Instruction CLSM Spectra Lab for PCA + clustering in XANES Wizard

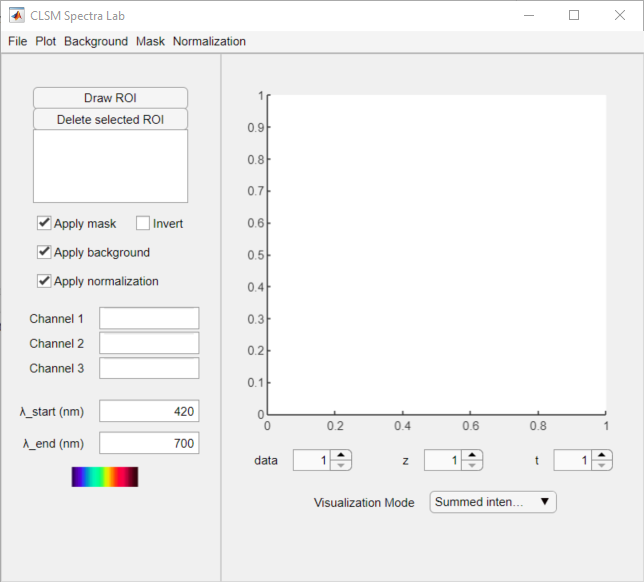
For version 0.25

# Motivation

The goal of this document is to provide a practical instruction to load images recorded on a confocal laser scanning microscope (CLSM) in the CLSM Spectra Lab App and prepare them for principal component analysis (PCA) + clustering in the XANES Wizard App. CLSM Spectra Lab is designed for CLSM data with a spectral component. The App has various plotting functions and can perform the computations “background correction”, “masking”, and “normalization”. These pre-treatment steps are essential for reliable clustering result of CLSM data, but are usually not included in the analysis software used in our group, because these were initially designed for other microscopy techniques. Ultimately, I hope that we can converge to a standardized pre-treatment procedure for CLSM data.

# Introduction

To start the app, open the CLSM Spectra Lab app via the “APPS” toolstrip. This should look like the figure below when freshly opened.



To load an image file, go to: “File” > “Load”. Multiple files can be selected if they are located in the same folder. If the images have been recorded in vastly different conditions, or have a different number of pixels, it is better to follow the workflow below for each set of ‘similar’ images. The reason is that some manipulations can only be done in the same way for all images in the data set. Even though the App can handle timeseries and z-stacks, it is not yet able export these to XANES Wizard.

Graphical user interface, application

Description automatically generated

The app has various features on the home screen that can be used right away (following the numbers in the figure above):

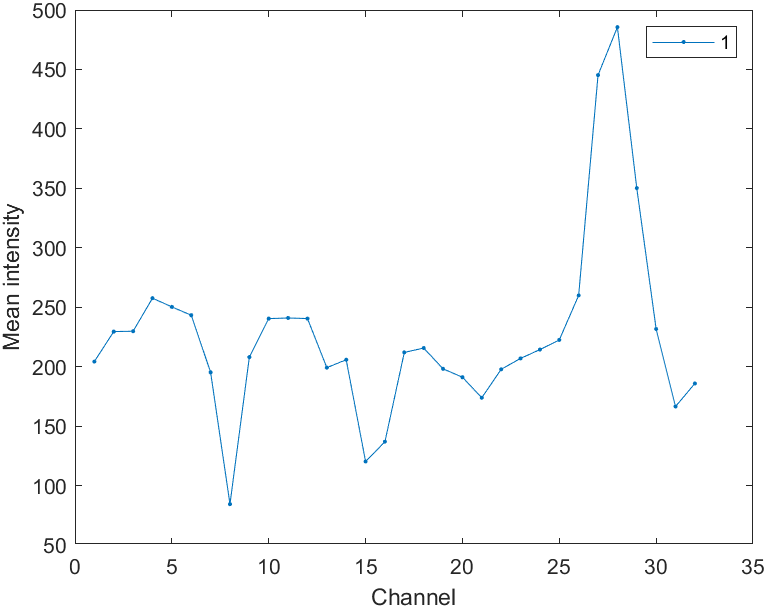
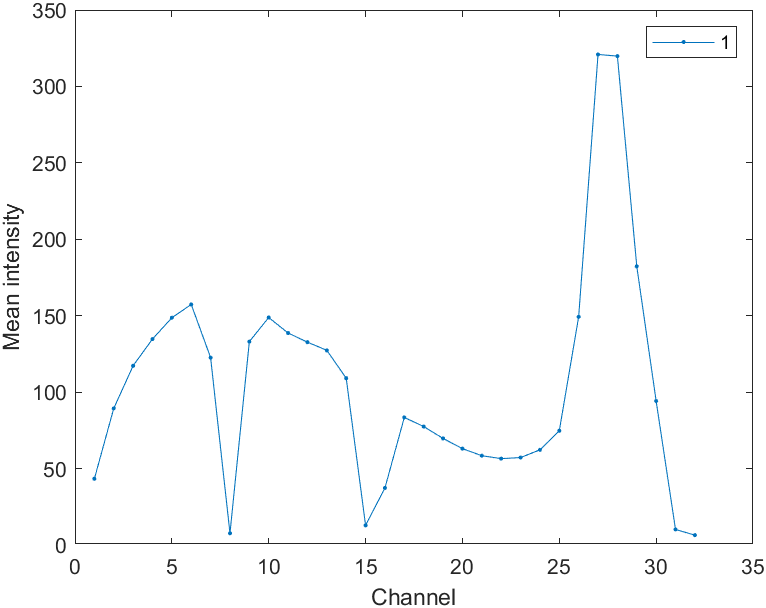
1. The user can navigate through the data, z-slices, and/or time steps via the buttons. A “please wait” box will appear until the image has been loaded.
2. The user can select the visualisation mode used to display the images in main window. The summed intensity over all spectral bins are displayed by default. The user can change the visualisation via the dropdown menu. “RGB blocks” divides the spectral bins in three equal blocks and displays each from low to high as red, green, and blue. Please note that this order is inverted with respect to the true colour, assuming that the increasing bin numbers represent longer wavelengths. To plot the approximate “true” colours, use the option “True colour”.
3. The wavelength of the first and last bin can be given in this box. These values are used for the “true colour” visualisation and are included in the export for XANES Wizard. The colour bar below displays the colour range of the bins.
4. Certain plot functions require the specification of bins as colour channel. This can be done in these boxes.
5. The mask, background correction, and normalisation can be applied for visualisation and computation. The visualisation is adapted automatically; however, computations have to be recomputed in order for the option to be applied.
6. Regions of interest (ROI) can be drawn and the spectra of the ROIs can be plotted. Click “Draw ROI” and draw the regions in the main visualisation window. Each ROI is assigned a unique number for identification. The corner points of the ROI can be interactively changed. The ROI can be deleted by selection of the ROI in the menu and pressing the “Delete selected ROI” button.

The images and their spectra can be plotted in the menu “Plot”.

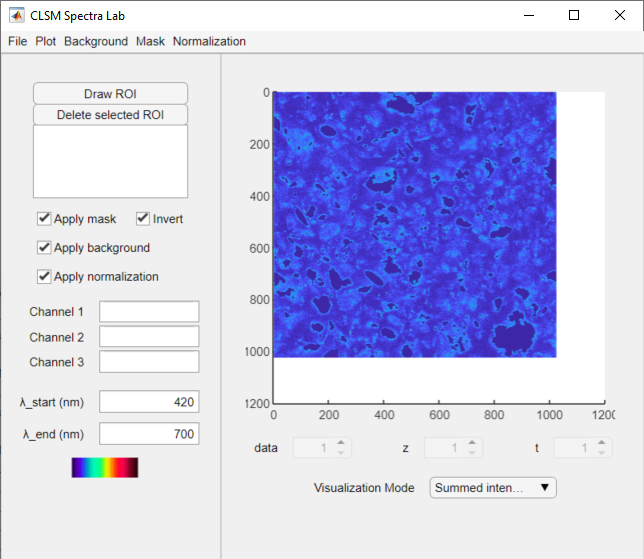
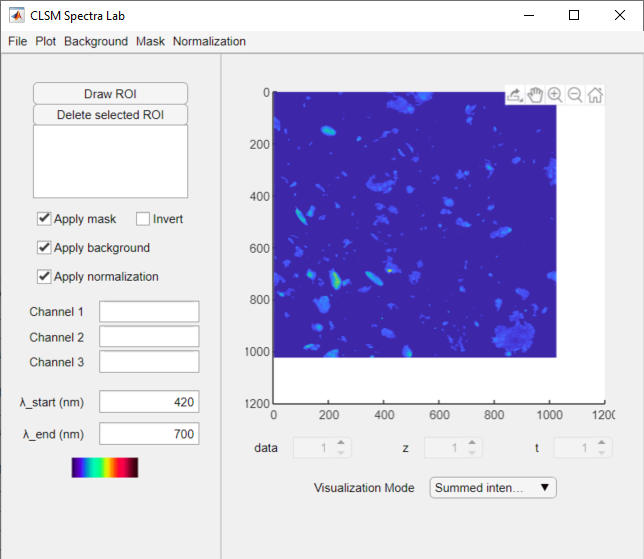
# Workflow

In this section, I would like to propose a workflow for the pre-treatment of CLSM data. This workflow should be polished based on user experience. I will explain the steps and how to perform them.

1. Press “Background” > “Compute background”. An input dialog opens: Compute per pixel (1=true/0=false) and fill in 1. A file selection dialog opens. The user should select a so-called “dark image”, which has been recorded with the shutter off or microscope to “AUX”. Please make sure that the dark image is recorded with the same settings as the image, that is same dwell time, HV, and pixels. This option subtracts the values in the dark image from the image. Inspect the result by plotting the spectrum of all pixels via “Plot” > “Plot Spectrum” > “All” both with and without “Apply background” marked “ON” and “OFF”. Before and after background correction:

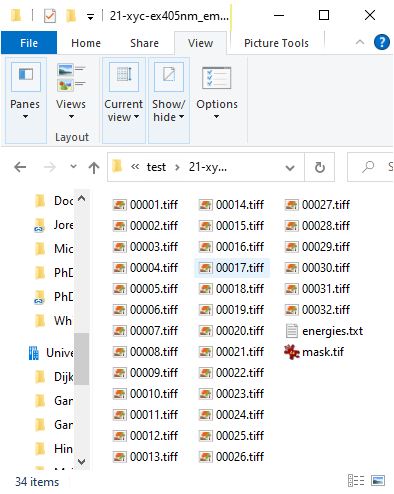
 

1. Please make sure that the “Apply background” is selected to do the masking with the background correction included. Press “Mask” > “First PCA” > “Compute mask”. A dialog box opens: Enter threshold value (empty for auto). Leave empty and press OK. Now image is displayed with the mask, where the dark blue regions are the background, and the mask can be inverted with the checkbox “Invert”:



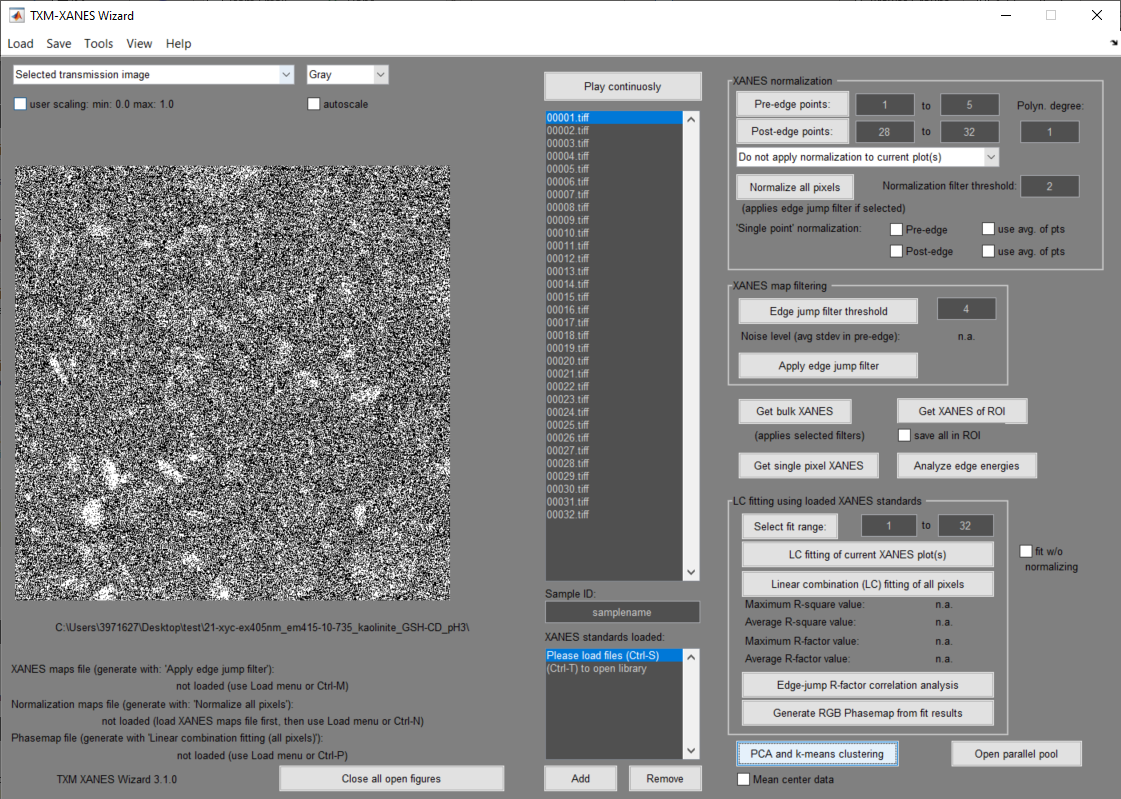
The value can be tuned for each image individually via “Mask” > “First PCA” > “Compute” for current data. The masking is done based on rescaled intensity values. A histogram of all rescaled intensity values can be plotted via “Mask” > “First PCA” > Plot histogram 1st eigen image”, which aids in selecting a good threshold value.

1. “Mask” > “Add sat. pixels”. Input dialog “Threshold counts”, keep the default value 4095. This function removes saturated pixels. A pixel is saturated when it has at least one bin in the spectrum with maximum value set by the user. Default is 4095, which is correct for the 12-bit format the Nikon A1 saves by default. Use 255 for 8-bit data and 65,535 for 16-bit images. The saturated pixels are removed from the current mask. This step is essential because the shape of the spectrum is distorted in the saturated pixels, because their true value is larger than the saved value in the image.
2. Please make sure that the “Apply background” and “Apply mask” are selected and “Invert” is deselected to do the normalisation with the background correction included. “Normalization” > “Max” > “Compute norm”. Use “Plot” > “Plot Spectrum” > “All” to inspect the result.
3. Please make sure that Apply background”, “Apply mask”, and “Apply normalization” are selected. Then, press “File” > “Export” > “XANES Wizard”, which opens a folder selection dialog box. Please select the folder where you want to save the exported images for XANES Wizard. In this folder, a new folder is created for each image in CLSM Spectra Lab. In this folder, ‘tiff’ images with an increasing number in the format “00001.tiff” are images of the individual spectral bins, “energies.txt” is a file containing the wavelength of the bins, and “mask.tif” is the mask of the image.



Now, we can import the files in XANES Wizard:

1. In XANES Wizard, go to “Load” > “Image stack” and select the numbered tiff images (not the mask!). An error will appear: “Energy could not be determined from the filename! […] List of energies?”. Press “yes” and load the file “energies.txt”. De images have now been loaded.



1. Perform PCA and Clustering without using XANES Wizard’s other functions via the button “PCA and k-means clustering”. A question dialog appears: “No normalization data file found. Do you […] filtered) data?”. Press “Yes, and apply external filter”. Press “save” and in the next window select “mask.tif” to “open”.
2. You’re ready to do the PCA and clustering. For more information about this procedure, please visit the course on the wiki <https://solisservices.sharepoint.com/sites/10857/icc_wiki> and go to “Knowledge” > “Lectures & Workshops” > “Principal component analysis (PCA) workshop”.

**Note:** If the data is noisy, this can lead to unexpected clustering results. Inspect the single-pixel spectra via “Plot” > “Plot spectrum” > “Single pixels”, click one or more pixels and press enter. The pixels in X and Y can be binned via “Tools” > “Bin input image” to reduce the noise level. A bin factor of 2, 3, or 4 is recommended. This should be done prior to the masking step.

The cluster results can be loaded in CLSM Spectra Lab to compute the average spectrum of the clusters without normalization. Because the clustering performed by XANES Wizard on the normalized hyperspectral image, the reported spectrum per cluster is also based on the normalized data. Better is to compute the average spectrum from the non-normalized data:

1. Go to “Plot” > “Plot spectrum” > “XANES Wizard clusters” and a file selection menu is opened requesting a “KmeansResults.mat” file, which is the default save name of XANES Wizard for clustering with K-means and/or EM-GMM. The file may be renamed; this does not affect the performance of CLSM Spectra Lab. The clustering result is plotted along with the average spectrum of the clusters given. If the mask, background, and/or normalization is/are applied, this is also taken for the computation of the average spectrum.
2. The spectrum plotted in (9) can be saved via “Tools” > “Save XANES Wizard spectrum”. The “KmeansResults.mat” file is requested and the spectrum per cluster is saved in a excel sheet in the same folder as ““KmeansResults.mat” file with the name “spectrum\_clustersXANESWiz\_bgc0\_msk0\_norm0.xlsx”. bgc indicates whether the background correction is applied; msk the mask is applied; and norm whether the normalization is applied. A “0” stands for “not applied” and “1” for “is applied”. If the mask/background correction/normalization is applied, but not computed, it will still be saved as “1”. The first column in the excel sheet is the spectrum of the first cluster, the second of the second cluster, etc. The first number is the number of pixels in the cluster, the others in the column are the spectrum intensity values per bin.

**Clustering**

As per version 0.30, clustering is available in CLSM Spectra Lab. Pixels with similar spectra properties can be grouped together in so-called “clusters”. Pixel clustering can be computed, plotted and analysed. There are two options to compute the clustering:

1. Perform PCA clustering analysis with XANESwizard and load the clustering result in CLSM Spectra Lab;
2. Compute the clustering based on a similarity fit to a set of reference spectra.

*Import from XANES Wizard:* The cluster results from XANES Wizard are exported to a file called “KmeanResults.mat”. This file can be selected for the current dataset via “Clustering” > “Import XANES Wizard” > “Set filename for current”. If this step is skipped, a file selection dialog box will appear one by one for all datasets that do not have a “KmeanResults.mat” selected. The data is loaded via the compute option: “Clustering” > “Import XANES Wizard” > “Compute”.

*Similarity fit:* CLSM Spectra Lab has the functionality to compute a similarity fit to a set of reference spectra. A least-squares residuals are computed for all individual pixels and the reference spectra. The pixel is assigned to the reference spectrum with the lowest residuals, and thus, the best fit. Because the area without fluorescent markers is characterised by a low emission intensity, rather than the dell-defined emission spectrum, the least squares fit is done in two separate groups: background and foreground. To decide what pixels are foreground and background, an intensity threshold is set. First, the reference spectra are loaded via “Clustering” > “Similarity fit” > “Load reference spectra”. The reference spectra must be saved in a delimited text or spreadsheet file. The data must be a matrix with the rows the intensity at each wavelength matching the to-be-analysed data and the columns the different clusters (i.e., background and/or materials). The similarity fit is computed via “Clustering” > “Similarity fit” > “Compute”. A dialog box is opened requiring the user to set the normalized background/foreground threshold as well as the number (index) of the reference spectra that are expected in the background. The other reference spectra are assumed to be in the foreground. Note that this background and foreground differs from the mask computed in CLSM Spectra Lab. The similarity fit is computed for all pixels, even when they are masked out.

The cluster result can be inspected in a few ways. “Clustering” > “Plot cluster map” plots the spatial map with the clusters colour coded. “Clustering” > “Plot spectrum clusters (all)” plots the spectrum of the clusters for the current data set (or for all data sets in one plot). Here, the mask is considered when applied to the current dataset. These plotted spectra can be copied to the clipboard via “Clustering” > “Copy spectrum clusters to clipboard (all)”.

To analyse the cluster maps, they are first processed via to remove small groups of pixels that are likely to be a false classification due to noise. The minimum number of adjacent pixels of the same cluster is set as well as their connectivity in “Clustering” > “Cluster map process settings”. The cluster maps before and after processing are shown (same can be plotted via “Clustering” > “Plot processed cluster maps”). As a result of the processing the number of pixels in the clusters can be larger/smaller than the number of pixels in the image. The cluster statistics are computed and exported to an Excel file via “Clustering” > “Save cluster statistics (all)”. The properties are computed for every particle in each cluster “*DATASET*\_Cluster*CLUSTERNUMBER*” and the mean properties per cluster are given in the tab “Mean”. The following properties of the “groups” of pixels are computed (think of as a particle or granule in sample):

1. Area: number of pixels in a “group”;
2. Centroid: x (\_1) and y (\_2) coordinates of a “group”;
3. Eccentricity: value between 0 (circle) and 1 (line) that describes the shape of the “group” (more info here: <https://ch.mathworks.com/help/images/ref/regionprops.html>);
4. FilledArea: same as Area but with all holes in “group” filled;
5. NN: distance to nearest neighbour. Please keep in mind when interpreting this value is very dependent on how close the “group” is to the edge of the image. Sufficient “groups” should be present in the map for this property to be meaningful.

Area, FilledArea, and NN are also computed in meters using the metadata in the CLSM images.