## data loading

```
import os
import shutil
import random
import numpy as np
import pandas as pd
import cv2
import matplotlib.pyplot as plt
from google.colab import drive
# Mount Google Drive if in Colab
try:
   drive.mount('/content/drive')
   print("Google Drive mounted successfully!")
   print("Running locally or Drive already mounted")
# Set paths
# Adjust these paths based on where you unzipped the LGG dataset
lgg_dataset_path = '/content/drive/MyDrive/ddd' # Path to the original LGG dataset
output_dir = '/content/drive/MyDrive/brain_tumor_dataset' # Where to save the organized dataset
# Create output directories
os.makedirs(os.path.join(output_dir, 'images'), exist_ok=True)
os.makedirs(os.path.join(output_dir, 'masks'), exist_ok=True)
# Function to extract and prepare dataset
def prepare_dataset(num_samples=10, random_selection=True):
   Prepare a subset of the LGG dataset for segmentation
   Parameters:
    -----
   num_samples : int
       Number of image-mask pairs to extract
   random\_selection : bool
       Whether to select random samples or the first ones
   # Get all case directories
   case_dirs = [d for d in os.listdir(lgg_dataset_path)
               if os.path.isdir(os.path.join(lgg_dataset_path, d))]
   print(f"Found {len(case_dirs)} case directories")
   if random_selection:
       # Randomly select cases
        selected_cases = random.sample(case_dirs, min(len(case_dirs), num_samples))
   else:
       # Take the first N cases
       selected cases = case dirs[:min(len(case dirs), num samples)]
   print(f"Selected {len(selected_cases)} cases")
   # Counter for processed images
   processed_count = 0
   # Process each selected case
   for case_id in selected_cases:
       case_dir = os.path.join(lgg_dataset_path, case_id)
       # Get all files in this case directory
        files = os.listdir(case_dir)
        # Get image files (without _mask suffix)
        image_files = [f for f in files if f.endswith('.tif') and '_mask' not in f]
        for img_file in image_files:
            # Get corresponding mask file
           mask_file = img_file.replace('.tif', '_mask.tif')
            if mask file in files:
               # Full paths
               img_path = os.path.join(case_dir, img_file)
                mask_path = os.path.join(case_dir, mask_file)
```

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# Read images
                img = cv2.imread(img_path)
                mask = cv2.imread(mask_path, cv2.IMREAD_GRAYSCALE)
                if img is not None and mask is not None:
                    # Convert BGR to RGB
                    img = cv2.cvtColor(img, cv2.COLOR_BGR2RGB)
                    # Take only the FLAIR channel (channel 1) from the original image
                    # In LGG dataset, FLAIR is the 2nd channel (index 1)
                    flair = img[:, :, 1]
                    # Output filenames
                    out_img_name = f"brain_tumor_{processed_count:03d}.png"
                    out_mask_name = f"brain_tumor_{processed_count:03d}_mask.png"
                   # Save paths
                   out_img_path = os.path.join(output_dir, 'images', out_img_name)
                    out_mask_path = os.path.join(output_dir, 'masks', out_mask_name)
                    # Save images
                    cv2.imwrite(out_img_path, flair)
                    cv2.imwrite(out mask path, mask)
                    processed_count += 1
                    print(f"Processed {processed_count}/{num_samples}: {out_img_name}")
                    if processed_count >= num_samples:
                        return
# Function to visualize the prepared dataset
def visualize_dataset(dataset_path, num_samples=5):
   Visualize the prepared dataset
   Parameters:
   dataset path : str
       Path to the prepared dataset
   num samples : int
       Number of samples to visualize
   images_dir = os.path.join(dataset_path, 'images')
   masks_dir = os.path.join(dataset_path, 'masks')
   # Get all image files
   image_files = sorted(os.listdir(images_dir))
   # Limit to the requested number of samples
   image_files = image_files[:min(len(image_files), num_samples)]
   # Create figure
   plt.figure(figsize=(12, 4 * len(image_files)))
   for i, img_file in enumerate(image_files):
        # Get corresponding mask file - handle different naming conventions
       if img_file.replace('.png', '_mask.png') in os.listdir(masks_dir):
           mask_file = img_file.replace('.png', '_mask.png')
        else:
           # Try alternative mask naming if needed
           mask_file = next((m for m in os.listdir(masks_dir) if m.startswith(img_file.split('.')[0])), None)
        if mask_file:
           # Read images
            img_path = os.path.join(images_dir, img_file)
           mask_path = os.path.join(masks_dir, mask_file)
            img = cv2.imread(img path, cv2.IMREAD GRAYSCALE) # Read as grayscale
           mask = cv2.imread(mask_path, cv2.IMREAD_GRAYSCALE)
            # Display image
           plt.subplot(len(image_files), 2, i * 2 + 1)
           plt.imshow(img, cmap='gray')
            plt.title(f"MRI Image: {img_file}")
           plt.axis('off')
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# птэһтай шаэк
             plt.subplot(len(image_files), 2, i * 2 + 2)
             plt.imshow(mask, cmap='gray')
             plt.title(f"Tumor Mask: {mask_file}")
             plt.axis('off')
    plt.tight_layout()
    plt.show()
# Execute dataset preparation
print("Preparing dataset...")
prepare_dataset(num_samples=10, random_selection=True)
print("Dataset preparation completed!")
# Visualize the prepared dataset
print("Visualizing prepared dataset...")
visualize_dataset(output_dir, num_samples=5)
# Print instructions for using the dataset with the segmentation code
print("\nDataset is ready to use with the brain tumor segmentation code!")
print(f"Images directory: {os.path.join(output_dir, 'images')}")
print(f"Masks directory: {os.path.join(output_dir, 'masks')}")
print("\nUpdate these paths in the main code:")
print("base_dir = '", output_dir, "'")
print("images_dir = os.path.join(base_dir, 'images')")
print("masks_dir = os.path.join(base_dir, 'masks')")
```

```
import os
import cv2
import numpy as np
import matplotlib.pyplot as plt
from google.colab import drive
from sklearn.model_selection import train_test_split
from \ sklearn. metrics \ import \ accuracy\_score, \ jaccard\_score, \ f1\_score, \ precision\_score, \ recall\_score
# Mount Google Drive (for Google Colab)
try:
    drive.mount('/content/drive')
    print("Google Drive mounted successfully!")
except:
    print("Running locally or Drive already mounted")
class BrainTumorSegmentation:
    def __init__(self, base_dir=None):
        Initialize the Brain Tumor Segmentation class
        base_dir : str
            Base directory containing the dataset
        self.base_dir = base_dir
        self.images = []
        self.masks = []
        self.processed_images = []
        self.segmented_masks = []
        self.metrics = {}
    def load_dataset(self, images_dir, masks_dir, max_samples=None):
        Load dataset images and their corresponding masks
        Parameters:
        images_dir : str
            Directory containing MRI scan images
        masks_dir : str
            Directory containing corresponding mask images
        max_samples : int, optional
            Maximum number of samples to load
        print("Loading dataset...")
        # List all files in the directories
        image_files = sorted(os.listdir(images_dir))
        # For LGG dataset, find corresponding mask files
        loaded count = 0
        for img_file in image_files:
            if not img_file.endswith(('.jpg', '.png', '.jpeg', '.tif')):
                continue
            # Construct mask filename based on LGG naming pattern
            if '_mask' not in img_file:
                mask_file = img_file.replace('.png', '_mask.png')
mask_file = mask_file.replace('.tif', '_mask.tif')
                mask_file = mask_file.replace('.jpg', '_mask.jpg')
                image_path = os.path.join(images_dir, img_file)
                mask_path = os.path.join(masks_dir, mask_file)
                 # Check if mask file exists
                 if not os.path.exists(mask_path):
                     print(f"Warning: No mask found for {img_file}")
                     continue
                 # Read image and mask
                 image = cv2.imread(image_path)
                mask = cv2.imread(mask_path, cv2.IMREAD_GRAYSCALE)
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if image is not None and mask is not None:
                # Handle grayscale vs color images
                if len(image.shape) == 3 and image.shape[2] == 3:
                    # For LGG dataset: take FLAIR channel (usually the most informative)
                    # FLAIR is typically in the middle channel (index 1)
                    gray_image = image[:, :, 1] # Extract FLAIR channel
                else:
                    gray_image = image.copy()
                # Convert to single channel and normalize
                if len(gray_image.shape) == 3:
                    gray_image = cv2.cvtColor(gray_image, cv2.COLOR_BGR2GRAY)
                # Normalize mask to binary (0 and 255)
                _, binary_mask = cv2.threshold(mask, 127, 255, cv2.THRESH_BINARY)
                # Add to the dataset
                self.images.append(gray_image)
                self.masks.append(binary_mask)
                loaded count += 1
                if max_samples is not None and loaded_count >= max_samples:
                    break
    print(f"Loaded {len(self.images)} images and {len(self.masks)} masks.")
def preprocess_images(self):
    """Preprocess the loaded MRI images"""
    print("Preprocessing images...")
    self.processed_images = []
    for image in self.images:
        # Ensure image is grayscale
        if len(image.shape) > 2:
            gray_image = cv2.cvtColor(image, cv2.COLOR_RGB2GRAY)
            gray_image = image.copy()
        # Normalize pixel values to range [0, 255]
        if gray_image.max() > 0:
            normalized = ((gray_image - gray_image.min()) /
                         (gray_image.max() - gray_image.min()) * 255).astype(np.uint8)
        else:
            normalized = gray_image
       # Apply CLAHE (Contrast Limited Adaptive Histogram Equalization)
        clahe = cv2.createCLAHE(clipLimit=2.0, tileGridSize=(8, 8))
       enhanced = clahe.apply(normalized)
        # Apply Gaussian blur to reduce noise
       blurred = cv2.GaussianBlur(enhanced, (5, 5), 0)
        # Store preprocessed image
        self.processed images.append(blurred)
    print(f"Preprocessed {len(self.processed_images)} images.")
def segment_tumors(self, method='watershed'):
    Segment tumor regions from preprocessed images
    Parameters:
    method : str
       Segmentation method to use ('threshold', 'watershed', 'kmeans')
    print(f"Segmenting tumors using {method} method...")
    self.segmented_masks = []
    for image in self.processed_images:
        if method == 'threshold':
           # Apply Otsu's thresholding
            _, segmented = cv2.threshold(image, 0, 255, cv2.THRESH_BINARY + cv2.THRESH_OTSU)
        elif method == 'watershed':
            # Watershed algorithm
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# Find sure background
            _, thresholded = cv2.threshold(image, 0, 255, cv2.THRESH_BINARY + cv2.THRESH_OTSU)
            # Noise removal with morphological operations
            kernel = np.ones((3, 3), np.uint8)
            opening = cv2.morphologyEx(thresholded, cv2.MORPH_OPEN, kernel, iterations=2)
            # Sure background area
            sure_bg = cv2.dilate(opening, kernel, iterations=3)
            # Finding sure foreground area
            dist_transform = cv2.distanceTransform(opening, cv2.DIST_L2, 5)
            _, sure_fg = cv2.threshold(dist_transform, 0.7*dist_transform.max(), 255, 0)
            # Finding unknown region
            sure_fg = np.uint8(sure_fg)
            unknown = cv2.subtract(sure_bg, sure_fg)
            # Marker labelling
            _, markers = cv2.connectedComponents(sure_fg)
            # Add one to all labels so that background is not 0, but 1
            markers = markers + 1
            # Mark the unknown region with 0
            markers[unknown == 255] = 0
            # Apply watershed
            markers = cv2.watershed(cv2.cvtColor(image, cv2.COLOR_GRAY2BGR), markers)
            segmented = np.zeros_like(image)
            segmented[markers > 1] = 255
        elif method == 'kmeans':
            # Apply K-means clustering
            image_data = image.reshape((-1, 1))
            image_data = np.float32(image_data)
            # Define criteria and apply kmeans
            criteria = (cv2.TERM_CRITERIA_EPS + cv2.TERM_CRITERIA_MAX_ITER, 100, 0.2)
            k = 3 # Number of clusters
            _, labels, centers = cv2.kmeans(image_data, k, None, criteria, 10, cv2.KMEANS_RANDOM_CENTERS
            # Find the cluster that corresponds to the tumor (usually the brightest)
            centers = np.uint8(centers)
            brightest_cluster = np.argmax(centers)
            # Create mask based on the brightest cluster
            segmented = np.zeros_like(image)
            segmented[labels.reshape(image.shape) == brightest_cluster] = 255
        else:
            raise ValueError(f"Unknown segmentation method: {method}")
        # Post-processing: fill holes and remove small objects
        segmented = self.post process mask(segmented)
        self.segmented_masks.append(segmented)
    print(f"Segmented {len(self.segmented_masks)} images.")
def post_process_mask(self, mask):
   Apply post-processing to improve the segmentation mask
   Parameters:
    mask : numpy.ndarray
       Binary segmentation mask
   Returns:
    numpy.ndarray
       Post-processed binary mask
   # Convert to binary
    if mask.dtype != np.uint8:
       mask = mask.astype(np.uint8)
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# Fill holes
    contours, _ = cv2.findContours(mask, cv2.RETR_EXTERNAL, cv2.CHAIN_APPROX SIMPLE)
    filled_mask = np.zeros_like(mask)
    cv2.drawContours(filled mask, contours, -1, 255, -1)
    # Remove small objects (noise)
    nb_components, output, stats, _ = cv2.connectedComponentsWithStats(filled_mask, connectivity=8)
    sizes = stats[1:, -1]
    min_size = 100  # Minimum size of objects to keep
    # Keep only components with size greater than min_size
    processed_mask = np.zeros_like(filled_mask)
    for i in range(1, nb_components):
       if sizes[i - 1] >= min_size:
           processed_mask[output == i] = 255
    return processed mask
def evaluate_segmentation(self, ground_truth_masks=None):
    Evaluate segmentation performance against ground truth masks
    Parameters:
    ground truth masks : list
       List of ground truth mask images
    -----
    dict
       Dictionary containing evaluation metrics
    if ground_truth_masks is None:
        ground_truth_masks = self.masks
    if len(ground_truth_masks) != len(self.segmented_masks):
       raise ValueError("Mismatch between number of ground truth masks and segmented masks")
    # Initialize metrics
    dice scores = []
    jaccard_scores = []
    precision_scores = []
    recall_scores = []
    for gt_mask, pred_mask in zip(ground_truth_masks, self.segmented_masks):
        # Binarize masks
        gt_binary = np.where(gt_mask > 0, 1, 0).flatten()
        pred_binary = np.where(pred_mask > 0, 1, 0).flatten()
        # Calculate Dice coefficient (F1 score)
       dice = f1_score(gt_binary, pred_binary, zero_division=1)
       dice_scores.append(dice)
        # Calculate Jaccard index (IoU)
        iou = jaccard score(gt binary, pred binary, zero division=1)
        jaccard_scores.append(iou)
        # Calculate precision and recall
       precision = precision_score(gt_binary, pred_binary, zero_division=1)
       recall = recall_score(gt_binary, pred_binary, zero_division=1)
       precision_scores.append(precision)
        recall_scores.append(recall)
    # Calculate average metrics
    self.metrics = {
        'dice_coefficient': np.mean(dice_scores),
        'jaccard_index': np.mean(jaccard_scores),
        'precision': np.mean(precision_scores),
        'recall': np.mean(recall_scores)
   }
    print("Segmentation Evaluation Metrics:")
    print(f" Dice Coefficient (F1-Score): {self.metrics['dice_coefficient']:.4f}")
   print(f" Jaccard Index (IoU): {self.metrics['jaccard_index']:.4f}")
    print(f" Precision: {self.metrics['precision']:.4f}")
   print(f" Recall: {self.metrics['recall']:.4f}")
```

```
return self.metrics
def visualize_results(self, num_samples=5):
    Visualize original images, ground truth masks, and segmented masks
   num samples : int
       Number of samples to visualize
    num_samples = min(num_samples, len(self.images))
    plt.figure(figsize=(15, 4 * num_samples))
    for i in range(num_samples):
        # Original image
       plt.subplot(num_samples, 3, i * 3 + 1)
       plt.imshow(self.images[i], cmap='gray')
       plt.title(f"Original Image {i+1}")
       plt.axis('off')
       # Ground truth mask
       plt.subplot(num samples, 3, i * 3 + 2)
       plt.imshow(self.masks[i], cmap='gray')
       plt.title(f"Ground Truth Mask {i+1}")
       plt.axis('off')
       # Segmented mask
       plt.subplot(num_samples, 3, i * 3 + 3)
       plt.imshow(self.segmented_masks[i], cmap='gray')
       plt.title(f"Segmented Mask {i+1}")
       plt.axis('off')
    plt.tight_layout()
   plt.show()
def overlay_results(self, num_samples=5):
   Overlay segmentation results on original images for visualization
    Parameters:
    num_samples : int
       Number of samples to visualize
    num_samples = min(num_samples, len(self.images))
    plt.figure(figsize=(12, 4 * num_samples))
    for i in range(num_samples):
        # Convert image to RGB for overlay
       display_img = cv2.cvtColor(self.images[i], cv2.COLOR_GRAY2RGB)
       # Original image with ground truth overlay
       plt.subplot(num_samples, 2, i * 2 + 1)
       # Create overlay with ground truth mask (green)
       overlay = display_img.copy()
        # Create green mask
        green_mask = np.zeros_like(overlay)
       green_mask[:, :, 1] = self.masks[i] # Green channel
        # Apply mask
       alpha = 0.5
        cv2.addWeighted(green_mask, alpha, overlay, 1 - alpha, 0, overlay)
       plt.imshow(overlay)
       plt.title(f"Original + Ground Truth {i+1}")
       plt.axis('off')
        # Original image with segmentation overlay
       plt.subplot(num_samples, 2, i * 2 + 2)
       # Create overlay with segmented mask (red)
```

```
overlay = display_img.copy()
           # Create red mask
           red_mask = np.zeros_like(overlay)
           red_mask[:, :, 0] = self.segmented_masks[i] # Red channel
           # Apply mask
           cv2.addWeighted(red_mask, alpha, overlay, 1 - alpha, 0, overlay)
           plt.imshow(overlay)
           plt.title(f"Original + Segmentation {i+1}")
           plt.axis('off')
        plt.tight_layout()
       plt.show()
   def run_full_pipeline(self, images_dir, masks_dir, max_samples=None, segmentation_method='watershed'):
        Run the full segmentation pipeline
       Parameters:
        images_dir : str
           Directory containing MRI scan images
        masks dir : str
           Directory containing corresponding mask images
        max_samples : int, optional
           Maximum number of samples to load
        segmentation_method : str
       Method to use for segmentation ('threshold', 'watershed', 'kmeans')
        # Load dataset
        self.load_dataset(images_dir, masks_dir, max_samples)
        # Preprocess images
        self.preprocess_images()
        # Segment tumors
        self.segment_tumors(method=segmentation_method)
        # Evaluate segmentation
        self.evaluate_segmentation()
        # Visualize results
        self.visualize_results()
        self.overlay_results()
        return self.metrics
# Execute the segmentation pipeline
if __name__ == "__main__":
   # Update these paths to your LGG dataset location
   base_dir = '/content/drive/MyDrive/brain_tumor_dataset'
   images_dir = os.path.join(base_dir, 'images')
   masks dir = os.path.join(base dir, 'masks')
   # Create segmentation object
   tumor_segmentation = BrainTumorSegmentation(base_dir)
   # Run full pipeline with all available segmentation methods
   methods = ['threshold', 'watershed', 'kmeans']
   results = {}
    for method in methods:
       print(f"\n{'-'*50}")
       print(f"Running segmentation with {method.upper()} method")
       print(f"{'-'*50}")
       metrics = tumor_segmentation.run_full_pipeline(
           images_dir=images_dir,
           masks_dir=masks_dir,
           max_samples=10, # Use 10 images as requested
           segmentation method=method
       )
        results[method] = metrics
```

# Compare results
print("\nComparison of Segmentation Methods:")

