### What is MEISTER?

MEISTER (Mass spEctrometry for Integrative Single cell and TissuE analysis with deep learning-based Reconstruction) is a data analysis framework that integrates a deep-learning-based reconstruction in order to accelerates high-mass-resolving MS, multimodal registration creating for 3D molecular images, and data integration of single cell mass spectra to 3D molecular images.

# Installation via Anaconda (recommended)

MEISTER is written in Python and can be installed by cloning the GitHub repository or by downloading the files. MEISTER uses several external dependences which are included the MEISTER environment. Before running the first time, install the MEISTER environment

- 1. Navigate to the MEISTER directory: cd MEISTER
- 2. Create the conda virtual environment: conda env create -f evironment.yml
- 3. Activate the virtual environment: conda activate MEISTER
- 4. Install Tensorflow separately into the environment to avoid potential conflicts, which will take a long time to solve: pip install tensorflow==2.10.1

## **Implementation**

### Training the Model

To begin, open terminal and navigate to the MEISTER repository. Be sure to activate the MEISTER environment.

```
conda activate MEISTER
```

Model training is performed via the script train\_sigma\_mode.py which is implemented using argparse for flexible implementation. To define model parameters, see examples in train\_prompt.txt, i.e.,

```
python train_signal_model.py --train_ROI ROO RO1 RO2 --path_file file_dir_coronal_3D_train.
    json --batch_size 256 --epochs_encoder 10 --epochs_regressor 30 --latent_dim 32
```

We will provide examples for how each of the above parameters can be set.

Starting with train: In this example, R00, R01, R02 are the images obtained at high-resolution. To obtain these labels, navigate to the Bruker .d file containing the high-resolution data. Open the XML Source File. From here, labels can be manually identified as shown:

The image source file will contain info for every FID obtained, so be sure to scroll to the bottom of the file to get labels for every image.

Next, navigate to your path\_file, here titled file\_dir\_coronal\_3D\_train.json

Specify your desired name for the project, and input the required parameters.

Next, we can specify the desired batch size, number of epochs for the encoder and regressor, and the number of latent dimensions. Note that omitting these parameters will use the default numbers as specified in train\_signal\_model.py. The batch size should be a power of two. The number of epochs can be increased or decreased during optimization if the model is underfit or overfit, respectively. Once all parameters are set, execute the training prompt in the terminal. For best performance, we recommend loading data from a SSD and running tensorflow on a GPU.

#### Reconstruction

When the encoder and regressor have completed, we can run the reconstruction which is executed with the bash script run\_deepmsi\_3Dcoronal\_slide1\_2.sh. Open this file to set the reconstruction parameters

```
MESTER > $ run_deepmais JOcoronal_slide1_2_lsh

# // Introduction | # // Introduction
```

Both decoder\_dir and regressor\_dir will be written to processed\_data when the training is complete. Like the high-resolution ROI, the low-resolution ROI labels can be found from the XML source file for the low-resolution data. Execution of this bash script in the terminal will run deep\_recon.py with the user-input parameters.