# VitisOmics

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# 1 Overview

This document contains the documentation for the "VitisOmics" project. This project aims at handling "omics" data in the genus Vitis (e.g. grapevine) in an open and reproducible way.

Such data are available from various places, Genoscope, URGI, NCBI, EBI, etc, and several committee from the IGGP (International Grape Genome Program) strive at improving interoperability. But my attempt, via the usage of git and GitHub, could prove for the community to be a useful addition to these efforts.

The project directory is organized as advised by Noble (PLoS Computational Biology 2009):

On any Unix-like system, it is easily done with the following commands:

```
touch AUTHORS COPYING README; mkdir -p doc data src bin results
```

On any Unix-like system, it can also be easily compressed and transferred (ignoring large data files):

```
cd ..; tar -czvf VitisOmics.tar.gz \
--exclude=VitisOmics/data --exclude=VitisOmics/results \
--exclude="*" --exclude=".*" VitisOmics
```

#### 1.1 Contributors

As of today: Timothée Flutre, Charles Romieu, Gautier Sarah

## 2 Data

```
cd data/
```

TODO: retrieve genome data from other cultivars than PN40024, e.g. Sultanina and Tannat

#### 2.1 URGI

• https://urgi.versailles.inra.fr/Species/Vitis/Data-Sequences/Genome-sequences

```
../../src/download_urgi.bash
```

When needed, the script decompresses zip files and compress them again but with gzip instead.

## 2.2 NCBI

- http://www.ncbi.nlm.nih.gov/genome/401
- ftp://ftp.ncbi.nlm.nih.gov/genomes/Vitis\_vinifera/

```
../../src/download_ncbi.bash
```

Note the important file scaffold\_names which provides the correspondence between original scaffold names (i.e. from the sequencing center) and various NCBI identifiers (RefSeq, GenBank, etc).

#### 2.3 EBI

12X.0 as well as soft-masking by RepeatMasker

```
../../src/download_ebi.bash
```

# 3 Results

```
cd results/
```

On a computer cluster, indexed files could be copied for usage by everyone, e.g. in /Genomics/Vitis if on the CIRAD cluster of the SouthGreen platform.

One needs to keep the info about the source (URGI or NCBI) because differences in terms of N spacers and additional scaffolds (from Aegilops) at the NCBI.

#### 3.1 **URGI**

```
cd urgi/
```

## 3.1.1 Reformat sequence headers for VITVI\_PN40024\_12x\_v0\_chroms\_URGI

Launch script:

Check:

```
zcat VV_chr12x.fsa.gz | wc -1 # 8240706
zcat VV_chr12x.fsa.gz | grep -c ">" # 33
zcat VITVI_PN40024_12x_v0_chroms_URGI.fa.gz | wc -1 # 8240706
zcat VITVI_PN40024_12x_v0_chroms_URGI.fa.gz | grep -c ">" # 33
diff <(zcat VV_chr12x.fsa.gz) <(zcat VITVI_PN40024_12x_v0_chroms_URGI.fa.gz)</pre>
```

Only the headers differ, not the sequences, so everything is fine.

Basic stats:

```
zcat VITVI_PN40024_12x_v0_chroms_URGI.fa.gz | md5sum # eff315994fafe35333462b9595e10ce5
```

## 3.1.2 Reformat sequence headers for VITVI\_PN40024\_12x\_v0\_scaffolds\_EMBL\_r102

Launch script:

Check:

Only the headers differ, not the sequences, so everything is fine.

Basic stats:

```
zcat VITVI_PN40024_12x_v0_scaffolds_EMBL_r102.fa.gz | md5sum # 4
fa2432d7a66c019c7cb41ee4d0cb7bc
```

3.1.3 Reformat sequence headers for VITVI\_PN40024\_12x\_v0\_contigs\_EMBL\_r102

TODO

3.1.4 Format VITVI\_PN40024\_12x\_v0\_chroms\_URGI for BLASTn

TODO: change Vvin to VITVI

```
../../src/format_Vvin-PN40024-12x-chr_blastn.bash
```

3.1.5 Index VITVI\_PN40024\_12x\_v0\_chroms\_URGI for BWA

TODO: change Vvin to VITVI

```
../../src/index_Vvin-PN40024-12x-chr_bwa.bash
```