

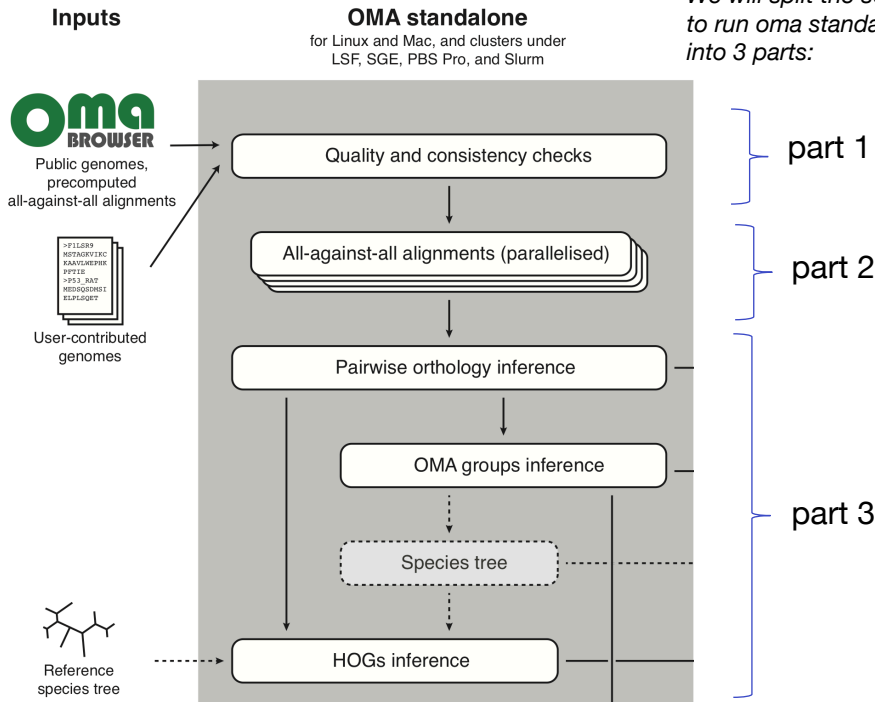
OMA Standalone CHEAT SHEET

OMA Standalone is a software for identifying homologs from complete genomes.

See <https://omabrowser.org/standalone/> for many more details!

Use this cheat sheet if:
 -You have your own sequenced genomes
 -You need to run it on the cluster (wally or axiom)

1. How it works



We will split the scripts to run oma standalone into 3 parts:

2. Get setup:

- Connect to cluster: `ssh <user>@wally-front1.unil.ch`
- Download software: `wget https://omabrowser.org/standalone/OMA.latest.tgz`
 - alternatively, install with homebrew: `brew tap brewsci/bio; brew install oma`
- Untar: `tar xvzf OMA.latest.tgz`
- Change directory: `cd OMA.latest`
- `mkdir logs`

3. Prepare genomes:

- Make sure all of your genomes meeting the following requirements:
- fasta files (1 for each genome)
 - protein sequences
 - the name of each fasta file is the name of the genome
 - all files must end in ".fa"
 - copy all genome fasta files into `OMA.latest/DB/`

4. Edit parameters.drw:

- Mostly can leave defaults
- Mandatory: specify outgroup species
- Optional: specify species tree (if not, the estimated species tree **must** be verified in the Output*!)

5. Prepare scripts:

oma_part1.sh

```
#!/bin/bash
#SBATCH --partition=wally
#SBATCH --nodes=1
#SBATCH --ntasks=1
#SBATCH --cpus-per-task=1
#SBATCH --mem=2GB
#SBATCH --job-name=oma1
#SBATCH --output=logs/oma1-%J.log
#SBATCH --export=None
#SBATCH --error=logs/oma1-%J.err

cd <full_path_to_OMA.latest>
./bin/oma -c
```

This is the database conversion part

oma_part2.sh

```
#!/bin/bash
#SBATCH --array=1-500
#SBATCH --partition=wally
#SBATCH --time=2:00:00
#SBATCH --nodes=1
#SBATCH --ntasks=1
#SBATCH --cpus-per-task=1
#SBATCH --mem=2GB
#SBATCH --job-name=oma2
#SBATCH --output=logs/%A.%a.log
#SBATCH --export=None
#SBATCH --error=logs/%A.a%.err

cd <full_path_to_OMA.latest>
export NR_PROCESSES=500
./bin/oma -s -W 7000
if [[ "$?" == "99" ]]; then
scontrol requeue /
${SLURM_ARRAY_JOB_ID}_${SLURM_ARRAY_TASK_ID}
fi
exit 0
```

This is the all-against-all, hence it is split into 500 parallelized jobs

oma_part3.sh

```
#!/bin/bash
#SBATCH --partition=wally
#SBATCH --nodes=1
#SBATCH --ntasks=1
#SBATCH --cpus-per-task=1
#SBATCH --mem=50GB
#SBATCH --job-name=oma3
#SBATCH --output=logs/%x_%J.log
#SBATCH --export=None
#SBATCH --error=logs/%x_%J.err

cd <full_path_to_OMA.latest>
./bin/oma
```

This is the orthology inference part

6. Run scripts:

- Run 1 at a time!
- `sbatch oma_part1.sh`
- `sacct` to check status
- If failed, check in `/logs`

7. Output:

Map-SeqNum-ID.txt
 OrthologousMatrix.txt
 OrthologousGroupsFasta
 OrthologousGroups.orthoxml
 OrthologousGroups.txt

Orthologous Groups, i.e. OMA Groups: useful for phylogenetic trees

OrthologousPairs.orthoxml
 PairwiseOrthologs/
 PhyleticProfileHOGs.txt
 PhyleticProfileOMAGroups.txt

useful for phylogenetic profiling

pairwise orthologs; useful for comparing 2 genomes

**Verify estimated species tree with phylo.io*

all HOGs at diff. taxonomic levels

EstimatedSpeciesTree.nwk
 EstimatedSpeciesTree.phyloxml
 HierarchicalGroups.orthoxml
 HOGFasta/
 Root HOGs i.e. gene families