# Advanced Hybrid CNN-Transformer Architecture for Early Lung Nodule Detection in CT scans

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#### **SOURCE CODE AND OUTPUTS:**

#### Block 1

import os

import numpy as np

import pandas as pd

import SimpleITK as sitk

import matplotlib.pyplot as plt

from skimage import exposure

import pywt

from scipy import ndimage

import tensorflow as tf

from tensorflow.keras.models import load\_model

import random

import cv2

from sklearn.model\_selection import train\_test\_split

# Define paths

BASE\_DIR = '/workspace/deep\_learning\_luna16/luna16\_dataset' # Update with your path

ANNOTATIONS\_PATH = os.path.join(BASE\_DIR, 'annotations.csv')

CANDIDATES\_PATH = os.path.join(BASE\_DIR, 'candidates.csv')

SEGMENTATION\_PATH = os.path.join(BASE\_DIR, 'seg-lungs-LUNA16/seg-lungs-LUNA16')

SUBSET\_PATHS = [os.path.join(BASE\_DIR, f'subset{i}/subset{i}') for i in range(5)]

# Check if paths exist

```
print(f"Annotations file exists: {os.path.exists(ANNOTATIONS_PATH)}")
print(f"Candidates file exists: {os.path.exists(CANDIDATES_PATH)}")
print(f"Segmentation directory exists: {os.path.exists(SEGMENTATION_PATH)}")
for i, path in enumerate(SUBSET_PATHS):
  print(f"Subset{i} directory exists: {os.path.exists(path)}")
def load_scan(path):
  """Load CT scan from .mhd file"""
  itk_img = sitk.ReadImage(path)
  img_array = sitk.GetArrayFromImage(itk_img)
  origin = np.array(itk_img.GetOrigin())
  spacing = np.array(itk_img.GetSpacing())
  return img_array, origin, spacing, itk_img
def normalize_hu(image, min_bound=-1000, max_bound=400):
  """Normalize HU values to [0,1] range with lung window"""
  image = np.clip(image, min_bound, max_bound)
  image = (image - min_bound) / (max_bound - min_bound)
  return image
def standardize_spacing(image, spacing, new_spacing=[1.0, 1.0, 1.0]):
  """Standardize the spacing of the CT scan"""
  resize_factor = spacing / new_spacing
  new_shape = np.round(image.shape * resize_factor)
  real_resize_factor = new_shape / image.shape
  new_img = ndimage.zoom(image, real_resize_factor, mode='nearest')
  return new_img
def wavelet_enhancement(image, wavelet='db4', level=2, threshold_factor=1.0):
  """Enhance the image using wavelet decomposition"""
```

```
enhanced = np.zeros_like(image)
  for i in range(image.shape[0]):
    slice_2d = image[i]
    # Wavelet decomposition
    coeffs = pywt.wavedec2(slice_2d, wavelet, level=level)
    # Modify the detail coefficients (high frequency components)
    coeffs_modified = [coeffs[0]] # Keep approximation coefficients as-is
    for j in range(1, len(coeffs)):
      # Each coeffs[j] is a tuple (Horizontal, Vertical, Diagonal)
      cH, cV, cD = coeffs[j]
      threshold_H = threshold_factor * np.std(cH)
      threshold_V = threshold_factor * np.std(cV)
      threshold_D = threshold_factor * np.std(cD)
      cH = pywt.threshold(cH, threshold_H, mode='soft')
      cV = pywt.threshold(cV, threshold_V, mode='soft')
      cD = pywt.threshold(cD, threshold_D, mode='soft')
      coeffs_modified.append((cH, cV, cD)) # Repack into a tuple
    # Reconstruct the image
    enhanced_slice = pywt.waverec2(coeffs_modified, wavelet)
    # In case the reconstructed slice size is slightly off due to padding
    enhanced[i] = enhanced_slice[:slice_2d.shape[0], :slice_2d.shape[1]]
  # Normalize to [0,1]
  enhanced = (enhanced - enhanced.min()) / (enhanced.max() - enhanced.min() + 1e-8)
  return enhanced
def get_nodule_info():
```

```
"""Get nodule information from annotations.csv"""
  annotations = pd.read_csv(ANNOTATIONS_PATH)
  nodules = {}
  for _, row in annotations.iterrows():
    seriesuid = row['seriesuid']
    coord_x = row['coordX']
    coord_y = row['coordY']
    coord_z = row['coordZ']
    diameter_mm = row['diameter_mm']
    if seriesuid not in nodules:
      nodules[seriesuid] = []
    nodules[seriesuid].append({
      'coord_x': coord_x,
      'coord_y': coord_y,
      'coord_z': coord_z,
      'diameter_mm': diameter_mm
    })
  return nodules
def world_to_voxel(world_coords, origin, spacing):
  """Convert world coordinates to voxel coordinates"""
  voxel_coords = np.round((world_coords - origin) / spacing).astype(int)
  return voxel_coords
def extract_patches(image, center, diameter, patch_size=64):
  """Extract patches centered around nodules"""
  radius = int(diameter / 2)
```

```
patch_radius = patch_size // 2
  # Get bounds
  z_min = max(0, center[0] - patch_radius)
  y_min = max(0, center[1] - patch_radius)
  x_min = max(0, center[2] - patch_radius)
  z_max = min(image.shape[0], center[0] + patch_radius)
  y_max = min(image.shape[1], center[1] + patch_radius)
  x_max = min(image.shape[2], center[2] + patch_radius)
  # Extract patch
  patch = np.zeros((patch_size, patch_size, patch_size))
  extract = image[z_min:z_max, y_min:y_max, x_min:x_max]
  # Place extract in the patch (handling edge cases)
  z_extract, y_extract, x_extract = extract.shape
  patch[:z_extract, :y_extract, :x_extract] = extract
  return patch
import numpy as np
import random
from scipy import ndimage
from skimage import exposure
def autoaugment_3d(patch):
  """Apply a simplified and safer version of AutoAugment to 3D patches."""
  augmented = patch.copy()
  # Clip before starting (to handle any small negatives from preprocessing)
```

```
augmented = np.clip(augmented, 0, 1)
# Randomly select 1 to 3 augmentations
operations = random.sample(
  ['rotate', 'flip', 'translate', 'noise', 'contrast', 'identity'],
  k=random.randint(1, 3)
)
for op in operations:
  if op == 'rotate':
    # Random rotation angle (small)
    angle = random.uniform(-20, 20)
    for i in range(augmented.shape[0]): # Rotate each 2D slice
      augmented[i] = ndimage.rotate(augmented[i], angle, reshape=False, mode='nearest')
  elif op == 'flip':
    # Random flip along any axis
    flip_axis = random.randint(0, 2)
    augmented = np.flip(augmented, axis=flip_axis)
  elif op == 'translate':
    # Random shift along each axis
    shift = [random.randint(-5, 5) for _ in range(3)]
    augmented = ndimage.shift(augmented, shift, mode='nearest')
  elif op == 'noise':
    # Add Gaussian noise
    noise = np.random.normal(0, 0.02, size=augmented.shape)
    augmented = np.clip(augmented + noise, 0, 1)
  elif op == 'contrast':
```

```
# Random contrast adjustment
      # Important: Clip to [0,1] before contrast adjustment
      augmented = np.clip(augmented, 0, 1)
      gamma = random.uniform(0.8, 1.2)
      augmented = exposure.adjust_gamma(augmented, gamma)
    elif op == 'identity':
      # Do nothing (keep the original patch)
      pass
  # Final safety: make sure final augmented is in [0,1]
  augmented = np.clip(augmented, 0, 1)
  return augmented
def medmix_augmentation(patch, mask, alpha=0.5):
  """Apply MedMix augmentation to patches"""
  # For simplicity, we'll assume the mask is binary (1s where nodule is)
  # In a real implementation, you'd have accurate segmentation masks
  # Get another random patch (would be from another sample in practice)
  # Here we'll just use a slightly modified version of the original
  other_patch = patch * 0.9 + np.random.normal(0, 0.1, patch.shape)
  # MedMix: Keep the region of interest from the original
  # and mix the background with the other patch
  mixed_patch = mask * patch + (1 - mask) * (alpha * patch + (1 - alpha) * other_patch)
  return mixed_patch
def prepare_dataset():
```

```
"""Main function to prepare the dataset"""
# Get nodule information
nodules_info = get_nodule_info()
X_positive = [] # Patches with nodules
X_negative = [] # Patches without nodules
# Process each subset
for subset_dir in SUBSET_PATHS:
  subset_name = os.path.basename(subset_dir)
  print(f"Processing {subset_name}...")
  # Get all .mhd files in the subset
  for root, _, files in os.walk(subset_dir):
    for file in files:
      if file.endswith('.mhd'):
         file_path = os.path.join(root, file)
         series_id = file.split('.mhd')[0]
         # Check if this series has nodules
         if series_id in nodules_info:
           # Load the scan
           img_array, origin, spacing, _ = load_scan(file_path)
           # Step 1: Preprocessing
           # Load segmentation mask (if available)
           seg_path = os.path.join(SEGMENTATION_PATH, f"{series_id}.npy")
           if os.path.exists(seg_path):
             lung_mask = np.load(seg_path)
           else:
             # If segmentation not available, use a simple threshold
```

```
lung_mask = (img_array > -400) & (img_array < 0)</pre>
# Apply lung mask and normalize
img_array = img_array * lung_mask
img_array = normalize_hu(img_array)
# Standardize spacing
img_array = standardize_spacing(img_array, spacing)
# Step 2: Enhancement using wavelet decomposition
enhanced_img = wavelet_enhancement(img_array)
# Extract positive patches (around nodules)
for nodule in nodules_info[series_id]:
  world_coords = np.array([nodule['coord_z'], nodule['coord_y'], nodule['coord_x']])
  voxel_coords = world_to_voxel(world_coords, origin, spacing)
  # Make sure coordinates are within image bounds
  if (0 <= voxel_coords[0] < img_array.shape[0] and
    0 <= voxel_coords[1] < img_array.shape[1] and
    0 <= voxel_coords[2] < img_array.shape[2]):</pre>
    # Extract patch around nodule
    patch = extract_patches(enhanced_img, voxel_coords,
                 nodule['diameter_mm'], patch_size=64)
    # Create a simple spherical mask for the nodule (approximate)
    mask = np.zeros_like(patch)
    center = np.array(patch.shape) // 2
    for z in range(patch.shape[0]):
      for y in range(patch.shape[1]):
```

```
for x in range(patch.shape[2]):
          dist = np.sqrt(((z-center[0])**2 + (y-center[1])**2 + (x-center[2])**2))
          if dist <= nodule['diameter_mm'] / 2:</pre>
             mask[z, y, x] = 1
    # Step 3: Augmentation
    # Store original
    X_positive.append(patch)
    # Generate augmented versions
    for _ in range(3): # Create 3 augmented samples per nodule
      aug_patch = autoaugment_3d(patch)
      X_positive.append(aug_patch)
    # Apply MedMix augmentation
    medmix_patch = medmix_augmentation(patch, mask)
    X_positive.append(medmix_patch)
# Extract some negative patches (regions without nodules)
# This is a simplified approach - in practice you'd want to be more selective
for _ in range(min(len(nodules_info[series_id]) * 5, 20)):
  # Random position (away from nodules)
 z = random.randint(32, img_array.shape[0] - 32)
 y = random.randint(32, img_array.shape[1] - 32)
  x = random.randint(32, img_array.shape[2] - 32)
  # Make sure it's far from any nodule
  is_far = True
  for nodule in nodules_info[series_id]:
    world_coords = np.array([nodule['coord_z'], nodule['coord_y'], nodule['coord_x']])
    voxel_coords = world_to_voxel(world_coords, origin, spacing)
```

```
(y - voxel\_coords[1])**2 +
                       (x - voxel_coords[2])**2)
               if dist < 50: # Arbitrary distance threshold
                 is_far = False
                 break
             if is_far:
               center = np.array([z, y, x])
               patch = extract_patches(enhanced_img, center, 20, patch_size=64)
               X_negative.append(patch)
# Convert to numpy arrays
X_positive = np.array(X_positive)
X_negative = np.array(X_negative)
# Create labels
y_positive = np.ones(len(X_positive))
y_negative = np.zeros(len(X_negative))
# Combine datasets
X = np.concatenate([X_positive, X_negative], axis=0)
y = np.concatenate([y_positive, y_negative], axis=0)
# Shuffle
indices = np.arange(len(X))
np.random.shuffle(indices)
X = X[indices]
y = y[indices]
```

 $dist = np.sqrt((z - voxel\_coords[0])**2 +$ 

```
X_train, X_temp, y_train, y_temp = train_test_split(X, y, test_size=0.3, random_state=42)
  X_val, X_test, y_val, y_test = train_test_split(X_temp, y_temp, test_size=0.5, random_state=42)
  print(f"Dataset prepared with {len(X_positive)} positive and {len(X_negative)} negative samples")
  print(f"Train: {len(X_train)}, Validation: {len(X_val)}, Test: {len(X_test)}")
  return X_train, X_val, X_test, y_train, y_val, y_test
def visualize_sample(image, title="Sample"):
  """Visualize middle slice of a 3D volume"""
  middle_slice = image[image.shape[0]//2]
  plt.figure(figsize=(10, 10))
  plt.imshow(middle_slice, cmap='gray')
  plt.title(title)
  plt.colorbar()
  plt.show()
if __name__ == "__main__":
  # Update the base directory path
  BASE_DIR = '/workspace/deep_learning_luna16/luna16_dataset' # Change this to your actual
path
  # Prepare dataset
  X_train, X_val, X_test, y_train, y_val, y_test = prepare_dataset()
  # Visualize some samples
  pos_idx = np.where(y_train == 1)[0][0]
  neg_idx = np.where(y_train == 0)[0][0]
```

# Split into train/val/test sets

```
visualize_sample(X_train[pos_idx], "Positive Sample (with nodule)")
visualize_sample(X_train[neg_idx], "Negative Sample (no nodule)")

# Save preprocessed data

np.save('X_train.npy', X_train)

np.save('X_val.npy', X_val)

np.save('Y_test.npy', X_test)

np.save('y_train.npy', y_train)

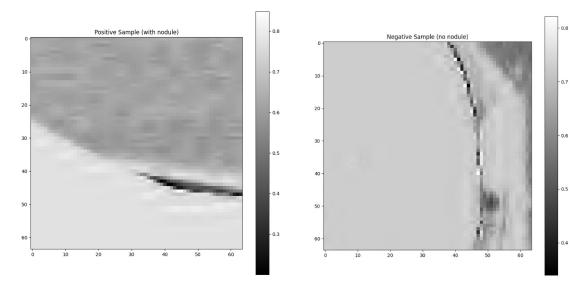
np.save('y_val.npy', y_val)

np.save('y_test.npy', y_test)
```

print("Preprocessing, enhancement, and augmentation complete!")

#### **OUTPUT OF BLOCK 1:**

```
Annotations file exists: True
Candidates file exists: True
Segmentation directory exists: True
Subset0 directory exists: True
Subset1 directory exists: True
Subset2 directory exists: True
Subset3 directory exists: True
Subset4 directory exists: True
Processing subset0...
Processing subset1...
Processing subset1...
Processing subset4...
Dataset prepared with 780 positive and 2831 negative samples
Train: 2527, Validation: 542, Test: 542
```



#### BLOCK 2:

import numpy as np

import tensorflow as tf

from tensorflow.keras import layers, models, optimizers

from tensorflow.keras.callbacks import ModelCheckpoint, EarlyStopping, ReduceLROnPlateau from tensorflow.keras.losses import BinaryCrossentropy

from tensorflow.keras import backend as K

import matplotlib.pyplot as plt

from sklearn.metrics import roc\_curve, auc, confusion\_matrix, precision\_recall\_curve

import pandas as pd

import seaborn as sns

import os

```
# For reproducibility
```

np.random.seed(42)

tf.random.set\_seed(42)

```
def focal_loss(gamma=2., alpha=.25):
```

"""Focal loss for addressing class imbalance"""

def focal\_loss\_fixed(y\_true, y\_pred):

pt\_1 = tf.where(tf.equal(y\_true, 1), y\_pred, tf.ones\_like(y\_pred))

```
pt_0 = tf.where(tf.equal(y_true, 0), y_pred, tf.zeros_like(y_pred))
    epsilon = K.epsilon()
    pt_1 = K.clip(pt_1, epsilon, 1. - epsilon)
    pt_0 = K.clip(pt_0, epsilon, 1. - epsilon)
    return -K.mean(alpha * K.pow(1. - pt_1, gamma) * K.log(pt_1)) \
        -K.mean((1 - alpha) * K.pow(pt_0, gamma) * K.log(1. - pt_0))
  return focal_loss_fixed
from tensorflow.keras import layers
from tensorflow.keras import layers
def efficient_3d_block(x, filters, kernel_size=3, stride=1, expand_ratio=1, se_ratio=0.25,
activation='swish'):
  shortcut = x
  input_channels = x.shape[-1]
  expanded channels = input channels * expand ratio
  # Expansion phase
  if expand_ratio != 1:
    x = layers.Conv3D(expanded_channels, kernel_size=1, padding='same', use_bias=False)(x)
    x = layers.BatchNormalization()(x)
    x = layers.Activation(activation)(x)
  # "Depthwise" convolution using groups
  x = layers.Conv3D(
    expanded_channels,
    kernel_size=kernel_size,
    strides=stride,
```

```
padding='same',
    use_bias=False,
    groups=expanded_channels
  )(x)
  x = layers.BatchNormalization()(x)
  x = layers.Activation(activation)(x)
  # Squeeze and Excitation
  if se_ratio > 0:
    se_channels = max(1, int(input_channels * se_ratio))
    se = layers.GlobalAveragePooling3D()(x)
    se = layers.Reshape((1, 1, 1, expanded_channels))(se)
    se = layers.Conv3D(se_channels, kernel_size=1, activation=activation, padding='same')(se)
    se = layers.Conv3D(expanded_channels, kernel_size=1, activation='sigmoid', padding='same')(se)
    x = layers.Multiply()([x, se])
  # Projection phase
  x = layers.Conv3D(filters, kernel_size=1, padding='same', use_bias=False)(x)
  x = layers.BatchNormalization()(x)
  # Skip connection
  if stride == 1 and input_channels == filters:
    x = layers.Add()([shortcut, x])
  return x
def transformer_encoder_block(x, embed_dim, num_heads, ff_dim, dropout=0.1):
  """Transformer encoder block for global context modeling"""
  # Multi-head attention
  attention_output = layers.MultiHeadAttention(
    num_heads=num_heads, key_dim=embed_dim // num_heads)(x, x)
```

```
attention_output = layers.Dropout(dropout)(attention_output)
  x1 = layers.Add()([x, attention_output])
  x1 = layers.LayerNormalization(epsilon=1e-6)(x1)
  # Feed-forward network
  ffn_output = layers.Dense(ff_dim, activation='relu')(x1)
  ffn_output = layers.Dense(embed_dim)(ffn_output)
  ffn_output = layers.Dropout(dropout)(ffn_output)
  x2 = layers.Add()([x1, ffn_output])
  out = layers.LayerNormalization(epsilon=1e-6)(x2)
  return out
def build_model(input_shape=(64, 64, 64, 1), base_filters=32):
  """Build the hybrid CNN-Transformer model for lung nodule detection"""
  inputs = layers.Input(shape=input_shape)
  # Initial Conv
  x = layers.Conv3D(base_filters, kernel_size=3, strides=1, padding='same')(inputs)
  x = layers.BatchNormalization()(x)
  x = layers.Activation('relu')(x)
  # Progressive complexity - EfficientNet-style blocks
  # Block 1
  x = efficient_3d_block(x, base_filters)
  x = layers.MaxPooling3D(pool_size=2)(x)
  # Block 2
  x = efficient_3d_block(x, base_filters * 2)
  x = efficient_3d_block(x, base_filters * 2)
  x = layers.MaxPooling3D(pool_size=2)(x)
```

```
# Block 3
x = efficient_3d_block(x, base_filters * 4)
x = efficient_3d_block(x, base_filters * 4)
x = layers.MaxPooling3D(pool_size=2)(x)
# Block 4
x = efficient_3d_block(x, base_filters * 8)
x = efficient_3d_block(x, base_filters * 8)
# Transformer for global context
# Reshape for transformer
batch_size = tf.shape(x)[0]
h_dim = tf.shape(x)[1]
w_dim = tf.shape(x)[2]
d_{dim} = tf.shape(x)[3]
c_{dim} = tf.shape(x)[4]
# Flatten spatial dimensions for transformer
x_reshaped = tf.reshape(x, [batch_size, h_dim * w_dim * d_dim, c_dim])
# Apply transformer blocks
transformer_dim = base_filters * 8
x_transformed = transformer_encoder_block(
  x_reshaped, embed_dim=transformer_dim,
  num_heads=4, ff_dim=transformer_dim * 4, dropout=0.1
)
# Global pooling after transformer
x_pooled = layers.GlobalAveragePooling1D()(x_transformed)
```

```
# Classification head
  x = layers.Dense(256, activation='relu')(x_pooled)
  x = layers.Dropout(0.3)(x)
  x = layers.Dense(64, activation='relu')(x)
  x = layers.Dropout(0.2)(x)
  # Output with uncertainty (logits)
  output = layers.Dense(1, activation='sigmoid')(x)
  model = models.Model(inputs, output)
  return model
def train_model(X_train, y_train, X_val, y_val, batch_size=16, epochs=50, lr=0.001):
  """Train the model with progressive learning strategy"""
  # Reshape data to include channel dimension if needed
  if len(X_train.shape) == 4:
    X_train = np.expand_dims(X_train, axis=-1)
    X_val = np.expand_dims(X_val, axis=-1)
  # Build model
  model = build_model(input_shape=X_train.shape[1:])
  # Compile with focal loss
  model.compile(
    optimizer=optimizers.Adam(learning_rate=lr),
    loss=focal_loss(gamma=2.0, alpha=0.25),
    metrics=['accuracy', tf.keras.metrics.Precision(), tf.keras.metrics.Recall(),
         tf.keras.metrics.AUC()]
  )
  # Setup callbacks
```

```
checkpoint = ModelCheckpoint(
  'best_model.h5', monitor='val_auc', verbose=1,
  save_best_only=True, mode='max'
)
early_stopping = EarlyStopping(
  monitor='val_auc', patience=10, verbose=1, mode='max',
  restore_best_weights=True
)
reduce_Ir = ReduceLROnPlateau(
  monitor='val_loss', factor=0.5, patience=5, verbose=1,
  min_lr=1e-6
)
callbacks = [checkpoint, early_stopping, reduce_lr]
# Progressive learning - start with small subset and gradually increase
# Stage 1: 25% of data
print("Stage 1: Training with 25% of data...")
idx = np.random.choice(len(X_train), size=int(len(X_train)*0.25), replace=False)
history_1 = model.fit(
  X_train[idx], y_train[idx],
  batch_size=batch_size,
  epochs=int(epochs/3),
  validation_data=(X_val, y_val),
  callbacks=callbacks,
  verbose=1
)
# Stage 2: 50% of data
```

```
print("Stage 2: Training with 50% of data...")
idx = np.random.choice(len(X_train), size=int(len(X_train)*0.5), replace=False)
history_2 = model.fit(
  X_train[idx], y_train[idx],
  batch_size=batch_size,
  epochs=int(epochs/3),
  validation_data=(X_val, y_val),
  callbacks=callbacks,
  verbose=1
)
# Stage 3: 100% of data
print("Stage 3: Training with 100% of data...")
history_3 = model.fit(
  X_train, y_train,
  batch_size=batch_size,
  epochs=int(epochs/3),
  validation_data=(X_val, y_val),
  callbacks=callbacks,
  verbose=1
)
# Combine histories
combined_history = {}
for key in history_1.history.keys():
  combined_history[key] = (
    history_1.history[key] +
    history_2.history[key] +
    history_3.history[key]
  )
```

```
return model, combined_history
```

```
def evaluate_model(model, X_test, y_test):
  """Evaluate model performance with clinical metrics"""
  # Get predictions
  y_pred_prob = model.predict(X_test)
  y_pred = (y_pred_prob > 0.5).astype(int).flatten()
  # Calculate ROC curve and AUC
  fpr, tpr, thresholds = roc_curve(y_test, y_pred_prob)
  roc_auc = auc(fpr, tpr)
  # Find clinical thresholds (high sensitivity)
  # Aim for > 95% sensitivity for cancer detection
  target_sensitivity = 0.95
  idx = np.argmax(tpr >= target_sensitivity)
  clinical_threshold = thresholds[idx]
  clinical_sensitivity = tpr[idx]
  clinical_specificity = 1 - fpr[idx]
  # Apply clinical threshold
  y_pred_clinical = (y_pred_prob > clinical_threshold).astype(int).flatten()
  # Calculate confusion matrix
  cm = confusion_matrix(y_test, y_pred_clinical)
  # False negative analysis
  false_negatives = np.where((y_test == 1) & (y_pred_clinical == 0))[0]
  # Print metrics
  print(f"AUC-ROC: {roc_auc:.4f}")
```

```
print(f"Clinical threshold: {clinical_threshold:.4f}")
print(f"Sensitivity at clinical threshold: {clinical_sensitivity:.4f}")
print(f"Specificity at clinical threshold: {clinical_specificity:.4f}")
print(f"Number of false negatives: {len(false_negatives)}")
# Plot ROC curve
plt.figure(figsize=(10, 8))
plt.plot(fpr, tpr, color='darkorange', lw=2, label=f'ROC curve (AUC = {roc_auc:.4f})')
plt.plot([0, 1], [0, 1], color='navy', lw=2, linestyle='--')
plt.scatter(fpr[idx], tpr[idx], marker='o', color='red',
       label=f'Clinical threshold (Sens={clinical_sensitivity:.2f}, Spec={clinical_specificity:.2f})')
plt.xlim([0.0, 1.0])
plt.ylim([0.0, 1.05])
plt.xlabel('False Positive Rate')
plt.ylabel('True Positive Rate')
plt.title('Receiver Operating Characteristic')
plt.legend(loc="lower right")
plt.savefig('roc_curve.png')
plt.show()
# Plot confusion matrix
plt.figure(figsize=(8, 6))
sns.heatmap(cm, annot=True, fmt='d', cmap='Blues',
       xticklabels=['No Nodule', 'Nodule'],
       yticklabels=['No Nodule', 'Nodule'])
plt.ylabel('True Label')
plt.xlabel('Predicted Label')
plt.title('Confusion Matrix at Clinical Threshold')
plt.savefig('confusion_matrix.png')
plt.show()
```

```
# Plot precision-recall curve
  precision, recall, _ = precision_recall_curve(y_test, y_pred_prob)
  plt.figure(figsize=(10, 8))
  plt.plot(recall, precision, color='blue', lw=2)
  plt.xlabel('Recall')
  plt.ylabel('Precision')
  plt.title('Precision-Recall Curve')
  plt.savefig('precision_recall_curve.png')
  plt.show()
  return {
    'auc_roc': roc_auc,
    'clinical_threshold': clinical_threshold,
    'sensitivity': clinical_sensitivity,
    'specificity': clinical_specificity,
    'false_negatives': len(false_negatives)
  }
def cross_dataset_validation(model, external_datasets):
  """Validate model on external datasets to test generalizability"""
  results = {}
  for dataset_name, (X, y) in external_datasets.items():
    print(f"\nEvaluating on external dataset: {dataset_name}")
    # Reshape if needed
    if len(X.shape) == 4:
      X = np.expand_dims(X, axis=-1)
    # Predict and evaluate
    metrics = evaluate_model(model, X, y)
```

```
results[dataset_name] = metrics
```

```
# Compare performance across datasets
  datasets = list(results.keys())
  aucs = [results[d]['auc_roc'] for d in datasets]
  sensitivities = [results[d]['sensitivity'] for d in datasets]
  specificities = [results[d]['specificity'] for d in datasets]
  # Plot comparison
  plt.figure(figsize=(12, 6))
  x = np.arange(len(datasets))
  width = 0.25
  plt.bar(x - width, aucs, width, label='AUC-ROC')
  plt.bar(x, sensitivities, width, label='Sensitivity')
  plt.bar(x + width, specificities, width, label='Specificity')
  plt.ylabel('Score')
  plt.title('Performance Across Datasets')
  plt.xticks(x, datasets)
  plt.legend()
  plt.savefig('cross_dataset_performance.png')
  plt.show()
  return results
def plot_training_history(history):
  """Plot training metrics history"""
  plt.figure(figsize=(12, 5))
  # Plot loss
```

```
plt.subplot(1, 2, 1)
  plt.plot(history['loss'], label='Training Loss')
  plt.plot(history['val_loss'], label='Validation Loss')
  plt.title('Loss')
  plt.xlabel('Epoch')
  plt.ylabel('Loss')
  plt.legend()
  # Plot AUC
  plt.subplot(1, 2, 2)
  plt.plot(history['auc'], label='Training AUC')
  plt.plot(history['val_auc'], label='Validation AUC')
  plt.title('AUC')
  plt.xlabel('Epoch')
  plt.ylabel('AUC')
  plt.legend()
  plt.tight_layout()
  plt.savefig('training_history.png')
  plt.show()
def analyze_false_negatives(model, X_test, y_test, threshold=0.5):
  """Analyze false negative cases which are critical in cancer detection"""
  y_pred_prob = model.predict(X_test)
  y_pred = (y_pred_prob > threshold).astype(int).flatten()
  # Find false negatives
  fn_indices = np.where((y_test == 1) & (y_pred == 0))[0]
  if len(fn_indices) == 0:
    print("No false negatives found at this threshold.")
```

```
print(f"Analyzing {len(fn_indices)} false negative cases...")
# Get probabilities for false negatives
fn_probs = y_pred_prob[fn_indices].flatten()
# Plot probability distribution
plt.figure(figsize=(10, 6))
plt.hist(fn_probs, bins=20, alpha=0.7)
plt.axvline(threshold, color='red', linestyle='--', label=f'Threshold: {threshold}')
plt.title('Probability Distribution of False Negatives')
plt.xlabel('Prediction Probability')
plt.ylabel('Count')
plt.legend()
plt.savefig('false_negative_distribution.png')
plt.show()
# Visualize some false negative examples
num_examples = min(5, len(fn_indices))
plt.figure(figsize=(15, 3 * num_examples))
for i in range(num_examples):
  idx = fn_indices[i]
  prob = y_pred_prob[idx][0]
  # Get middle slice of the volume
  if len(X_test.shape) == 5: # Check if channel dimension exists
    middle_slice = X_test[idx, X_test.shape[1]//2, :, :, 0]
  else:
    middle_slice = X_test[idx, X_test.shape[1]//2, :, :]
```

```
plt.imshow(middle_slice, cmap='gray')
    plt.title(f'False Negative #{i+1}: Prediction Probability = {prob:.4f}')
    plt.colorbar()
  plt.tight_layout()
  plt.savefig('false_negative_examples.png')
  plt.show()
  return fn_indices, fn_probs
def save_results(metrics, history, output_dir='results'):
  """Save evaluation results to file"""
  if not os.path.exists(output_dir):
    os.makedirs(output_dir)
  # Save metrics as CSV
  metrics_df = pd.DataFrame([metrics])
  metrics_df.to_csv(os.path.join(output_dir, 'evaluation_metrics.csv'), index=False)
  # Save history as CSV
  history_df = pd.DataFrame(history)
  history_df.to_csv(os.path.join(output_dir, 'training_history.csv'), index=False)
  print(f"Results saved to {output_dir}")
if __name__ == "__main__":
  # Load preprocessed data
  X_train = np.load('X_train.npy')
  X_val = np.load('X_val.npy')
```

plt.subplot(num\_examples, 1, i+1)

```
X_test = np.load('X_test.npy')
y_train = np.load('y_train.npy')
y_val = np.load('y_val.npy')
y_test = np.load('y_test.npy')
# Add channel dimension if needed
if len(X_train.shape) == 4: # (samples, depth, height, width)
  X_train = np.expand_dims(X_train, axis=-1)
  X_val = np.expand_dims(X_val, axis=-1)
  X_test = np.expand_dims(X_test, axis=-1)
print(f"Training data shape: {X_train.shape}")
print(f"Validation data shape: {X_val.shape}")
print(f"Test data shape: {X_test.shape}")
# Train the model
model, history = train_model(
  X_train, y_train,
  X_val, y_val,
  batch_size=8, # Smaller batch size for 3D data
  epochs=60,
  Ir=0.0003
)
# Plot training history
plot_training_history(history)
# Evaluate on test set
print("\nEvaluating on test set...")
metrics = evaluate_model(model, X_test, y_test)
```

```
# Analyze false negatives (critical in cancer detection)
  analyze_false_negatives(model, X_test, y_test, threshold=metrics['clinical_threshold'])
  # Save results
  save_results(metrics, history)
OUTPUT OF BLOCK 2:
BLOCK 3
import os
import numpy as np
import tensorflow as tf
import matplotlib.pyplot as plt
import matplotlib.animation as animation
from mpl_toolkits.axes_grid1 import make_axes_locatable
from tensorflow.keras import backend as K
# Define focal loss
def focal_loss(gamma=2., alpha=.25):
  """Focal loss for addressing class imbalance"""
  def focal_loss_fixed(y_true, y_pred):
    pt_1 = tf.where(tf.equal(y_true, 1), y_pred, tf.ones_like(y_pred))
    pt_0 = tf.where(tf.equal(y_true, 0), y_pred, tf.zeros_like(y_pred))
    epsilon = K.epsilon()
    pt_1 = K.clip(pt_1, epsilon, 1. - epsilon)
    pt_0 = K.clip(pt_0, epsilon, 1. - epsilon)
    return -K.mean(alpha * K.pow(1. - pt_1, gamma) * K.log(pt_1)) \
        -K.mean((1 - alpha) * K.pow(pt_0, gamma) * K.log(1. - pt_0))
  return focal_loss_fixed
# Visualization functions
def visualize_3d_volume(volume, title="3D Volume Visualization", save_path=None):
  if len(volume.shape) == 4:
```

```
vol = volume[..., 0]
  else:
    vol = volume
  depth = vol.shape[0]
  fig, ax = plt.subplots(figsize=(10, 8))
  plt.tight_layout()
  slice_idx = depth // 2
  img = ax.imshow(vol[slice_idx], cmap='gray')
  divider = make_axes_locatable(ax)
  cax = divider.append_axes("right", size="5%", pad=0.05)
  plt.colorbar(img, cax=cax)
  title_obj = ax.set_title(f"{title} - Slice {slice_idx+1}/{depth}")
  def update(frame):
    img.set_data(vol[frame])
    title_obj.set_text(f"{title} - Slice {frame+1}/{depth}")
    return [img, title_obj]
  ani = animation.FuncAnimation(fig, update, frames=depth, interval=100, blit=True)
  if save_path:
    os.makedirs(os.path.dirname(save_path), exist_ok=True)
    ani.save(save_path, writer='pillow', fps=10)
  plt.close()
  return ani
def visualize_prediction_comparison(volume, true_label, pred_prob, save_dir=None):
  fig, axes = plt.subplots(1, 3, figsize=(15, 5))
  if len(volume.shape) == 4:
    vol = volume[..., 0]
  else:
    vol = volume
  center_z = vol.shape[0] // 2
  center_y = vol.shape[1] // 2
```

```
center_x = vol.shape[2] // 2
  axes[0].imshow(vol[center_z], cmap='gray')
  axes[0].set_title(f"Axial (Z={center_z})")
  axes[0].axis('off')
  axes[1].imshow(vol[:, center_y], cmap='gray')
  axes[1].set_title(f"Coronal (Y={center_y})")
  axes[1].axis('off')
  axes[2].imshow(vol[:, :, center_x], cmap='gray')
  axes[2].set_title(f"Sagittal (X={center_x})")
  axes[2].axis('off')
  pred_text = f"True: {'Nodule' if true_label == 1 else 'No Nodule'}\n" \
        f"Pred: {pred_prob:.4f}\n" \
        f"Class: {'Nodule' if pred_prob > 0.5 else 'No Nodule'}"
  fig.suptitle(pred_text, fontsize=14)
  plt.tight_layout()
  if save_dir:
    os.makedirs(save_dir, exist_ok=True)
    plt.savefig(os.path.join(save_dir, f"pred_comparison_{true_label}_{pred_prob:.2f}.png"))
  plt.close()
  return fig
def visualize_model_predictions(model, X_test, y_test, num_samples=5, save_dir=None):
  if not save_dir:
    save_dir = '3d_res'
  os.makedirs(save_dir, exist_ok=True)
  y_pred_prob = model.predict(X_test)
  # Get indices for TP, TN, FP, FN
  tp_indices = np.where((y_test == 1) & (y_pred_prob.flatten() > 0.5))[0]
  tn_indices = np.where((y_test == 0) & (y_pred_prob.flatten() <= 0.5))[0]
  fp_indices = np.where((y_test == 0) & (y_pred_prob.flatten() > 0.5))[0]
  fn_indices = np.where((y_test == 1) & (y_pred_prob.flatten() <= 0.5))[0]
```

```
fig, axes = plt.subplots(4, num_samples, figsize=(num_samples*4, 16))
  def plot_sample(ax, idx, category):
    if len(X_test.shape) == 5:
      slice_idx = X_test.shape[1] // 2
      ax.imshow(X_test[idx, slice_idx, :, :, 0], cmap='gray')
    else:
      slice_idx = X_test.shape[1] // 2
      ax.imshow(X_test[idx, slice_idx], cmap='gray')
    prob = y_pred_prob[idx][0]
    ax.set_title(f"{category}\nProb: {prob:.2f}")
    ax.axis('off')
  categories = ['True Positive', 'True Negative', 'False Positive', 'False Negative']
  index_lists = [tp_indices, tn_indices, fp_indices, fn_indices]
  for i, (category, indices) in enumerate(zip(categories, index_lists)):
    count = min(len(indices), num_samples)
    for j in range(count):
      plot_sample(axes[i, j], indices[j], category)
    for j in range(count, num_samples):
      axes[i, j].axis('off')
      axes[i, j].set_title(f"{category}\nN/A")
  plt.tight_layout()
  plt.savefig(os.path.join(save_dir, 'prediction_summary.png'))
  plt.close()
def generate_feature_maps(model, sample, layer_names=None, save_dir=None):
  if not save_dir:
    save_dir = '3d_res/feature_maps'
  os.makedirs(save_dir, exist_ok=True)
  if not layer_names:
    layer_names = [layer.name for layer in model.layers if 'conv' in layer.name.lower()]
  feature_models = []
```

```
for layer_name in layer_names:
    feature_model = tf.keras.Model(inputs=model.input,
outputs=model.get_layer(layer_name).output)
    feature_models.append((layer_name, feature_model))
  if len(sample.shape) == 4:
    sample = np.expand_dims(sample, axis=0)
  for layer_name, feature_model in feature_models:
    print(f"Generating feature maps for layer: {layer_name}")
    feature_maps = feature_model.predict(sample)
    if len(feature_maps.shape) == 5:
      n_features = feature_maps.shape[-1]
      middle_slice = feature_maps.shape[1] // 2
      grid_size = int(np.ceil(np.sqrt(min(n_features, 16))))
      fig, axes = plt.subplots(grid_size, grid_size, figsize=(12, 12))
      fig.suptitle(f"Feature Maps - Layer: {layer_name}", fontsize=16)
      feature_idx = 0
      for i in range(grid_size):
         for j in range(grid_size):
           if feature idx < min(n features, 16):
             ax = axes[i, j] if grid_size > 1 else axes
             im = ax.imshow(feature_maps[0, middle_slice, :, :, feature_idx], cmap='viridis')
             ax.set_title(f"Feature {feature_idx+1}")
             ax.axis('off')
             feature idx += 1
           else:
             if grid_size > 1:
               axes[i, j].axis('off')
             else:
               axes.axis('off')
      plt.tight_layout()
      plt.savefig(os.path.join(save_dir, f"feature_maps_{layer_name}.png"))
```

```
plt.close()
  print(f"Feature maps saved to {save_dir}")
# Pipeline
def run_pipeline(base_dir, output_dir='3d_res'):
  print("="*80)
  print("LUNG CANCER DETECTION PIPELINE")
  print("="*80)
  # Create output directory
  os.makedirs(output_dir, exist_ok=True)
  # Step 1: Load preprocessed data
  data_dir = '.' # Try working directory first
  if not all(os.path.exists(os.path.join(data_dir, f)) for f in
        ['X_train.npy', 'X_val.npy', 'X_test.npy', 'y_train.npy', 'y_val.npy', 'y_test.npy']):
    data_dir = base_dir # Fall back to luna16_dataset
  print(f"Loading preprocessed data from {data_dir}...")
  try:
    X_train = np.load(os.path.join(data_dir, 'X_train.npy'))
    X_val = np.load(os.path.join(data_dir, 'X_val.npy'))
    X_test = np.load(os.path.join(data_dir, 'X_test.npy'))
    y_train = np.load(os.path.join(data_dir, 'y_train.npy'))
    y_val = np.load(os.path.join(data_dir, 'y_val.npy'))
    y_test = np.load(os.path.join(data_dir, 'y_test.npy'))
  except FileNotFoundError as e:
    raise FileNotFoundError(f"Preprocessed data not found in {data_dir}. Ensure X_train.npy, etc.,
are present: {str(e)}")
  # Add channel dimension if needed
  if len(X_train.shape) == 4:
```

```
X_train = np.expand_dims(X_train, axis=-1)
  X_val = np.expand_dims(X_val, axis=-1)
  X_test = np.expand_dims(X_test, axis=-1)
print(f"Data shapes:")
print(f" Training: {X_train.shape}, {y_train.shape}")
print(f" Validation: {X_val.shape}, {y_val.shape}")
print(f" Testing: {X_test.shape}, {y_test.shape}")
# Step 2: Load model
print("\nLoading model...")
try:
  model = tf.keras.models.load_model(
    os.path.join(data_dir, 'best_model.h5'),
    custom_objects={'focal_loss_fixed': focal_loss(gamma=2.0, alpha=0.25)}
  )
except FileNotFoundError:
  raise FileNotFoundError(f"Model file 'best_model.h5' not found in {data_dir}.")
# Step 3: Evaluate the model (skipped)
print("\nEvaluation already completed, proceeding to visualization...")
# Step 4: Analyze false negatives (skipped)
print("\nFalse negative analysis already completed, proceeding to visualization...")
# Step 5: Visualize Predictions
print("\nVisualizing model predictions...")
vis_dir = output_dir
# Get predictions
y_pred_prob = model.predict(X_test)
```

```
# Get indices for TP, TN, FP, FN
tp_indices = np.where((y_test == 1) & (y_pred_prob.flatten() > 0.5))[0]
tn_indices = np.where((y_test == 0) & (y_pred_prob.flatten() <= 0.5))[0]
fp_indices = np.where((y_test == 0) & (y_pred_prob.flatten() > 0.5))[0]
fn_indices = np.where((y_test == 1) & (y_pred_prob.flatten() <= 0.5))[0]
# 1. Visualize 3D volumes (animations for 2 positive and 2 negative samples)
print("Generating 3D volume animations...")
for idx in tp_indices[:2]:
  visualize_3d_volume(
    X_test[idx],
    title=f"Positive Sample (Nodule) - Index {idx}",
    save_path=os.path.join(vis_dir, 'true_positive', f"volume_positive_{idx}.gif")
  )
for idx in tn_indices[:2]:
  visualize_3d_volume(
    X_test[idx],
    title=f"Negative Sample (No Nodule) - Index {idx}",
    save_path=os.path.join(vis_dir, 'true_negative', f"volume_negative_{idx}.gif")
  )
# 2. Visualize prediction comparisons
print("Generating prediction comparison plots...")
categories = ['true_positive', 'true_negative', 'false_positive', 'false_negative']
index_lists = [tp_indices, tn_indices, fp_indices, fn_indices]
for category, indices in zip(categories, index_lists):
  count = min(len(indices), 3)
  for j in range(count):
    idx = indices[j]
    visualize_prediction_comparison(
```

```
X_test[idx],
      y_test[idx],
      y_pred_prob[idx][0],
      save_dir=os.path.join(vis_dir, category)
    )
# 3. Visualize model predictions (summary grid)
print("Generating prediction summary grid...")
visualize_model_predictions(
  model=model,
  X_test=X_test,
  y_test=y_test,
  num_samples=5,
  save_dir=vis_dir
)
# 4. Generate feature maps (for one positive and one negative sample)
print("Generating feature maps...")
if tp_indices.size > 0:
  generate_feature_maps(
    model,
    X_test[tp_indices[0]],
    save_dir=os.path.join(vis_dir, 'feature_maps_positive')
  )
if tn_indices.size > 0:
  generate_feature_maps(
    model,
    X_test[tn_indices[0]],
    save_dir=os.path.join(vis_dir, 'feature_maps_negative')
  )
```

```
print(f"Visualizations saved in {vis_dir}")

print("\nPipeline completed successfully!")

print(f"Results saved to {output_dir}")

return model, {}

if __name__ == "__main__":

base_dir = '/workspace/deep_learning_luna16/luna16_dataset'

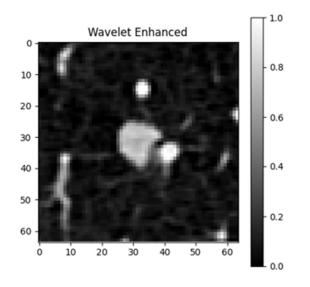
model, metrics = run_pipeline(base_dir)
```

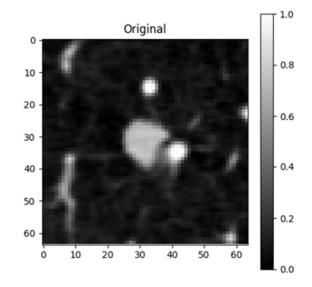
#### **OUTPUT OF BLOCK 3**

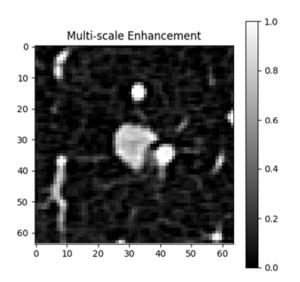
```
LUNG CANCER DETECTION PIPELINE
Loading preprocessed data from ....
Data shapes:
 Training: (2527, 64, 64, 64, 1), (2527,)
 Validation: (542, 64, 64, 64, 1), (542,)
 Testing: (542, 64, 64, 64, 1), (542,)
Loading model...
Evaluation already completed, proceeding to visualization...
False negative analysis already completed, proceeding to visualization...
Visualizing model predictions...
                                 ==] - 2s 92ms/step
17/17 [=====
Generating 3D volume animations...
Generating prediction comparison plots...
Generating prediction summary grid...
17/17 [===============] - 1s 75ms/step
Generating feature maps...
Generating feature maps for layer: conv3d_7
1/1 [-----] - 0s 159ms/step
Generating feature maps for layer: conv3d_8
Visualizations saved in 3d_res
Pipeline completed successfully!
Results saved to 3d res
```

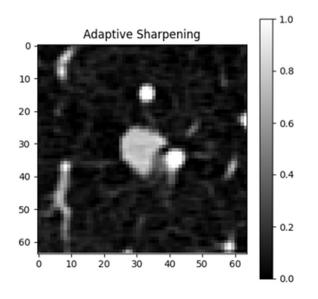
# OUTPUTS(RESULTS)

# • Enhancement:

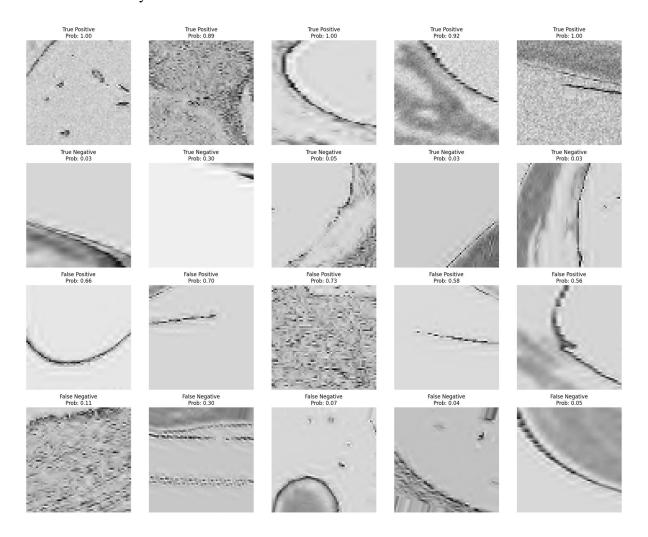




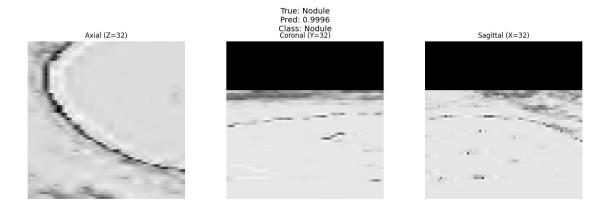




# • Prediction Summary

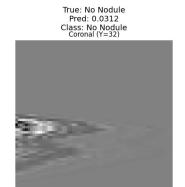


# • True positive sample:



# • True Negative sample:

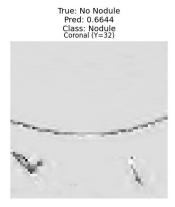


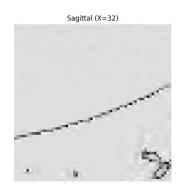




#### • False Positive







# • False Negative:

