Proteomics data analysis in cancer biology with Matlab

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- **Date**: 08/14/2017
- **Github repository**: https://github.com/friveramariani/Proteomic-Examples

Summary

This report represents an example Matlab proteomic data analysis. The dataset analyzed in this report can be found here, which is the FDA-NCI Clinical Proteomics Program Databank. The samples downloaded from the FDA-NIC Proteomics Programa Databank correspond to SELDI Mass-Spec profiles of ovarian cancer samples: **Cancer Group** vs **Normal Group**. The study related to this dataset was published, in 2004, in the Endocrine Related Cancer journal. **Briefly**, after *transforming* the Mass-Spec data, some *variables were initialized* to facilitate the downstream workflow, *visualization of Mass-Spec profiles* performed, and lastly the *features ranked* with t-test statistic.

Loading pre-processed dataset

After preprocessing the dataset into .mat format (find the code here, the dataset was loaded.

```
load OvarianCancerQAQCdataset
whos
```

```
  Name                Size                Bytes  Class
                    Attributes

  Cidx              216x1                   216  logical

  Cvec              121x1                   968  double
```

```
  MZ              15000x1                120000  double

  N                   1x1                     8  double

  Nidx              216x1                   216  logical

  Nvec               95x1                   760  double

  Y               15000x216              25920000  double

  ans                1x10                    80  double

  feat              100x1                   800  double

  grp               216x1                 26784  cell

  hC                 10x1                     0
 matlab.graphics.chart.primitive.Line
  hN                 10x1                     0
 matlab.graphics.chart.primitive.Line
  max_C           15000x1                120000  double

  max_N           15000x1                120000  double

  mean_C          15000x1                120000  double

  mean_N          15000x1                120000  double

  min_C           15000x1                120000  double

  min_N           15000x1                120000  double

  sig_Masses        100x1                   800  double

  stat            15000x1                120000  double

  xAxisLabel          1x17                   34  char

  yAxisLabel          1x13                   26  char
```

Initializing variables

A set of vector variables, which will be used in the downstream workflow, are initialized.

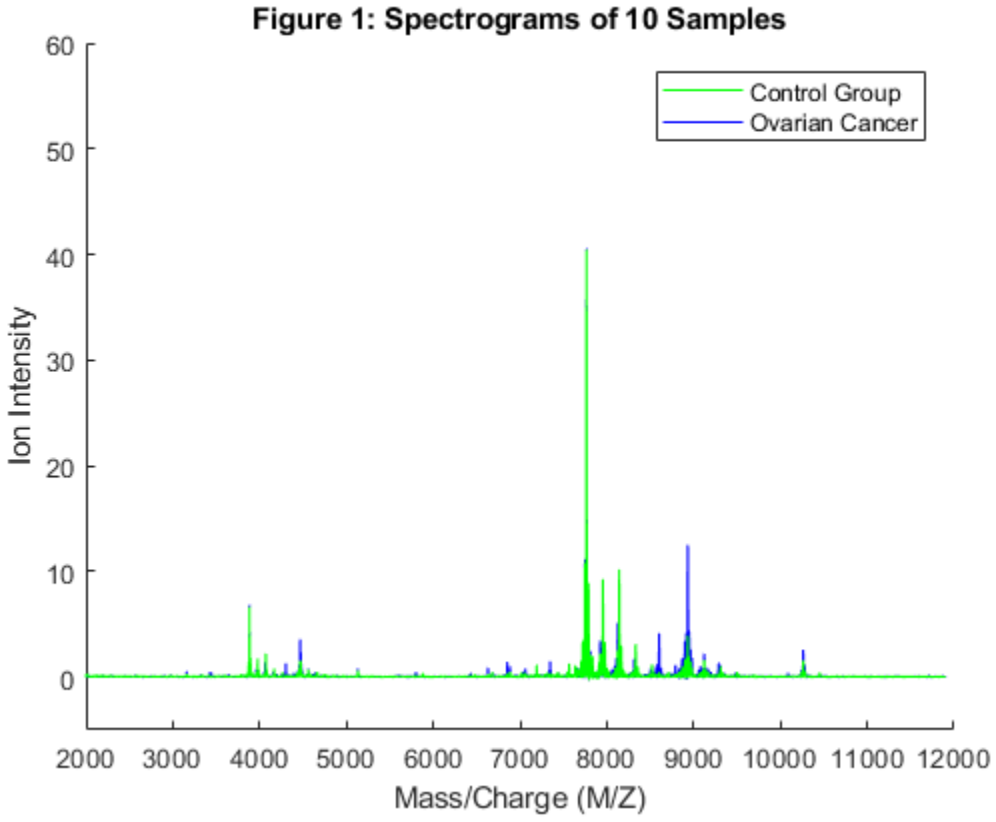
```
N = numel(grp);                          % vector of number of samples
Cidx = strcmp('Cancer',grp);             % logical index vector for
 Cancer samples' group
Nidx = strcmp('Normal',grp);             % logical index vector for
 Normal samples' group
Cvec = find(Cidx);                       % index vector for Cancer
 samples
```

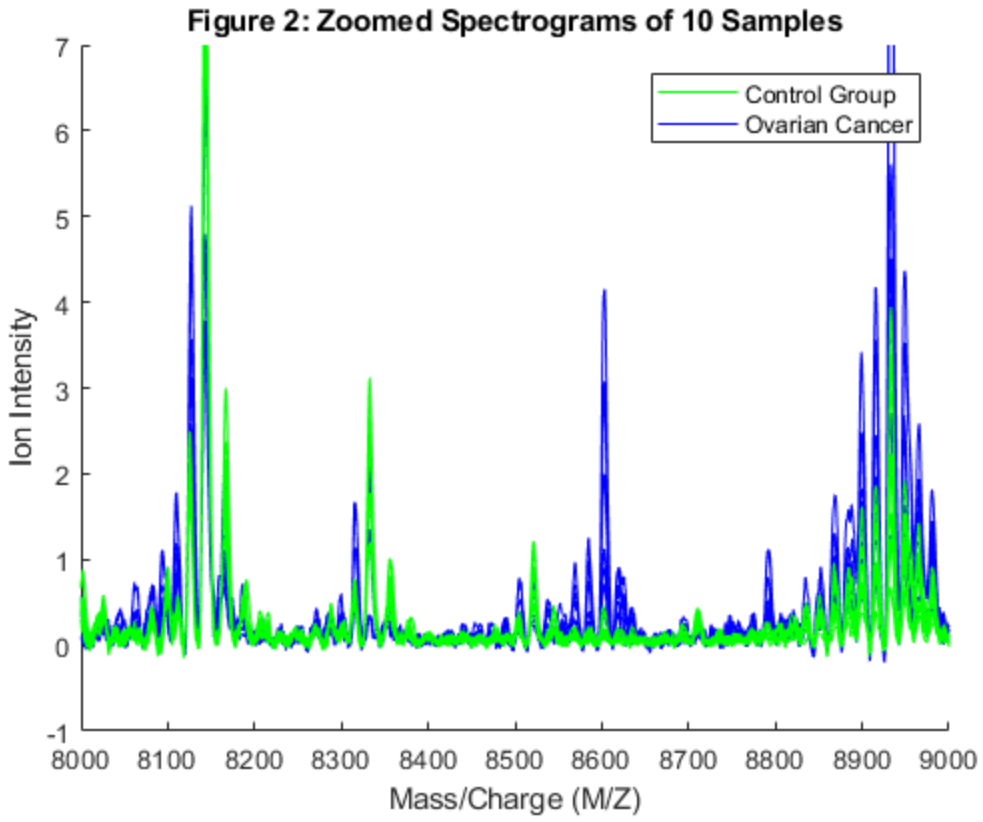
```
Nvec = find(Nidx);                          % index vector for Normal
 samples
xAxisLabel = 'Mass/Charge (M/Z)';           % x-axis label for plots
yAxisLabel = 'Ion Intensity';               % y-axis label for plots
```

Visualizing a set of the samples

Fine below the spectogram of 10 samples. Figure 1 corresponds to original spectogram, while figure 2 to a zoomed spectrogram.

```
figure; hold on;
hC = plot(MZ,Y(:,Cvec(1:10)),'b');
hN = plot(MZ,Y(:,Nvec(1:10)),'g');
xlabel(xAxisLabel); ylabel(yAxisLabel);
axis([2000 12000 -5 60])
legend([hN(1),hC(1)],{'Control Group','Ovarian Cancer'})
title('Figure 1: Spectrograms of 10 Samples')

figure; hold on;
hC = plot(MZ,Y(:,Cvec(1:10)),'b');
hN = plot(MZ,Y(:,Nvec(1:10)),'g');
xlabel(xAxisLabel); ylabel(yAxisLabel);
axis([8000 9000 -1 7])
legend([hN(1),hC(1)],{'Control Group','Ovarian Cancer'})
title('Figure 2: Zoomed Spectrograms of 10 Samples')
```





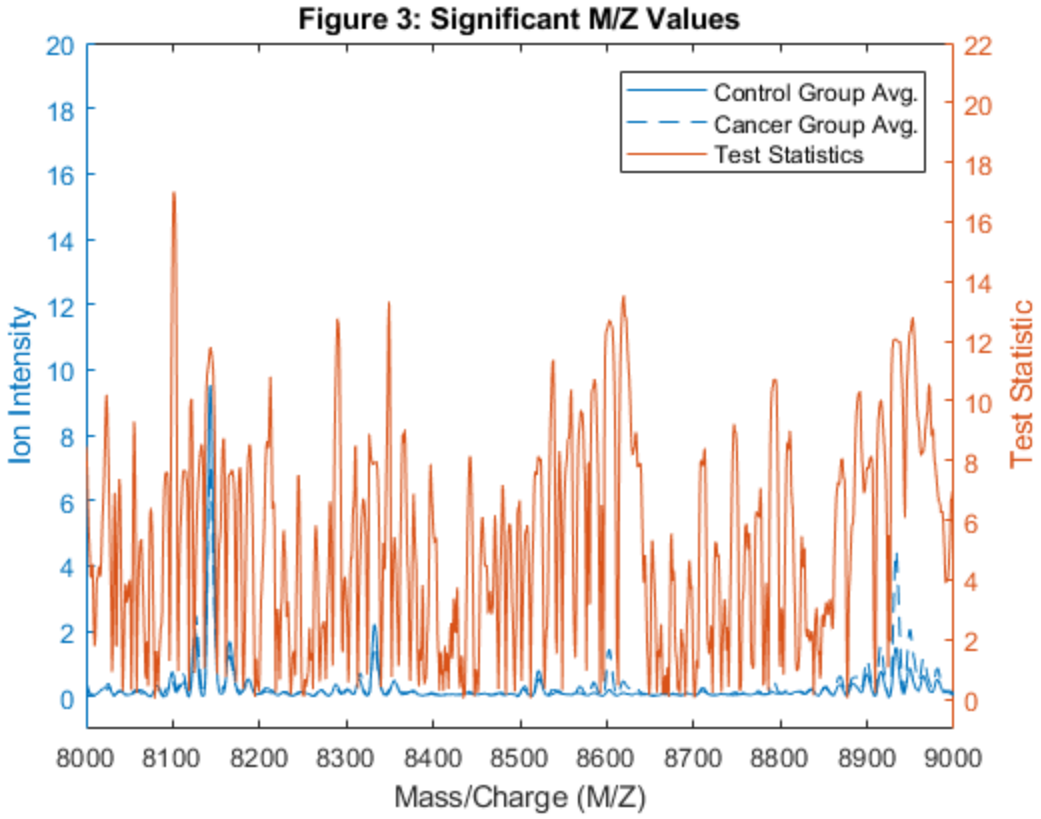
Ranking features

Significant masses were identified using a two-way t-statistic. After ranking the features, a set of variables were initialized to generate the plot (figure 3) for the spectogram with two-way t-statistic.

```
[feat,stat] = rankfeatures(Y,grp,'CRITERION','ttest','NUMBER',100);
sig_Masses = MZ(feat);
sig_Masses(1:10)' %display the first 10 significant masses

mean_N = mean(Y(:,Nidx),2);  % group average for control samples
max_N = max(Y(:,Nidx),[],2); % top envelopes of the control samples
min_N = min(Y(:,Nidx),[],2); % bottom envelopes of the control samples
mean_C = mean(Y(:,Cidx),2);  % group average for cancer samples
max_C = max(Y(:,Cidx),[],2); % top envelopes of the control samples
min_C = min(Y(:,Cidx),[],2); % bottom envelopes of the control samples

figure;

yyaxis left
plot(MZ, [mean_N mean_C]);
ylim([-1,20])
xlim([8000,9000])
title('Figure 3: Significant M/Z Values')
xlabel(xAxisLabel);
ylabel(yAxisLabel);
```

```
yyaxis right
plot(MZ,stat);
ylim([-1,22])
ylabel('Test Statistic');

legend({'Control Group Avg.','Cancer Group Avg.', 'Test Statistics'})
```

```
ans =

   1.0e+03 *

  Columns 1 through 7

    8.1009    8.1016    8.1024    8.1001    8.1032    7.7366    7.7359

  Columns 8 through 10

    7.7374    7.7253    7.7245
```



Other possible approaches

Although not performed in this example proteomic data analysis, other approaches that could have been added include:

1. identify the amino acid sequences of the statistically significant features

2. and identify the proteins by matching amino acid sequences to databases.

In future *"omics" data analysis in Matlab, as well as in R and Python more thorough and detailed workflow will be shared.

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