iDiv

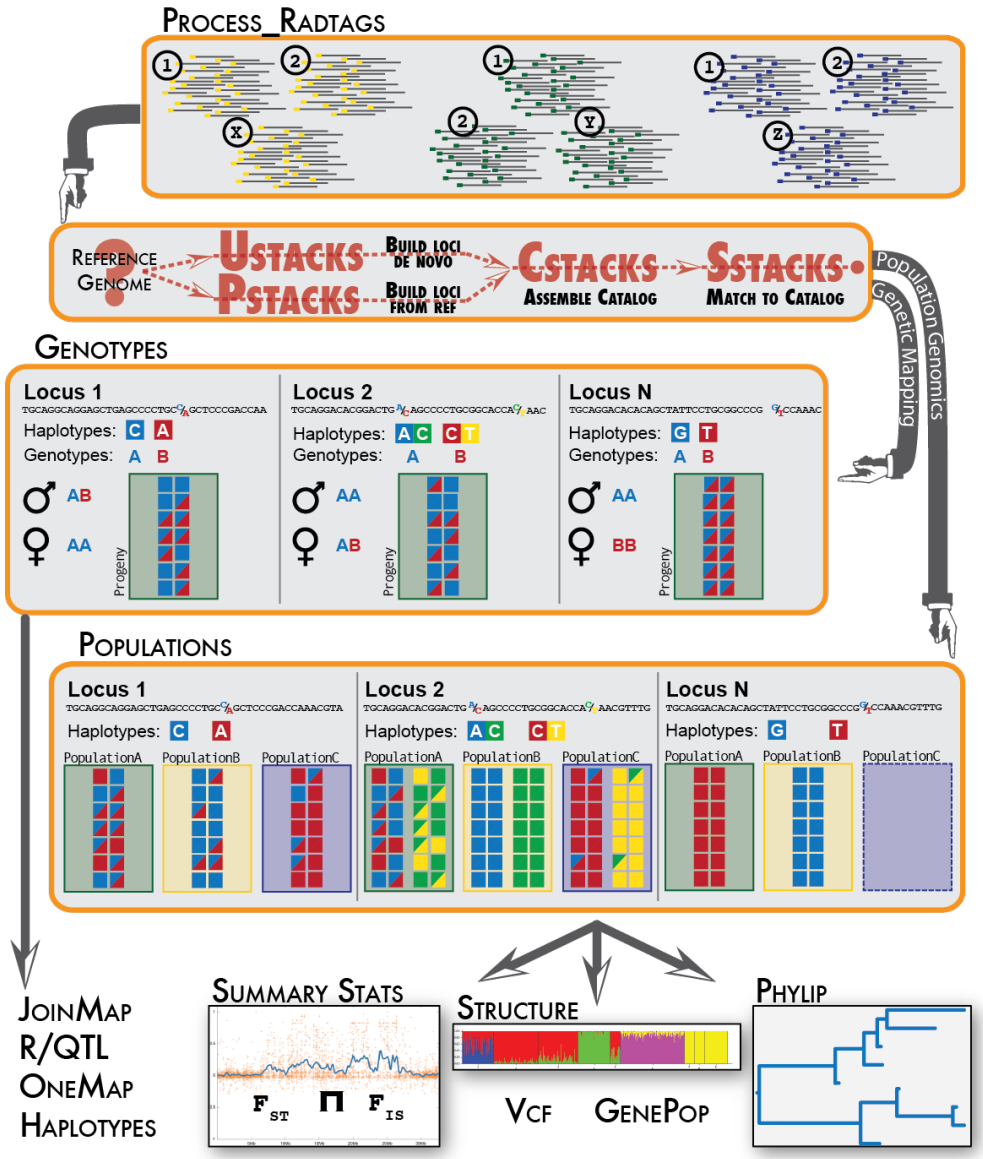
Estimating genetic diversity and population information from short read (ddRAD-seq) type data

03 – denovo_map

Chris Barratt (sDiv / Evolution and Adaptation)

Laura Mendez (Evolution and Adaptation)

Assistant: Dimas Calderon (Evolution and Adaptation)



Rochette, Rivera-Colón & Catchen, 2019 - Mol. Ecology

ddRAD sequencing

ddRADseq Data

HPC cluster (EVE) - UFZ

Stacks v2.61

process_radtags	Clean and demultiplex the data
ustacks	Building loci *de novo* for each sample
cstacks	Creates a *catalog* of all loci across the populations according to sequence similarity
sstacks	Match each sample against the catalog
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gstacks	A contig is assembled from pair-end reads and overlapped with the single-end locus. SNP calling
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Output files for population structure analyses, phylogenetics, demographic history…

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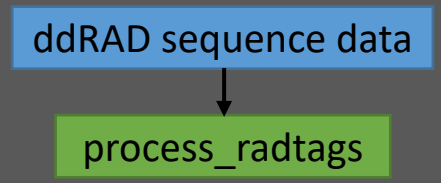
Bioinformatics Workflow



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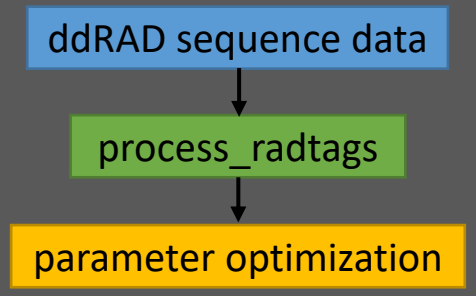

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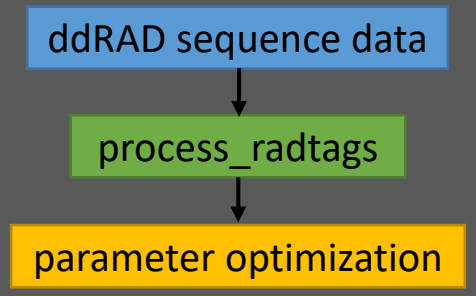

Methods in Ecology and Evolution

BRITISH ECOLOGICAL SOCIETY

Methods in Ecology and Evolution 2017, **8**, 1360–1373 doi: 10.1111/2041-210X.12775

Lost in parameter space: a road map for STACKS

Josephine R. Paris[1], Jamie R. Stevens[1] and Julian M. Catchen*[,2]

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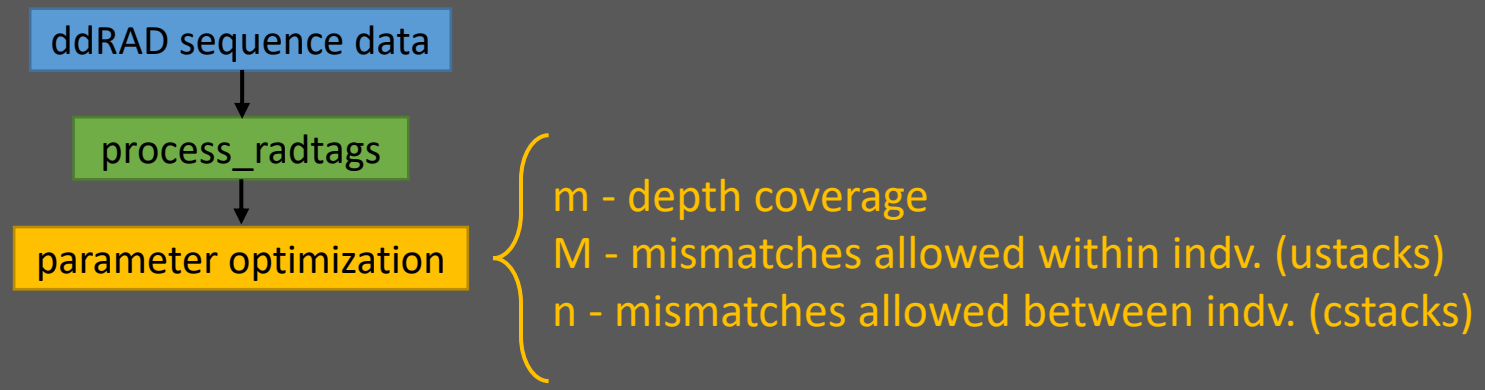

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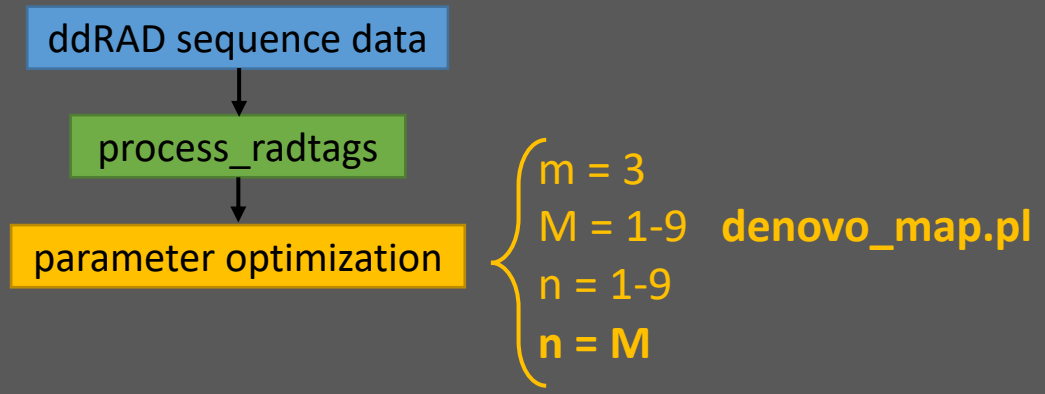

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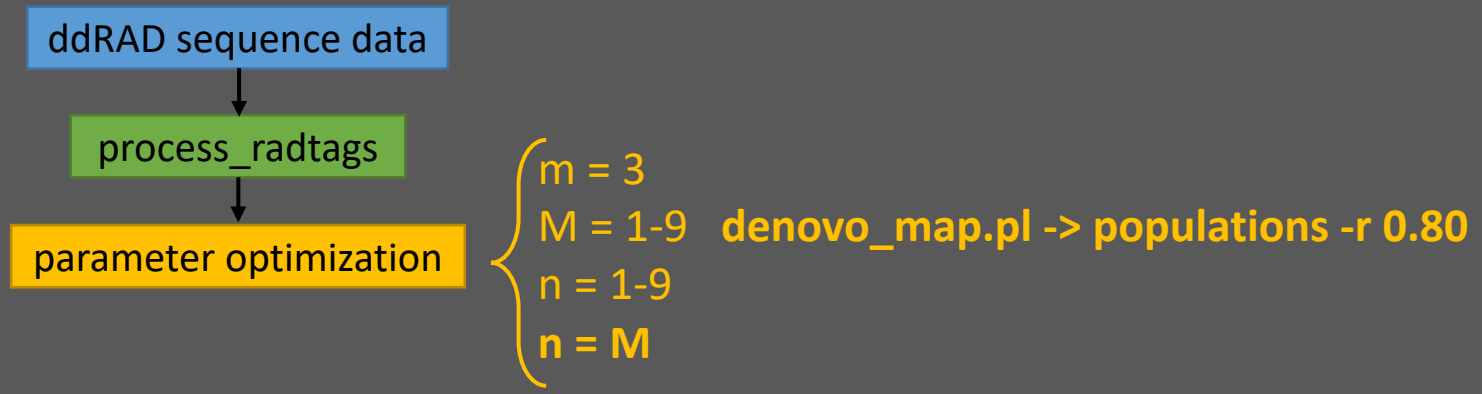

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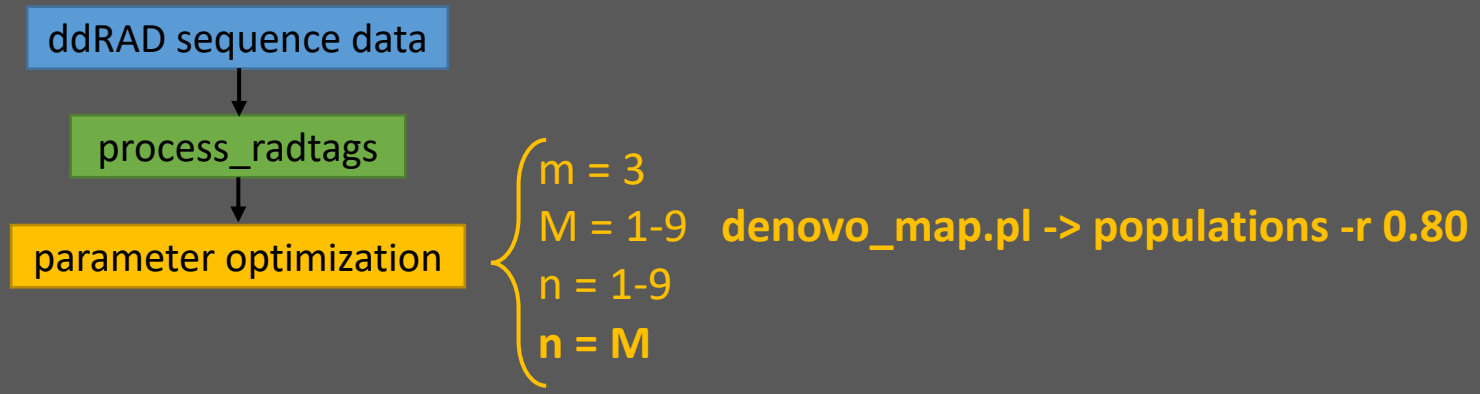
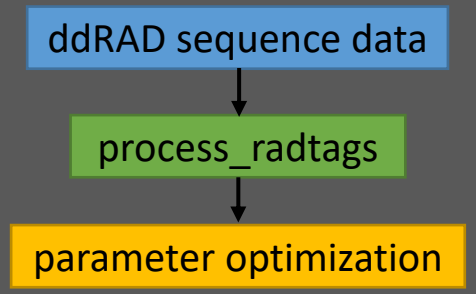


"a locus must be found in 80% of individuals of a single population to be processed"

Bioinformatics Workflow



- Number of r80 polymorphic loci
- Number of new polymorphic found for each iteration of M

01_denovo_map_test.parameters

```
#!/bin/bash

#SBATCH -J denovo_map_test.parameters
#SBATCH --mail-user=YOUREMAIL@gmail.com
#SBATCH --mail-type=BEGIN,END,FAIL,TIME_LIMIT
#SBATCH --output=/work/%u/%x-%j.out
#SBATCH --error=/work/%u/%x-%j.err
#SBATCH --cpus-per-task=20
#SBATCH --mem-per-cpu=8G
#SBATCH -t 48:00:00

# Set the requested number of cores to the number of Threads your app should use
export OMP_NUM_THREADS=${SLURM_CPUS_PER_TASK:-1}

# Paths and filenames for this analysis

M_values="1 2 3 4 5 6 7 8 9"

WORK_DIR="/work/$USER/ddRAD-seq_workshop"
popmap="$WORK_DIR/data/Exercise_3/popmaps/test.popmap.txt"
OUT_DIR="$WORK_DIR/outputs/Exercise_3/test.denovo"

mkdir "$OUT_DIR"

# Create subdirectories
cd "$OUT_DIR" || exit

for M in $M_values
do
    mkdir stacks.M"$M"
done

## Load modules and activate software

module purge
module load Anaconda3
source activate /gpfs0/global/apps/stacks_2.61

# denovo_map.pl - it will execute the Stacks pipeline by running each of the Stacks components individually: ustacks, cstacks, sstacks, tsv2bam, gstacks and populations.
# We are doing this to select the parameters M (ustacks) and n (cstacks) which optimal value depends on the amount of genetic diversity within the species and with the quality of the raw data as well.
# Therefore this has to be done with every species separately, with only a subset of samples from all the populations. This subset is written in the test.popmap files and therefore Stacks will only
# run the analyses over those samples specified. We will vary M and n (M=n) from 1 to 9, and set m = 3.

# -samples = file path to the samples (samples will be read from population map)
# --popmap = file path to the population map (<sample name><TAB><population>)
# -o = file path to write the pipeline output files
# -X = additional options for specific pipeline components, e.g. -X "populations: --min-maf 0.05". We will run populations separately afterwards
# -M = number of mismatches allowed between stacks within individuals (for ustacks)
# -n =number of mismatches allowed between stacks between individuals (for cstacks)
# -m = Minimum depth of coverage required to create a stack (default 3)
# --paired = after assembling RAD loci, assemble contigs for each locus from paired-end reads
# --rm-pcr-duplicates = remove all but one set of read pairs of the same sample that have the same insert length
# -r = minimum percentage of individuals in a population required to process a locus for that population (for populations; default: 0)
# -T = the number of threads/CPUs to use (default: 1)

# Run denovo_map on the subset of samples told by the popmap

for M in $M_values
do
    out_dir="$OUT_DIR/stacks.M$M"
    reads_dir="$WORK_DIR/data/Exercise_3/demultiplexed_data/HC"
    log_file="$out_dir"/denovo_map.oe
    denovo_map.pl --samples "$reads_dir" --popmap "$popmap" -o "$out_dir" -T "$SLURM_CPUS_PER_TASK" -M "$M" -n "$M" -m 3 --paired &> "$log_file"
done

# Run populations with '-r 0.80' (loci present in 80% of samples)

for M in $M_values
do
    stacks_dir=stacks.M"$M"
    out_dir="$stacks_dir"/populations.r80
    mkdir "$out_dir"
    log_file="$out_dir"/populations.oe
    populations -P "$stacks_dir" -O "$out_dir" -t "$SLURM_CPUS_PER_TASK" -r 0.80 &> "$log_file"
done
```

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M_values="1 2 3 4 5 6 7 8 9"

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popmap="$WORK_DIR/data/Exercise_3/popmaps/test.popmap.txt"
OUT_DIR="$WORK_DIR/outputs/Exercise_3/test.denovo"

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# Create subdirectories
cd "$OUT_DIR" || exit

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do
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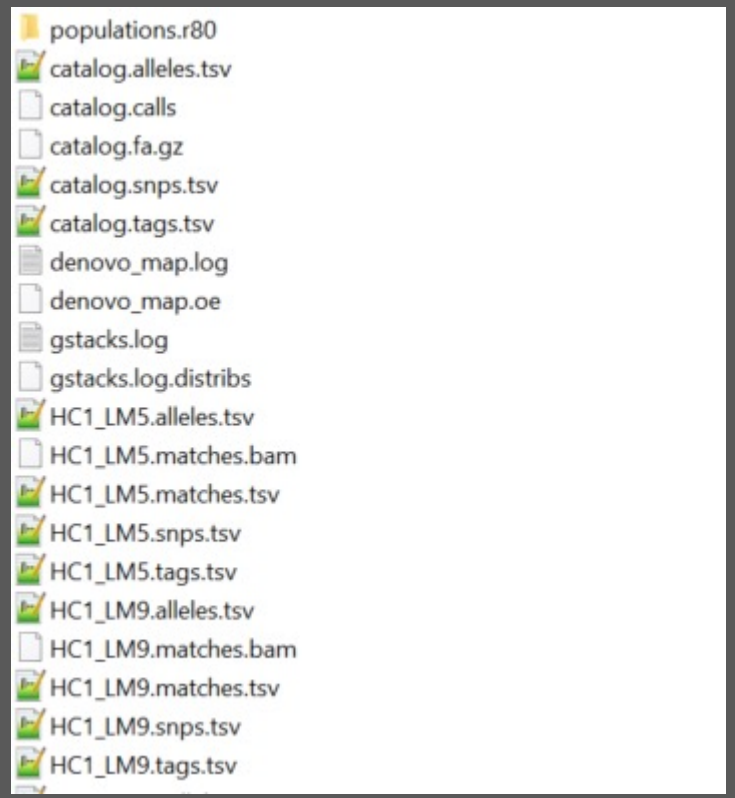
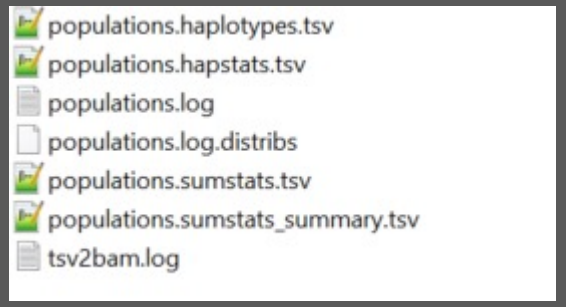
    out_dir="$stacks_dir"/populations.r80
    mkdir "$out_dir"
    log_file="$out_dir"/populations.oe
    populations -P "$stacks_dir" -O "$out_dir" -t "$SLURM_CPUS_PER_TASK" -r 0.80 &> "$log_file"
done
```


stacks.M1 File folder 15-Jun-22 16:19:34
stacks.M2 File folder 15-Jun-22 16:19:08
stacks.M3 File folder 15-Jun-22 16:19:13
stacks.M4 File folder 15-Jun-22 16:19:03
stacks.M5 File folder 15-Jun-22 16:19:24
stacks.M6 File folder 15-Jun-22 16:18:59
stacks.M7 File folder 15-Jun-22 16:19:39
stacks.M8 File folder 15-Jun-22 16:19:29
stacks.M9 File folder 15-Jun-22 16:19:18


populations.r80
catalog.alleles.tsv
catalog.calls
catalog.fa.gz
catalog.snps.tsv
catalog.tags.tsv
denovo_map.log
denovo_map.oe
gstacks.log
gstacks.log.distribs
HC1_LM5.alleles.tsv
HC1_LM5.matches.bam
HC1_LM5.matches.tsv
HC1_LM5.snps.tsv
HC1_LM5.tags.tsv
HC1_LM9.alleles.tsv
HC1_LM9.matches.bam
HC1_LM9.matches.tsv
HC1_LM9.snps.tsv
HC1_LM9.tags.tsv


populations.haplotypes.tsv
populations.hapstats.tsv
populations.log
populations.log.distribs
populations.sumstats.tsv
populations.sumstats_summary.tsv
tsv2bam.log

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- populations.r80
- catalog.alleles.tsv
- catalog.calls
- catalog.fa.gz
- catalog.snps.tsv
- catalog.tags.tsv
- denovo_map.log
- denovo_map.oe
- gstacks.log
- gstacks.log.distribs
- HC1_LM5.alleles.tsv
- HC1_LM5.matches.bam
- HC1_LM5.matches.tsv
- HC1_LM5.snps.tsv
- HC1_LM5.tags.tsv
- HC1_LM9.alleles.tsv
- HC1_LM9.matches.bam
- HC1_LM9.matches.tsv
- HC1_LM9.snps.tsv
- HC1_LM9.tags.tsv

- populations.haplotypes.tsv
- populations.hapstats.tsv
- populations.log
- populations.log.distribs
- populations.sumstats.tsv
- populations.sumstats_summary.tsv
- tsv2bam.log

- Number of r80 polymorphic loci
- Number of new polymorphic found for each iteration of M

02_extract_results

```
#!/bin/bash

#SBATCH -J extract_results
#SBATCH --mail-user=YOUREMAIL@gmail.com
#SBATCH --mail-type=BEGIN,END,FAIL,TIME_LIMIT
#SBATCH --output=/work/%u/%x-%j.out
#SBATCH --error=/work/%u/%x-%j.err
#SBATCH --mem-per-cpu=4G
#SBATCH -t 1:00:00

# Paths and filenames for this analysis

M_values="1 2 3 4 5 6 7 8 9"

WORK_DIR="/work/$USER/ddRAD-seq_workshop/outputs/Exercise_3/test.denovo"

## Load modules and activate software

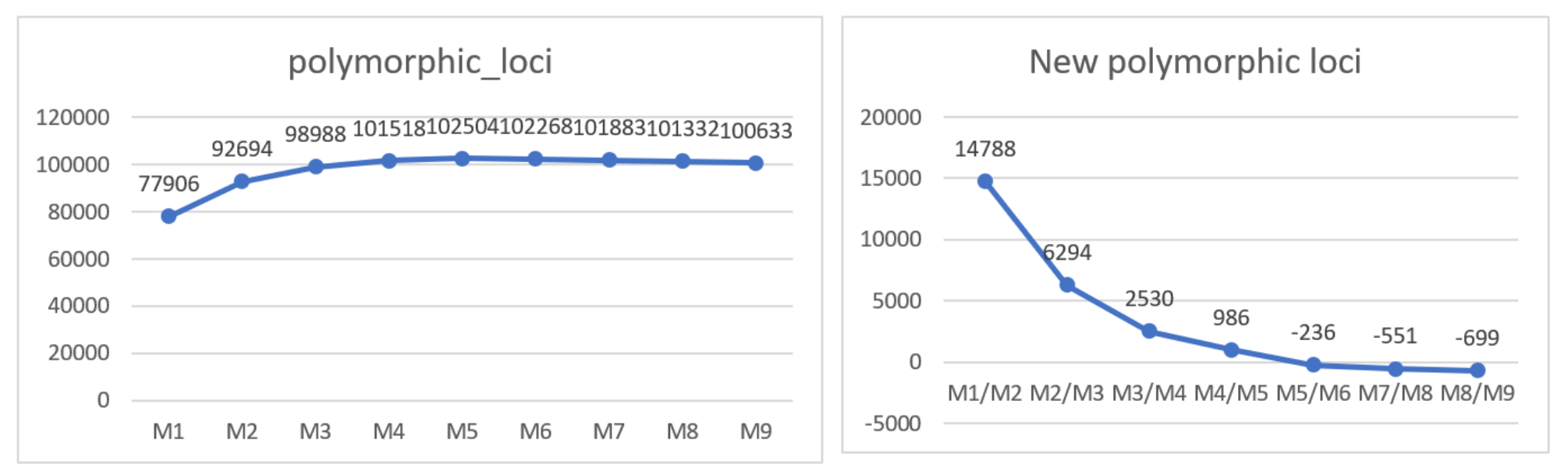
module purge
module load Anaconda3
source activate /gpfs0/global/apps/stacks_2.61

cd "$WORK_DIR" || exit
mkdir "$WORK_DIR/results"


for M in $M_values
do
stacks-dist-extract stacks.M"$M"/populations.r80/populations.log.distribs snps_per_loc_postfilters >> results/M"$M"_snp_distribution.tsv
cat stacks.M"$M"/populations.r80/populations.sumstats.tsv | grep -v "^#" | cut -f 1 | sort -n | uniq | wc -l >> results/M"$M"_r80.polymorphicLOCI.tsv
awk 'NR == 6 {print $5}' stacks.M"$M"/populations.r80/populations.sumstats_summary.tsv >> results/M"$M"_r80.polymorphicLOCI_summary.tsv
cat results/*.polymorphicLOCI.tsv >> results/all.polymorphicLOCI.tsv
cat  results/*.polymorphicLOCI_summary.tsv > results/all.polymorphicLOCI_summary.tsv
done
```

Bioinformatics

Bismarckia nobilis



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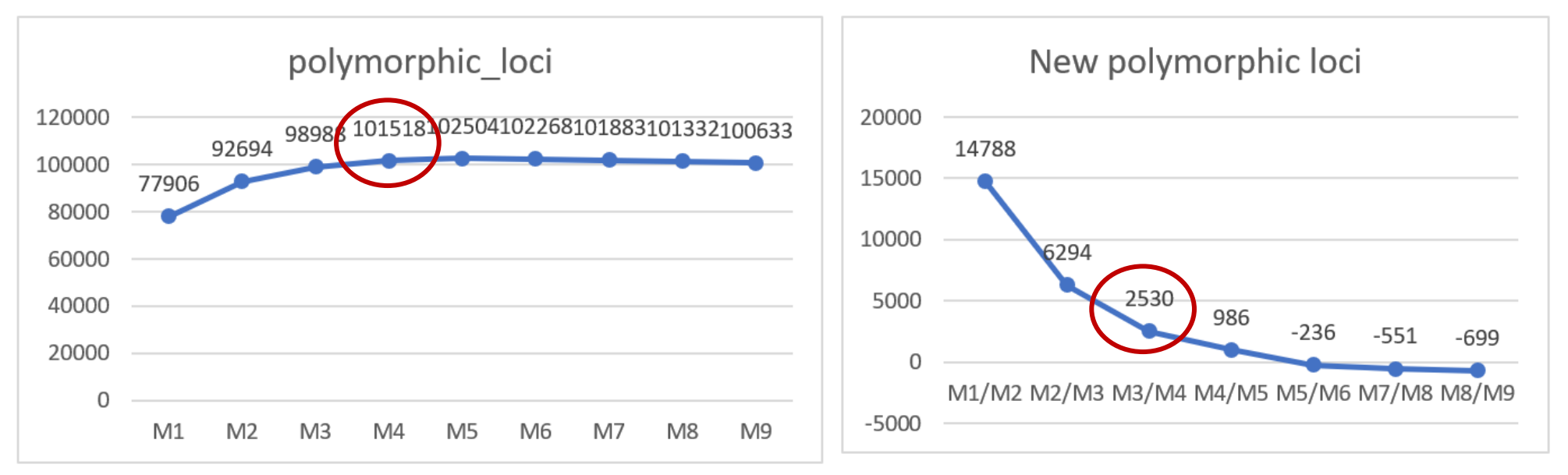

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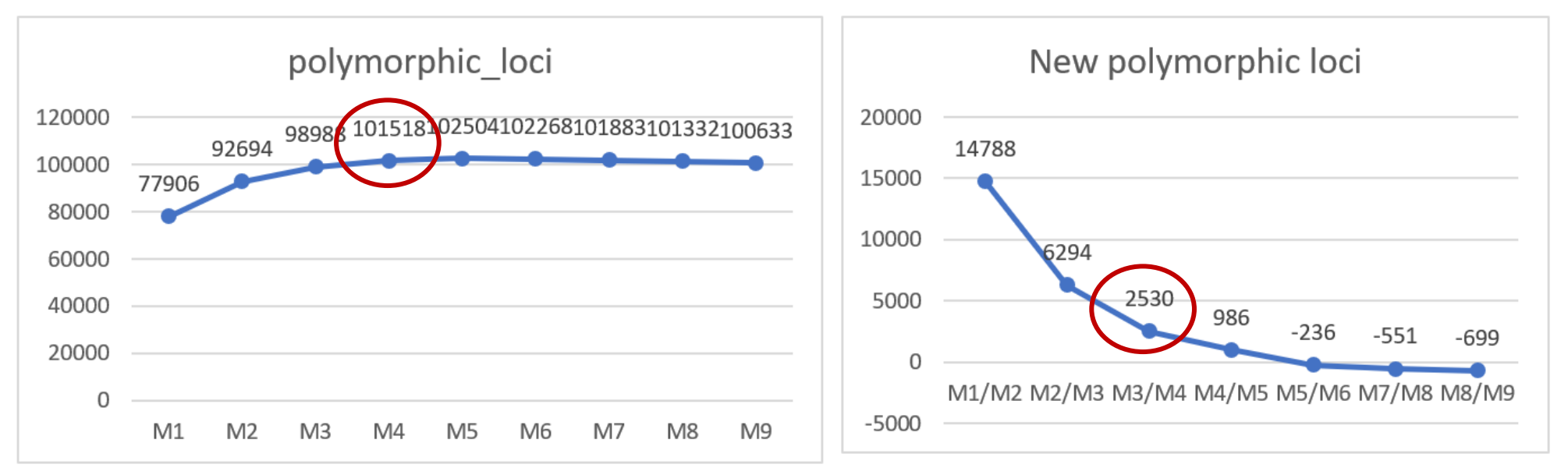
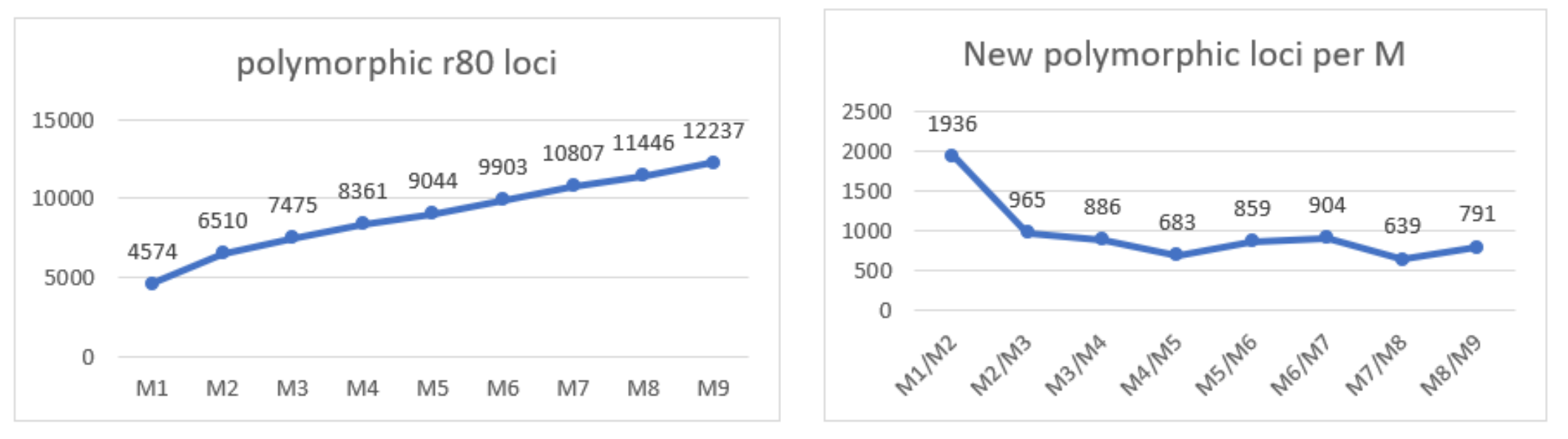


M = 4

M = n

Bioinformatics

Dypsis pinnatifrons



Bioinformatics

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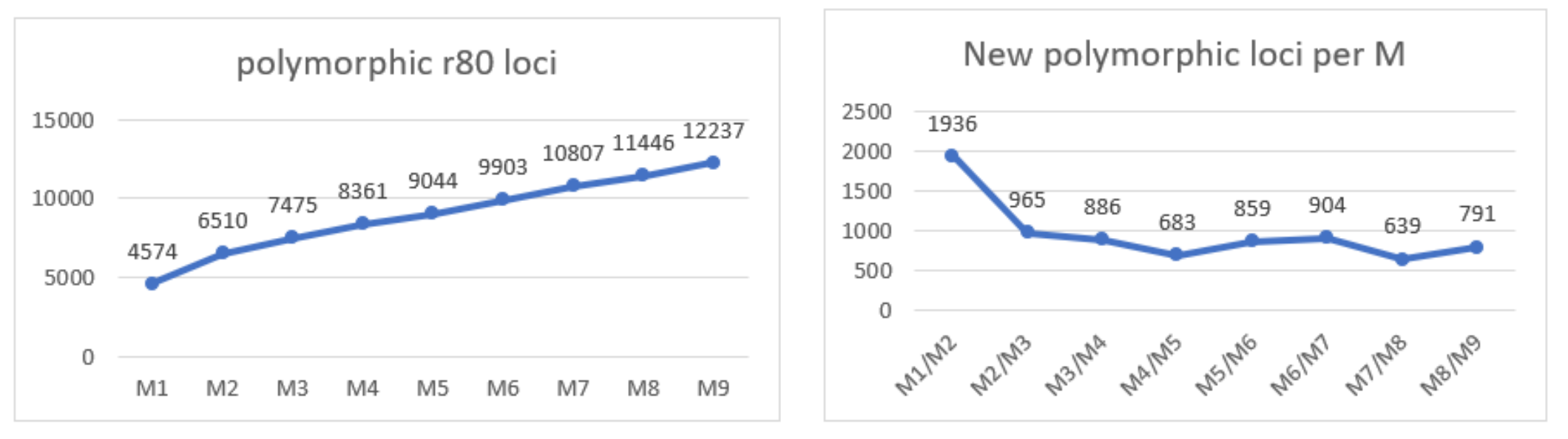
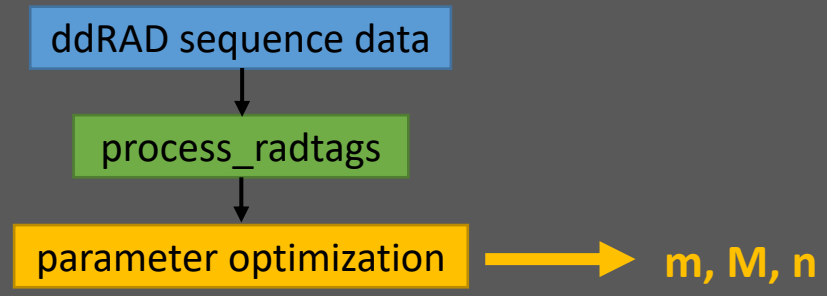


M = 5
M = n

Bioinformatics Workflow



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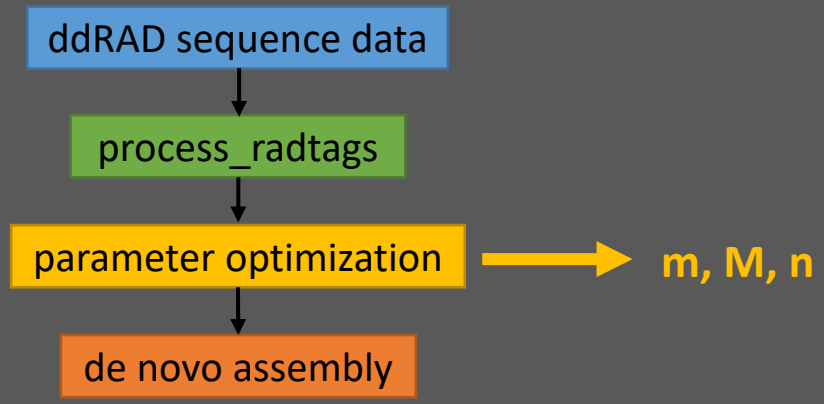

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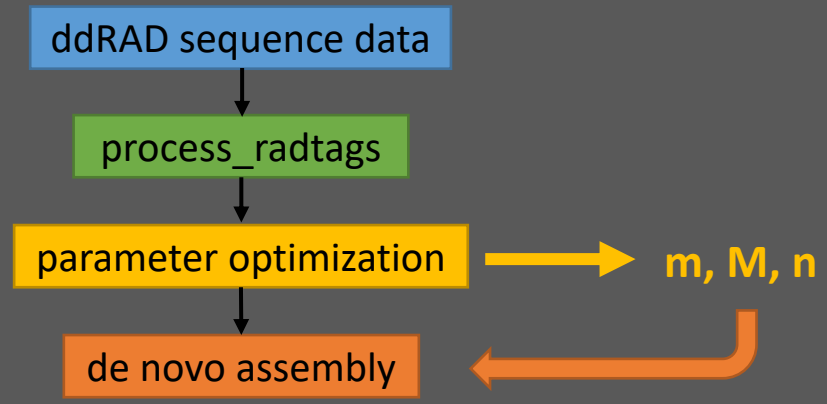

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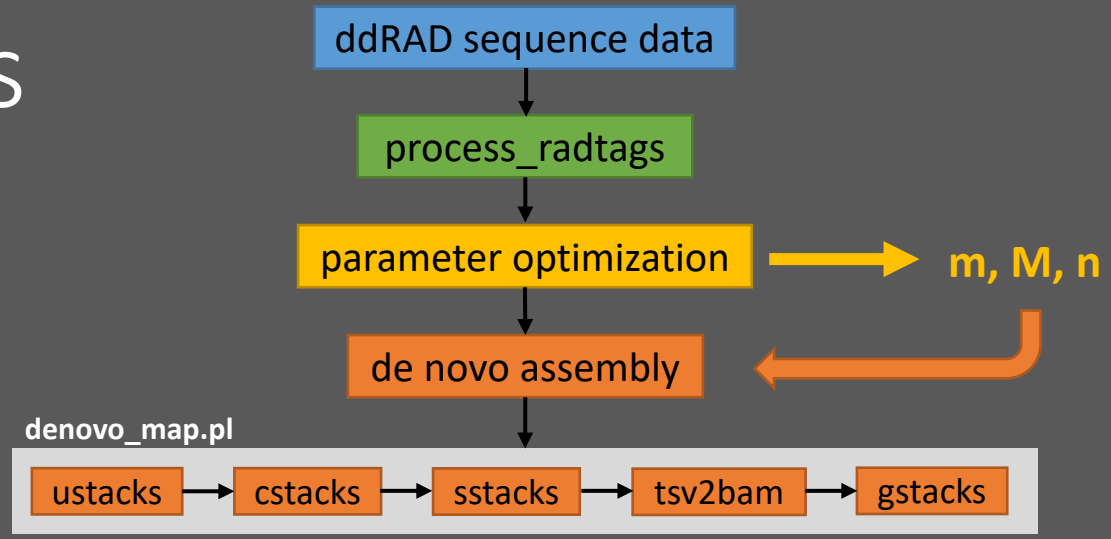

Rochette, Rivera-Colón & Catchen, 2019 - Mol. Ecology

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populations	Population level statistics and output in different formats. Possibility for further filtering

Output files for population structure analyses, phylogenetics, demographic history…

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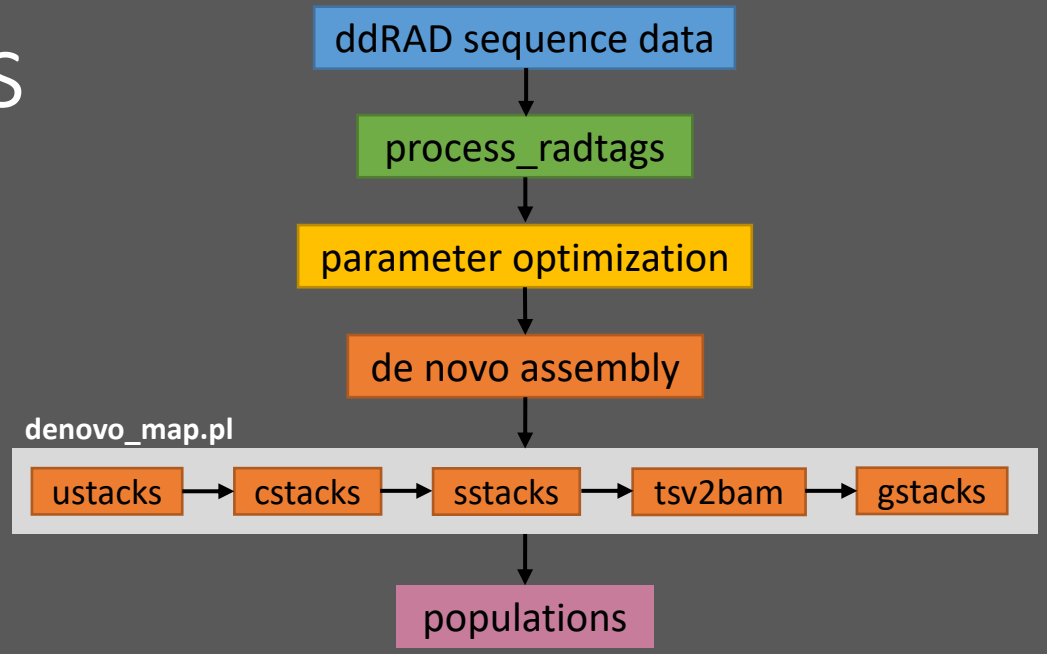

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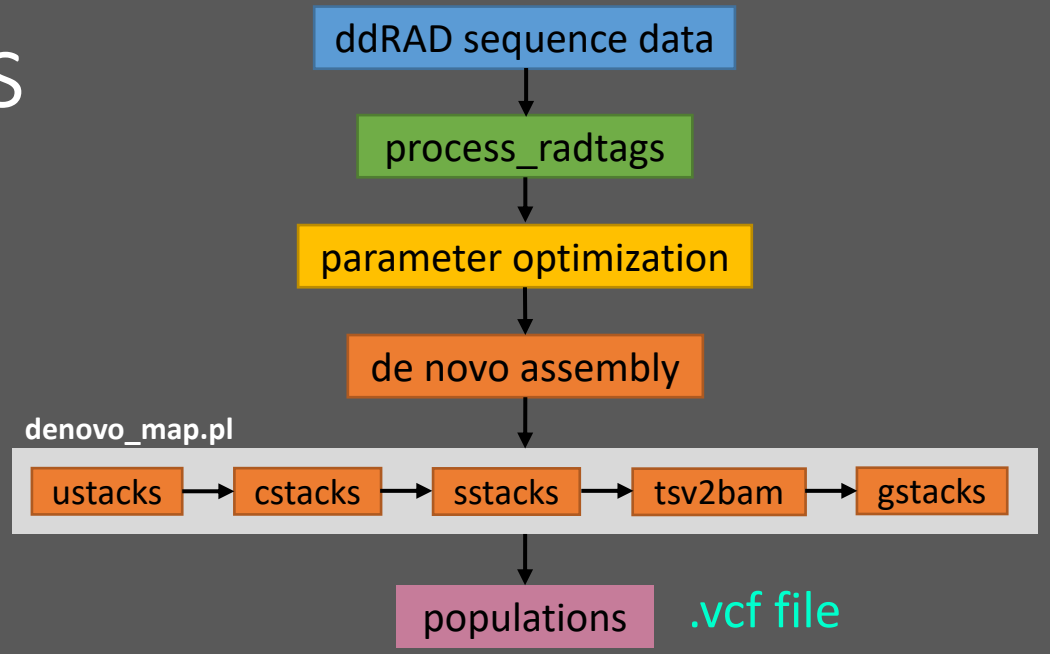

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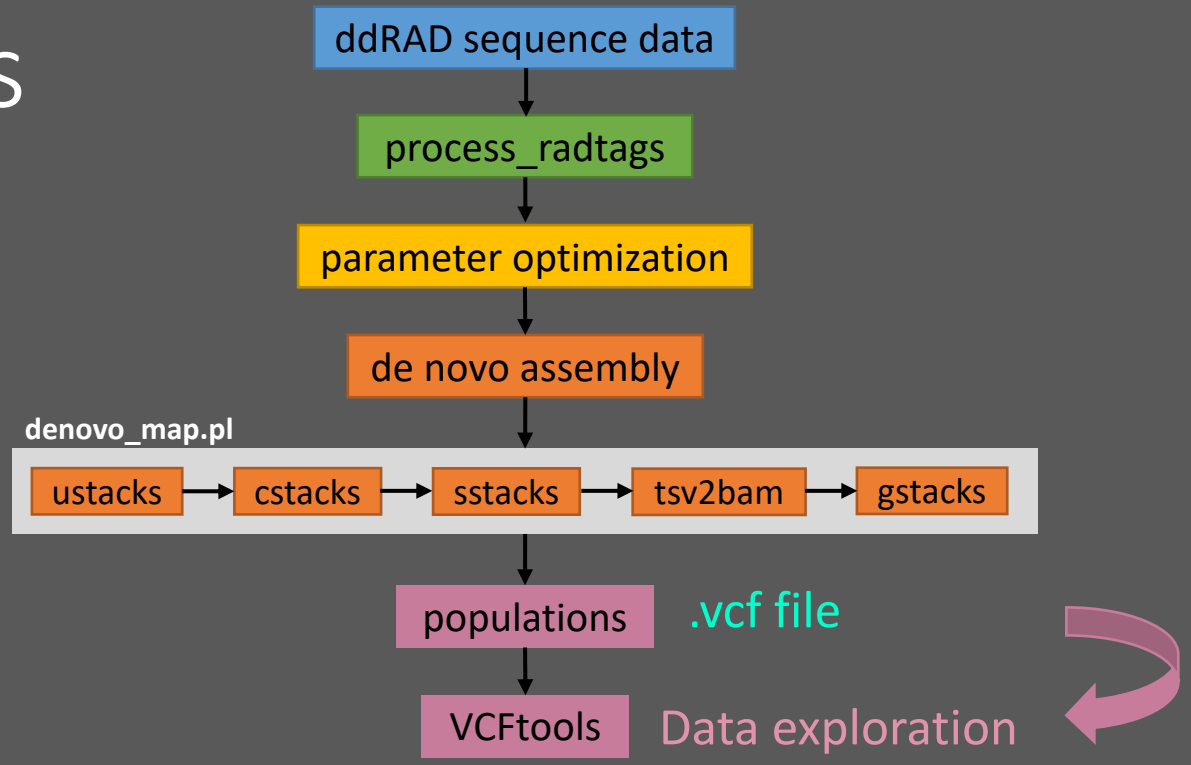

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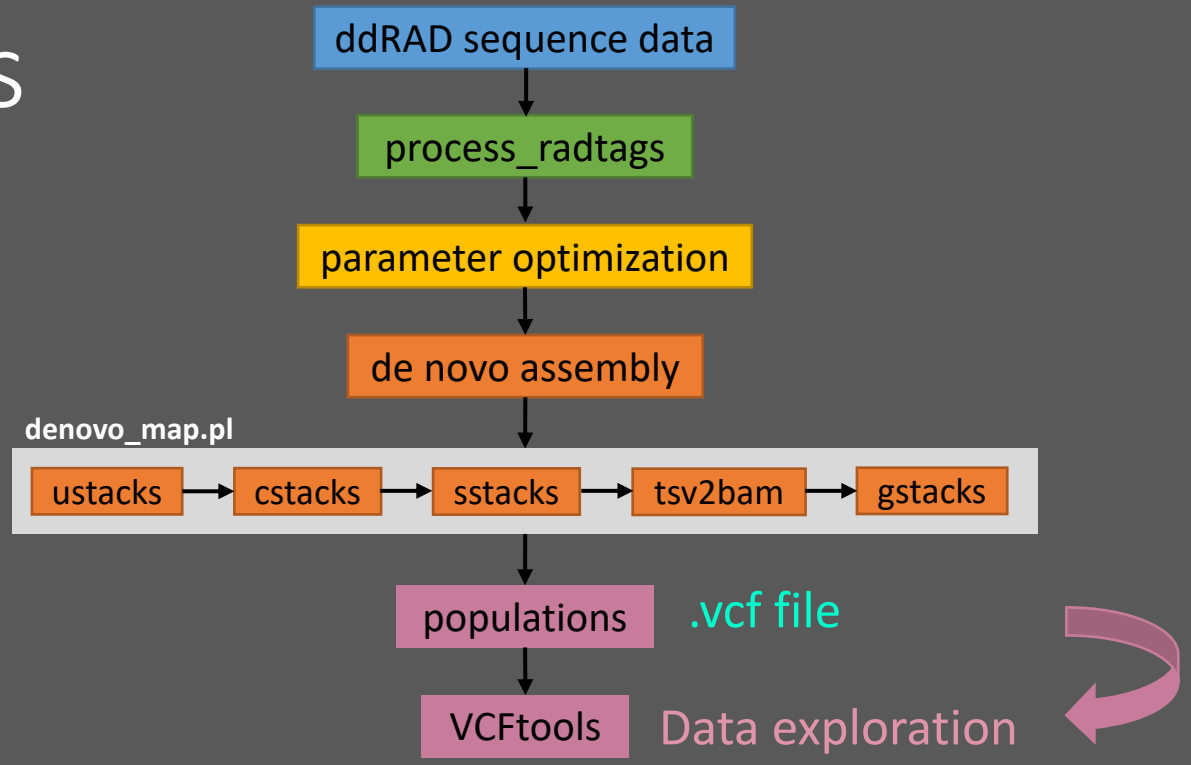


--freq2 = allele frequency

--depth = mean depth per individual

--site-mean-depth = mean depth per site

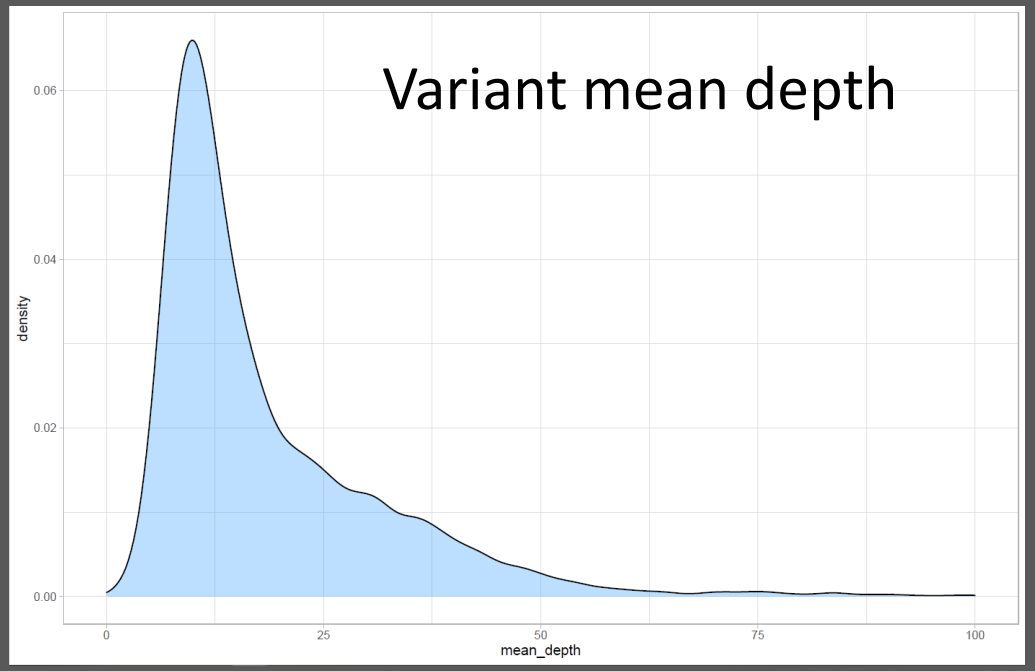
--missing-indv = proportion of missing data per individual

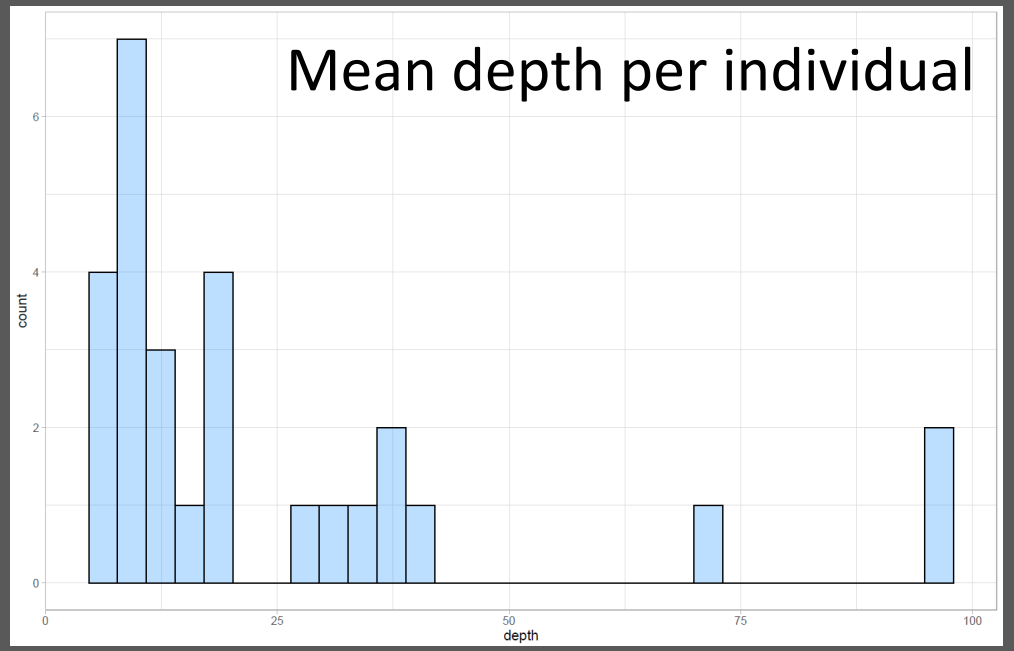
--missing-site = proportion of missing data per site

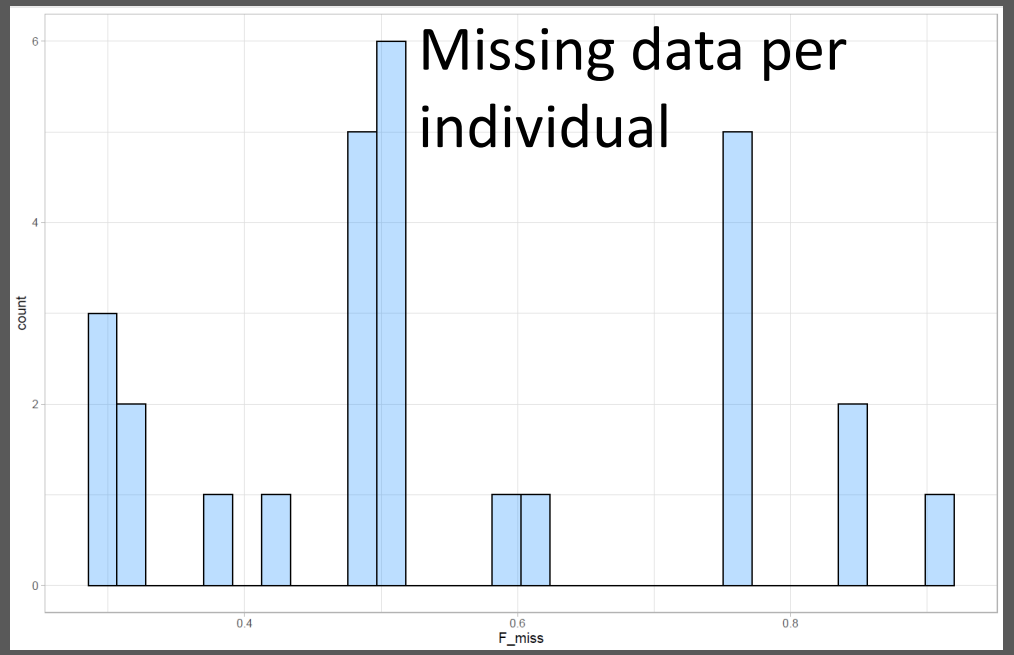
--het = heterozygosity and inbreeding coefficient per individual

Bioinformatics

Borassus madagascariensis
28 individuals



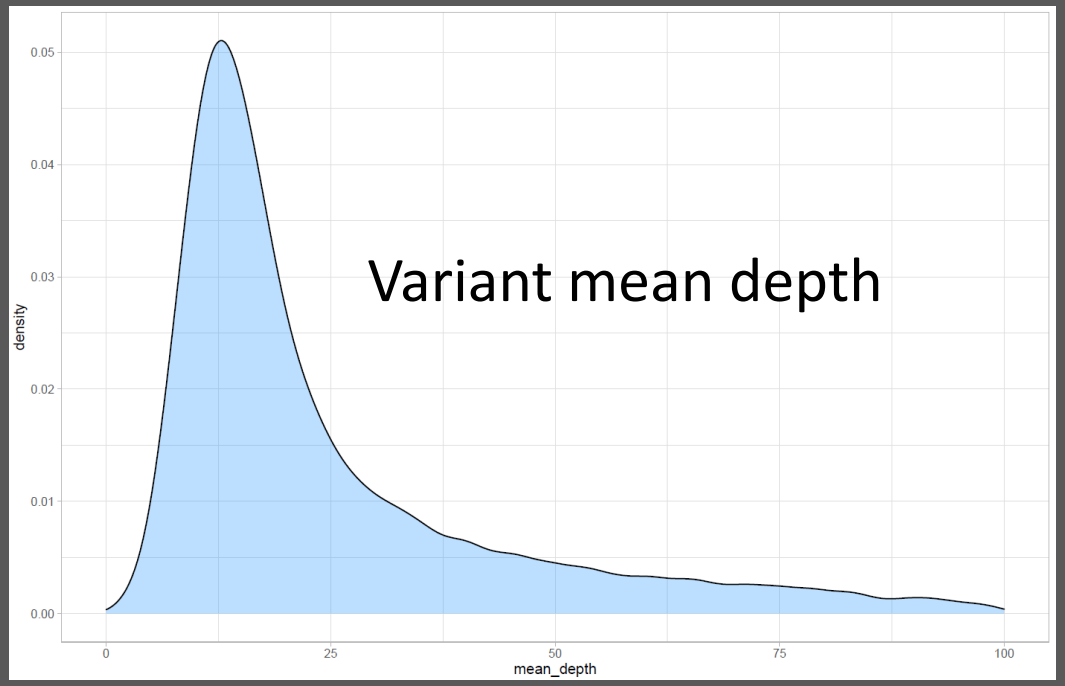


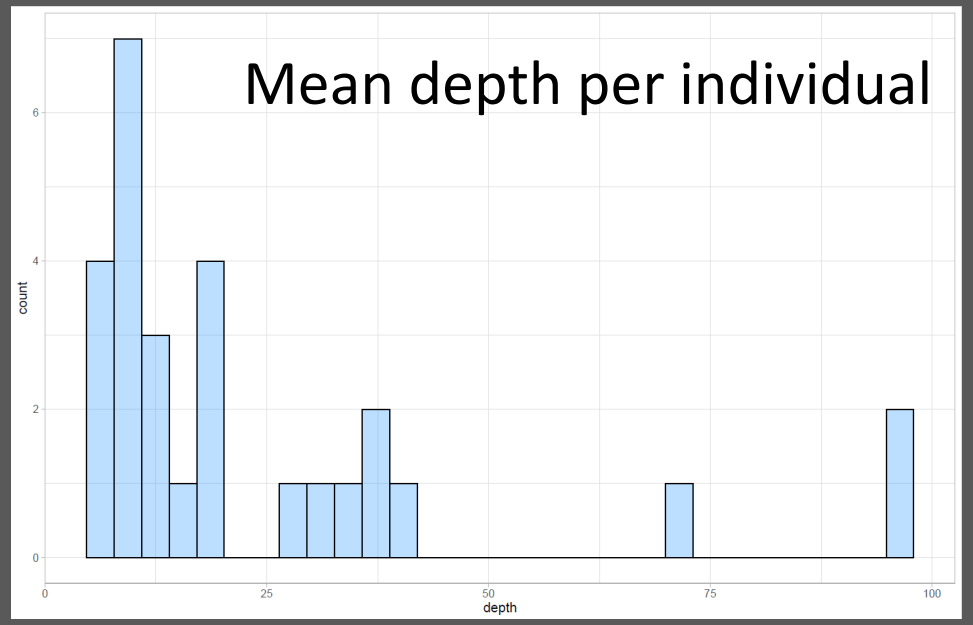


Bioinformatics

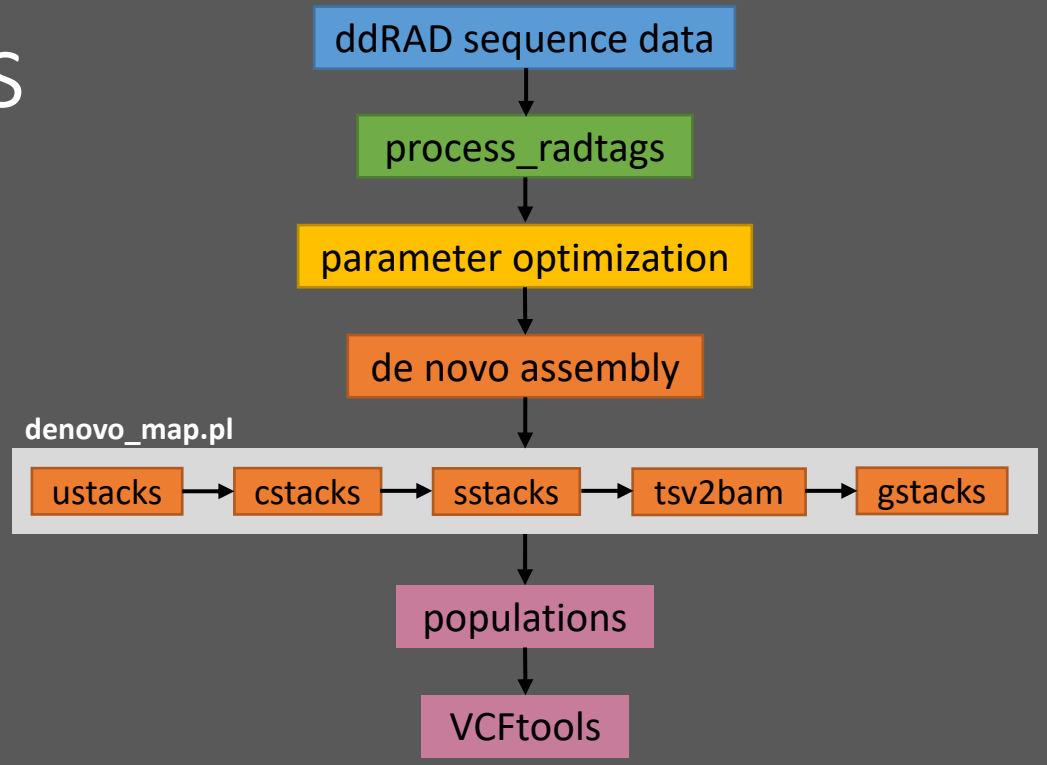
Bismarckia nobilis

63 individuals

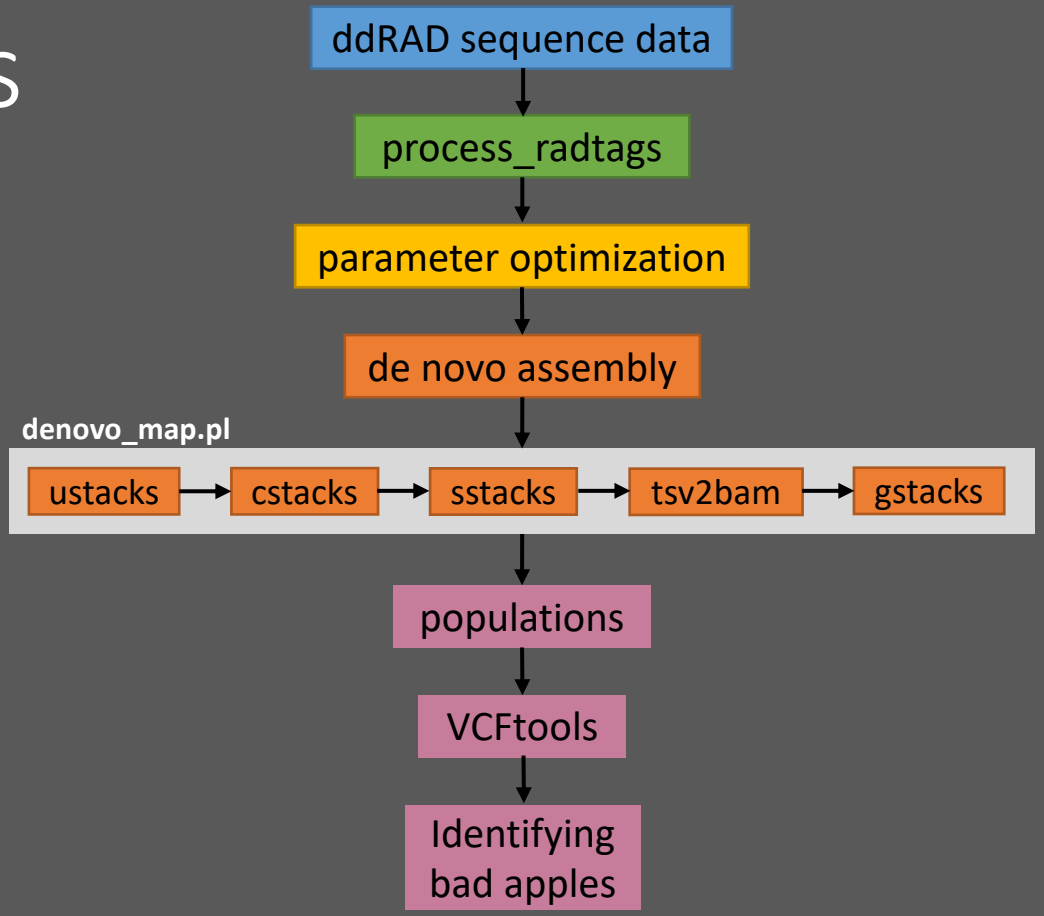




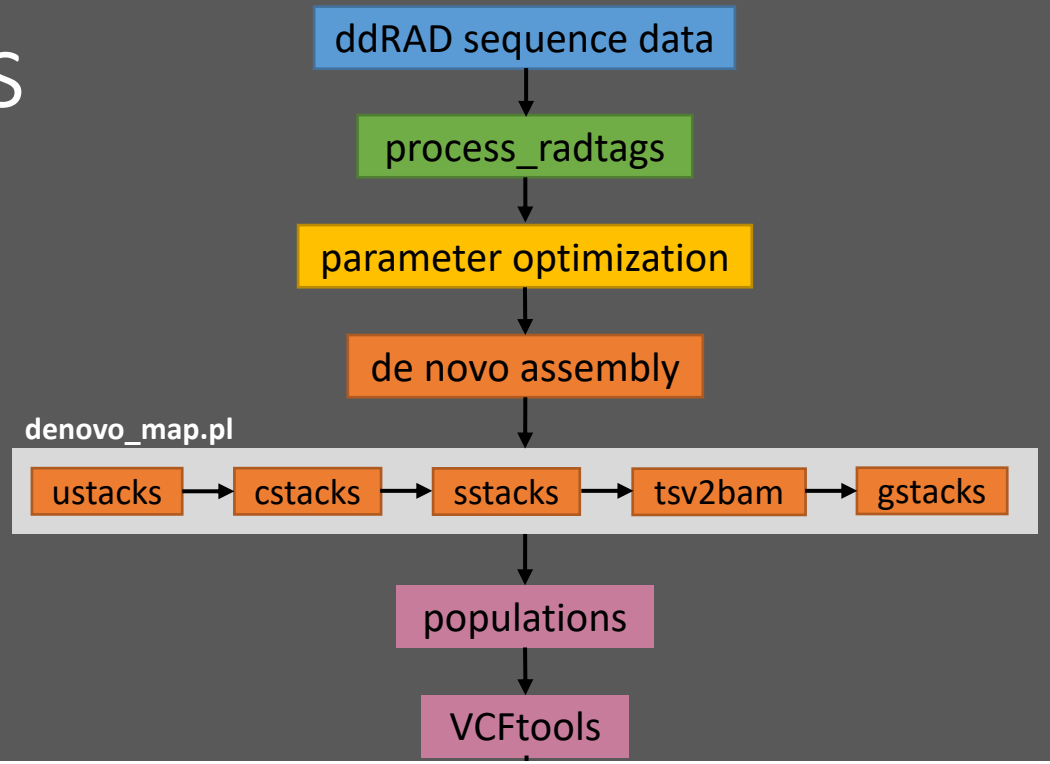
Bioinformatics Workflow



Bioinformatics Workflow



Bioinformatics Workflow



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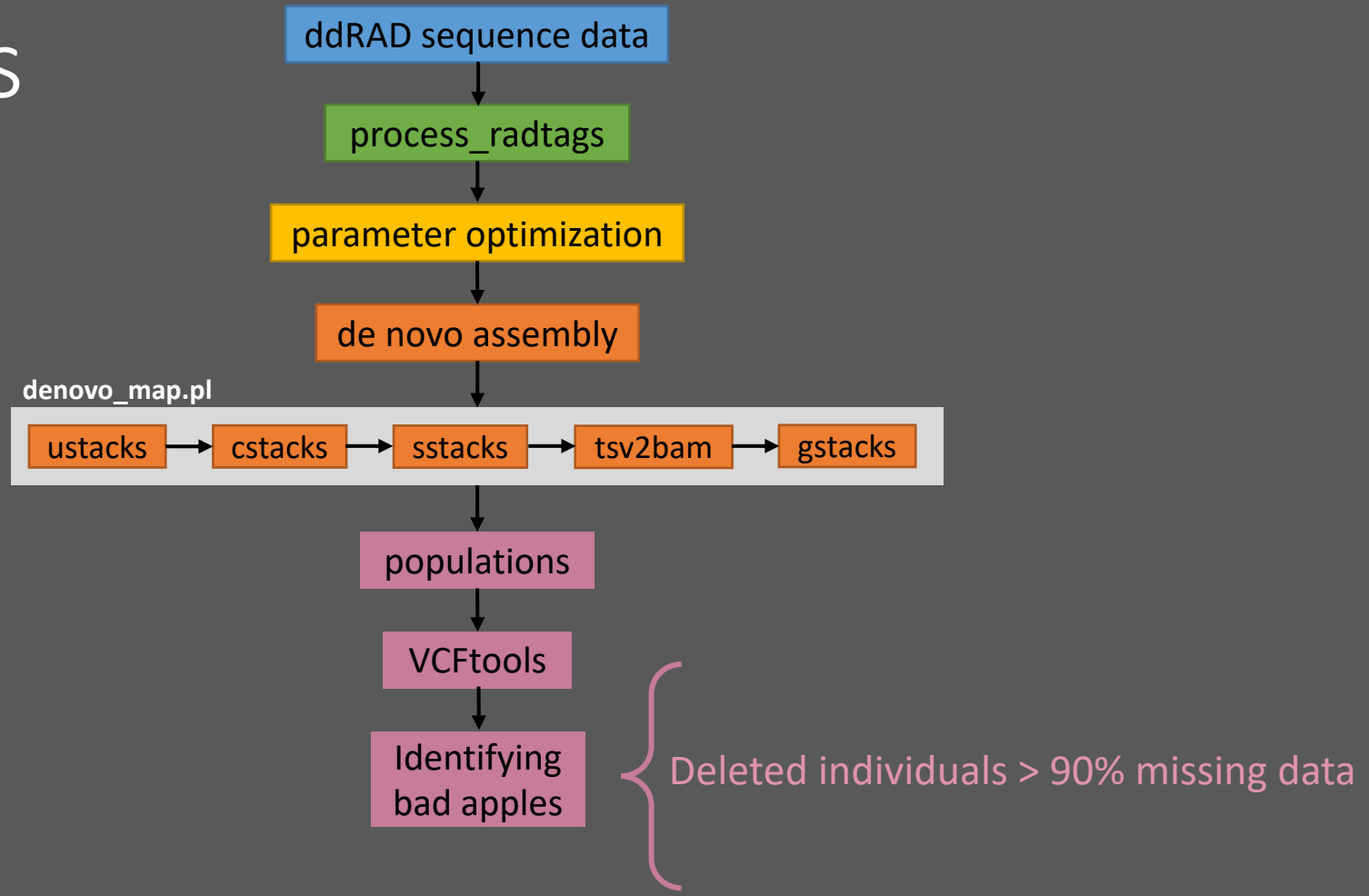
RESEARCH ARTICLE

Methods in Ecology and Evolution BRITISH ECOLOGICAL SOCIETY

Removing the bad apples: A simple bioinformatic method to improve loci-recovery in de novo RADseq data for non-model organisms

José Cerca[1,2,3] | Marius F. Maurstad[1,4] | Nicolas C. Rochette[5,6] | Angel G. Rivera-Colón[5] | Niraj Rayamajhi[5] | Julian M. Catchen[5] | Torsten H. Struck[1]

Bioinformatics Workflow



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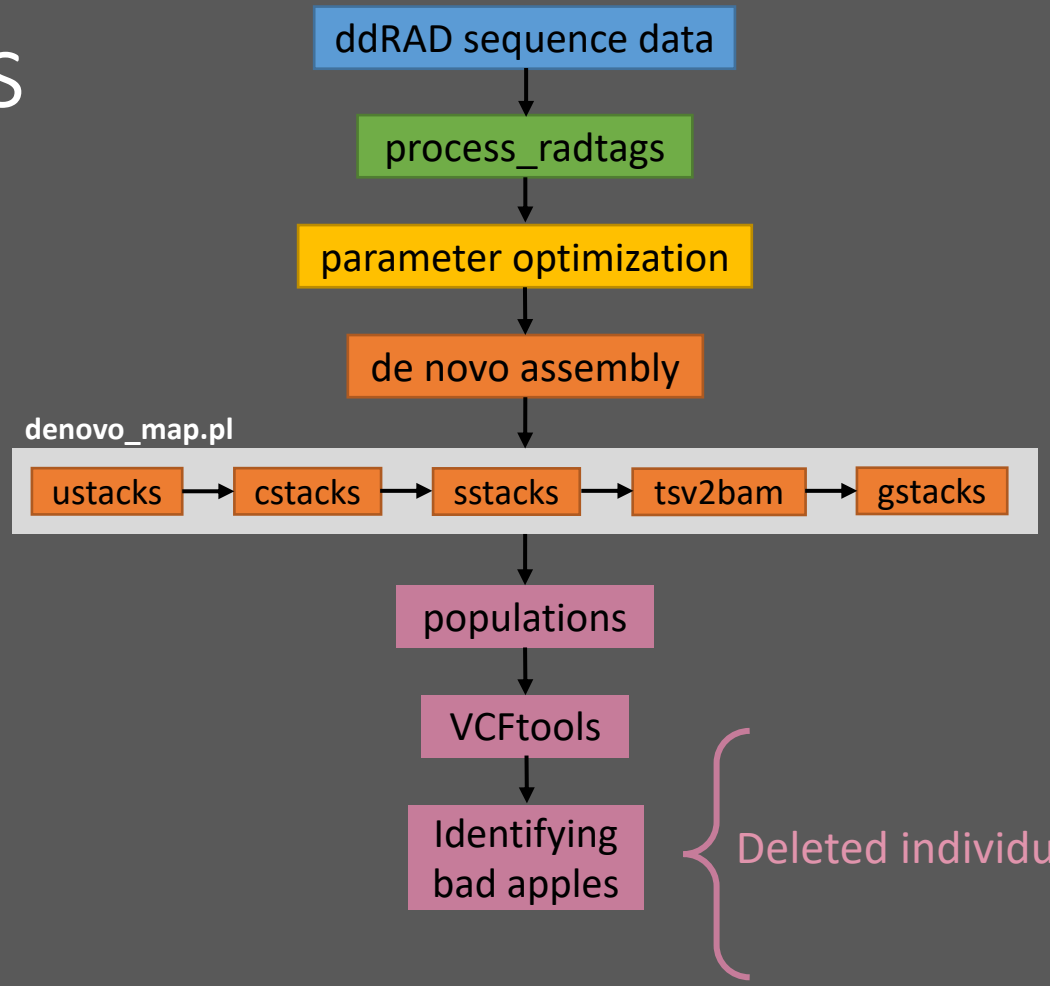

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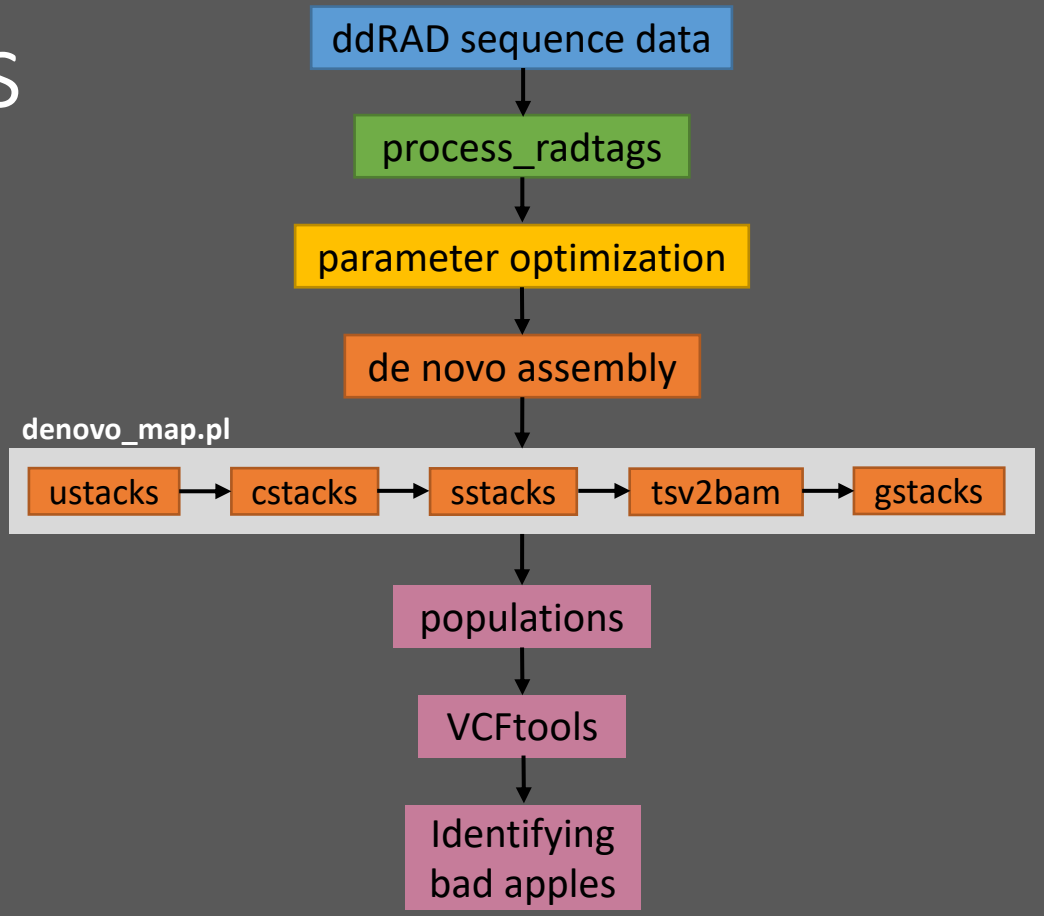

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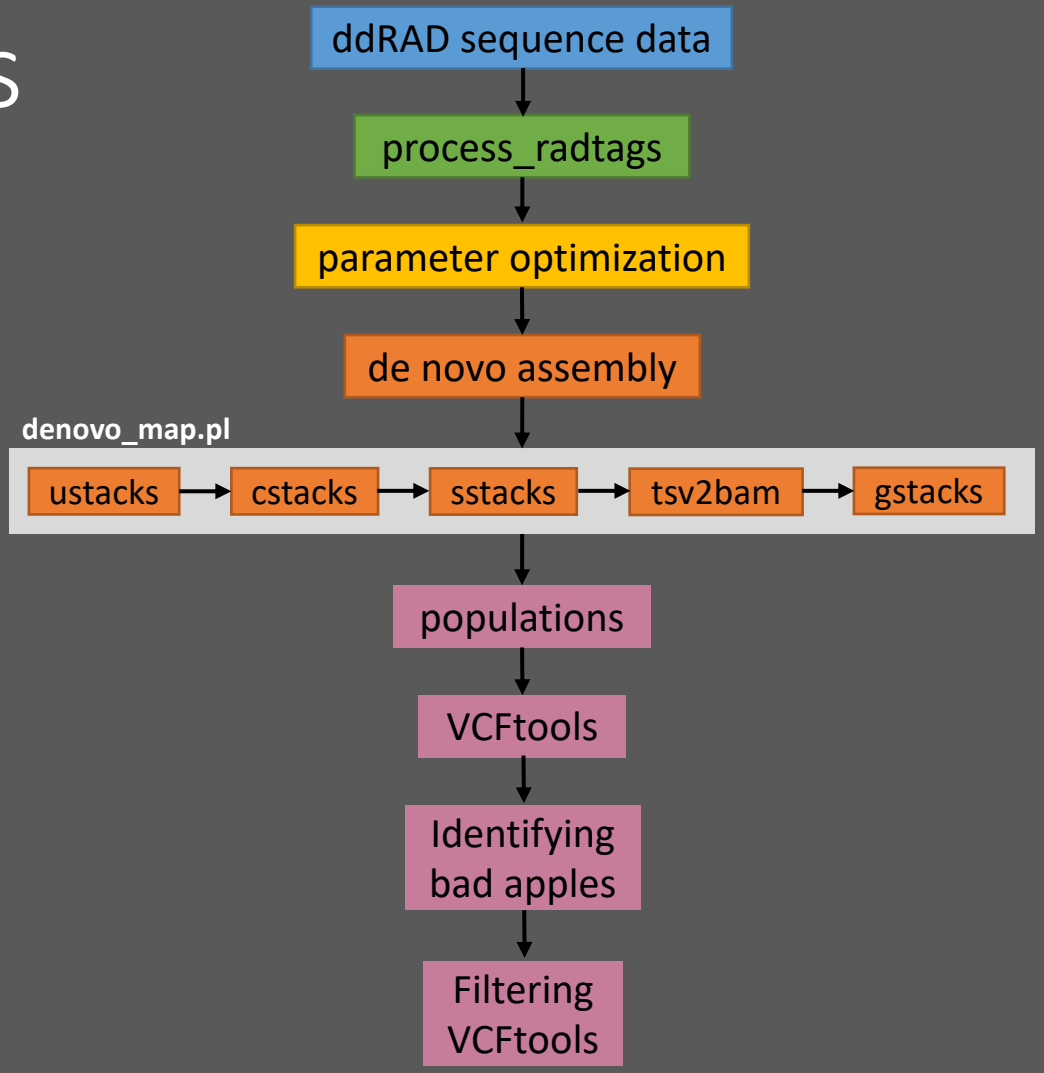

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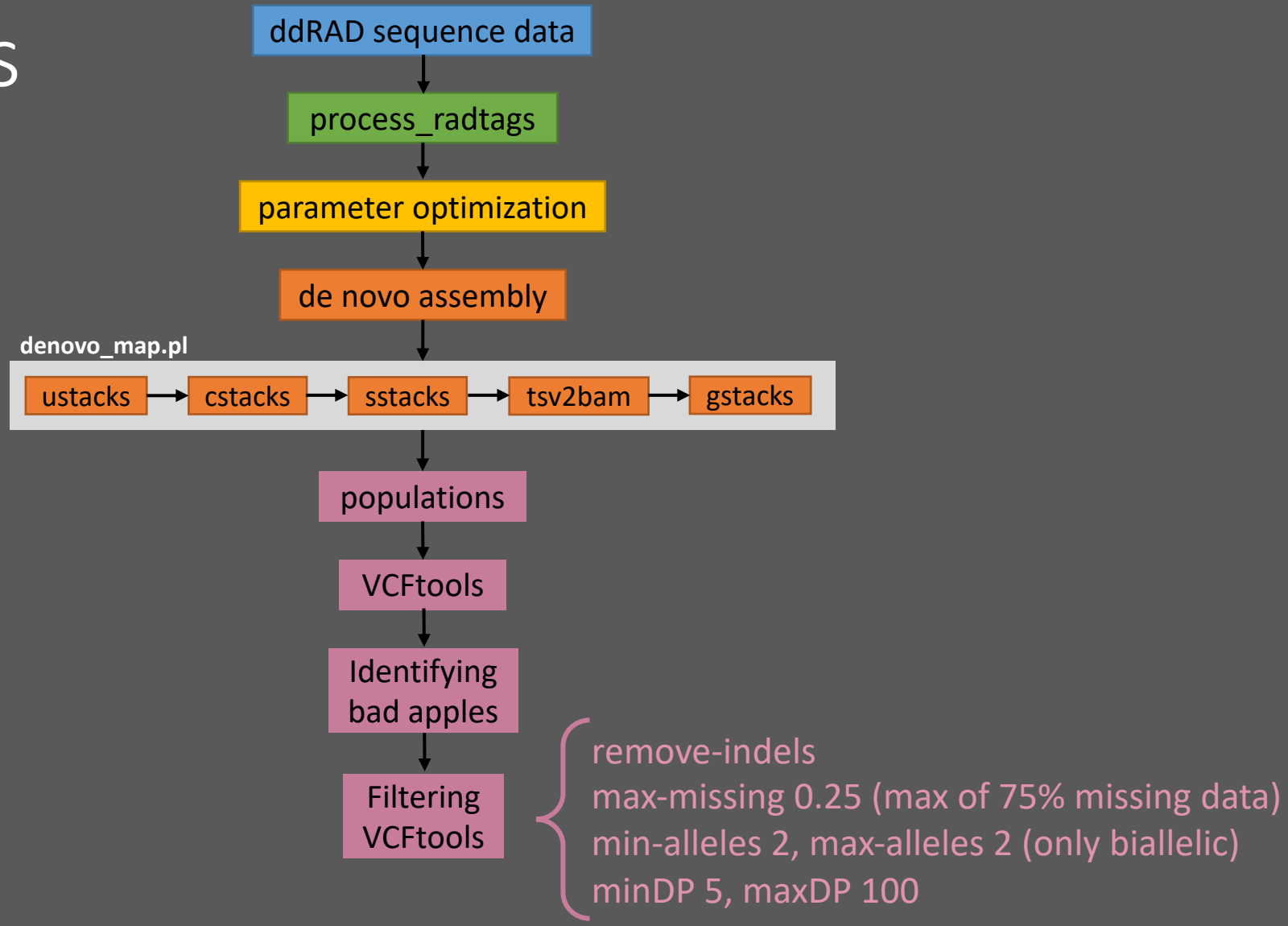

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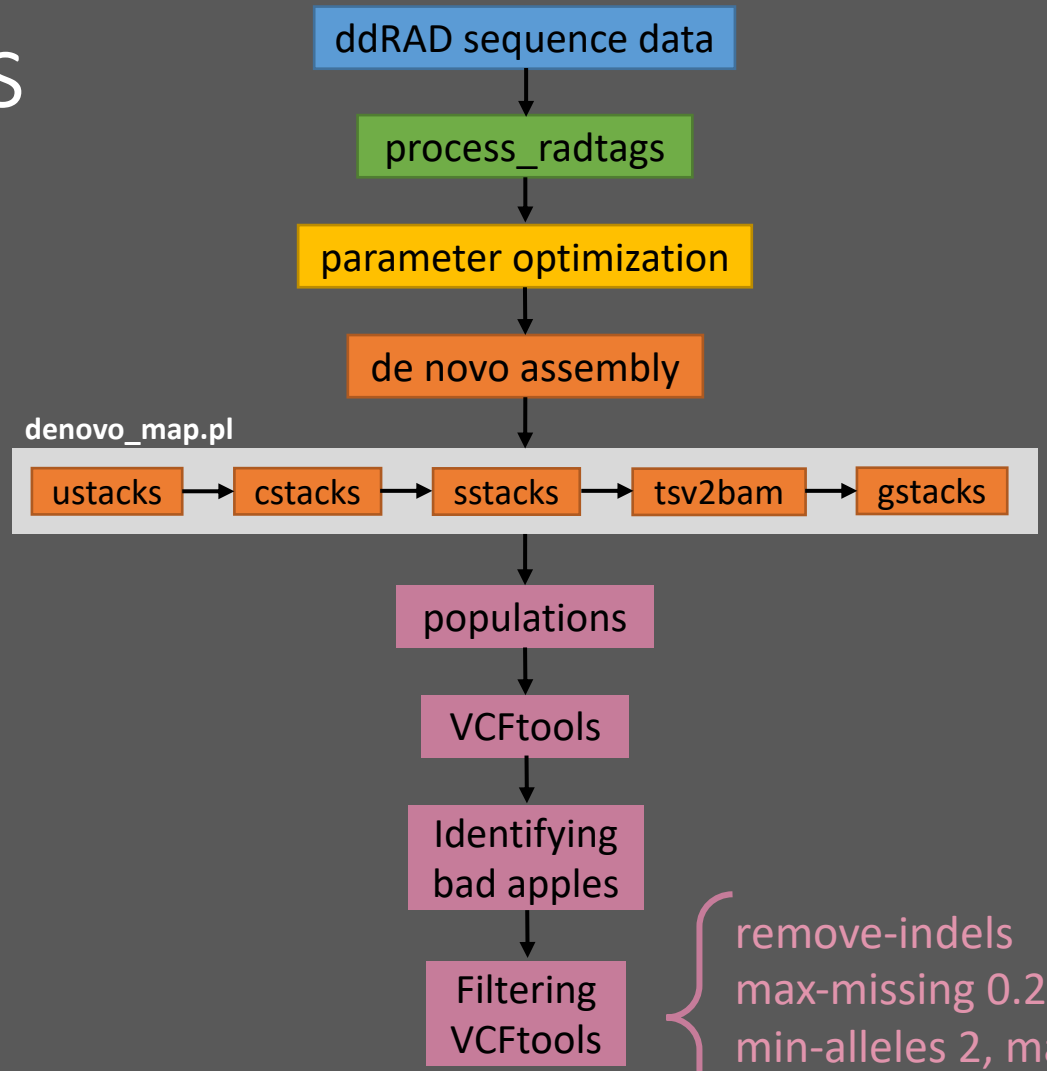
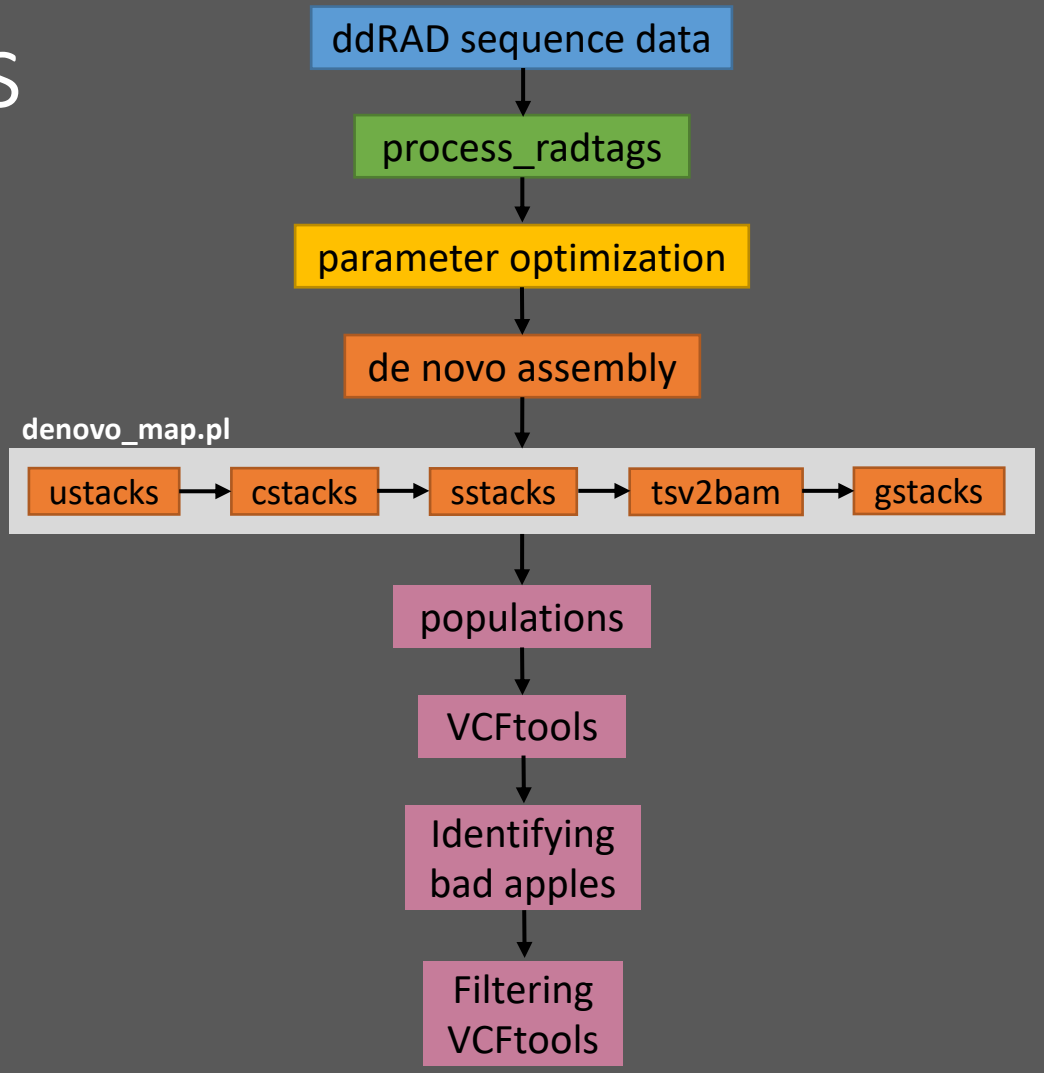


To control for false-positive loci

remove-indels
max-missing 0.25 (max of 75% missing data)
min-alleles 2, max-alleles 2 (only biallelic)
minDP 5, maxDP 100

Bioinformatics Workflow



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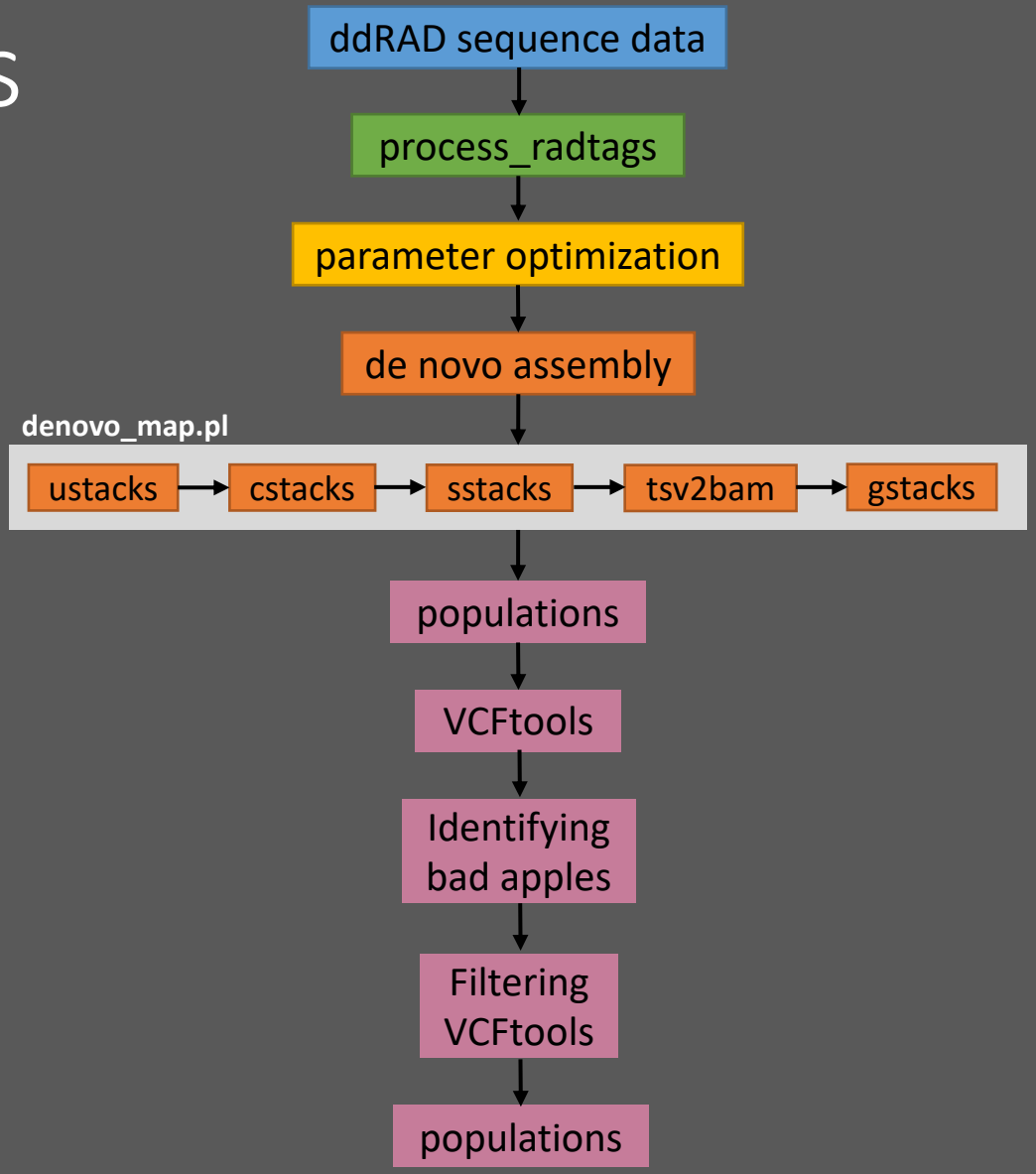
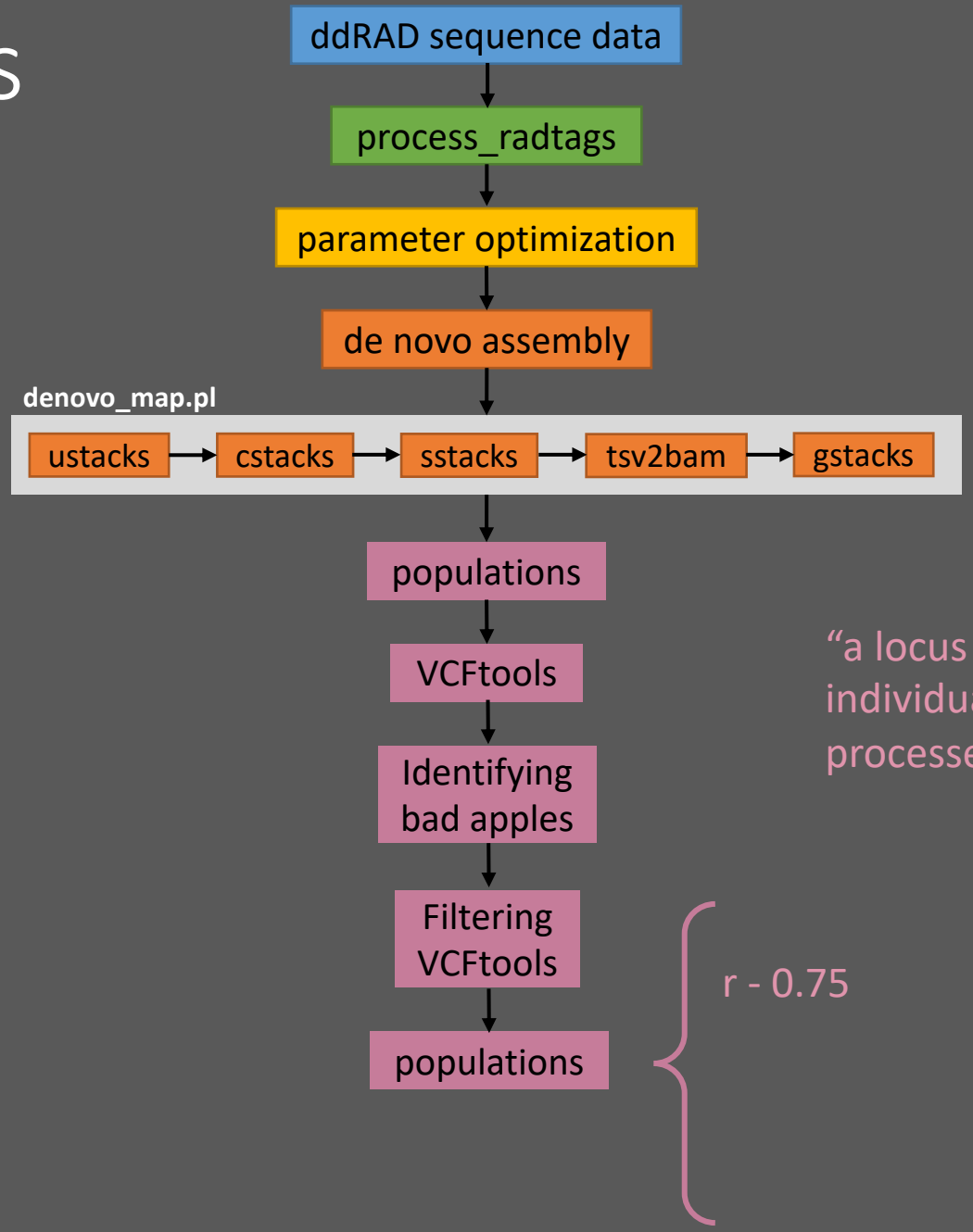
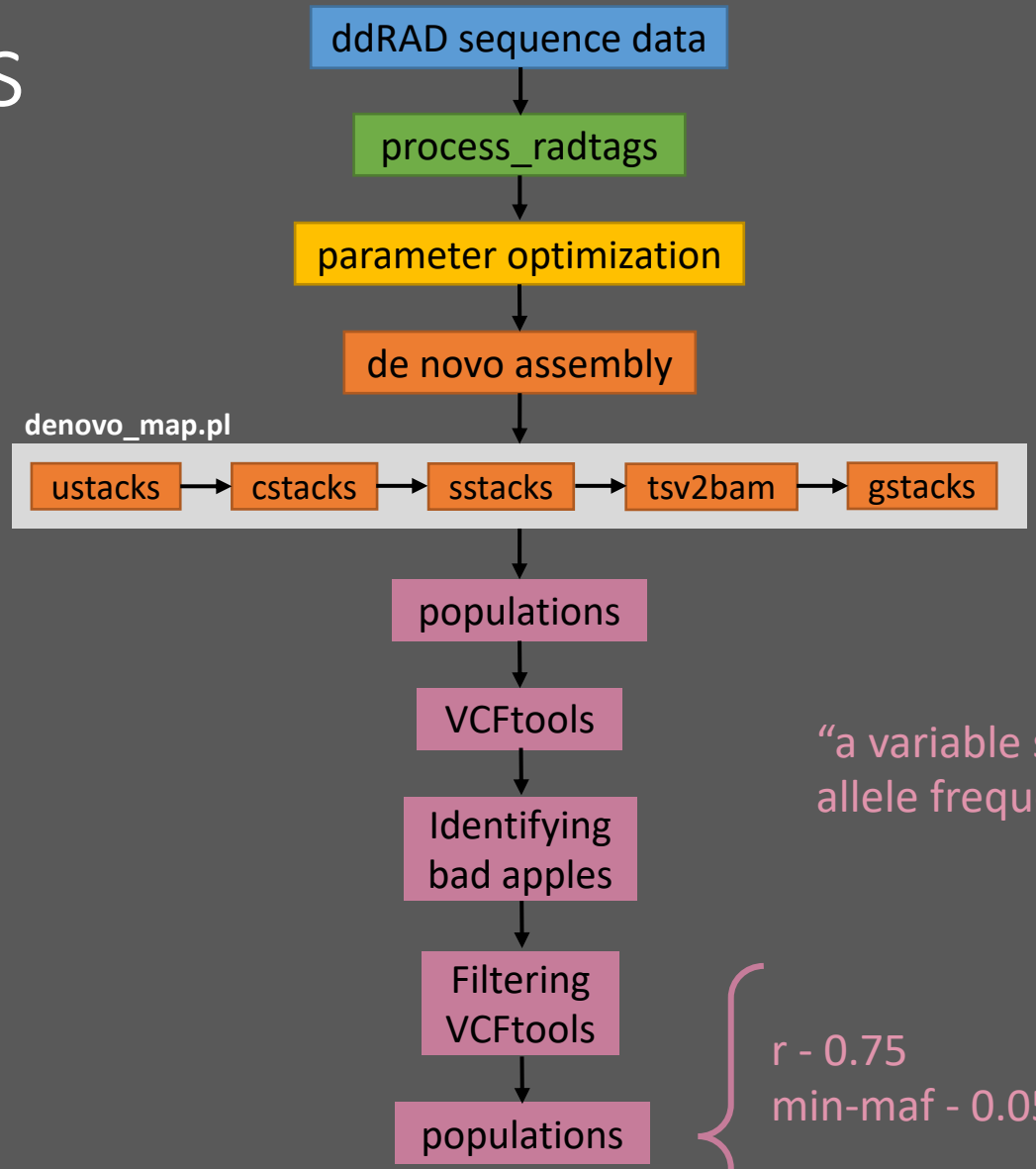


Bioinformatics Workflow



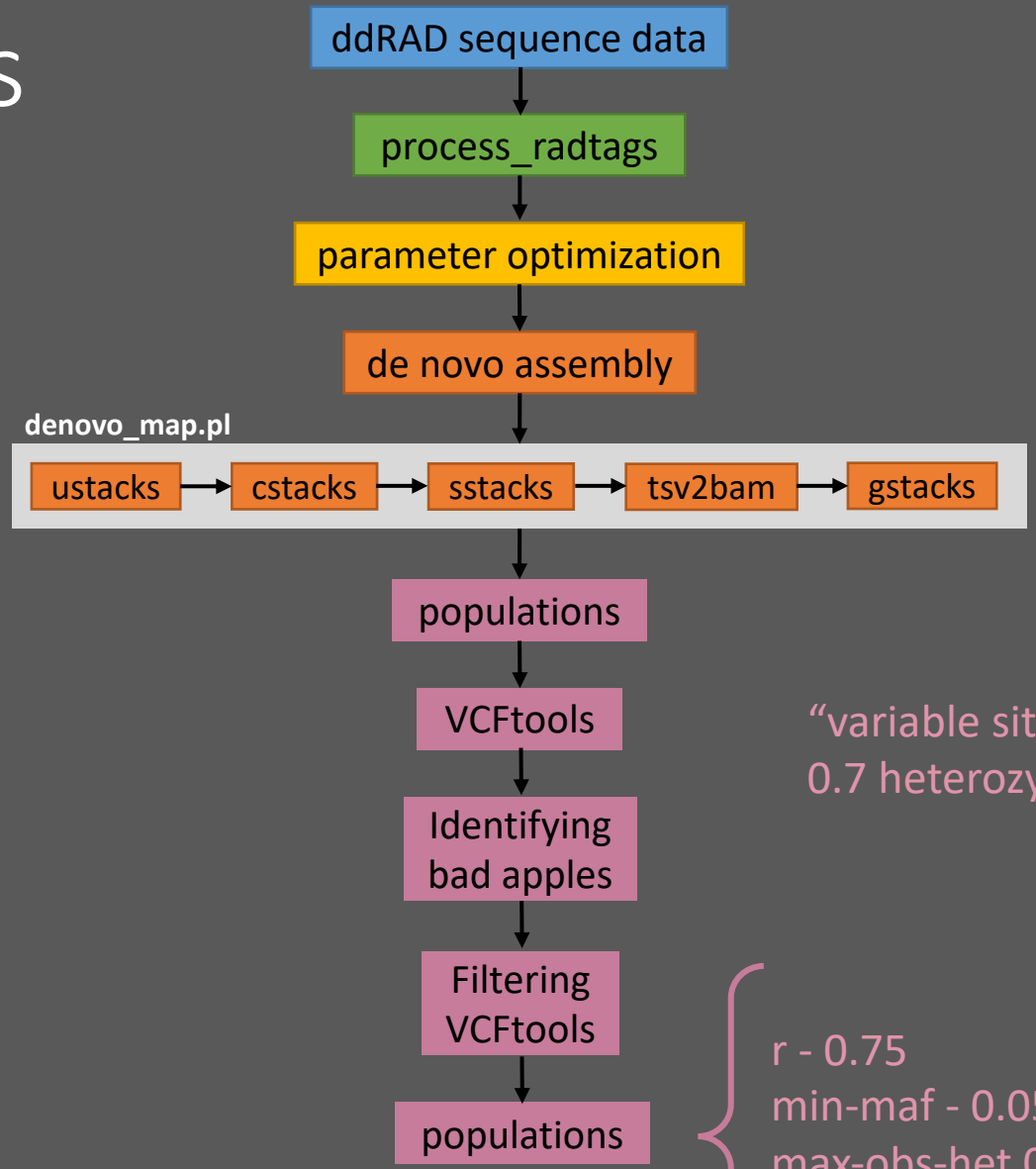
"a locus must be found in 75% of individuals of a single population to be processed"

Bioinformatics Workflow



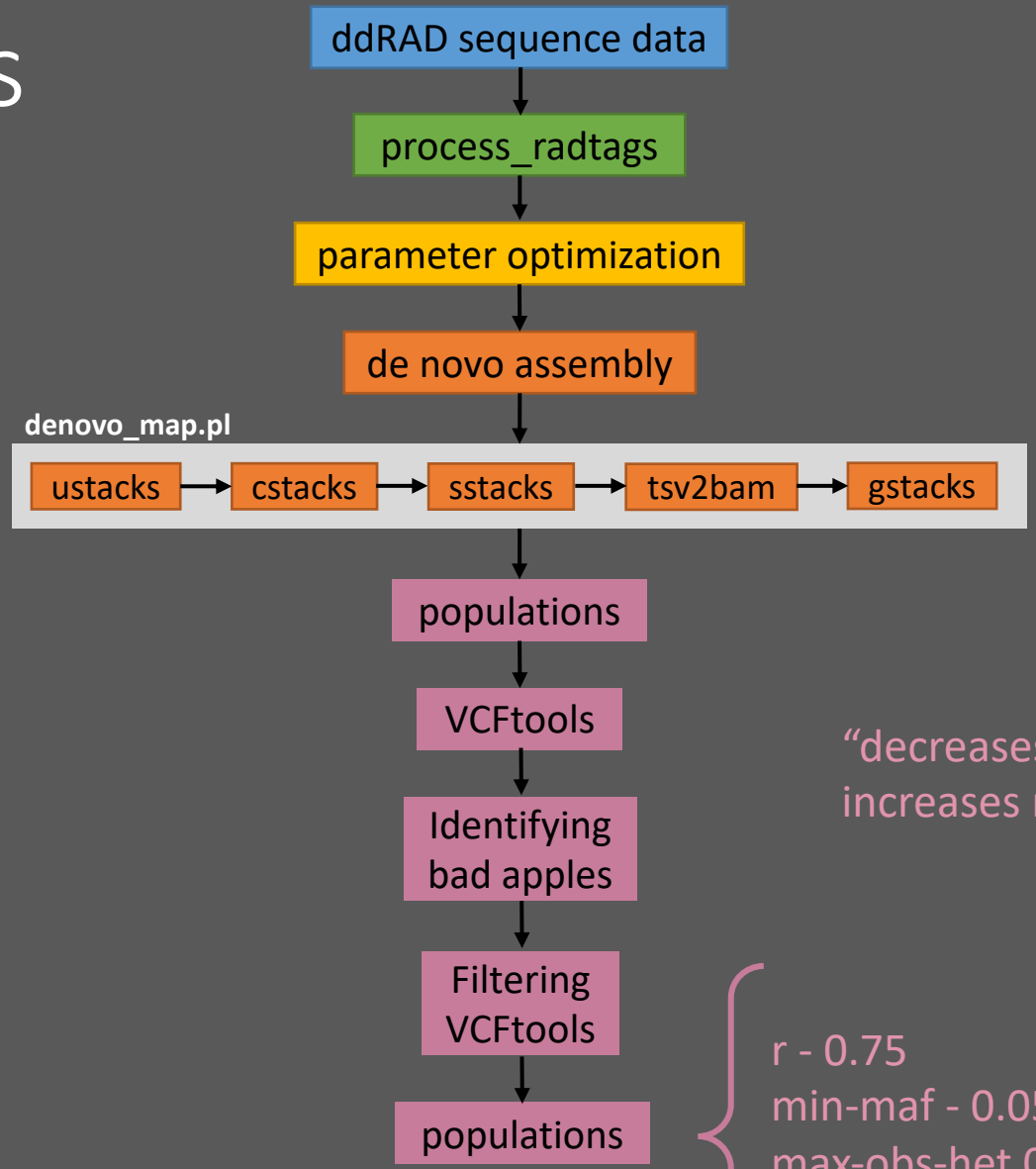
"a variable site must possess a minimum allele frequency of 5% to be included"

Bioinformatics Workflow



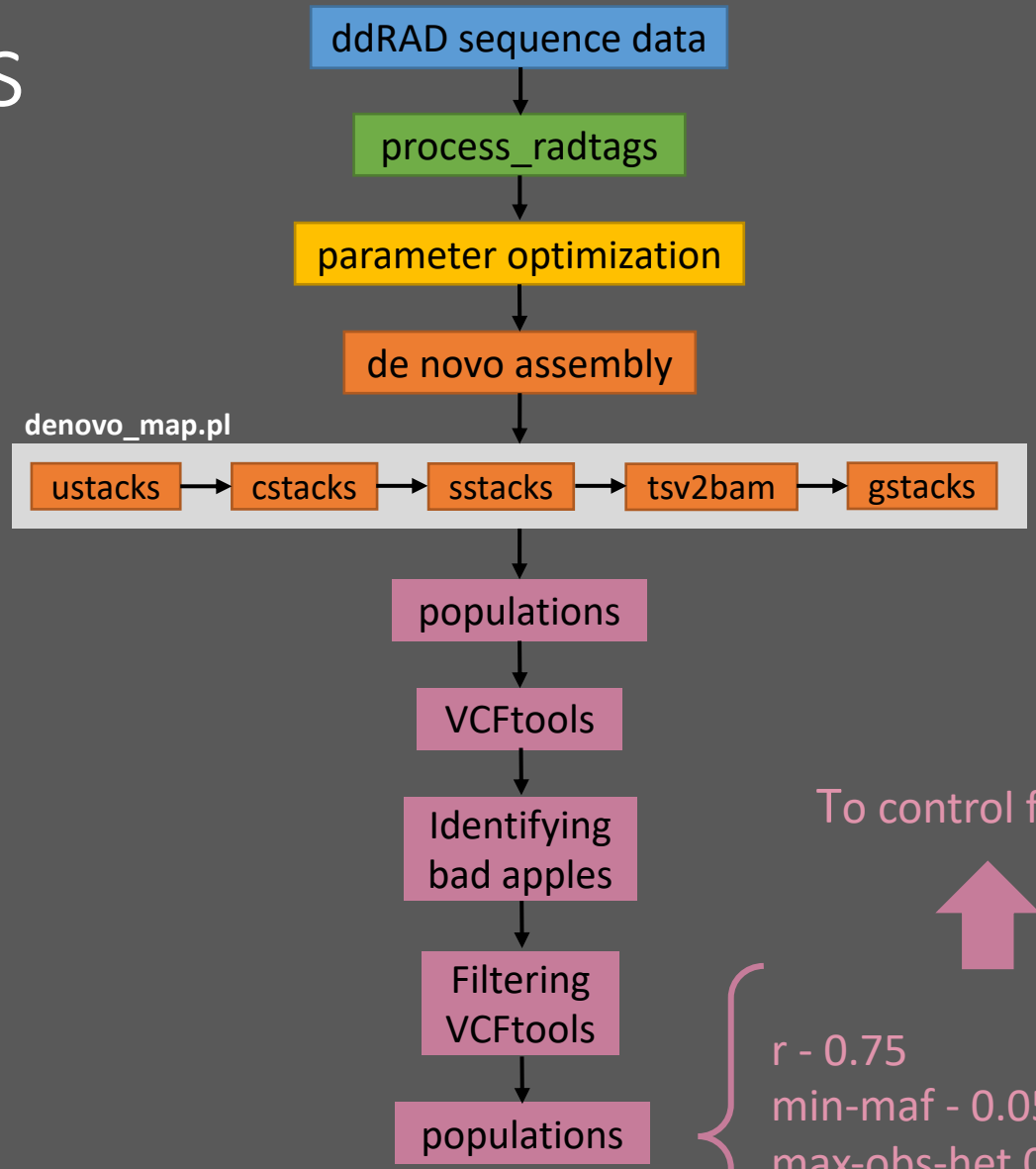
"variable sites with a maximum level of 0.7 heterozygosity will be included"

Bioinformatics Workflow

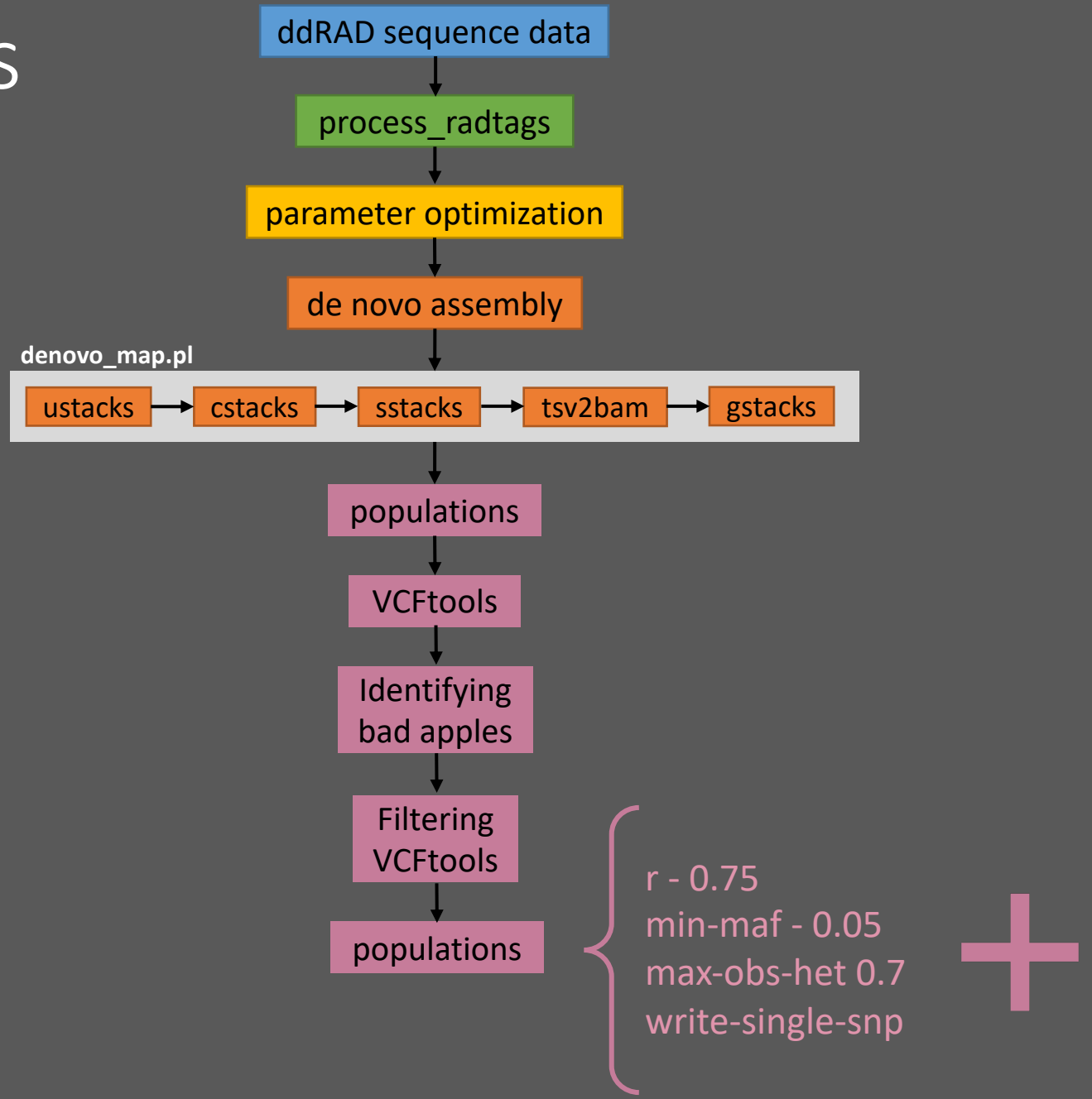


"decreases linkage disequilibrium and increases reproducibility"

Bioinformatics Workflow



Bioinformatics Workflow



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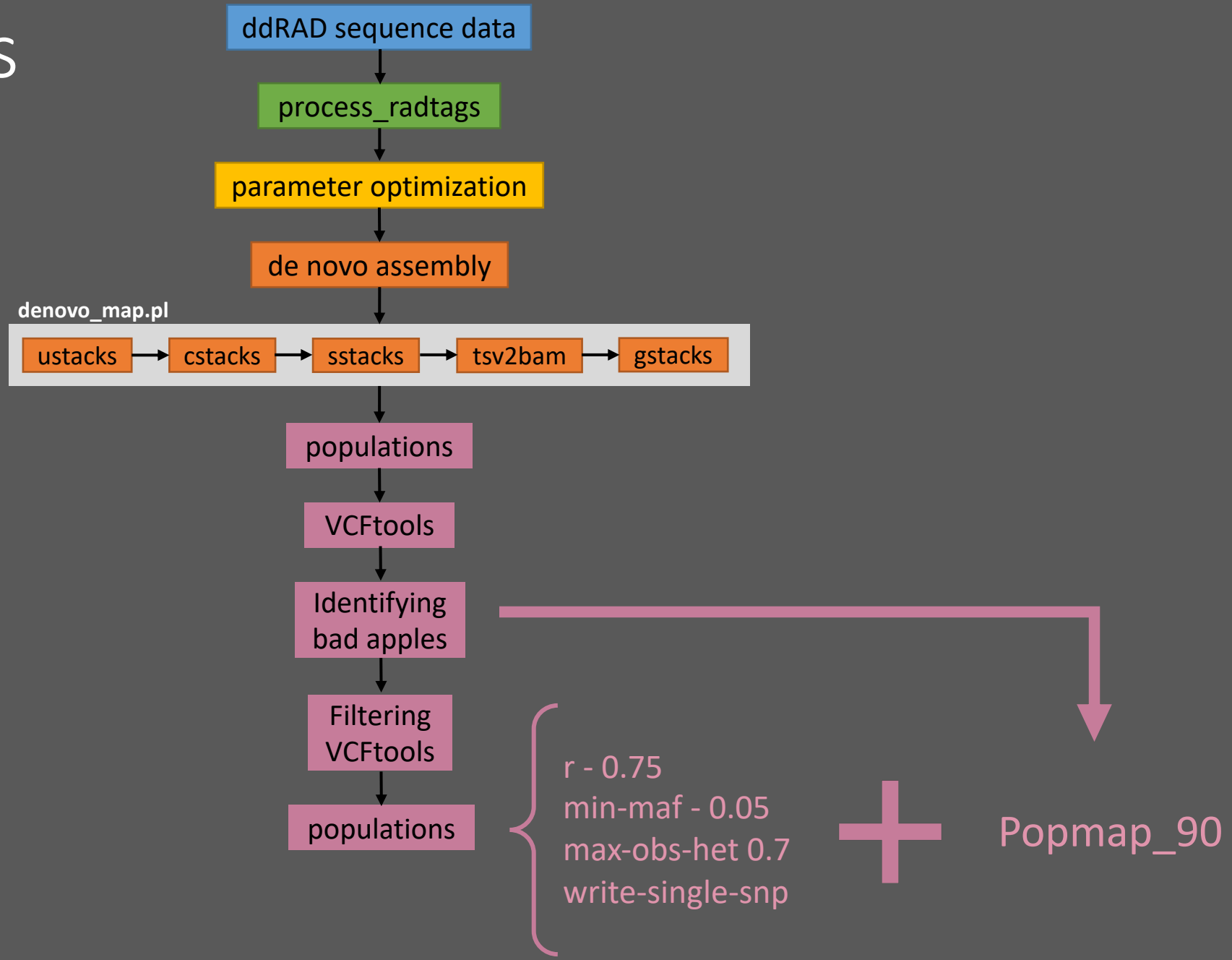
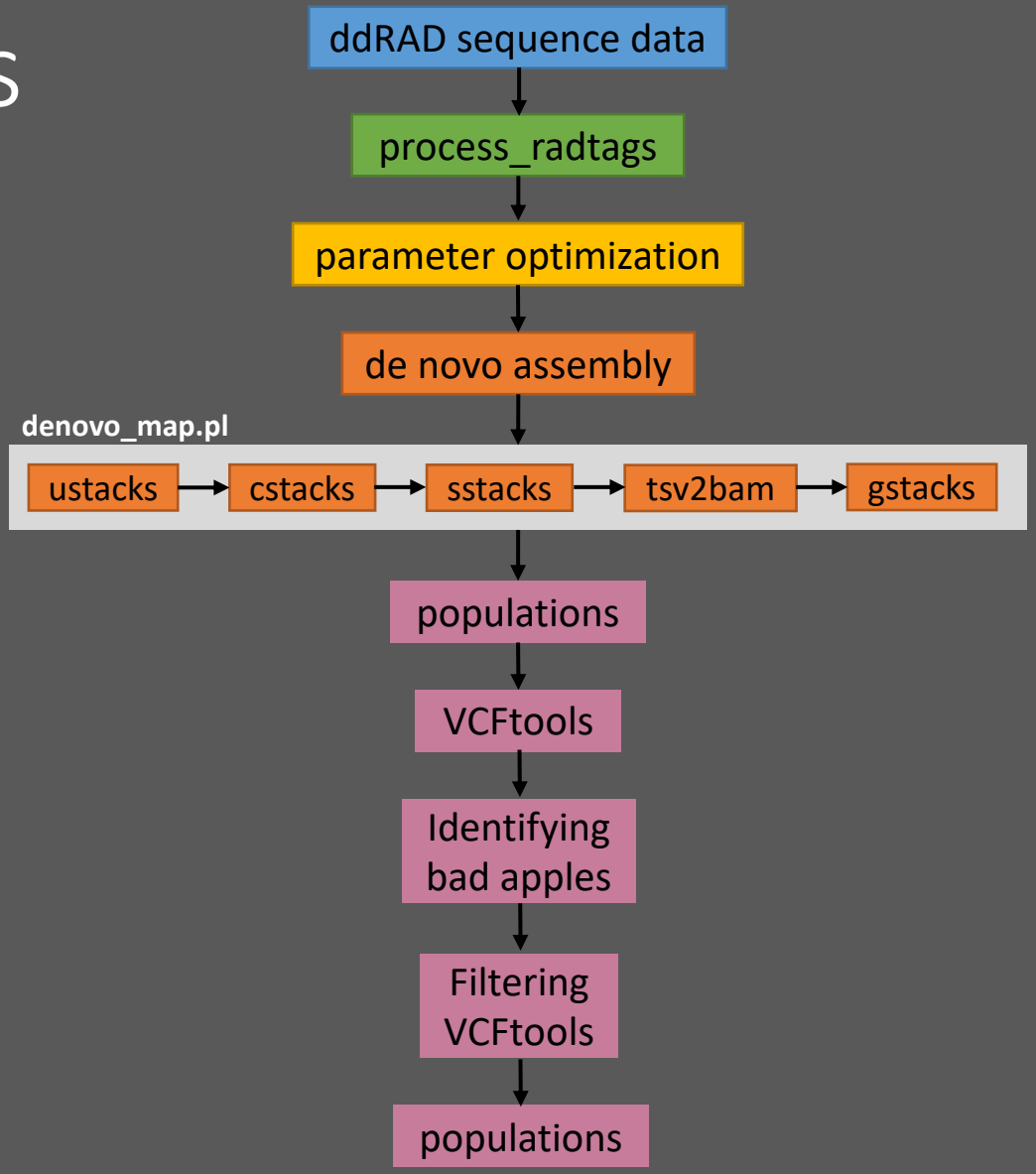


Bioinformatics Workflow



Bioinformatics Workflow