From Movie to 3D Space…

You should have a folder full of DV files.

Folder starting with 1,

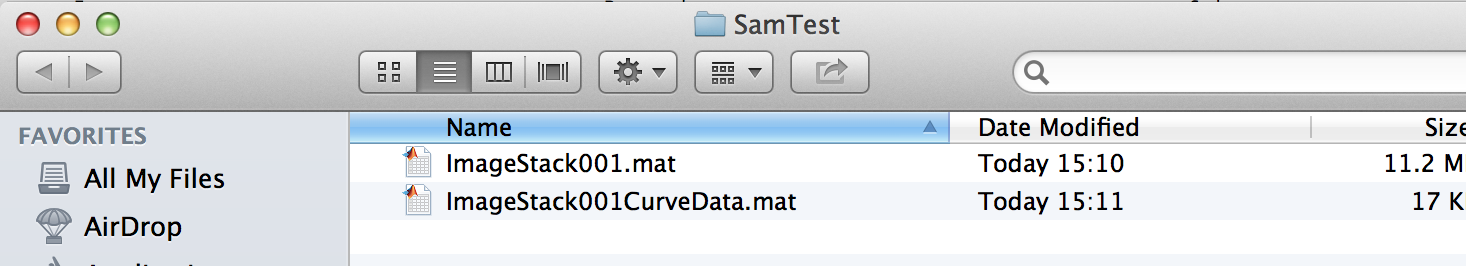
RunMeFirst.

* Index of file (which of the files you want to do
* Start of frame, end of frame
* Path (that can be reg expression)

DV becomes a matlab variables; This one does the segmentation . which can take quite a while. As a rule of thumb, it 1500

* RunMeFirst(1, 1,10, '/Users/iasmam/Desktop/SamJ\_Code\_and\_Data/DemoData/GLA6\_exp3\_220713\_untreated01\_114\_R3D.dv','/Users/iasmam/Desktop/SamTest' )
* Movies should have all the same number of frames (assumptions)

After this step, we have for each file the following files



# Macintosh HD:Users:iasmam:Desktop:Screen Shot 2015-02-06 at 15.28.52.png

This are the two parameters directly in the code, that influence how the segmenation is done.

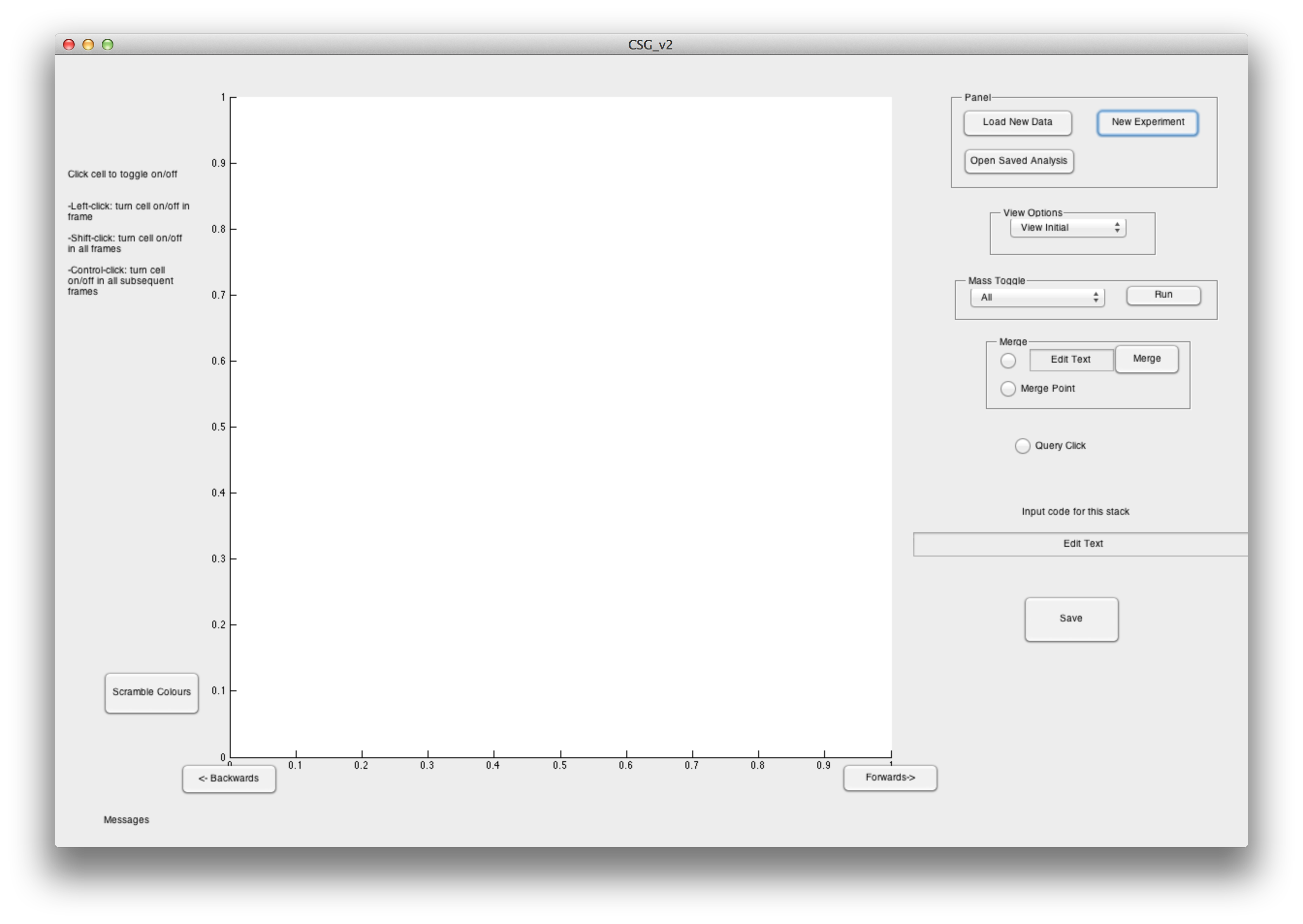
Rangeband= how much noise it is picking up  
spagtialband= how large a cell is (say 3 is the readious… if a cell is large, say always larger than 10, you might want 10, as the smallest cluster).

We can give a good default (as shown in the image).

The next step involves opening a guid

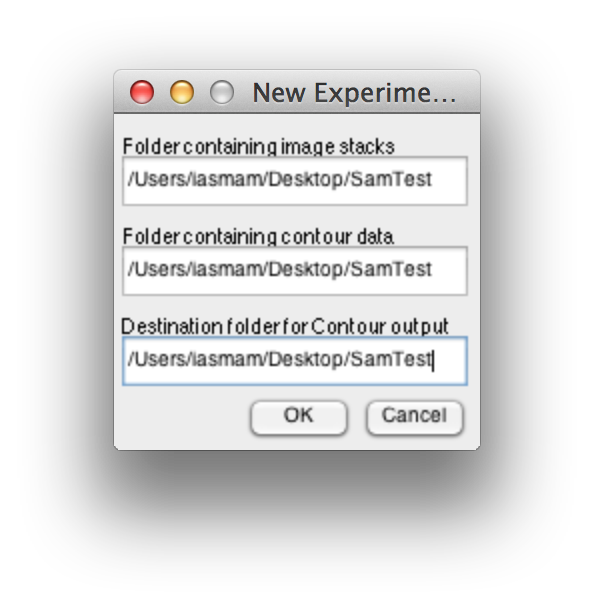
In the folder 1-ImageSementationFull there is a file called “CSG\_v2.m”

Open in Matlab this file and start it. You should see:



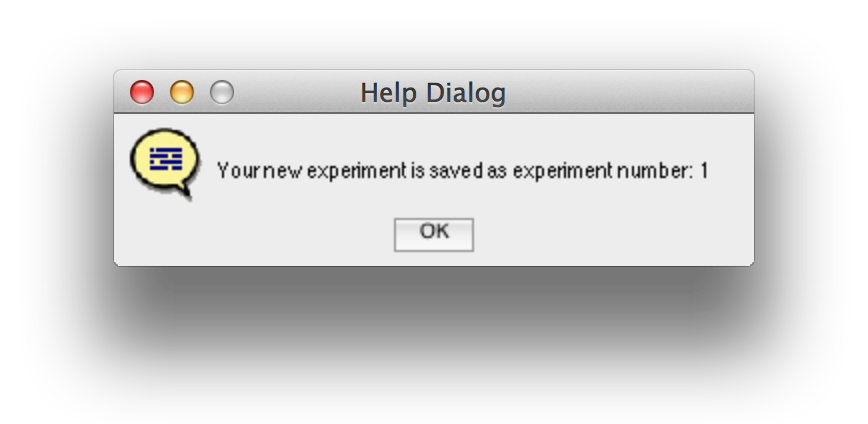
Click on “New Experiment”

And enter the path, where you have been saving things, in all three boxes.



If you are a programmer, the 3 paths would make sense, or if you want to do some advanced output. Otherwise, if you have just been following this tutorial, you have to put 3 times the same path in.

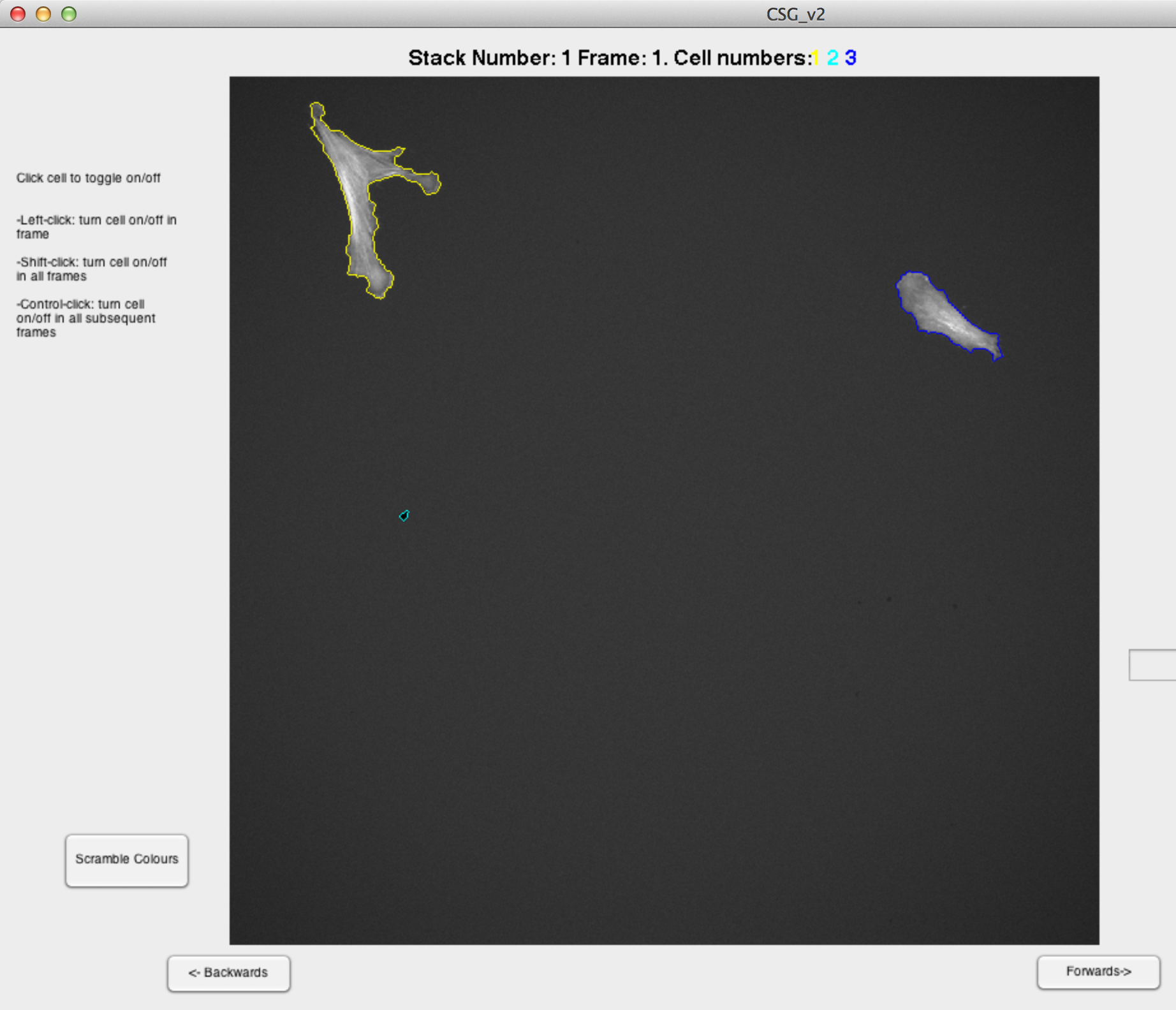
If you were successful, you should see a dialog window, that should look like this: Important: it should have a number!



# Now to open a stack. You have to enter the experiment number and the number of the stack (see below)

# Macintosh HD:Users:iasmam:Desktop:Screen Shot 2015-02-07 at 14.42.13.png

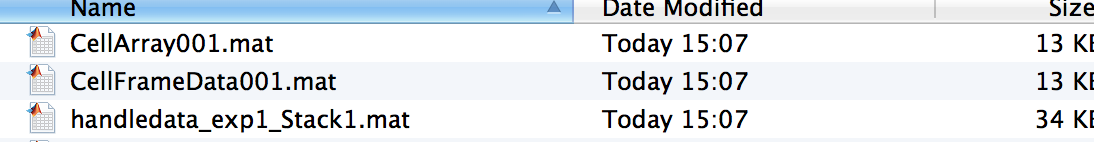
# If everthing was working out. You should see your stack, with the outline of the cells.



This interface permits you to manually correct things. Say I would like to remove the small detected cell here, I would Shft+Click on it, and then save the data. Saving the data, means that I generate 3 new files for the stack. CewllArray001.mat

CellFrameData001.mat

Handledata\_exp1.Stack.mat



# Options in the GUI

* You can delete cell shapes (also froma specific point)
* You can merge cells (even with merge point)
* There are instruction on the movie.

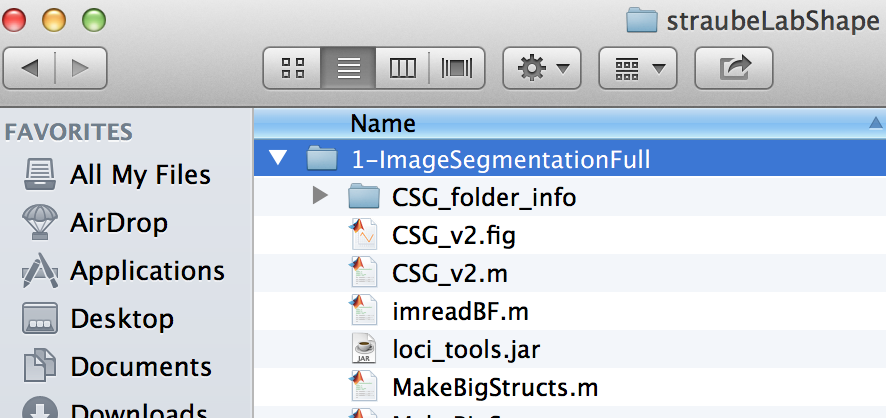
You can toggle off the short lived ones, and you can toggle of the small ones. Be careful as the two annul each other. Which is pretty annoying.

You can only load one stack at the time.  
You save them in your folder.

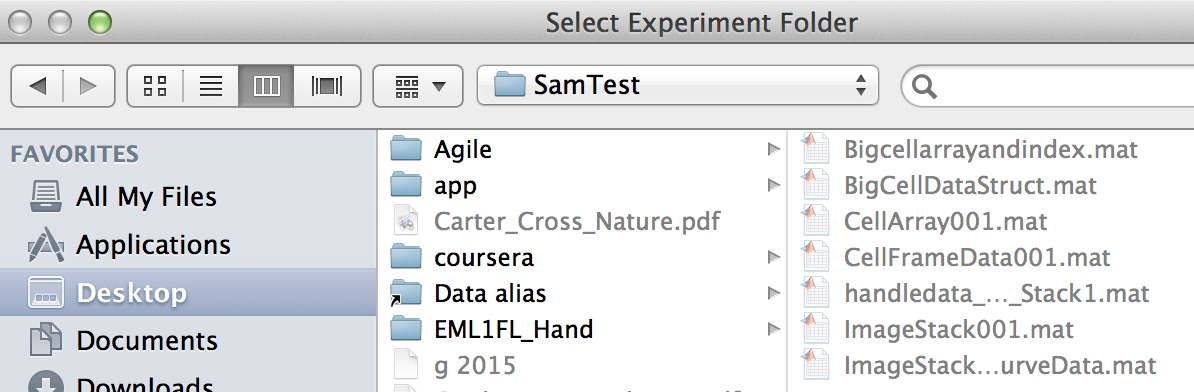
# Once all your files are done

Say, you have corrected all files, then you are ready for the next step: in Folder 1-ImageSegmantation Full, there is the file MakeBigStructs.m

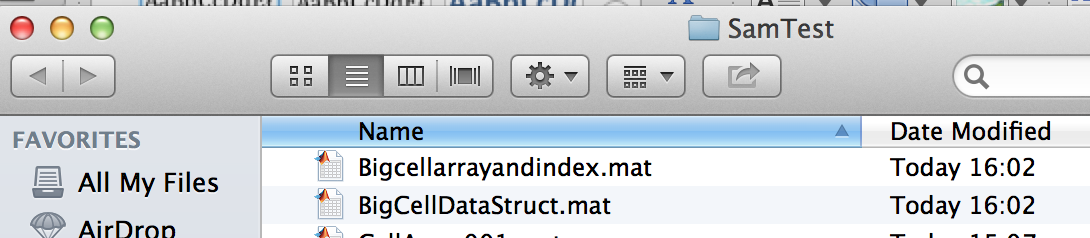
Open it in Matlab and click on the “Run” Button.



There should be an explorer that opens, and you should enter as the directory, the experminet directory you have been using.

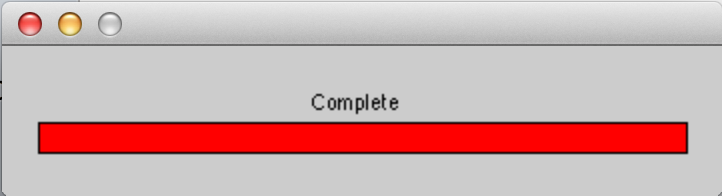


After selecting the folder, the program should create 2 new files in the folder, namely Bigcellarrayandindex.mat and BigCellDataStruct.mat (see below).



Now you are ready for the next processing step. You go to the Folder   
“2-ShapeManifoldEmbedding”

And open the script “RunShapeManifoldEmbedding.m” in Matlab. Click on the “Run” Button in Matlab, and the script should ask you for experiment folder (as before). Select the folder and when the script complets correctly, you should see the following dialog box.



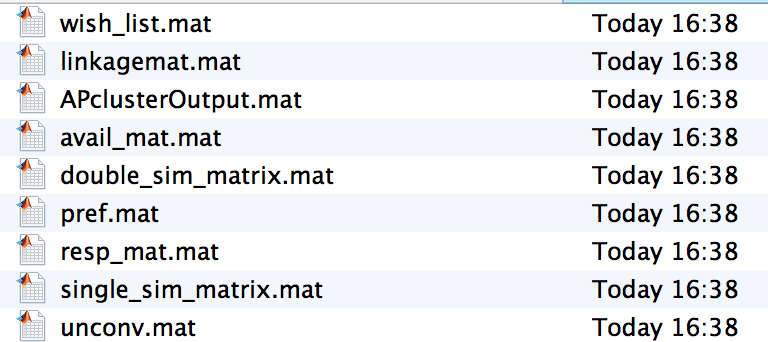
Also, if the program run correctly, you should have the file “CellShapeData.mat” in your experiment folder.

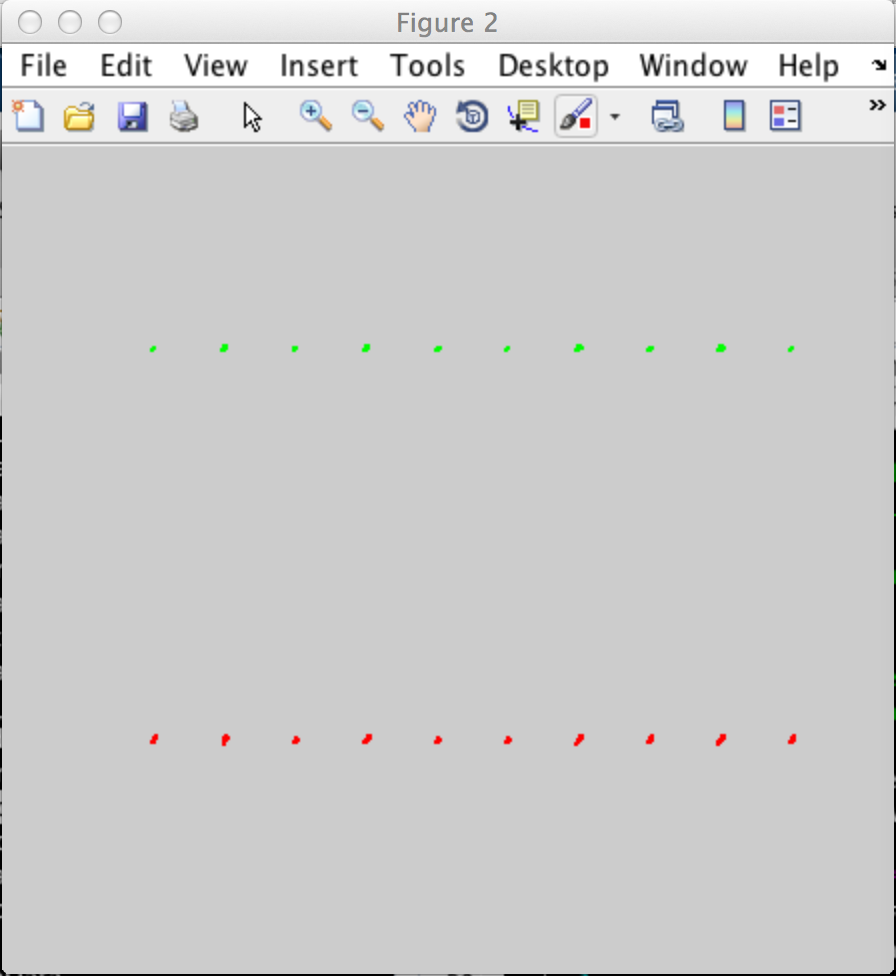
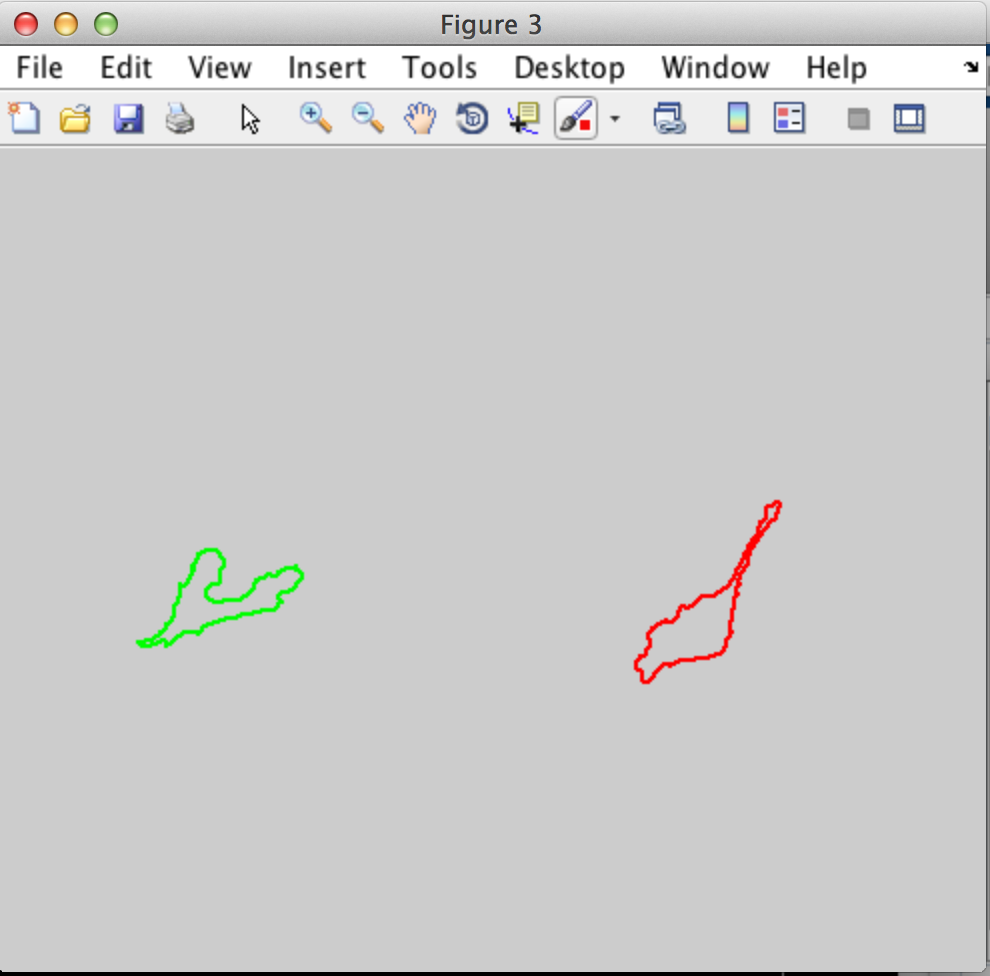
Now you are ready to move to the steps in Folder

“3-Extended\_Affinity\_Propagation”.

Open the program: Run\_Affinnity\_Progagation.m in Matlab and click “Run”. It should ask for your ExperimentFolder. After selecting, the programs will run through (this might take quite a while, please be patient.)

IF the program was successful, there should be the following files there:



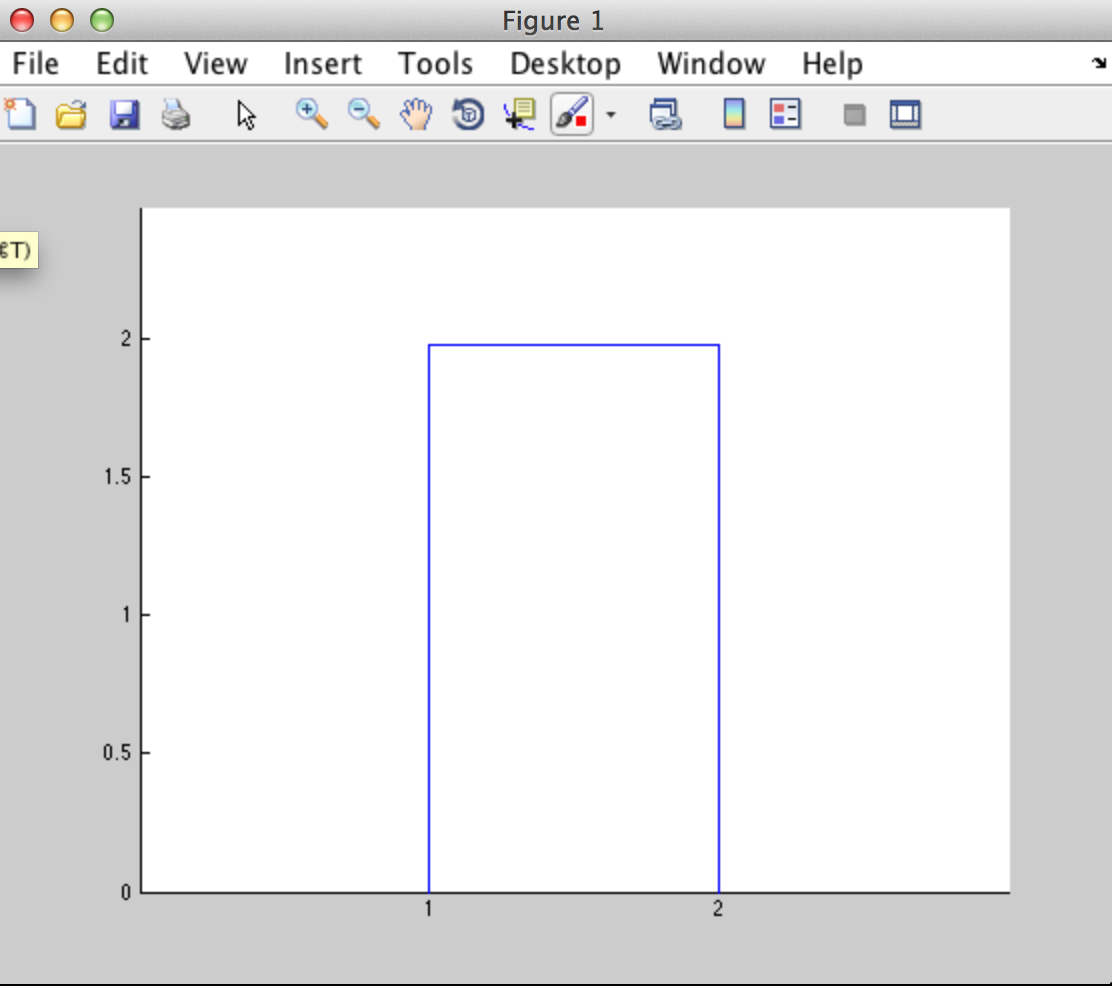
Open “unordered\_list.m” and click on Run. Again select your experiment folder, and then the program should run through. You should get 2 figures. 

You can save vector images by clicking on File>Save As…

Ordered List.

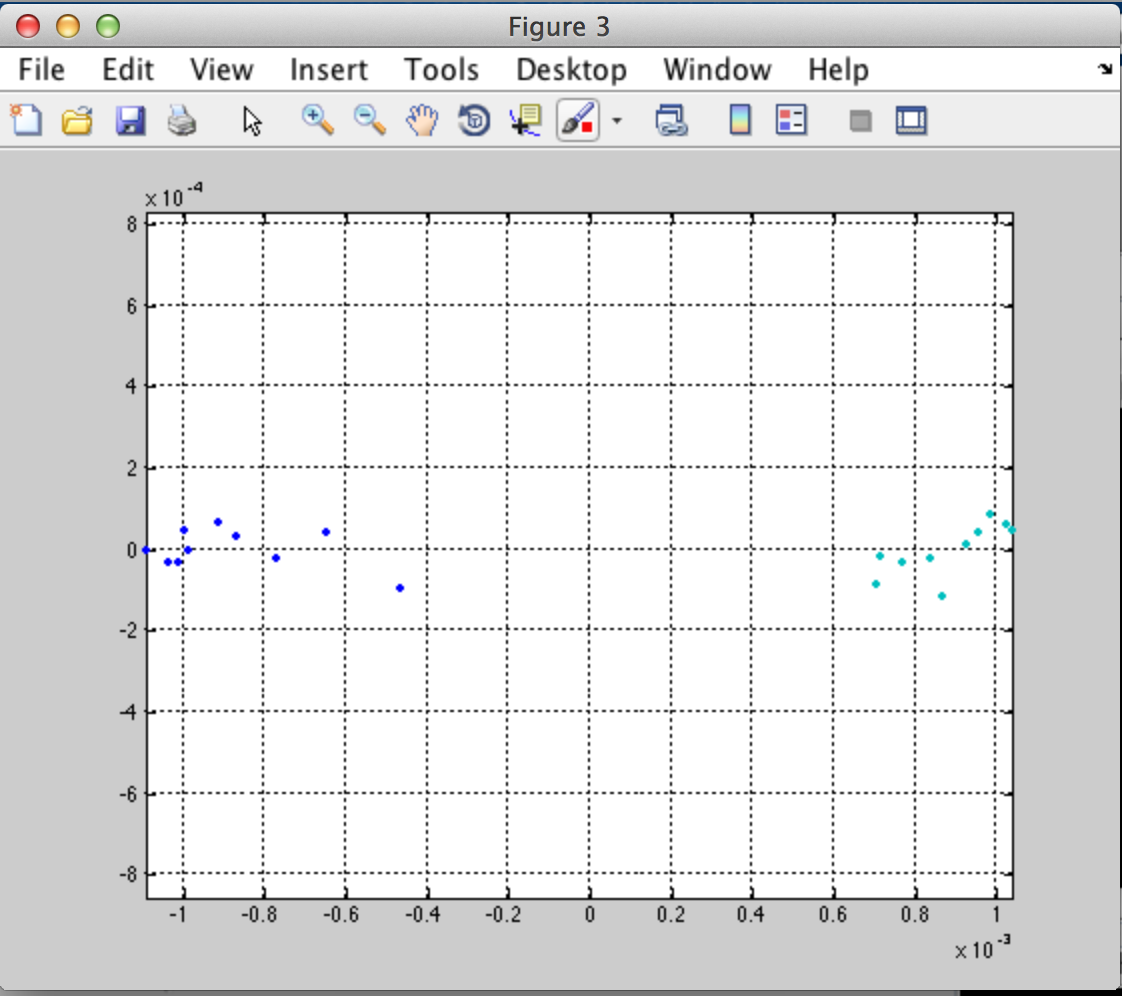
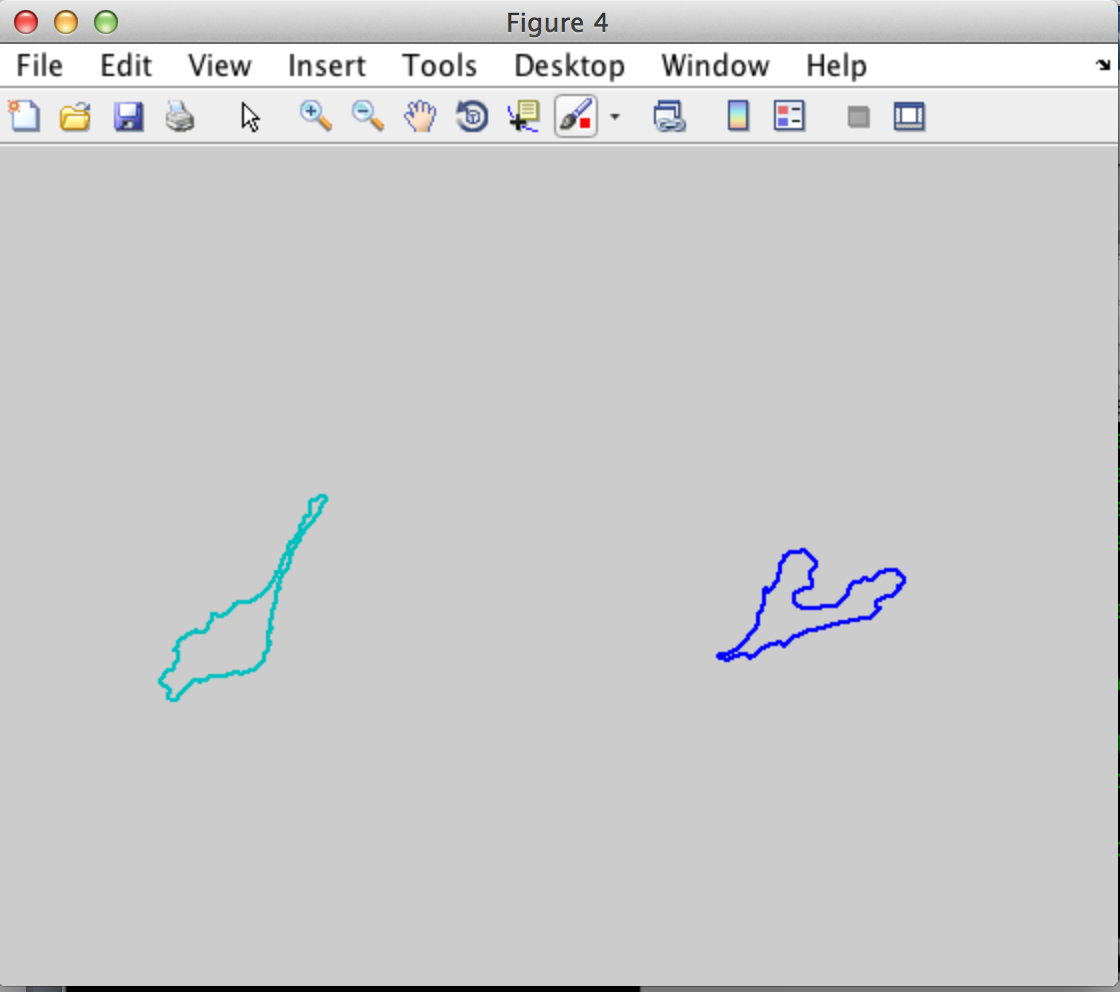
Sometimes, the unordered list might produce too many classes. Than, we first look at the dendogram, by entering zero to the file.

ordered\_list(0,CellShapeData,APe\_output\_foldername)



Now, if you enter a number, you get additional output: namely the following diagrams.

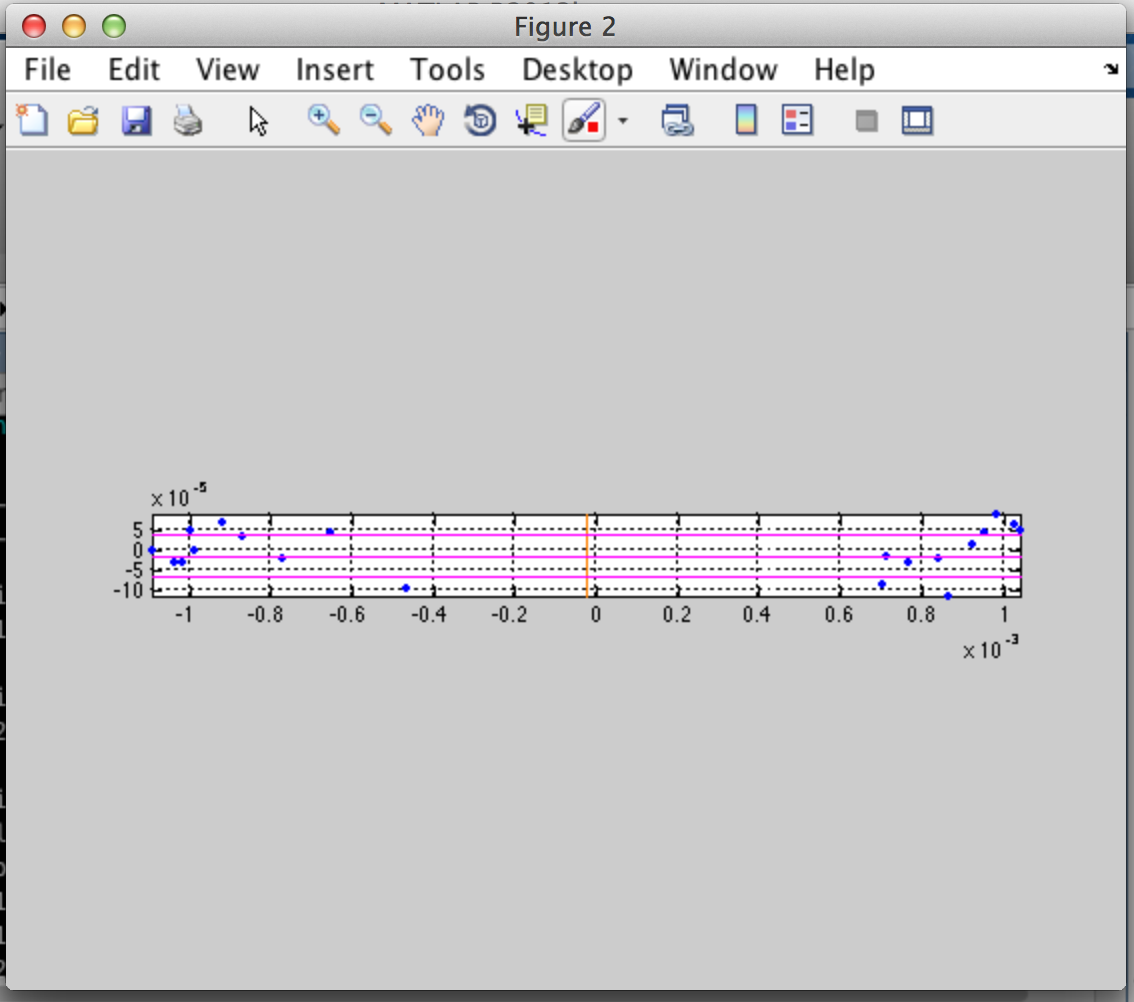
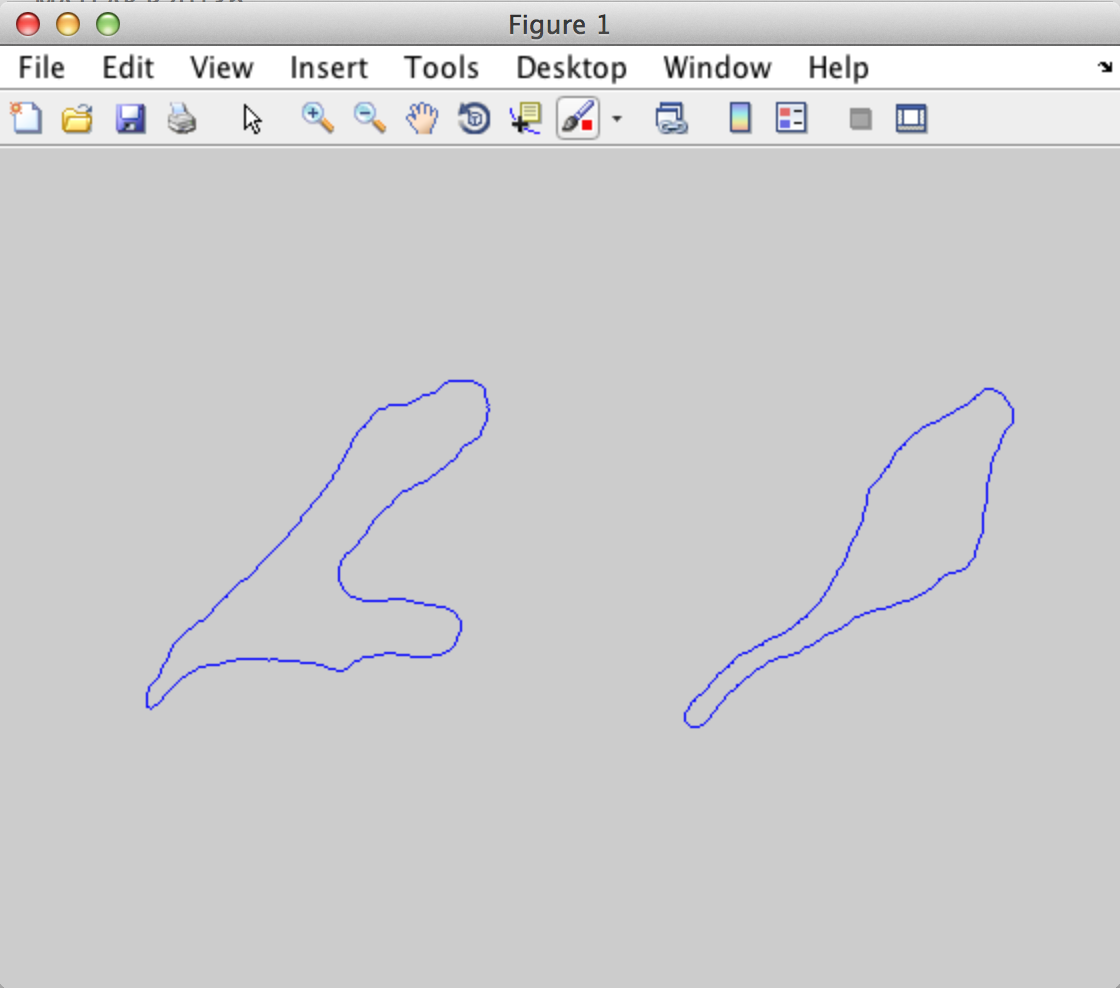
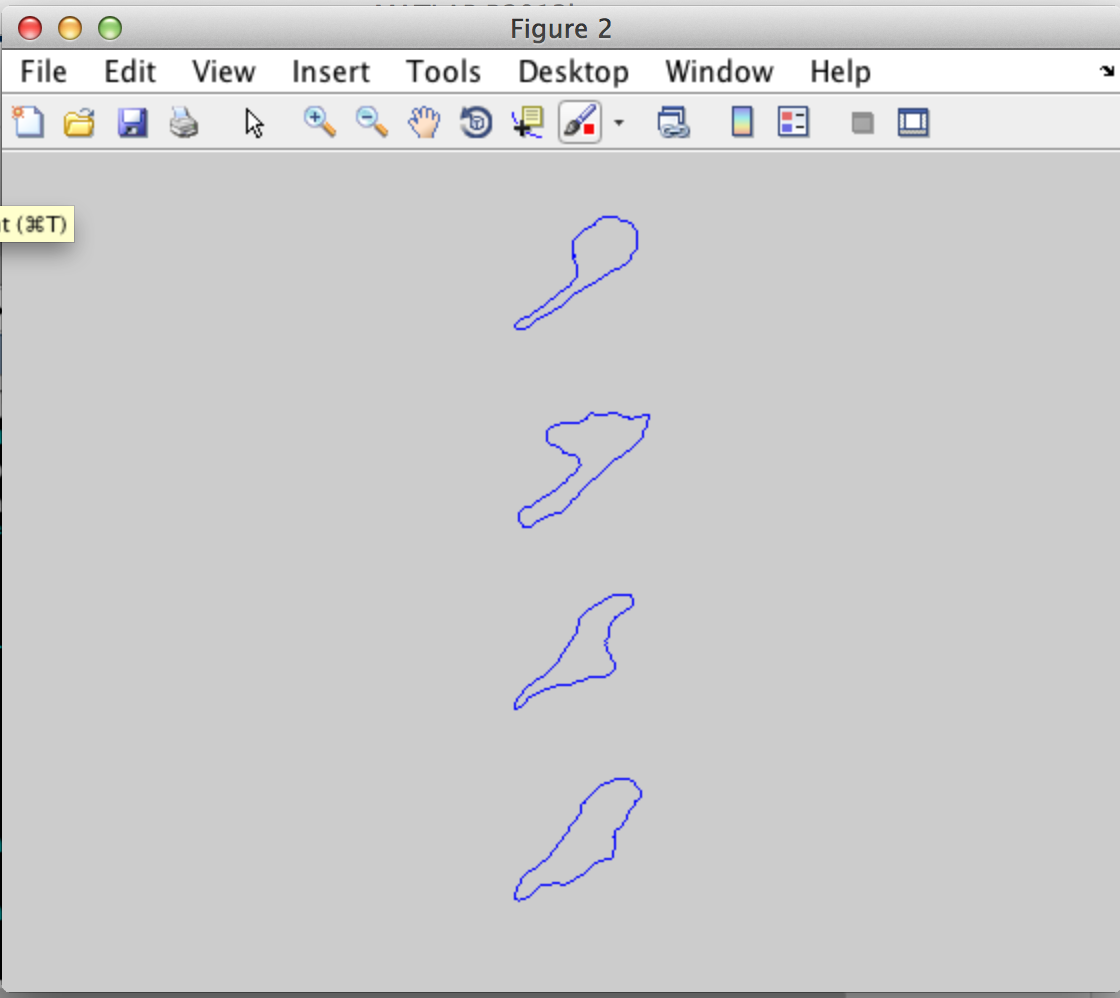
ordered\_list(2,CellShapeData,APe\_output\_foldername)



# 4 Shape Slicer

To run shape sliser, go to Folder 4

Run “Run\_SpaceSlicer( )” it asks you for the experiment directory. Select the folder and rund the program. You should get the following 3 figures.



The figure sam has on page 45 in the thesis draft Anne showed me, this figure is larger. Also, on the third images, there are figure shapes. They were done with illustrator, and for the method paper, this probably would need to change. But for now it is running.