# TP seg MEDIMA 204 STUDENTS

January 18, 2024

# 1 IMA204 Practical Session - Segmentation of medical images

### 2 Introduction

Images to use are provided in a zip file called **data.zip** to install in your local directory from where you are running this notebook.

The goal of this practical session is to push K-means segmentation method as a pre-segmentation tool on different types of medical images: CT scans for the segmentation of kidneys and tumors, MRI for the corpus callosum in the brain , temporal sequences of MRI images for segmentation of the myocardium.

You are provided with pre-processing ideas. You will have to adjust a pipeline for **AT LEAST** one application (kidney/tumor, corpus callosum or heart).

You have to submit your code and comment your results.

**Deadline**: You will have to upload a single jupyter notebook .ipynb with your answers (code + text) before the deadline (please check on Ecampus/Moodle).

The uploaded file should be named 'TP\_SegMedImage\_YOURSURNAME.ipynb'.

# RUN THE WHOLE NOTEBOOK FIRST TO GET A FULL OVERVIEW OF USE-CASES AND THE NOTION OF HYPERPARAMETERS.

```
[]: import os
  import numpy as np
  import matplotlib.pyplot as plt
  from matplotlib.colors import ListedColormap, LinearSegmentedColormap

# For ploting utilities of contours on images
  from mpl_toolkits.axes_grid1 import AxesGrid
  from mpl_toolkits.axes_grid1 import make_axes_locatable

# For listing files in local foler
  import glob

# For loading .mat files
  from scipy.io import loadmat

import skimage
```

```
from skimage.io import imread
from skimage import morphology
from skimage.segmentation import watershed
from skimage.filters import rank
from skimage.util import img_as_ubyte
from skimage.morphology import disk
from scipy import ndimage
from skimage.measure import find_contours
import skimage.morphology as morpho
from skimage import data
from skimage import color
from skimage import morphology
from skimage import segmentation
from skimage.filters import gaussian
# For Kmeans
import cv2
Working_directory = os.getcwd()
print('Working_directory = ',Working_directory)
```

```
[]: def my_kmeans(image,k):
         \#k = number of clusters
         # Reshaping the image
        pixel_vals = image.reshape((-1,1))
        # Convert to float type only for supporting cv2.kmean
        pixel vals = np.float32(pixel vals)
        criteria = (cv2.TERM CRITERIA EPS + cv2.TERM CRITERIA MAX ITER, 100, 0.
      ⇔85) #criteria
        retval, labels, centers = cv2.kmeans(pixel_vals, k,
                     None, criteria, 10, cv2.KMEANS_PP_CENTERS)
         centers = np.uint8(centers) # convert data into 8-bit values
         segmented_data
                        = centers[labels.flatten()] # Mapping labels to center_
      ⇔points( RGB Value)
         segmented image = segmented data.reshape((image.shape)) # reshape data__
      ⇒into the original image dimensions
         segmented_labels = labels # Mapping labels to center points( RGB Value)
         segmented_labels = segmented_labels.reshape((image.shape)) # reshape data__
      ⇒into the original image dimensions
        return segmented_image,segmented_labels
```

```
def my_colormap_white_bkg(Colormap_name,numLabels):
    #numLabels = number of colors

Colormap = plt.get_cmap(Colormap_name, numLabels)
    newcolors = Colormap(np.linspace(0, 1, numLabels))
    bkg_color = np.array([256/256, 256/256, 256/256, 1])
    max_color = np.array([0/256, 0/256, 0/256, 1])
    newcolors[numLabels-1, :] = max_color
    newcolors[0, :] = bkg_color
    newcmp = ListedColormap(newcolors)
    return newcmp
```

#### 2.1 Abdominal CT

You have at your disposal **6 abdominal CT scans** of different subjects. Subjects may have renal tumor. You also have the manual segmentations for both kidney and, when present, the tumor.

Variables defined: \* abdominalCT\_path \* listImagesabdCT \* Img\_abdo\_ex \* Seg abdo kidney ex, Seg abdo tumor ex

```
[]: abdominalCT_path = Working_directory + '/data/abdominalCT'
     os.listdir(abdominalCT_path)
     listImagesabdCT=glob.glob(abdominalCT_path + '/*-seg.tiff')
     print('There are', len(listImagesabdCT), 'abdomical CT images')
     # Choose a figure and plot it with the ground truth segmentation
     indexIm=0 # between 0 and 5
     # Abdominal CT
     filename_Segmentation = listImagesabdCT[indexIm]
     Labels_abdo_ex = imread(filename_Segmentation)
                      = filename_Segmentation[:-9] + '.tiff'
     filename
     Img_abdo_ex
                      = imread(filename)
     print('Reading image ', filename)
     print(np.unique(Labels_abdo_ex))
     if Img_abdo_ex.shape != Labels_abdo_ex.shape:
       raise NameError('image and mask should have the same shape, problem...')
     # In Labels_abdo_ex we may have two values: 127 is for kidney and 255 for renal_
      \hookrightarrow tumor
     Seg_abdo_kidney_ex=Labels_abdo_ex==127
     if np.sum(Seg_abdo_kidney_ex)==0:
       print('There is no kidney')
     Cont_abdo_kidney_ex = find_contours(Seg_abdo_kidney_ex, 0.5)
```

```
Seg_abdo_tumor_ex=Labels_abdo_ex==255
if np.sum(Seg_abdo_tumor_ex)==0:
  print('There is no tumor')
Cont_abdo_tumor_ex = find_contours(Seg_abdo_tumor_ex, 0.5)
fig = plt.figure(figsize=(17, 7))
grid = AxesGrid(fig, 111,
                nrows ncols = (2, 3),
                axes_pad = 0.5)
grid[0].imshow(Img abdo ex, cmap='gray')
grid[0].axis('off')
grid[0].set_title("Original image")
grid[1].imshow(Seg_abdo_kidney_ex,cmap='gray')
grid[1].axis('off')
grid[1].set_title("Segmentation Mask Kidney")
grid[2].imshow(Seg_abdo_tumor_ex,cmap='gray')
grid[2].axis('off')
grid[2].set_title("Segmentation Mask Tumor")
grid[3].imshow(Img_abdo_ex, cmap='gray')
for contour in Cont_abdo_kidney_ex:
  grid[3].plot(contour[:, 1], contour[:, 0], linewidth=2, c='r')
grid[3].axis('off')
grid[3].set title("Image + contour kidney")
grid[4].imshow(Img_abdo_ex, cmap='gray')
for contour in Cont abdo tumor ex:
  grid[4].plot(contour[:, 1], contour[:, 0], linewidth=2, c='r')
grid[4].axis('off')
grid[4].set_title("Image +contour tumor")
```

### 3 Brain MRI

Here you can select medial slices of the brain of **4 different subjects**. You also have manual segmentations of the corpus callosum.

Variables defined: \* brainMRI\_path \* listImagesbrainMRI \* Seg\_brain\_ex \* Img\_brain\_ex

```
[]: brainMRI_path = Working_directory + '/data/brainMRI'
    os.listdir(brainMRI_path)
    listImagesbrainMRI=glob.glob(brainMRI_path + '/*-seg.png')
    print('There are', len(listImagesbrainMRI), 'brain MRI images')
    print(listImagesbrainMRI)

# Choose a brain MRI and plot it with the ground truth segmentation
    indexIm = 3 # between 0 and 3
    filename_seg = listImagesbrainMRI[indexIm]
    Seg_brain_ex = imread(filename_seg)
```

```
filename
            = filename_seg[:-8] + '.png'
Img brain ex = imread(filename)
print('Reading image ', filename)
if Img_brain_ex.shape != Seg_brain_ex.shape:
  raise NameError('image and mask should have the same shape, problem...')
# In Im Seg we have masks of the corpus callosum
            = Seg_brain_ex==255
contourMask = find contours(maskCC, 0.5)
fig = plt.figure(figsize=(17, 7))
grid = AxesGrid(fig, 111,
                nrows_ncols = (1, 3),
                axes_pad = 0.5)
grid[0].imshow(Img_brain_ex, cmap='gray')
grid[0].axis('off')
grid[0].set_title("Original image")
grid[1].imshow(maskCC,cmap='gray')
grid[1].axis('off')
grid[1].set_title("Segmentation Mask\n Corpus Callosum");
grid[2].imshow(Img_brain_ex, cmap='gray')
for contour in contourMask:
  grid[2].plot(contour[:, 1], contour[:, 0], linewidth=2, c='r')
grid[2].axis('off')
grid[2].set_title("Image with segmentation\n corpus callosum")
```

### 4 Cardiac MRI

The last section is about MRI sequences of the heart. You are provided with a **single use case**. Your goal is to segment the left ventricule. Be careful, the segmentation is not a make but a series of points (landmarks). To obtain a binary mask, you should first interpolate the points (using for instance a spline).

The structure of this image data is more complex and needs some coding to load single slices to segment.

Variables defined: \* MRIheart path \* Img cardiac ex \* Seg cardiac ex

```
[]: # Read one case
MRIheart_path = Working_directory + '/data/MRIheart/'
    os.listdir(MRIheart_path)
    data = loadmat(MRIheart_path + 'dataMRIheart.mat')
    data = data['data']
    seg = loadmat(MRIheart_path + 'segMRIheart.mat')
    seg = seg['seg']
```

```
Ex_index_select = 6
Img_cardiac_ex = data[:,:,Ex_index_select,1] # can be index 4,5,6,...
Cont_cardiac_ex = seg[Ex_index_select,1][:]
print('MRI volume of the heart composed of', data.shape[2], 'slices along the z_{\sqcup}
 ⇔axis and', data.shape[3],
'temporal frames. Each slice is an image ', data.shape[0], ' x ', data.
 \hookrightarrowshape[1])
print('For each slice and at each time frame we have a manual segmentation ⊔
 print('Be careful, some slices do not contain the left ventricle myocardium and \Box

→the manual segmentation is not simply empty but it contains the value:',

□

 \hookrightarrowseg[0,0])
plt.figure(figsize=(20, 9))
plt.suptitle('Sample slices at a single time frame with the red manual ⊔
 ⇔segmentation at the bottom')
plt.subplot(2, 5, 1)
plt.imshow(data[:,:,4,1],cmap="gray")
plt.gca().invert_yaxis()
plt.subplot(2, 5, 2)
plt.imshow(data[:,:,5,1],cmap="gray")
plt.gca().invert_yaxis()
plt.subplot(2, 5, 3)
plt.imshow(data[:,:,6,1],cmap="gray")
plt.gca().invert_yaxis()
plt.subplot(2, 5, 4)
plt.imshow(data[:,:,7,1],cmap="gray")
plt.gca().invert_yaxis()
plt.subplot(2, 5, 5)
plt.imshow(data[:,:,8,1],cmap="gray")
plt.gca().invert_yaxis()
plt.subplot(2, 5, 6)
plt.imshow(data[:,:,4,1],cmap="gray")
plt.gca().invert_yaxis()
plt.scatter(seg[4,1][:,0], seg[4,1][:,1], c='r',alpha=0.1)
plt.subplot(2, 5, 7)
plt.imshow(data[:,:,5,1],cmap="gray")
plt.gca().invert_yaxis()
```

```
plt.scatter(seg[5,1][:,0], seg[5,1][:,1], c='r',alpha=0.1)

plt.subplot(2, 5, 8)
plt.imshow(data[:,:,6,1],cmap="gray")
plt.gca().invert_yaxis()
plt.scatter(seg[6,1][:,0], seg[6,1][:,1], c='r',alpha=0.1)

plt.subplot(2, 5, 9)
plt.imshow(data[:,:,7,1],cmap="gray")
plt.gca().invert_yaxis()
plt.scatter(seg[7,1][:,0], seg[7,1][:,1], c='r',alpha=0.1)

plt.subplot(2, 5, 10)
plt.imshow(data[:,:,8,1],cmap="gray")
plt.gca().invert_yaxis()
plt.scatter(seg[8,1][:,0], seg[8,1][:,1], c='r',alpha=0.1);
```

#### 4.1 For cardiac MRI: how to create a binary mask from the provided contours

### 5 Preliminaries

### 5.1 Mathematical Morphology

Incitations to use morphological operators seen during the previous lectures to segment the provided images.

Think about the structural elements and the hyper-parameters ... We typically adapt their values to the image resolution and type of structures targeted in our segmentation (eg. bright or dark).

```
[]: #Select input image
     \# Img\_test = Img\_cardiac\_ex
\# Img\_test = Img\_brain\_ex
     Img_test
                      = Img_abdo_ex ; # With: Seg_abdo_kidney_ex, Seg_abdo_tumor_ex
     # Define Element
     Radius
     se
                      = disk(Radius)
     # Morpho closing
     Img_test_close = morpho.closing(Img_test,se)
     # Morpho Opening
     Img_test_open
                    = morpho.opening(Img_test,se)
     # Morpho Gradient
     Img_test_grad = morpho.dilation(Img_test,se)-morpho.erosion(Img_test,se)
     #Figure display
     fig, axes
                      = plt.subplots(2,2, figsize=(10, 10))
                      = axes.ravel()
     ax
     ax[0].imshow(Img_test,cmap='gray')
     ax[0].set title("ORI")
     ax[1].imshow(Img_test_close,cmap='gray')
     ax[1].set title("Closing")
     ax[2].imshow(Img_test_open,cmap='gray')
     ax[2].set_title("Opening")
     ax[3].imshow(Img_test_grad,cmap='gray')
     ax[3].set_title("Grad")
     plt.show()
```

# 6 Preliminaries

#### 6.1 Regular filtering

```
[]: img_histeq = skimage.exposure.equalize_adapthist(Img_test, clip_limit=0.03)
    img_median = ndimage.median_filter(Img_test, size=4)
    img_histeqmed = ndimage.median_filter(img_histeq, size=4)

fig, axes = plt.subplots(2,2, figsize=(10, 10))
    ax = axes.ravel()
    ax[0].imshow(Img_test,cmap='gray')
    ax[0].set_title("ORI")
    ax[1].imshow(img_histeq,cmap='gray')
```

```
ax[1].set_title("HistEq")
ax[2].imshow(img_median,cmap='gray')
ax[2].set_title("Median")
ax[3].imshow(img_histeqmed,cmap='gray')
ax[3].set_title("Combined");
```

### 7 ASSIGNMENT:

Question Develop a segmentation pipeline building upon **kmeans** as proposed below for **at least one application** and report quality of segmentation results comparing your results to the provided ground truth on multiple images.

We want you to focus on **pre-processing** your image AND **post process** your segmentation result to extract **THE structure of interest**. For the postprocessing you can rely on extraction of connected components and apply to criteria (size, shape, position,..) to extract the composant that most likely corresponds to the structure of interest.

When processing medical images, and given that you are provided with several ground-truth (gt) segmentations you can push the exercice to consider: \* Cropping the field of view to remove the background (a common issue in medical images) \* Target a range of intensity values based on learning from the gt masks and corresponding images \* Learn priors on shape/intensity statistics from the provided gt masks

# 7.1 Kmeans routine to segment your images

Kmeans is very often used as pre-segmentation to initialise a finer segmentation.

Provides a segmentation of the image using k-means clustering. Be careful: Kmeans uses random initialisation and is therefore different at each run and randomly assigns labels to clusters (0,1,2,...,K).

```
Img_test > Target_value_thresh,
                            min_size=500,connectivity=1),
                             area_threshold=500)
Seg_thresh = morphology.opening(Seg_thresh, morphology.disk(3))
Seg_thresh = Seg_thresh.astype('uint8')
# [2] Example of segmentation via kmeans
Seg_km,Seg_km_labels = my_kmeans(Img_test,nber_clusters)
# get discrete colormap to display results
Colormap = plt.get_cmap('nipy_spectral', nber_clusters)
# Display results
fig, axes = plt.subplots(2,2, figsize=(5, 5))
         = axes.ravel()
ax[0].imshow(Img_test, cmap='gray')
ax[0].set_title('Original image')
ax[1].imshow(Seg_thresh, cmap='gray')
ax[1].set_title('Thresholding')
          = ax[2].imshow(Seg_km, cmap='nipy_spectral')
tmp
divider = make_axes_locatable(ax[2])
         = divider.append_axes('right', size='5%', pad=0.05)
ax[2].set_title('kmeans Centers')
fig.colorbar(tmp, cax=cax, orientation='vertical')
       = ax[3].imshow(Seg_km_labels, cmap=Colormap)
tmp
divider = make_axes_locatable(ax[3])
       = divider.append_axes('right', size='5%', pad=0.05)
fig.colorbar(tmp, cax=cax, orientation='vertical')
ax[3].set_title('kmeans labels')
fig.tight_layout()
plt.show();
# Get mean pixel intensity values under all Kmeans labels
Img_label_means = np.empty(nber_clusters)
for i in range(0,nber clusters):
    Img_label_mask = Seg_km_labels==i
    Img_label_mask = Img_label_mask.astype('uint8')
                   = cv2.mean(Img_test, Img_label_mask)
   Img_label_means[i] = tmp[0]
#print(Img_label_means)
# Plot histograms of Kmeans clusters
Bins = np.sort(np.concatenate((0,Img_label_means), axis=None))
```

```
hist_kmeans, bins_kmeans = np.histogram(Seg_km.flatten(),
                                         bins=Bins)
bar_width = 5
         = plt.figure(figsize=(2, 1))
fig
         = fig.add_axes([0,0,1,1])
ax.bar(bins_kmeans[1:-1],(hist_kmeans[1:]),bar_width);
plt.show()
# Set a target intensity value and get the Kmeans label closest to it
Target_value = cv2.mean(Img_test, Seg_thresh)
Target value = Target value[0]
Diff = np.absolute(Img_label_means-Target_value*np.ones(nber_clusters))
Label select = np.argmin(Diff)
print('Your Target_value = ', np.round(Target_value).astype(int))
print('Distance of your Target_value to Centers = ', np.round(Diff).astype(int))
print('Your selected label (starting with 0)= ', Label_select)
# Filter the selected label and get a binary segmentation mask
Img_label_select = Seg_km_labels==Label_select
Img_label_select = Img_label_select.astype('float64')
# Examples to refine your segmentation mask
Img_label_select_smooth = gaussian(Img_label_select, 2,
                            preserve_range=True)
Img_label_select_smooth_clean = morphology.remove_small_holes(
   morphology.remove_small_objects(
        Img_label_select_smooth>0.25, min_size=50,connectivity=1),
   area_threshold=50)
fig, axes = plt.subplots(2,2, figsize=(5, 5))
          = axes.ravel()
ax[0].imshow(Img label select, cmap='gray')
ax[1].imshow(Img_label_select_smooth, cmap='gray')
ax[2].imshow(Img label select smooth clean, cmap='gray')
ax[3].imshow(Img_seg_gt, cmap='gray')
fig.tight_layout()
plt.show()
```

# 7.2 From Kmeans to Connected Components

This is an example on how to exploit the results from Kmeans. Note that this example uses several hard-coded **hyperparameters** which is not appropriate to segment robustly several cases.

If you reuse this piece of code for the question below, propose some approaches to set some of the hyperparameter values automatically (eg using the known average size of the structure of interest from the gt segmentation you have)

```
[]: Img label select filter = gaussian(Img label select, 2,
                                 preserve_range=True)
     Img_label_select_smooth = Img_label_select_filter>0.25
     Img_label_select_smooth_clean = morphology.remove_small_holes(
         morphology.remove small objects(
             Img_label_select_smooth, min_size=150,connectivity=1),
         area_threshold=150)
     # Extract connected components
     Thresh = Img_label_select_smooth_clean
     output = cv2.connectedComponentsWithStats(Thresh.astype(np.uint8))
     (numLabels, labels, stats, centroids) = output
     print('Max value in labels = ',labels.max())
     print('Number of connected components = ',numLabels)
     # Extract 1 connected component
     thresh = labels==1
     # plots results
     Colormap = my colormap white bkg('nipy spectral',numLabels)
     fig, axes = plt.subplots(2,2, figsize=(5, 5))
              = axes.ravel()
     ax
            = ax[0].imshow(labels, cmap=Colormap)
               = make_axes_locatable(ax[0])
     divider
               = divider.append_axes('right', size='5%', pad=0.05)
     ax[0].set_title('Connected Components')
     ax[0].axis('off')
     fig.colorbar(tmp, cax=cax, orientation='vertical')
     ax[1].imshow(thresh, cmap=Colormap)
     ax[1].set_title('Selected label')
     ax[1].axis('off')
     ax[2].imshow(Img label select smooth, cmap=Colormap)
     ax[2].set_title('Img_label_select_smooth')
     ax[2].axis('off')
     ax[3].imshow(Img_label_select_smooth_clean, cmap=Colormap)
     ax[3].set_title('Img_label_select_smooth_clean');
     ax[3].axis('off');
```

# 7.3 Implement your full segmentation pipeline

#### 7.3.1 TO DO:

- Implement a full pipeline to segment TWO EXAMPLES from a single use case. Your code must run over the two cases in a loop. We need to be able to run the code without any adjustment to local path to access the input data.
- Implement and compute a quality metric (eg Dice, overlap, relative area differences, errors between max diameters (something used to measure tumors), distances between contours) that compares your final segmentation with the ground-truth.
- Extra point if you provide results on more use-cases or more than 2 examples per use case.
- Feel free to also test your segmentation pipeline on images degraded by noise for example.
- Your final solution can involve registration or active contours from previous lectures. Just make sure to include the required functions and imports in your final notebook.

#### 7.3.2 If you use the cardiac MRI dataset:

- Extra point because extra difficulties to load and prepare the data.
- In this case the data preparation is part of your code below as you can only rely on the input data provided as it is.
- Use any slice from any case you want as your "learning" ground truths on which you can learn object size or shape characteristics for example.
- Test your segmentation on any slice from any case, as long as the slices were not part of your "learning" ground-truth.
- See if you can get your solution to not detect anything on slices that don't contain the left ventricle myocardium.

### 7.3.3 Important instructions:

- You cannot use the ground truth of the images you segment to segment them! But you can use the ground-truths of other examples of the same use-case to learn a priori knowledge such as size or average pixel intensity.
- List all your hard-coded hyperparameters at the beginning of your code, indicating a variable name and its value. You will be penalised if you leave any hard-coded hyperparameter values inside your code.

Nb: Dice = np.sum([seg==gt])\*2.0/(np.sum(seg)+np.sum(gt)) #seg is the segmentation and gt is the ground truth. Both are of same size

[]: