This tutorial explains building a **ColorModel** using the *TASBEFlowAnalytics* software package. A **ColorModel** improves the precision of flow cytometry measurements by translating raw fcs data to a comparable unit data. A previous understanding of Flow Cytometry is assumed within this tutorial. However, resources are available for users new to the subject and are available here.

Creating A Color Model

As a starting point for users, a MATLAB script with an example color model is available here. The script is labeled as *make_color_model.m*. Users should proceed to download and open this file using MATLAB or Octave. Additionally, example data files are also provided to create the **ColorModel** and are made available here.

Within the script, the first step is to set up paths for a few different values. The first path should be directed to where the fcs data files exist. Currently, the script is coded to include the path to the example files that are provided. The second and third path are set to point towards the *bead* and *blank* file path. This process of setting up paths is entirely optional and exist solely to prevent from having to use full paths when referring to the data files.

```
1 stem0312 = '../example_controls/2012-03-12_';
2 beadfile = [stem0312 'Beads_P3.fcs'];
3 blankfile = [stem0312 'blank_P3.fcs'];
```

Autogating

Next step is to initialize Autogating. First, the autogating filters need to be initilized as shown in line 1. Some key default filter parameters are already set, but they can be modified as needed by the user. For example, the default number of gaussian components which are searched for are two. The components are set to search for one single-cell component and the second component is typically a non-cell or clump component. Line 2 shows how to change the number of searched components.

Additionally, the default channel names are set as FSCA,SSCA,FSCH,FSCW,SSCH,SSCW. These channel names can be modified to add more channel or fewer as needed as shown in line 3. Please see the API to see the complete list of filter parameters available.

Once the parameters are set, we can initialize *Autogating* as shown in line 5. The inputs include the location of the blankfile, the parameters, and output directory where graphical figures will be stored.

Creating Channels and ColorPairs

A useful function of the TASBEFlowAnalytics package is its ability for using multi-color controls to convert other colors in MEFL units. In order to begin this process, channels are created for each color being measured. There are three channels being created, FITC-A, PE-Tx-Red-YG-A, and Pacific Blue-A. As shown in line 4, a channel requires a name, wavelength, center filter, and width filter. Additionally, the directory for where the color file exists must be specified as shown in line 6. Functions exist to add details by which a channel can be identified. For example, setPrintName and setLineSpec are used to set the name the channel is known by and the color which will appear on graphs when referring to that channel. These are optional functions to use.

```
1
   channels = {}; colorfiles = {};
2
3
   channels\{1\} = Channel('FITC-A', 488, 515, 20);
   colorfiles{1} = [stem0312 'EYFP_P3.fcs'];
5
   channels{1} = setPrintName(channels{1}, 'EYFP');
   channels{1} = setLineSpec(channels{1}, 'y');
7
8
   channels\{2\} = Channel('PE-Tx-Red-YG-A', 561, 610, 20);
9
   colorfiles{2} = [stem0312 'mkate_P3.fcs'];
10
   channels{2} = setPrintName(channels{2}, 'mKate');
11
   channels{2} = setLineSpec(channels{2}, 'r');
12
13
14
   channels{3} = Channel('Pacific Blue-A', 405, 450, 50);
15
   colorfiles{3} = [stem0312 'ebfp2_P3.fcs'];
   channels{3} = setPrintName(channels{3}, 'EBFP2');
17
   channels{3} = setLineSpec(channels{3}, 'b');
```

The next step is to create the *colorpairfiles* which specify which colors are being converted. Line 3 shows how to create an instance of a color pairing. The first two parameters are the colors which are being converted into one another. We see that *FITC-A* and *PE-Tx-Red-YG-A* are the colors being converted. The third parameter is an alternate color which is used to assist in estimating the conversion which in this case is *Pacific Blue-A*.

Creating The ColorModel

All of the inputs necessary to make a color model have been created. The next step is to create the model. As shown in line 1, the parameters needed are the bead file, blank file, the created channels, colorfiles, and colorpairfiles. Additionally, some additional functions have been pre-set for the user. In particular, details for the beads used have been specified through the functions set_bead_model and set_bead_batch. Additionally, line 12 shows the function to limit what bead data values are ignored. Currently, the minimum is set such that any data below 10² is considered too smeared and is discarded. The peak threshold determines the minimum count per bin for something to be considered part of a peak. This value can be adjusted using set_bead_peak_threshold as shown in line 13.

```
1
  CM = ColorModel(beadfile, blankfile, channels, colorfiles, colorpairfiles);
  CM=set_bead_plot(CM, 2);
  CM=set_translation_plot(CM, true);
  CM=set_noise_plot(CM, true);
  CM=set_bead_model(CM,'SpheroTech RCP-30-5A');
6
7
  CM=set_bead_batch(CM,'Lot AA01, AA02, AA03, AA04, AB01, AB02, AC01, GAA01-R');
8
9
   CM=set_FITC_channel_name(CM, 'FITC-A');
10
   CM=set_translation_channel_min(CM,[2,2,2]);
11
  CM=set_bead_min(CM, 2);
12
13
   CM=set_bead_peak_threshold(CM, 1000);
14
15
  settings = TASBESettings();
16
  settings = setSetting(settings, 'path', 'plots');
17
18 | CM = add_filter(CM, autogate);
```

Discussion of Graphs











