## **Quality Control Processing Preprocessing** 1. Normalize data 1. Checking the percentage of peptides normalyse() in library detected 1. Preprocessing proteomics 2. Remove outliers - qc\_check() - outliers() - prepare peaks() 3. Data Imputation 2. Identifying the components of - prepare\_pd() impute() Peptide Library Not Detected. 4. Cleavage motif determination mspms() - find\_nd\_peptdes() - join\_with\_library() cterm\_cleavage() 3. Visualizing QC Results add cleavages() 2. Preprocessing peptide library - nterm\_cleavage() - plot\_qc\_check() 5. Consolidating data - calculate\_all\_cleavages() - plot\_nd\_peptides() - prepare\_for\_stats()

## **Statistics / Visualization Preparation**

## 4. Count of cleavages by position

- plot\_rt\_qc()

- count\_of\_cleavages\_per\_pos()
- 5. iceLogo

1. Log2 fold change

2. T- tests

3. Anova

- mspms\_log2fc()

mspms\_t\_test()

-log2fct condition()

- mspms\_anova()

-log2fc\_t\_test()

-log2fct time()

3. Log2 fold change + t-tests

- calc\_AA\_count\_of\_motif()
- calc\_prop\_of\_motif()
- calc AA motif zscore()
- calc\_sig\_zscores()
- calc\_percent\_difference()
- calc\_AA\_fold\_change()
- prepare\_sig\_p\_diff()
- prepare\_pd()
- prepare\_fc()
- extract\_re()

prepare\_icelogo\_data()

## **Visualizations**

- 1. Principle component analysis
- plot\_pca()
- 2. Heatmap/unsupervised hierarchical clustering
  - plot\_heatmap()

- polish()

- 3. Volcano plot
  - plot\_volcano()
- 4. Cleavage position plot
  - plot\_cleavage\_pos()
- 5. Icelogo plots
  - plot\_icelogo()

  - plot\_all\_icelogos()

Integrated in Integrated in Shiny App generate\_report()