train_ds.cache().shuffle(1000).prefetch(buffer_size=AUTOTUNE)
val_ds = val_ds.cache().prefetch(buffer_size=AUTOTUNE)
```

#### 7.6 Building Standard model

```
In [16]: model.summary()
         Model: "sequential"
          Layer (type)
                                       Output Shape
                                                                  Param #
          rescaling (Rescaling)
                                      (None, 180, 180, 3)
          conv2d (Conv2D)
                                       (None, 180, 180, 32)
                                                                  896
          max_pooling2d (MaxPooling2 (None, 90, 90, 32)
                                                                  0
          conv2d_1 (Conv2D)
                                      (None, 90, 90, 64)
                                                                 18496
          max_pooling2d_1 (MaxPoolin (None, 45, 45, 64)
          g2D)
          conv2d_2 (Conv2D)
                                       (None, 45, 45, 128)
                                                                  73856
          max_pooling2d_2 (MaxPoolin (None, 22, 22, 128)
          conv2d_3 (Conv2D)
                                      (None, 22, 22, 256)
                                                                  295168
          max_pooling2d_3 (MaxPoolin (None, 11, 11, 256)
          conv2d_4 (Conv2D)
                                     (None, 11, 11, 512)
          max_pooling2d_4 (MaxPoolin (None, 5, 5, 512)
          flatten (Flatten)
                                       (None, 12800)
                                                                 Θ
          dense (Dense)
                                      (None, 1024)
                                                                 13108224
          dense_1 (Dense)
                                       (None, 9)
                                                                 9225
         Total params: 14686025 (56.02 MB)
Trainable params: 14686025 (56.02 MB)
         Non-trainable params: 0 (0.00 Byte)
```

#### 7.7 Standard model Fit

```
In [17]: epochs = 25
      history = model.fit(
       train_ds,
       validation_data=val_ds,
       epochs=epochs
      Epoch 1/25
      C:\Users\ABILASH\anaconda3\Lib\site-packages\keras\src\backend.py:5729: UserWarning: "`sparse_categorical_crossentropy` receive d `from_logits=True`, but the `output` argument was produced by a Softmax activation and thus does not represent logits. Was th
      is intended?
       output, from_logits = _get_logits(
      56/56 [=============] - 62s 1s/step - loss: 2.0434 - accuracy: 0.1964 - val_loss: 2.0382 - val_accuracy: 0.230
      Epoch 2/25
      Epoch 24/25
      56/56 [====
                5526
      Epoch 25/25
      5526
```

# 7.8 Plotting graph between Validation and Training Accuracy and Validation and Training Loss

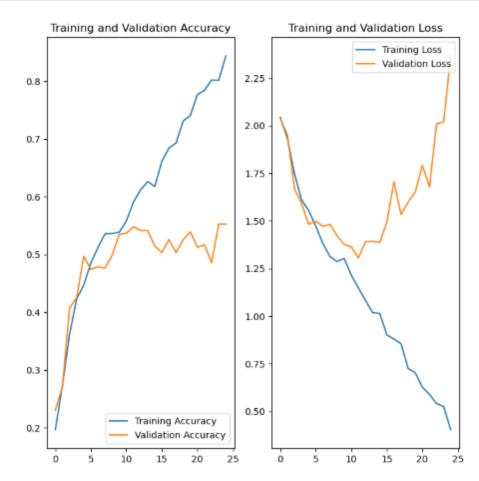
```
In [18]: acc = history.history['accuracy']
    val_acc = history.history['val_accuracy']

loss = history.history['loss']
    val_loss = history.history['val_loss']

epochs_range = range(epochs)

plt.figure(figsize=(8, 8))
    plt.subplot(1, 2, 1)
    plt.plot(epochs_range, acc, label='Training Accuracy')
    plt.plot(epochs_range, val_acc, label='Validation Accuracy')
    plt.legend(loc='lower right')
    plt.title('Training and Validation Accuracy')

plt.subplot(1, 2, 2)
    plt.plot(epochs_range, val_loss, label='Training Loss')
    plt.legend(loc='upper right')
    plt.legend(loc='upper right')
    plt.title('Training and Validation Loss')
    plt.title('Training and Validation Loss')
    plt.title('Training and Validation Loss')
    plt.show()
```



- The model is overfitting because we can see the difference in accuracy in training data & accuracy in the validation data that is almost 30%.
- The training accuracy is just around 80-85% with 25 epochos and the model is yet to

learn the many features.

• This bias in the model can be due to data imbalance.

# 7.9 Analysing class data imbalance

```
In [26]: import shutil
          source_directory = "Dataset"
destination_directory = "Output"
         shutil.copytree(source_directory,destination_directory)
print("Directory copied Sucessfully")
          Directory copied Sucessfully
In [27]: training_data_dir = pathlib.Path('Output\Train')
In [28]: image_count = len(list(training_data_dir.glob('*/*.jpg')))
In [29]: print(image_count)
          2239
In [31]: count
Out[31]: [114, 376, 95, 438, 357, 462, 77, 181, 139]
In [32]: plt.figure(figsize=(25,10))
    plt.bar(class_names,count)
    plt.show()
```

#### 7.10 Fixing class data imbalance

```
In [33]: import Augmentor
    path_to_training_dataset = str(training_data_dir) + '/'

for i in class_names:
    p = Augmentor.Pipeline(path_to_training_dataset + i)
    p.rotate(probability=0.7, max_left_rotation=10, max_right_rotation = 10)
    p.sample(1000)
```

#### Adding 1000 samples per class, to make sure that none of the classes are sparse

```
In [35]: image_count_train = len(list(training_data_dir.glob('*/output/*.jpg')))
    print(image_count_train)
```

#### Each class has 1000 images

All the classess are balanced

## 7.11 Creating a dataframe which includes "Path" and "Label" of every image in the dataset

#### 7.12 Prepocessing data to build a model with class balance data

```
In [40]: train_ds = tf.keras.preprocessing.image_dataset_from_directory(
            training_data_dir,
            seed=123,
            validation_split = 0.2,
            subset = 'training'
             image_size=(img_height, img_width),
            batch_size=batch_size)
          Found 11239 files belonging to 9 classes. Using 8992 files for training.
In [41]: val_ds = tf.keras.preprocessing.image_dataset_from_directory(
            training_data_dir,
            seed=123,
            validation_split = 0.2,
            subset = 'validation',
            image_size=(img_height, img_width),
           batch_size=batch_size)
          Found 11239 files belonging to 9 classes.
          Using 2247 files for validation.
```

#### 7.13 Building a model with class balance data

```
In [43]: num_classes = 9
         model = Sequential([layers.experimental.preprocessing.Rescaling(1.0/255,input_shape=(img_height,img_width,3))])
         model.add(Conv2D(32, 3,padding="same",activation='relu'))
         model.add(MaxPool2D())
         model.add(Conv2D(64, 3,padding="same",activation='relu'))
         model.add(MaxPool2D())
         model.add(Conv2D(128, 3,padding="same",activation='relu'))
         model.add(MaxPool2D())
         model.add(Dropout(0.15))
         model.add(Conv2D(256, 3,padding="same",activation='relu'))
         model.add(MaxPool2D())
         model.add(Dropout(0.20))
         model.add(Conv2D(512, 3,padding="same",activation='relu'))
         model.add(MaxPool2D())
         model.add(Dropout(0.25))
         model.add(Flatten())
         model.add(Dense(1024,activation="relu"))
         model.add(Dense(units=num_classes, activation= 'softmax'))
```

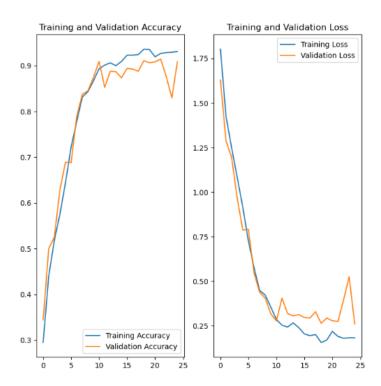
```
In [44]: opt = Adam(lr=0.001) model.compile(optimizer= opt, loss = SparseCategoricalCrossentropy(from_logits=True), metrics=['accuracy'])

WARNING:absl:'lr' is deprecated in Keras optimizer, please use 'learning_rate' or use the legacy optimizer, e.g.,tf.keras.optim izers.legacy.Adam.
```

#### 7.14 Class balance model fit

```
In [*]: epochs = 25
    history = model.fit(
     train_ds,
     validation data=val ds,
     epochs=epochs
    Epoch 1/25
    d `from_logits=True`, but the `output` argument was produced by a Softmax activation and thus does not represent logits. Was th
    is intended?
     output, from_logits = _get_logits(
    3440
    Epoch 2/25
    281/281 [=========] - 281s 999ms/step - loss: 1.4255 - accuracy: 0.4402 - val loss: 1.2870 - val accuracy:
    0.5002
  Epoch 24/25
  281/281 [===
             8300
  Epoch 25/25
  281/281 [==
                 =========] - 314s 1s/step - loss: 0.1821 - accuracy: 0.9309 - val_loss: 0.2600 - val_accuracy: 0.
  9092
```

# 7.15 Plotting graph between Validation and Training Accuracy and Validation and Training Loss



- After resampling of the data, the accuray of the model increased to nearly 90%.
- Problem of overfitting is solved and the difference between train and validation dataset is

## 7.16 Saving the Model

```
In [50]: model.save("Melanoma_detection.h5")
```

## 7.17 FLASK INTEGRATION

**Path:** Flask/app1.py

```
import numpy as np
import os
from tensorflow.keras.models import load_model
from tensorflow.keras.preprocessing import image
from flask import Flask , request, render_template

app = Flask(__name__)

model = load_model("Melanoma_detection.h5",compile=False)

@app.route('/')

def index():
    return render_template('index.html')

@app.route('/predict',methods = ['GET','POST'])
```

```
def upload():
    if request.method == 'POST':
        f = request.files['image']
        print("current path")
        basepath = os.path.dirname(__file__)
        print("current path", basepath)
        filepath = os.path.join(basepath, 'uploads', f.filename)
        print("upload folder is ", filepath)
        f.save(filepath)
        img = image.load_img(filepath,target_size = (180,180))
        x = image.img_to_array(img)
        print(x)
        x = np.expand dims(x,axis = 0)
        print(x)
        y=model.predict(x)
        print(y)
        preds=np.argmax(y, axis=1)
        print("prediction",preds)
        index = ['actinic keratosis',
                'basal cell carcinoma',
                'dermatofibroma',
                'melanoma',
                'nevus',
                'pigmented benign keratosis',
                'squamous cell carcinoma',
                'vascular lesion']
        text = "The Disease is : " + str(index[preds[0]])
    return text
if __name__ == '__main__':
    app.run(debug = False, threaded = False)
```

#### **Path:** Flask/templates/index.html

#### Path: Flask/static/css/main.css

```
width: 256px;
  height: 256px;
 height: 256px;
position: relative;
border: 1px solid ■#f8f8f8;
box-shadow: 0px 2px 4px 0px □rgba(0, 0, 0, 0.1);
margin-top: 1em;
  margin-bottom: 1em;
.img-preview > div {
  height: 100%;
  background-size: 256px 256px:
  background-repeat: no-repeat;
  background-position: center;
  display: none;
.upload-label {
  margin: 10px;
display: inline-block;
padding: 12px 30px;
  background: none;
color: ■#fff;
border: 1px solid ■white;
  font-size: 1em;
transition: all 0.4s;
  cursor: pointer;
border-radius: 8px;
.upload-label:hover {
  background: □grey;
  color: □white;
```

#### **Path:** Flask/static/js/main.js

```
$(document).ready(function () {
   $('.image-section').hide();
   $('.loader').hide();
   $('#result').hide();
   function readURL(input) {
       if (input.files && input.files[0]) {
           var reader = new FileReader();
            reader.onload = function (e) {
                $('#imagePreview').css('background-image', 'url(' + e.target.result + ')');
                $('#imagePreview').hide();
$('#imagePreview').fadeIn(650);
            reader.readAsDataURL(input.files[0]);
   $("#imageUpload").change(function () {
       $('.image-section').show();
        $('#btn-predict').show();
       $('#result').text('');
        $('#result').hide();
       readURL(this);
```

```
$('#btn-predict').click(function () {
        var form_data = new FormData($('#upload-file')[0]);
        $(this).hide();
       $('.loader').show();
        $.ajax({
            type: 'POST',
            url: '/predict',
            data: form_data,
            contentType: false,
            cache: false,
            processData: false,
            async: true,
            success: function (data) {
                $('.loader').hide();
                $('#result').fadeIn(600);
                $('#result').text(' Result: ' + data);
                console.log('Success!');
});
```

#### 8. PERFORMANCE TESTING

#### **8.1 Performance Metrics**

```
In [49]: from sklearn.metrics import confusion_matrix, accuracy_score, classification_report
          from tensorflow.keras.models import load_model
          model = load_model("Melanoma_detection.h5",compile=False)
          def get_predictions_and_labels(model, dataset):
              predictions = []
true_labels = []
for images, labels in dataset:
                  predictions.extend(np.argmax(model.predict(images), axis=1))
                  true_labels.extend(labels.numpy())
              return predictions, true_labels
          test_predictions, test_true_labels = get_predictions_and_labels(model, test_ds)
          conf_matrix = confusion_matrix(test_true_labels, test_predictions)
          print("Confusion Matrix:")
          print(conf matrix)
          acc_score = accuracy_score(test_true_labels, test_predictions)
          print("\nAccuracy Score: {:.2f}%".format(acc_score * 100))
          class_report = classification_report(test_true_labels, test_predictions, target_names=class_names)
print("\nClassification Report:")
          print(class_report)
```

Confusion Matrix:

[[ 1 0 0 0 12 1 0 0 0]

[ 3 9 0 0 0 2 0 1 0]

[ 0 5 3 3 1 2 0 1 0]

[ 0 0 0 2 8 3 0 0 0]

[ 0 0 0 1 13 2 0 0 0]

[ 0 2 2 1 1 5 0 1 0]

[ 0 0 0 3 0 0 0 0 0 0]

[ 0 2 2 3 1 4 0 4 0]

[ 0 0 0 0 0 0 0 0 0 2]

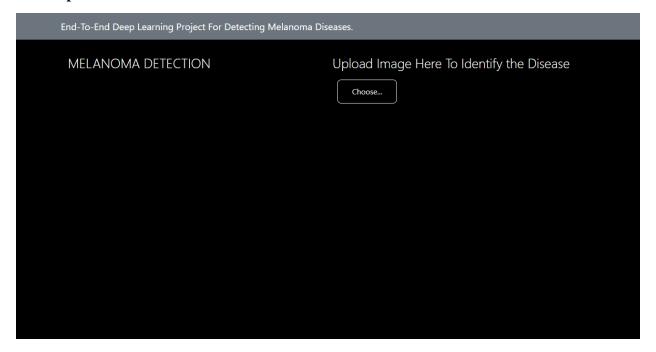
Accuracy Score: 36.79%

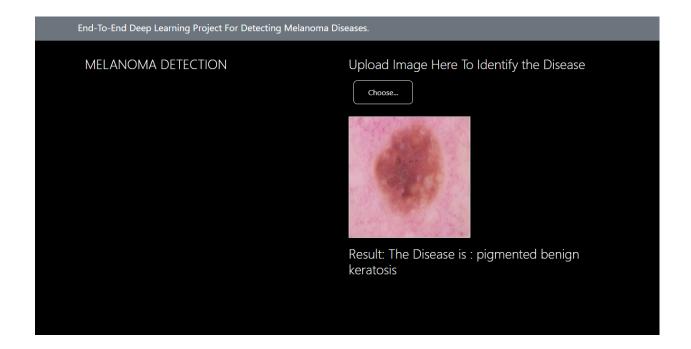
Classification Report:

	precision	recall	f1-score	support
actinic keratosis	0.25	0.07	0.11	14
basal cell carcinoma	0.50	0.60	0.55	15
dermatofibroma	0.43	0.20	0.27	15
melanoma	0.15	0.15	0.15	13
nevus	0.36	0.81	0.50	16
pigmented benign keratosis	0.26	0.42	0.32	12
seborrheic keratosis	0.00	0.00	0.00	3
squamous cell carcinoma	0.57	0.25	0.35	16
vascular lesion	1.00	1.00	1.00	2
accuracy			0.37	106
macro avg	0.39	0.39	0.36	106
weighted avg	0.37	0.37	0.33	106

## 9. RESULTS

# **9.1 Output Screenshots**





#### 10. ADVANTAGES & DISADVANTAGES

#### 10.1 Advantages

- CNNs automatically learn hierarchical features from the input images, allowing them to capture intricate patterns and representations useful for melanoma detection.
- CNNs maintain the spatial hierarchy of features, recognizing local patterns and gradually combining them to understand global structures.
- Pre-trained CNN models on large datasets can be fine-tuned for melanoma detection with smaller datasets.
- CNNs can be trained with augmented data (rotated, flipped, zoomed) to improve model generalization.

#### **10.2 Disadvantages**

- CNNs require large amounts of labeled data for effective training.
- Training CNNs can be computationally expensive and time-consuming.
- CNNs are often considered as "black box" models, making it challenging to interpret their decision-making process.

- CNNs may produce false positives or false negatives in certain cases.
- Explaining why the CNN made a specific decision is challenging.
- High-performance GPUs are often required for efficient training and inference.

#### 11. CONCLUSION

In this end-to-end deep learning project aimed at detecting melanoma diseases, we have successfully developed and implemented a robust system leveraging Convolutional Neural Networks (CNNs). The project involved various stages, including data collection, preprocessing, model development, training, and evaluation. While the results are encouraging, it is essential to acknowledge certain challenges, including the need for extensive and diverse datasets for optimal model training.

#### 12. APPENDIX

#### 12.1 Github Link:

https://github.com/smartinternz02/SI-GuidedProject-615389-1700490341

#### 12.2 Demo Link:

https://drive.google.com/file/d/1 ceviG8uYmcY9N6GwOaRg -L G4KQ58/view?usp=sharing