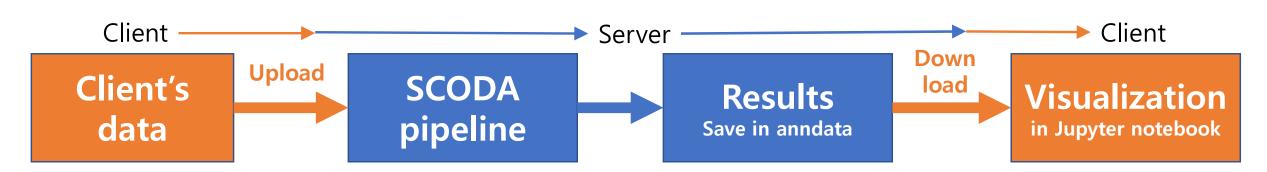
#### What is SCODA?

- Web-based, Fully-automated, all-in-one computing service for Single-Cell (transcript)Omics Data Analysis (single-cell RNA-seq)
- ☐ Especially useful for
  - O In-vivo Tissue/tumor micro environment (TME) study.
  - O Immune cell profiling in, for example, autoimmune disease & cancer
- ☐ Simply uploading your count data, you get standardized functional analysis results in ten-minutes or so (depending on your data size)
- Results include
  - O High precision cell-type annotation (major-type, minor-type and subset)
  - O Condition specific DEG and GSA results per-cell-type
  - O Possible cell-cell interactions between different cell-types per-condition
  - O CNV estimates and tumor-cell identification results



# SCODA pipeline overview (Single-Cell Omics Data Analysis pipeline)



#### **Supported format**

- 1. Compressed 10x mtx files sets + (optional) meta data (csv file)
- 2. Compressed CSV (cell-by-gene matrix)+ (optional) meta data (csv file)
- 3. Compressed h5ad file

#### **Functions**

- 1. Celltype annotation using HiCAT (v0.6.7)
- 2. DEG & GSA/GSEA per minor celltype
- 3. Inference of cell-cell interaction using CellPhoneDB (v2.1.7)
- 4. (optional) Tumor cell identification using InferCNV (v0.4.2) + icnv addon
- 5. More will be added soon

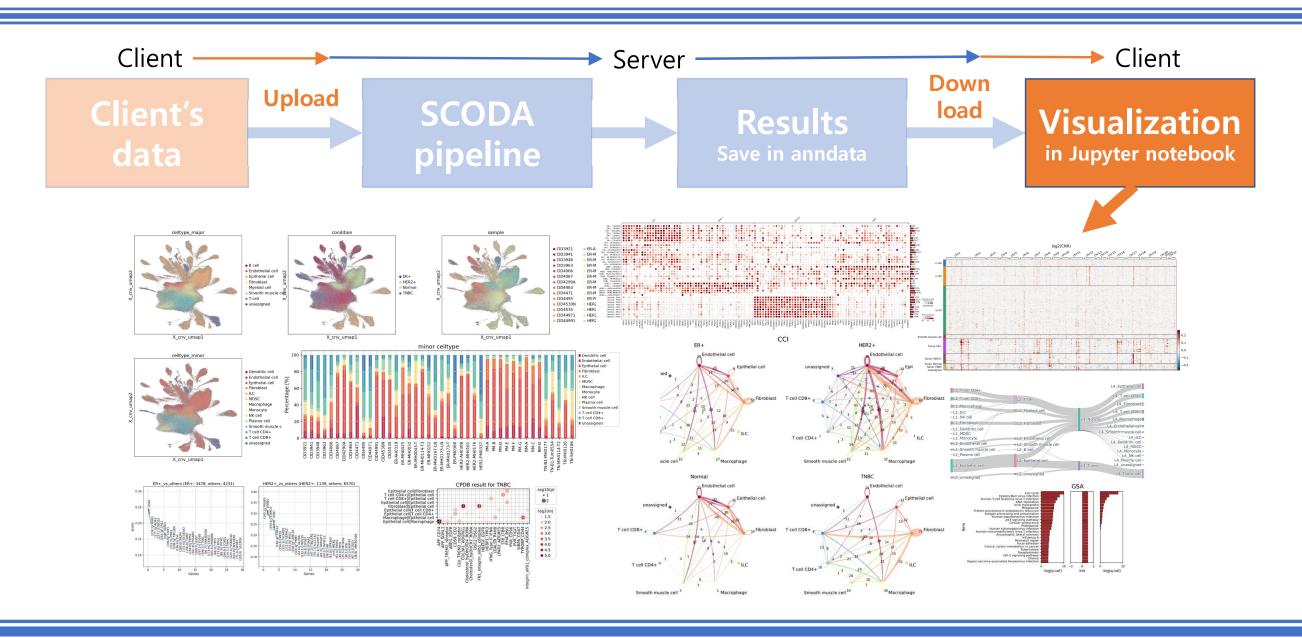
### Results saved in anndata format (v0.8.0)

- 1. Celltype annotation in obs field
- 2. DEG & GSA/GSEA results for each celltype in uns field
- 3. CellPhoneDB results in uns field
- 4. (optional) Tumor cell identification results in obs field

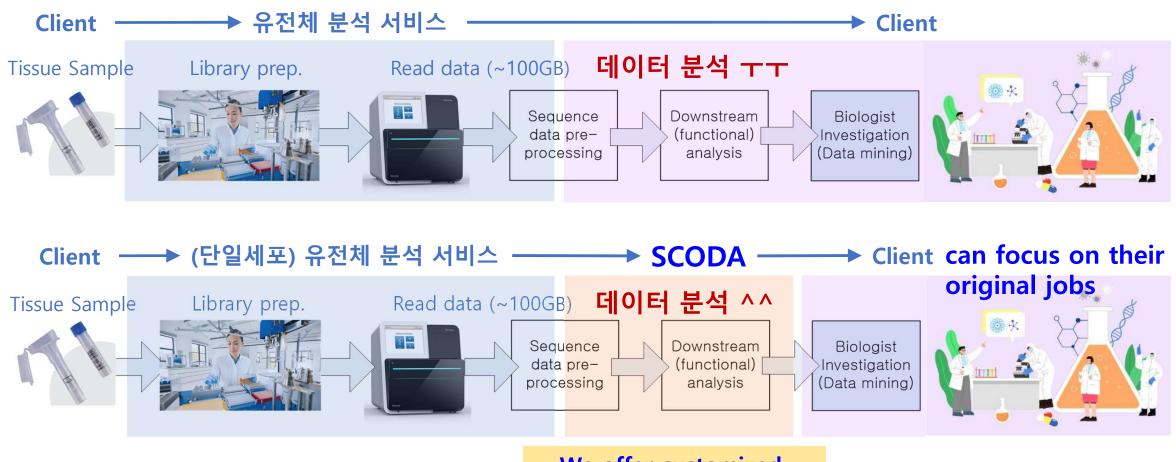
#### **Exploring the results**

- 1. Use SCANpy & SCODA-viz package (open source)
- 2. Example Jupyter notebook provided
- 3. With a little bit of programming, users can get deeper insight into the data by comparing cells in different condition

## SCODA-viz: visualization lib. (free open source)



# Why SCODA?



We offer customized functional analysis service

#### How to use SCODA?

- ☐ With your raw datasets (e.g., CellRanger v3 count data)
- 1. Go to SCODA home <a href="https://mlbi-lab.net">https://mlbi-lab.net</a> to upload and get results.
- 2. Get Jupyter notebook to explore the results: <a href="https://github.com/combio-dku/scoda\_explorer">https://github.com/combio-dku/scoda\_explorer</a>
- 3. Mine the SCODA results in Jupyter notebook on your own computer (or in Google Colab)
- 4. Contact us if you have larger datasets
- Contact: <u>mlbi.dku@gmail.com</u> for any inquiries or online workshop for using SCODA-viz

# Why SCODA?

- □ Trend in bio/medical research: convergence of data science & bio/medicine.
  □ Requires complicated communications between biologist and informatician.
  □ SCODA is an automated one-stop computing service using verified, standardized functional analysis tools for single-cell omics data.
  □ With SCODA, you don't need to spend your time to do something that are not your field of expertise, e.g.,
   for complicated computing environment setup
   develop pipelines to preprocess your datasets
- ☐ Can accelerate the research with single-cell RNA-seq data.
- The only thing you need is a little-bit of programming skill, e.g., handling data frame, for your own data mining.
- ☐ It provides you with a much higher degree of freedom for your research.

### What is SCODA?

- ☐ The computing server accepts 10x mtx format (CellRanger v3 output), csv or anndata formatted h5ad file (optionally with meta data)
- SCODA pipeline utilizes verified, suitably licensed (or free) open source SWs (HiCAT v0.6.12, CellPhoneDB v2.1.7, InferCNV v0.4.2, anndata v0.8.0)
- ☐ Free visualization library is provided for your own data mining (along with scanpy, searborn, matplotlib packages)
- ☐ You only need a little bit of programming skill (in python or R) to explore the SCODA results by yourself to mine biological implications.
- ☐ Small group workshop (training) for data mining is available.
- □ With SCODA, you can focus more on your original jobs!!! (i.e., biology and/or medical research)

### Try SCODA now

☐ Are you ready to unravel the mysteries of single-cell gene expression? And gain deeper insights into cellular heterogeneity? Look no further! Our single-cell RNA-seq data analysis pipeline was designed to empower your research with state-of-the-art computational capabilities. The visualization and data mining is fully customizable with a little bit of programming skill. ☐ You can tailor the analysis pipeline to your specific needs. With optional analysis parameter settings, you have control to adapt workflows for your specific requirement. ☐ Don't hesitate to contact us (mlbi.dku@gmail.com). We're here to help you.