

Input files
FASTQ
unaligned BAM

Configure Parameters

Compile Run Directives

- *Input*
- *Metadata*
- *Tokenization*
- *Barcode sets*
- *Prior normalization*
- *Output*

Validate Configuration

For each read

Prepare Segment Output

- *Extract tokens*
- *Assemble transforms*

For each barcode

(sample, cellular, molecular)

PAMLD

or

MDD

or

Other

Output files

- *Classified*
 - *Unclassified*
- FASTQ/BAM/CRAM*

**Demux
Report**
JSON