



## Interview exercise

## Requirements:

1. Single cell data: <a href="https://github.com/jkubis96/Interview\_exercises.git">https://github.com/jkubis96/Interview\_exercises.git</a>

2. R softwere (recommended IDE: RStudio)

2. Seurat library for analysis: <a href="https://satijalab.org/seurat/articles/install.html">https://satijalab.org/seurat/articles/install.html</a>

3. Other useful libraries: tidyverse, readxl

## **Exercise I:**

- 1. Download necessary data
- 2. Load data to R and run the analysis using the Seurat package
- 3. Conduct basic visualization: count ~ genes (scatter plot), number of genes (violin plot), percent [%] of mitochondrial genes, etc
- 4. Conduct data normalization [explain this type of normalization]
- 5. Find the most variable marker genes in the data set and visualize them by plot
- 6. Conduct scaling of the data and determine principal components (PC) [It will be necessary for the clustering]
- 7. Choose the right principle components for the next analysis (use Elbow plot, JackStraw plot, or both for visualization)
- 8. Conduct clustering using SNN/KNN algorithms based on previously chosen PCs
- 9. Run tSNE or UMAP and visualize obtained clusters
- 10. Find marker gene specific to obtained clusters based on average FC and p-val < 0.05
- 11. Select top 10 markers for each cluster based on avg\_log2FC and save into \*.xlsx format