An Interdisciplinary Summer School on Mining of Biological Data for Master and PhD students,

Norwegian University of Life Sciences, As, Norway.

Tutorial for module 8: "Clustering", rooms U223 and U226, 15:30-16:45, Monday, 13 of August, 2018. **Mentor:** Petr V. Nazarov, Ph.D., Researcher of Proteome and Genome Research Unit, Department of Oncology, Luxembourg Institute of Health, Strassen, Luxembourg.

Online materials: http://edu.sablab.net/nmbu2018/

Data for tutorial: http://edu.sablab.net/data/txt/mRNAIFNg.txt

It contains annotated genes in rows and samples in columns, values are log2 transformed expressions.

NOTE: before starting, you may go through the code used for the lecture: <u>http://edu.sablab.net/nmbu2018/lecture.html</u>

NOTE: depending on the task you can consider samples as objects (gene expression are features then) or genes as objects (expression in samples are features).

Tasks

- 1. Install required packages (follow the online materials)
- 2. Import the data (follow the online materials)
 - a. Prepare **matrix X** with gene expressions removing lowly expressed and nonannotated features (GeneSymbol is "")
 - b. Create standardized gene expression matrix Z, so that all genes have mean = 0, st.dev. = 1
 - c. Perform and plot PCA of samples and genes (use both X and Z)
- 3. Cluster the samples (expected outputs are presented in online materials)
 - a. Use pheatmap() to make bi-cluster of genes and samples (for X and Z)
 - b. Cluster the samples using k-means or PAM and define the reasonable number of clusters. Visualize in PCA plot. Any difference when using X or Z matrices?
- 4. Cluster the genes (expected outputs are presented in online materials)
 - Use k-means or PAM methods to cluster the genes from standardized Z matrix. Visualize them on corresponding PCA plot (genes as objects, samples as features).