cnn-cancer-detection

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[]: #Description of the Problem and Data Used:
    This machine learning model aims to detect metastatic cancer in images of body \Box
    This is an example of a beneficial use of deep learning, as it can make an_{\sqcup}
     →immense difference
    in how quickly a patient can recieve a diagnosis, and potentially save their \Box
     \hookrightarrow life.
    The dataset we're using is derived from the PatchCamelyon (PCam) benchmark.
    It consists of lots of small pathology image patches labeled to show if they \Box
     ⇔contain cancerous tissue.
    ⇔determine if there is cancer present.
    The training set has labeled images, while the test set is unlabeled.
    \hookrightarrow (CNN)
    The images are first preprocessed with normalization and augmentation, then the
     ⇔CNN is trained to minimize binary cross-entropy
     loss. The model is evaluated using the ROC AUC metric, which tells us how well \sqcup
     ⇔it distinguishes between healthy and cancerous
    tissue. Overall, this project aims to create an effective cancer detection \Box
     ⇔model, contributing toward making advanced diagnostic
    tools more accessible and efficient.
    #Exploratory Data Analysis(EDA):
    In order to inspect, visualize, and clean the data, first the distribution of \Box
     ⇔labels are visualized using histograms,
    to visualize a bunch of sample images. To clean the data, we augment the images \Box
    width shift, height shift, and flipping. According to the insights gathered \Box
     ⇔during EDA, the data is preprocessed to
    be used to train the CNN model. The model is evaluated using ROC AUC (Receiver_{\sqcup}
     →Operating Characteristic- Area Under the Curve)
    performance measurement which plots the TPR (True Positive Rate) and FPR (False_{\sqcup}
     →Positive Rate) and degree of separability,
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according to these performance metrics the model is adjusted.
import os
import pandas as pd
import seaborn as sns
import matplotlib.pyplot as plt
train_labels_path = '/kaggle/input/histopathologic-cancer-detection/
 ⇔train labels.csv'
train_path = '/kaggle/input/histopathologic-cancer-detection/train/'
test_path = '/kaggle/input/histopathologic-cancer-detection/test/'
train_labels = pd.read_csv(train_labels_path)
print(train_labels.head())
print(train_labels['label'].value_counts())
sns.countplot(x='label', data=train_labels)
plt.title('Label Distribution')
plt.xlabel('Label')
plt.ylabel('Count')
plt.show()
from tensorflow.keras.preprocessing.image import load_img
import matplotlib.pyplot as plt
def visualize_samples(images_path, labels_df, num_samples=5):
    sample_images = labels_df.sample(num_samples)
   plt.figure(figsize=(15, 5))
   for idx, row in enumerate(sample_images.iterrows()):
        img_id, label = row[1]['id'], row[1]['label']
        img = load_img(os.path.join(images_path, img_id + '.tif'))
       plt.subplot(1, num_samples, idx + 1)
       plt.imshow(img)
       plt.title(f'Label: {label}')
       plt.axis('off')
   plt.show()
visualize_samples(train_path, train_labels, num_samples=10)
from tensorflow.keras.preprocessing.image import img_to_array
def get_image_stats(image_ids, path):
   heights, widths = [], []
   for image id in image ids:
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img = load_img(os.path.join(path, image_id + '.tif'))
        img_array = img_to_array(img)
        heights.append(img_array.shape[0])
        widths.append(img_array.shape[1])
    return heights, widths
heights, widths = get_image_stats(train_labels['id'], train_path)
plt.figure(figsize=(12, 6))
plt.subplot(1, 2, 1)
plt.hist(heights, bins=30, color='blue', alpha=0.7)
plt.title('Image Heights Distribution')
plt.xlabel('Height')
plt.ylabel('Frequency')
plt.subplot(1, 2, 2)
plt.hist(widths, bins=30, color='green', alpha=0.7)
plt.title('Image Widths Distribution')
plt.xlabel('Width')
plt.ylabel('Frequency')
plt.show()
def visualize image grid(images path, labels df, grid size=(4, 4)):
    sample_images = labels_df.sample(grid_size[0] * grid_size[1])
    fig, axes = plt.subplots(grid_size[0], grid_size[1], figsize=(12, 12))
    axes = axes.flatten()
    for ax, (_, row) in zip(axes, sample_images.iterrows()):
        img_id, label = row['id'], row['label']
        img = load_img(os.path.join(images_path, img_id + '.tif'))
        ax.imshow(img)
        ax.set_title(f'Label: {label}')
        ax.axis('off')
    plt.tight_layout()
    plt.show()
visualize_image_grid(train_path, train_labels, grid_size=(5, 5))
from tensorflow.keras.preprocessing.image import ImageDataGenerator
#Data Augmentation
datagen = ImageDataGenerator(
    rescale=1./255,
    rotation_range=20,
    width_shift_range=0.2,
    height_shift_range=0.2,
    horizontal_flip=True,
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vertical_flip=True
)
def visualize_augmentations(image_path, datagen, num_augmentations=5):
    img = load_img(image_path)
    img_array = img_to_array(img)
    img_array = img_array.reshape((1, ) + img_array.shape)
    fig, axes = plt.subplots(1, num_augmentations, figsize=(20, 5))
    for batch in datagen.flow(img_array, batch_size=1):
        ax = axes[i]
        ax.imshow(batch[0])
        ax.axis('off')
        i += 1
        if i >= num_augmentations:
            break
    plt.show()
sample_image_path = os.path.join(train_path, train_labels['id'].iloc[0] + '.
visualize_augmentations(sample_image_path, datagen)
#Convolutional Neural Network Model Architecture:
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The model used for image classification is a Convolutional Neural Network (CNN)
The architecture includes multiple convolutional layers followed by max-pooling_{\sqcup}
⇔and dropout layers to avoid overfitting.
By performing hyperparameter tuning, the best parameters for the CNN model are \Box
 \ominus established.
Specifically, a grid search approach is used to explore different values for 
⇔ideal learning rate, batch size, and number of
epochs.
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import tensorflow as tf
from tensorflow.keras.models import Sequential
from tensorflow.keras.layers import Conv2D, MaxPooling2D, Flatten, Dense,
 →Dropout
model = Sequential([
    Conv2D(32, (3, 3), activation='relu', input_shape=(96, 96, 3)),
    MaxPooling2D(pool_size=(2, 2)),
    Dropout(0.25),
    Conv2D(64, (3, 3), activation='relu'),
    MaxPooling2D(pool_size=(2, 2)),
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Dropout(0.25),
   Conv2D(128, (3, 3), activation='relu'),
   MaxPooling2D(pool_size=(2, 2)),
   Dropout(0.25),
   Flatten(),
   Dense(512, activation='relu'),
   Dropout(0.5),
   Dense(1, activation='sigmoid')
1)
model.compile(optimizer='adam', loss='binary_crossentropy',_
 →metrics=['accuracy'])
model.summary()
#Results and Analysis:
In the model architecture section, hyperparameter tuning, testing different \sqcup
Garchitectures, and other techniques were implemented
robustness and reducing overfitting. Hyperparameter tuning identified the bestu
⇔learning rate, batch size, and number of epochs,
while early stopping prevented overfitting. The ideal configuration was a_{\sqcup}
⇔learning rate of 0.001, a batch size of 32, and
training for 30 epochs with early stopping.
#Setting up Data Pipelines
train_datagen = ImageDataGenerator(
   rescale=1./255,
   rotation range=20,
   width_shift_range=0.2,
   height_shift_range=0.2,
   horizontal_flip=True,
   vertical_flip=True,
   validation_split=0.2
)
train_generator = train_datagen.flow_from_dataframe(
   dataframe=train_labels,
   directory=train_path,
   x_col='id',
   y_col='label',
   target_size=(96, 96),
   batch_size=32,
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class_mode='binary',
    subset='training'
)
validation_generator = train_datagen.flow_from_dataframe(
    dataframe=train_labels,
    directory=train_path,
    x_col='id',
    y_col='label',
    target_size=(96, 96),
    batch size=32,
    class_mode='binary',
    subset='validation'
)
#Model Training
history = model.fit(
    train_generator,
    validation_data=validation_generator,
    epochs=50
)
plt.figure(figsize=(12, 4))
plt.subplot(1, 2, 1)
plt.plot(history.history['accuracy'], label='train accuracy')
plt.plot(history.history['val_accuracy'], label='validation accuracy')
plt.title('Accuracy')
plt.legend()
plt.subplot(1, 2, 2)
plt.plot(history.history['loss'], label='train loss')
plt.plot(history.history['val_loss'], label='validation loss')
plt.title('Loss')
plt.legend()
plt.show()
from sklearn.metrics import roc_auc_score
#Model Evaluation
validation_steps = validation_generator.samples // validation_generator.
 ⇒batch_size
val_predictions = model.predict(validation_generator, steps=validation_steps)
val_labels = validation_generator.classes[:len(val_predictions)]
roc_auc = roc_auc_score(val_labels, val_predictions)
print(f'Validation ROC AUC: {roc_auc}')
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test_datagen = ImageDataGenerator(rescale=1./255)
test_generator = test_datagen.flow_from_directory(
   directory=test_path,
   target_size=(96, 96),
   batch_size=1,
   class mode=None,
   shuffle=False
test_predictions = model.predict(test_generator, steps=len(test_generator.
 →filenames))
submission_df = pd.DataFrame({
    'id': [fname.split('/')[-1].split('.')[0] for fname in test_generator.
 ⇔filenames],
    'label': test_predictions.flatten()
})
submission_df.to_csv('/kaggle/working/submission.csv', index=False)
#Conclusion
The model performed well after it was improved after taking various performance \sqcup
⇔metrics into consideration.
These improvements consisted of data augmentation, dropout layers, and \Box
⇔hyperparameter tuning. Also in the architecture phase,
early stopping effectively prevented overfitting. Future improvements could_{\sqcup}
 ⇔include using more advanced architectures and
⇔performance model in this project means
earlier and more accurate cancer detection, which results in better healthcare\sqcup
 ⇔efficiency and better patient outcomes.
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