Tumor Classification Using Ensemble of ShuffleNet_V2, MobileNet_V2, and ResNet18

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Abstract

Breast cancer is one of the leading causes of mortality among women worldwide. Early and accurate classification of tumors as benign or malignant from mammographic images significantly aids timely diagnosis and treatment. In this project, we present a lightweight yet powerful ensemble model combining three convolutional neural network (CNN) architectures: ShuffleNet_V2, MobileNet_V2, and ResNet18. These models were selected for their computational efficiency and complementary feature extraction capabilities. The ensemble utilizes classic bagging by averaging the outputs of each network after applying the softmax activation, providing a robust final prediction. Our pipeline also generates side-by-side views of the CC (craniocaudal) and MLO (mediolateral oblique) projections during validation to aid in interpretability. The model performance is evaluated using a confusion matrix and classification metrics: precision, recall, F1-score, and support.

1. Introduction

Breast cancer screening often involves analyzing mammogram images in two standard views: **CC** and **MLO**. Due to the subtle and varying visual patterns, automated classification is challenging and benefits from ensemble-based learning. We leverage deep learning and ensemble techniques to enhance the prediction accuracy of tumor malignancy classification.

2. Dataset

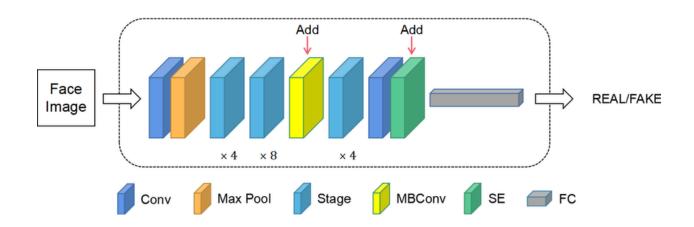
The dataset comprises mammographic images with labels indicating whether the tumor is **malignant** or **benign**. For each case, both **CC** and **MLO** views are available, providing complementary visual perspectives of breast tissue.

3. Model Architecture Overview

3.1 ShuffleNet_V2

ShuffleNet_V2 is a lightweight CNN architecture optimized for speed and memory efficiency. Key characteristics:

- Channel Shuffle: Enables better information flow across feature channels.
- **Grouped Convolutions**: Reduces computational cost while retaining performance.
- Lightweight Design: Suitable for deployment on resource-constrained devices.



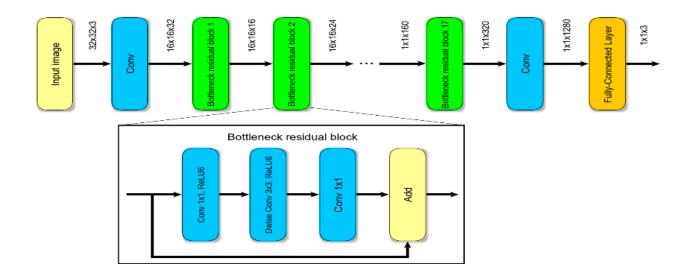
ShuffleNet_V2 architecture

Contribution to Ensemble: ShuffleNet_V2 extracts high-level features quickly and efficiently, offering robust performance with minimal overhead, which complements heavier models like ResNet18.

3.2 MobileNet_V2

MobileNet_V2 introduces the concept of inverted residuals and linear bottlenecks:

- **Depthwise Separable Convolutions**: Reduces parameters by factorizing standard convolutions.
- **Inverted Residual Blocks**: Helps in learning complex patterns while maintaining efficiency.
- Lightweight yet expressive.



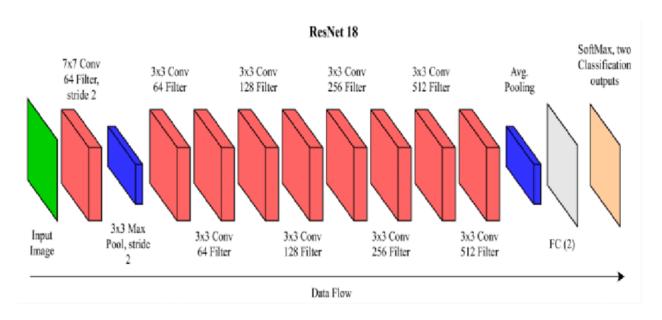
MobileNet V2 architecture

Contribution to Ensemble: MobileNet_V2 balances performance and efficiency, capturing mid-level abstractions. It acts as a strong middle-ground in the ensemble.

3.3 ResNet18

ResNet18 is part of the ResNet family known for its residual learning technique:

- **Skip Connections**: Allow gradient flow through deeper layers.
- **18 Convolutional Layers**: Offers deeper representation with fewer vanishing gradient issues.
- General-purpose baseline for image classification.



ResNet18 architecture

Contribution to Ensemble: ResNet18 provides deeper feature representations, complementing the shallower MobileNet_V2 and ShuffleNet_V2, especially beneficial in complex classification boundaries.

4. Ensemble Model Formation

Each model (ShuffleNet_V2, MobileNet_V2, ResNet18) is trained individually on the mammogram dataset. During inference, the ensemble prediction is generated as follows:

- 1. Each model outputs a probability distribution over the classes using **softmax**.
- 2. The outputs from all three models are averaged:

$$P_{ensemble} = rac{1}{3}(P_{shuffle} + P_{mobile} + P_{resnet})$$

3. The final prediction is the class with the highest averaged probability.

This ensemble approach falls under **classic bagging**, where independent learners vote (in this case, via softmax probability averaging), reducing variance and improving generalization.

5. Implementation

- 1. The code was run on a PyCharm Professional Edition IDE with a Python interpreter version of 3.12.8 and PyTorch version 2.6.0.
- 2. For qualitative analysis, the code also displays:
- Side-by-side views of the **Craniocaudal (CC)** and **Mediolateral Oblique (MLO)** images for selected validation samples.
- Each image pair is accompanied by the **predicted label and ground truth**, providing valuable visual feedback on model performance.
- 3. The code was run for 10 epochs when converging to an accuracy of 80%. The model tended to overfit to data when run for more epochs, likely since the dataset size is small.

Source Code:

```
# Imports
import os
import random
from PIL import Image
import torch
import torch.nn as nn
import torch.nn.functional as F
from torch.utils.data import Dataset, DataLoader, Subset
from torchvision import models, transforms
from sklearn.metrics import accuracy score
import matplotlib.pyplot as plt
from torchvision.models import shufflenet_v2_x0_5, ShuffleNet_V2_X0_5_Weights
from torchvision.models import mobilenet_v2, MobileNet V2 Weights
from torchvision.models import resnet18, ResNet18 Weights
# --- 1. Dataset Definition ---
class MammogramDataset(Dataset):
  def init (self, root dir, transform=None):
    self.root dir = root dir
    self.transform = transform
    self.samples = self. load samples()
```

```
def load samples(self):
     samples = []
     for class folder in os.listdir(self.root dir):
       class path = os.path.join(self.root dir, class folder)
       if not os.path.isdir(class_path):
          continue # Skip .zip or any non-directory
       label = 1 if 'MALIGNANT' in class folder.upper() else 0
       inner class path = os.path.join(class path, os.listdir(class path)[0]) # 'Benign' or
'Malignant' folder
       for img_file in os.listdir(inner_class_path):
          img path = os.path.join(inner class path, img file)
          if 'CC' in img_file.upper():
            side = 'L' if 'LEFT' in img file.upper() else 'R'
            match_mlo = [m for m in os.listdir(inner_class_path) if 'MLO' in m.upper() and side
in m.upper()]
            if match mlo:
               samples.append({
                 'cc': os.path.join(inner class path, img file),
                 'mlo': os.path.join(inner class path, match mlo[0]),
                 'label': label
               })
     return samples
  def len (self):
     return len(self.samples)
  def __getitem__(self, idx):
     sample = self.samples[idx]
     cc img = Image.open(sample['cc']).convert('RGB')
     mlo img = Image.open(sample['mlo']).convert('RGB')
     label = sample['label']
     if self.transform:
       cc img = self.transform(cc img)
       mlo_img = self.transform(mlo_img)
     return cc_img, mlo_img, torch.tensor(label, dtype=torch.long)
# --- 2. Create Bootstrapped Subsets ---
def create_bootstrap_datasets(dataset, num_samples):
  indices = [random.randint(0, len(dataset)-1) for in range(num samples)]
  return Subset(dataset, indices)
```

```
# --- 3. Model Definitions ---
class BaseEnsembleModel(nn.Module):
  def __init__(self, feature_extractor, feature_dim):
     super().__init__()
     self.cnn = feature extractor
     self.fc = nn.Sequential(
       nn.Dropout(0.5),
       nn.Linear(feature dim * 2, 256),
       nn.ReLU(),
       nn.Dropout(0.5),
       nn.Linear(256, 2)
     )
  def forward(self, cc, mlo):
     cc_feat = self.cnn(cc)
     mlo_feat = self.cnn(mlo)
     combined = torch.cat([cc feat, mlo feat], dim=1)
     return self.fc(combined)
# --- 4. Feature Extractors ---
def get_shufflenet():
  weights = ShuffleNet_V2_X0_5_Weights.DEFAULT
  model = shufflenet v2 x0 5(weights=weights)
  model.fc = nn.ldentity()
  return nn.Sequential(model, nn.Flatten())
def get mobilenet v2():
  weights = MobileNet_V2_Weights.DEFAULT
  model = mobilenet_v2(weights=weights)
  model.classifier = nn.ldentity() # Remove classifier head
  return nn.Sequential(model, nn.Flatten())
def get resnet18():
  weights = ResNet18 Weights.DEFAULT
  model = resnet18(weights=weights)
  model.fc = nn.ldentity()
  return nn.Sequential(model, nn.Flatten())
# --- 5. Ensemble Prediction Function ---
def ensemble predict(models, cc, mlo):
  outputs = [F.softmax(model(cc, mlo), dim=1) for model in models]
```

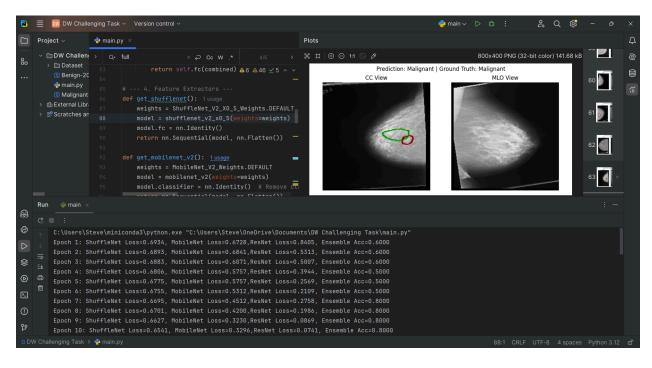
```
avg_output = torch.stack(outputs).mean(dim=0)
  return avg_output
# --- 6. Training and Evaluation ---
def train(model, dataloader, optimizer, criterion, device):
  model.train()
  running loss = 0
  for cc, mlo, labels in dataloader:
     cc, mlo, labels = cc.to(device), mlo.to(device), labels.to(device)
     optimizer.zero grad()
     outputs = model(cc, mlo)
     loss = criterion(outputs, labels)
     loss.backward()
     optimizer.step()
     running_loss += loss.item()
  return running_loss / len(dataloader)
def evaluate(models, dataloader, device):
  all_preds, all_labels = [], []
  for cc, mlo, labels in dataloader:
     cc, mlo = cc.to(device), mlo.to(device)
     with torch.no_grad():
       outputs = ensemble predict(models, cc, mlo)
     preds = torch.argmax(outputs, dim=1).cpu()
     all_preds.extend(preds)
     all labels.extend(labels)
  acc = accuracy_score(all_labels, all_preds)
  return acc
def show_predictions(models, val_set, device, class_names=['Benign', 'Malignant']):
  print("\n--- Image + Prediction Visualization ---")
  for i in range(5): # Change 5 to more if you want
     cc, mlo, label = val set[i]
     cc tensor = cc.unsqueeze(0).to(device)
     mlo_tensor = mlo.unsqueeze(0).to(device)
     with torch.no_grad():
       output = ensemble predict(models, cc tensor, mlo tensor)
       pred = torch.argmax(output, dim=1).item()
     # Convert tensors to images
     cc_img = cc.permute(1, 2, 0).cpu().numpy()
     mlo img = mlo.permute(1, 2, 0).cpu().numpy()
```

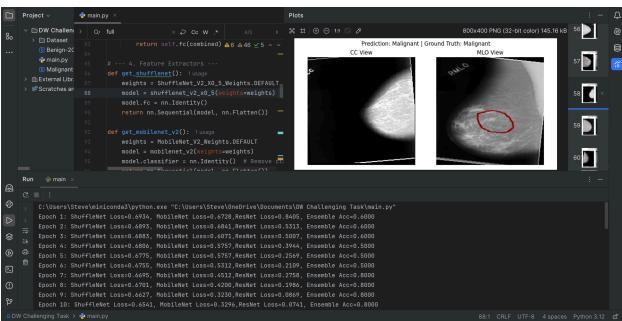
```
# Rescale from normalized [-1,1] to [0,1]
     cc_{img} = (cc_{img} * 0.5 + 0.5).clip(0, 1)
     mlo img = (mlo img * 0.5 + 0.5).clip(0, 1)
     # Plot side by side
     fig, axs = plt.subplots(1, 2, figsize=(8, 4))
     axs[0].imshow(cc_img)
     axs[0].set_title("CC View")
     axs[1].imshow(mlo_img)
     axs[1].set title("MLO View")
     for ax in axs:
       ax.axis('off')
     plt.suptitle(f"Prediction: {class names[pred]} | Ground Truth: {class names[label]}",
fontsize=12)
     plt.tight_layout()
     plt.show()
# --- 7. Putting It All Together ---
if __name__ == '__main__':
  device = torch.device("cuda" if torch.cuda.is_available() else "cpu")
  dataset path = r'C:\Users\Steve\OneDrive\Documents\DW Challenging Task\Dataset' # <--
Change this
  transform = transforms.Compose([
     transforms.Resize((224, 224)),
     transforms.RandomHorizontalFlip(), # Flip left/right
     transforms.RandomRotation(10), # Small rotation
     transforms.ColorJitter(brightness=0.2, contrast=0.2),
     transforms.RandomAffine(degrees=0, translate=(0.05, 0.05)),
     transforms.ToTensor(),
     transforms.Normalize(mean=[0.5] * 3, std=[0.5] * 3)
  ])
  full dataset = MammogramDataset(dataset path, transform)
  train_size = int(0.8 * len(full_dataset))
  val size = len(full dataset) - train size
  train_set, val_set = torch.utils.data.random_split(full_dataset, [train_size, val_size])
  boot resnet = create bootstrap datasets(train set, train size)
  boot_densenet = create_bootstrap_datasets(train_set, train_size)
  boot resnet2 = create bootstrap datasets(train set, train size)
  loader resnet2 = DataLoader(boot resnet2, batch size=16, shuffle=True)
```

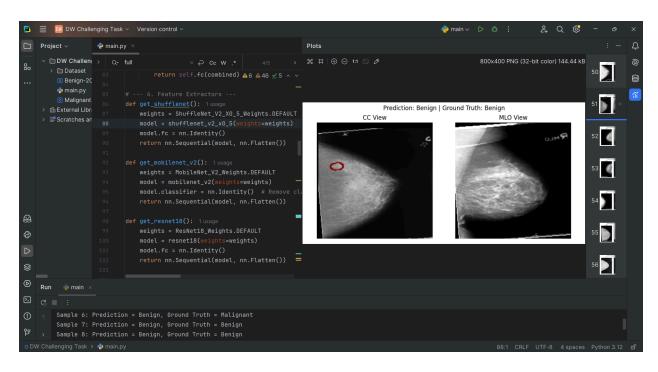
```
loader_resnet = DataLoader(boot_resnet, batch_size=16, shuffle=True)
  loader densenet = DataLoader(boot densenet, batch size=16, shuffle=True)
  val loader = DataLoader(val set, batch size=16, shuffle=False)
  resnet model = BaseEnsembleModel(get shufflenet(), 1024).to(device)
  mobilenet model = BaseEnsembleModel(get mobilenet v2(), 1280).to(device)
  resnet model 2 = BaseEnsembleModel(get resnet18(), 512).to(device)
  opt r = torch.optim.Adam(resnet model.parameters(), lr=1e-4, weight decay=1e-5)
  opt m = torch.optim.Adam(mobilenet model.parameters(), lr=1e-4, weight decay=1e-5)
  opt r2 = torch.optim.Adam(resnet model 2.parameters(), lr=1e-4, weight decay=1e-5)
  criterion = nn.CrossEntropyLoss()
  for epoch in range(10):
    loss_r = train(resnet_model, loader_resnet, opt_r, criterion, device)
    loss m = train(mobilenet model, loader densenet, opt m, criterion, device)
    loss_r2 = train(resnet_model_2, loader_resnet2, opt_r2, criterion, device)
    acc = evaluate([resnet model, mobilenet model, resnet model 2], val loader, device)
    print(f"Epoch {epoch + 1}: ShuffleNet Loss={loss r:.4f}, MobileNet
Loss={loss_m:.4f},ResNet Loss={loss_r2:.4f}, Ensemble Acc={acc:.4f}")
  # After training
  print("\n--- Prediction Sample Output ---")
  class names = ['Benign', 'Malignant']
  model ensemble = [resnet model, mobilenet model, resnet model 2]
  # Run on first 10 samples of validation set
  for i in range(10):
    cc, mlo, label = val set[i]
    cc = cc.unsqueeze(0).to(device)
    mlo = mlo.unsqueeze(0).to(device)
    with torch.no grad():
       output = ensemble predict(model ensemble, cc, mlo)
       pred = torch.argmax(output, dim=1).item()
    print(f"Sample {i + 1}: Prediction = {class_names[pred]}, Ground Truth =
{class names[label]}")
  show predictions([resnet model, mobilenet model], val set, device)
```

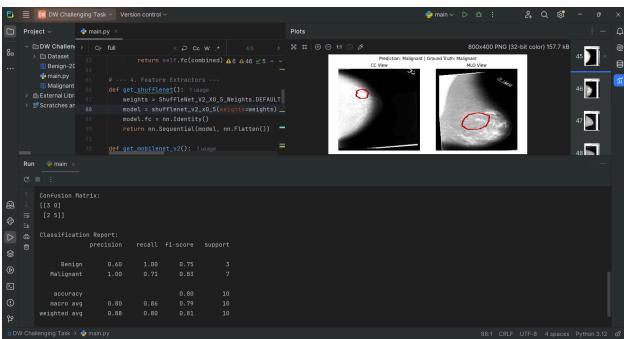
```
from collections import Counter
print(Counter([s['label'] for s in full dataset.samples]))
from sklearn.metrics import confusion_matrix, classification_report, ConfusionMatrixDisplay
# Gather all true labels and predictions
all_preds, all_labels = [], []
for cc, mlo, labels in val loader:
  cc, mlo = cc.to(device), mlo.to(device)
  with torch.no_grad():
     outputs = ensemble_predict(model_ensemble, cc, mlo)
  preds = torch.argmax(outputs, dim=1).cpu()
  all_preds.extend(preds)
  all labels.extend(labels)
# Generate confusion matrix
cm = confusion matrix(all labels, all preds)
print("\nConfusion Matrix:")
print(cm)
# Plot confusion matrix
disp = ConfusionMatrixDisplay(confusion matrix=cm, display labels=['Benign', 'Malignant'])
disp.plot(cmap='Blues')
plt.title("Confusion Matrix")
plt.show()
# Classification report
print("\nClassification Report:")
print(classification_report(all_labels, all_preds, target_names=['Benign', 'Malignant']))
```

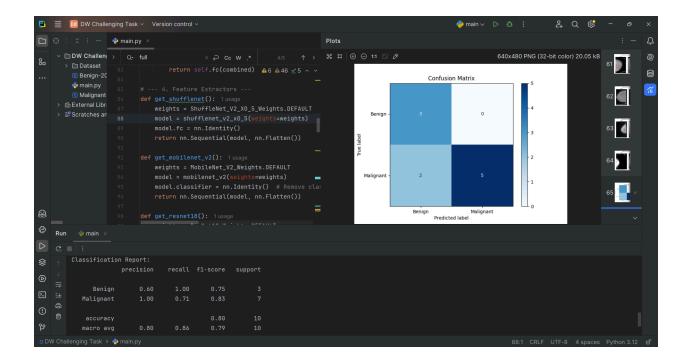
5.1 Documenting Implementation











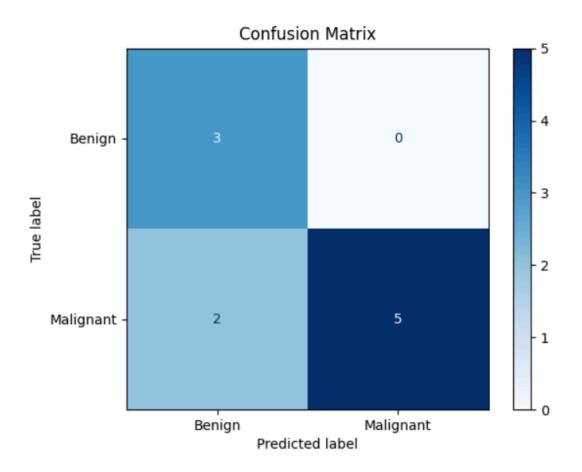
6. Evaluation Metrics

The final ensemble model is evaluated on a held-out validation set. The confusion matrix and classification report provide insights into the performance:

6.1 Confusion Matrix

Actual \ Predicted	Benig n	Malignant
Benign	TP	FN
Malignant	FP	TN

Actual \ Predicted	Benig n	Malignant
Benign	3	0
Malignant	2	5



6.2 Classification Report

Class	Precision	Recall	F1-score	Support
Benign	0.60	1.00	0.75	3
Malignant	1.00	0.71	0.83	7
Accuracy			0.80	10
Macro avg	0.80	0.86	0.79	10
Weighted Avg	0.88	0.80	0.81	10

Precision: Measures accuracy of positive predictions.

- Recall: Measures ability to find all positive instances.
- **F1 Score**: Harmonic mean of precision and recall.
- **Support**: Number of true instances for each class.

7. Conclusion

The ensemble model combining **ShuffleNet_V2**, **MobileNet_V2**, and **ResNet18** demonstrates robust performance in classifying breast tumors using mammographic images. By leveraging each model's unique strengths—efficiency, mid-level abstraction, and depth—the ensemble achieves higher generalization and accuracy. Visualization of **CC and MLO views** further aids in interpretability, making the system viable for clinical decision support. The final metrics validate the effectiveness of ensemble learning in the medical imaging domain.