SUBTYPING PIPELINE

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Quick Guide

Important!

Before you start the pipeline, make absolutely sure that all file names, sequence names, structures of input tables are correct and **follow exactly the patterns** as described in examples.

Pipeline Directory

Locate to Pipeline directory:

```
$ cd ~/rki_subtyping/Pipeline
```

Open IDE (Visual Code Studio)

```
$ code ../
```

Input Folders

Make sure you have 5 directories:

AllSeqsCO20 Folder

Provide this folder with .xlsx files as listed (from NGS pipeline):

```
AllSeqsC020/

— MS95_Seqs_ENV_C020_V5.xlsx

— MS95_Seqs_INT_C020_V5.xlsx

— MS95_Seqs_PRRT_C020_V5.xlsx
```

Refer to Supplementary part of the guide, if you need more information (Page 21).

InputFasta Folder

Provide this folder with files as listed (from NGS pipeline):

```
InputFasta/
— MS95_ENV_20.fasta
— MS95_INT_20.fasta
— MS95_PRRT_20.fasta
```

Refer to Supplementary part of the guide, if you need more information (Page 22).

Conda Environment

Activate subtyping_pipeline environment.

\$ conda activate subtyping_pipeline

Be sure you have change in prompt:

(subtyping_pipeline) beast2@Beast2:~/rki_sybtyping/Pipeline\$

Pipeline with --outdir parameter

Parameter —outdir determines a name of an ouput folder. The command will generate four enumerated output folders within Results folder. Without specifying an output folder you can get a warning message.

\$ nextflow Scripts/subtyping_pipeline.nf --outdir Results

ManualRega Folder (1)

Provide the folder with .csv files (separator: comma) generated by Rega using marked .fasta files from the folder procuded by the pipeline:

```
~/rki_subtyping/Pipeline/Results/1_marked_fasta
```

These .fasta files have M at the end of the file name:

ManualRega Folder (2)

Rega online tool always generates files with the same name results, e.g. results.csv.

Rename these files accordingly, using the pattern as in the example below:

```
ManualRega/
— Manual_Rega_MS95_ENV_20M.csv
— Manual_Rega_MS95_INT_20M.csv
— Manual_Rega_MS95_PRRT_20M.csv
```

Refer to Supplementary part of the guide, if you need more information (Page 23).

Pipeline with --fullpipeline parameter

Repeat the previous command with ——fullpipeline parameter and —resume flag. The latter allows for generating an output up to 12_mafft folder. The complete processes are cached.

```
$ nextflow Scripts/subtyping_pipeline.nf --outdir Result --fullpipeline -resume
```

Check the output of 12_mafft folder before you run iqtree analysis (msa files - multiple sequence alignments)!

Pipeline with --iqtree parameter

Parameter —iqtree allows for running the iqtree process that produces 13_iqtree folder within Results. The folder contains .iqtree, .treefile, and .log files. The parameter can be added at this point, as the last command with report and plot outputs being produced or not added at all (no 13_iqtree folder then).

```
$ nextflow Scripts/subtyping_pipeline.nf --outdir Results --fullpipeline --iqtree -resume
```

You can monitor the log file while running iqtree within work folder using respective process ID (68), e.g [68/72f0eb].

Decision

Manually modify files (see below) which contain Manual tag in PRRT_Subpype, INT_Subtype, and ENV_Subtype columns. Save changes and close xlsx files.

```
9_joint_with_tags/
— full_MS95_ENV_20M.xlsx
— full_MS95_INT_20M.xlsx
— full_MS95_PRRT_20M.xlsx
```

Report and Plot

```
$ nextflow Scripts/subtyping_pipeline.nf --outdir Results --fullpipeline --iqtree -resume
```

Repeating the command above generates 14_report folder with MS95_subtype_uploads.xlsx report file.

Repating it again generates a MS95_subtype_counts.png plot and adds it to the 14_report folder.

Clean Up

Once the pipeline has generated Results folder with all desired output files (save all needed outputs first), the input files can be removed from the input folders.

```
$ rm -rf InputFasta/* AllSeqsC020/* ManualRega/* Results/
```

The same is true for dot and nextflow temp files/folders:

```
$ rm -rf .nextflow work .nextflow.log .nextflow.log.*
```

Processes Overview

```
[1b/f2f10a] process > mark fasta (2)
                                                 [100%] 3 of 3, cached: 3 \checkmark
                                                 [100%] 3 of 3, cached: 3 ✓
[a9/dd644a] process > get tags (3)
                                                 [100%] 3 of 3, cached: 3 \checkmark
[97/70bbdd] process > comet (3)
[73/a28f41] process > stanford (3)
                                                 [100%] 2 of 2, cached: 2 \checkmark
                                                 [100%] 2 of 2, cached: 2 \checkmark
[e6/e4af1d] process > json to csv (3)
[65/e0eb90] process > clean rega (3)
                                                 [100%] 3 of 3, cached: 3 \checkmark
                                                 [100%] 1 of 1, cached: 1 \checkmark
[87/4d2fcf] process > join env (1)
[34/36991e] process > join int (1)
                                                 [100%] 1 of 1, cached: 1 \checkmark
[62/e59285] process > join prrt (1)
                                                 [100%] 1 of 1, cached: 1 \checkmark
[8d/dad394] process > make decision (1)
                                                 [100%] 1 of 1, cached: 1 \checkmark
[d8/983216] process > join with tags
                                                 [100%] 1 of 1, cached: 1 \checkmark
[e6/ceaa42] process > fasta for mafft (2)
                                                 [100%] 3 of 3, cached: 3 \checkmark
                                                 [100%] 1 of 1 cached: 1 ✓
[54/89322b] process > env concat panel (1)
[a4/b7aaee] process > int concat panel (1)
                                                 [100%] 1 of 1, cached: 1 \checkmark
[f7/9e1ccf] process > prrt concat panel (1)
                                                 [100%] 1 of 1, cached: 1 \checkmark
[c0/786bcd] process > mafft (3)
                                                 [100%] 3 of 3, cached: 2 \checkmark
[68/72f0eb] process > igtree (3)
                                                 [100%] 3 of 3, cached: 3 \checkmark
[3c/0fb71f] process > report
                                                 [100%] 1 of 1, cached: 1 \checkmark
[c5/462a18] process > countplot (1)
                                                 [100%] 1 of 1 🗸
```

Supplementary

Example of .xlsx within AllSeqsCO20

Example of .fasta within InputFasta

>20-02955_ENV_20

Example of .csv within ManualRega

An example of a csv file produced by Rega online tool (names of columns and only one sample for demonstration)

```
"name","length","assignment","rule","support","begin","end","type","pure",
"pure_support","pure_inner","pure_outer","scan_best_support","scan_assigned_support",
"scan_assigned_nosupport","scan_best_profile","scan_assigned_profile","crf",
"crf_support","crf_inner","crf_outer","crfscan_best_support",
"crfscan_assigned_support","crfscan_assigned_nosupport","crfscan_best_profile",
"crfscan_assigned_profile","major_id","minor_id"
"20-02944_PRRT_20","1026.0","HIV-1 CRF 06_CPX","4","98.0","1823.0","2848.0","Human
immunodeficiency virus 1","HIV-1 Subtype G","93.0","0.0","93.0","0.5","0.357","0.643",
"G K A1 A1 A1 A1 G A1 A1 G G G G G","G - - - - - - G G G G -","HIV-1 CRF 06_CPX",
"98.0","0.0","98.0","1.0","1.0","0.0","06_CPX 06_CPX 06_CP
```

References Folder

This folder contains reference panels and does not need any change unless reference panels should be replaced.

```
References/
— Reference_ENV_Panel_Stanford.fas
— Reference_INT_Panel_Stanford.fas
— Reference_PRRT_Panel_Stanford.fas
```

Scripts Folder

This folder contains the scripts and does not need any change.

```
Scripts/
  — comet_rest.py
  – decision.py
   fasta_for_mafft.py
   full_join.py
   json_parser.py
   nexflow.config
   plot.py
   rega_cleanup.py
    repeat_marking.py
   report.py
    subtyping_pipeline.nf
    tag parser.py
```

Conda

Conda Info

List available conda environments.

Conda Version

Pipeline's version of conda 4.14.0

\$ conda --version

Deactivation of Environment

This command is used to deactivate the current invironment.

\$ conda deactivate

Be sure you have change in prompt:

(base) beast2@Beast2:~/rki_sybtyping/Pipeline\$

GitHub Repo

Repo Link

The project is hosted here. Use this link to clone the repo in case of data loss.

How to Clone

Locate to home directory

\$ cd

Clone the repo

\$ git clone https://github.com/vera-rykalina/rki_subtyping

Modify path of ProjectDir within subtyping_pipeline.nf

projectDir = "/home/beast2/rki_subtyping/Pipeline"

Notes

Keep in Mind (1)

- The pipeline does not take into account subsubtypes. If there are subsubtypes they are converted to subtypes. For instance, A1 is converted to A, F2 is converted to F etc.
- The pipeline does not perform a full quality check of

 fasta sequences. Illegal characters should be excluded
 (the pipeline takes care only of underscores so far).

 Sequences with illegal characters are not accepted by Rega online tool.

Keep in Mind (2)

Make sure that sample names do not exceed 30 characters in length. Long sample names get shorten by Rega online tool that can cause issues. E.g, this sequence name PK105_F482_23_MiS84_S86_20consensus_PRRT_20 is too long and gets shortern by Rega to PK105_F482_23_MiS84_S86_20cons. In such cases a manual change is necessary.

Keep in Mind (3)

- Be sure you are connected to the Internet
- You can always delete the whole Results folder or individual subfolder/subfolders within Results and repeat the commnand with -resume.