- « Classical » analysis
- Clusters analysis
- Optional step
- TODO →
 modifications to
 do
- RNA-seq data

Pre-processing:

- Demultiplexing
- Quality reports
- · Filtering of reads with bad IQF
- Filtering of PCR duplicates
- Extract Unique Molecular Identifier TODO
- · Adapters trimming
- Size selection

For each sample

Pre-processing:

- Pool all sequences / sample
- Pool all sequences over all samples

Mapping:

- Filtering of reads mapped to rRNA
- · Mapping to reference genome
- · Filtering of multi-reads
- Mapping output sorted & indexed

All sequences availables over all samples

PCR duplicates removing TODO

Metagene analysis

P-site offsets analysis

Multi-mapped annotations clustering

Triplet-periodicity analysis TODO

<u>Isoform-level</u> estimation

Expression estimation considering:

- Longest CDS
- Splicing-isoform-level assessing
- Clusters

Differenretial analysis:

- Genes/Clusters translation
- Regulation level