Dependencies

Snakemake

Python

R

Installation

1.
2. Snakemake
   Installation via Conda:
   a. conda install -n base -c conda-forge mamba
   b. mamba create -c conda-forge -c bioconda -n snakemake snakemake
   c. conda activate snakemake
   Find other options in https://snakemake.readthedocs.io/en/stable/getting_started/installation.html

Pipeline Description

The pipeline structure follows the Snakemake modularization. Modules are set in a main snake file which will be called for the workflow execution (Figure 1). A module consists of one or more rules which decompose the workflow into small steps by specifying how to create sets of output files from sets of input files. A rule can not be run if the input files are not found, also if an error occurs during the execution the output files created are deleted to keep consistency in the workflow in case a corrupted output is created. A rule also could include configurable parameters such as params, shell, messages, among others (Figure 2). File names, paths and parameters can be set in specific configuration files for easy modifications and administration (Figure 3 and Figure 4).

```
# Config files containing hard coded variables
configfile: os.path.abspath(os.path.join(os.path.dirname(workflow.basedir), 'snakefiles/config.json'))

# Runtime variables
include: "variables.py"

# rule all
rule all:
    input:
        dupconcordant = config['data_dup_path'] + '/' + config['dup_concordant_file'],
        dupdiscordant = config['data_dup_path'] + '/' + config['dup_disconcordant_file']
        samplespassmerged = config['sample_merged_file']


# Include modules
include: "module1-data-conversion.snake"
include: "module2-data-calling.snake"
include: "module3-data-clean.snake"
```

Figure 1. Snakemake file. This file includes the main rule “all” (by default snakemake executes this rule), and all the modules which will be executed in the pipeline workflow. In this file you can define your pipeline adding or removing any module allowing a flexible workflow. The file config.json contains the paths names to access the files described in each rule. And, the variable.py file contains the path that should be created for the workflow structure results. This file also can define links to external tools, libraries and resources which could be called in any rule. And, it also can define global parameters and external parameters for external tools.

```
# Run PennCNV to detec CNVs
# 1. Input: PFB and GCmodel files
# 2. output: list of calls by sample in *.rawcn file
rule DETECT_CNVs:
    input:
        pbffile = config['pfb_file'],
        gcmodelfile = config['gcmodel_file']
    output:
       call_raw = expand(config['data_calling_path'] + '/' + '{prefix}' + '.rawcn', prefix=calling_prefix),
       call_log = expand(config['data_calling_path'] + '/' + '{prefix}' + '.log', prefix=calling_prefix)
    params:
        signalintensitydir =  config['data_intensity_path'],
        outputdir = config['data_calling_path'],
        hmmfile = config['hmm_file'],
        listfile = config['list_signal_files_file'],
        logfile =  config['log_path'] + '/' + 'data_calling.log',
    message: "Calling for CNV detection from {input} in {output}"

        shell:
            """
            #Calling from PennCNV command
            detect_cnv.pl \
                -test \
                -conf \
                -hmm {params.hmmfile} \
                -pfb {input.pbffile} \
                -directory {params.signalintensitydir} \
                -list {params.listfile} \
                -log {output.call_log} \
                -out {output.call_raw} \
                -gcmodel {input.gcmodelfile}

            #login
            samples_with_calls=`awk '{{print $5}}'  {output.call_raw} | sort -u | wc -l`
            total_calls=`wc -l  {output.call_raw} | cut -f1 -d' '`
            echo "Were found $samples_with_calls with $total_calls calls" >> {params.logfile}
```

Figure 2. Module and rule structure. This figure shows a Module composed by one rule. Most commonly, rules consist of a name, input files, output files, and a shell command to generate the output from the input. This rule also contains the *params* specification which defines the alternative parameters used in this rule. Labels in red specify the path name for a file or directory. They are defined in the config.json file.

```
{
    "final_report_file": "/path_to/GSA2016_308_025_FinalReport.txt",
    "signal_intensity_file": "/path_to/signal_intensity.txt",
    "list_signal_files_file": "/path_to/data_conversion/list.txt",
    "map_file": "/path_to/data_conversion/sample_map.txt",
    "snp_file": "/path_to/data_conversion/SNPfile.txt",
    "pfb_file": "/path_to/data_conversion/addison.pfb",
    "gcmodel_file": "/path_to/data_conversion/hg19.gcmodel",
    "gc_content_file": "/path_to/resources/gc5Base.sorted.txt",
    "hmm_file": "/path_to/resources/hhall.hmm",


    "data_conversion_path": "/path_to/data_conversion",
    "data_intensity_path" :  "/path_to/data_conversion/data_intensity",
    "data_calling_path": "/path_to/data_calling",
    "data_clean_path": "/path_to/data_clean",
    "graphic_path": "/path_to/graphic",
    "graphic_qc_path": "/path_to/graphic/qc",
    "log_path": "/path_to/logs",
    "data_dup_path": "/path_to/data_clean/DUP"

}
```

Figure 3. Config file. This file contains file and directory paths for input and output data. Any change in the file location just has to be modified in this file. Similarly, the output directory can be changed just by modifying this file. Changes are broadcast to all modules and rules.

```
### snakemake_workflows initialization ######################################
libdir = os.path.abspath(os.path.join(os.path.dirname(workflow.basedir), '../lib'))
resourcesdir = os.path.abspath(os.path.join(os.path.dirname(workflow.basedir), '../resources'))

### programs
plink = "plink"
shapeit = "path_to/shapeit.v2.904.3.10.0-693.11.6.el7.x86_64/bin/shapeit"
vcf_sort = "/path_to/vcftools_0.1.13/perl/vcf-sort"
bcftools = "/path_to/bcftools/bcftools-1.8/bin/bcftools"
### prefix
### module 1,2 and 3
signal_prefix = "split"
calling_prefix = "sampleall"
plink_prefix = 'study_raw'

### Workflow parameters ###################################
PLINK_EXT =['.bed','.bim','.fam']
TPLINK_EXT =['.tped','.tfam']

### variables #############################################
### PennCNV #######
qcnumcnv = "50"
wf = "0.05"
qcbafdrift = "0.01"
qclrrsd = "0.3"

### Create paths if don't exist ###########################################
### data_conversion and data_preparation modules ###
if not os.path.exists(config['log_path']):
    os.makedirs(config['log_path'])
if not os.path.exists(config['data_conversion_path']):
    os.makedirs(config['data_conversion_path'])
if not os.path.exists(config['data_intensity_path']):
    os.makedirs(config['data_intensity_path'])
if not os.path.exists(config['data_calling_path']):
    os.makedirs(config['data_calling_path'])
if not os.path.exists(config['data_clean_path']):
    os.makedirs(config['data_clean_path'])
if not os.path.exists(config['data_dup_path']):
    os.makedirs(config['data_dup_path'])
if not os.path.exists(config['graphic_path']):
    os.makedirs(config['graphic_path'])
if not os.path.exists(config['graphic_qc_path']):
    os.makedirs(config['graphic_qc_path'])
```

Figure 4. The variable.py file is a configuration file which contains the path structure created for the workflow to store results. This file also can define links to external tools, libraries and resources which could be called in any rule. And, it also can define global parameters and external parameters for external tools.

Calling and quality control pipeline

1. **Module Data Conversion**

Rule CREATE_INTENSITY_FILES

Description: Create the individual signal intensity files for use with PennCNV calling. Also create the map file containing the original ID sample with the new name assigned by PennCNV (split* when using the –numeric option to hide the original ID for sensitive data).

Input: signal intensity and final report file from GenomeStudio (Figure 5 and Figure 6).

Output: a file with the list of names for each intensity file created, the SNPs list file (name, Chr, Position), and the individual signal intensity (Figure 7).

A log file is generated inside the logs directory with the module name (logs/data_conversion.log) showing useful information for future reports.

Figure 5. Signal intensity file format. This figure shows the extraction of one patient intensity signal for the first four markers.

Name	Chr	Position	Patient1.GType	Patient1.B Allele Freq	Patient1.Log R Ratio
1:100292476	1	100292476	AA	0,00	-0,3792592
1:101064936	1	101064936	AA	0,01713777	-0,1650909
1:103380393	1	103380393	BB	0,95952	-0,02254537
1:104303716	1	104303716	BB	0,9974257	-0,09440669
1:104864464	1	104864464	BB	1	-0,08369645

Figure 6. Final report from Genome studio format

```
GSGT Version    2.0.4
Processing Date 1/17/2020 11:48 AM
Content         GSAMD-24v1-0_20011747_A4.bpm
Num SNPs        700078
Total SNPs      700078
Num Samples     6112
Total Samples   6112
[Data]
SNP Name        Sample ID       B Allele Freq   Log R Ratio
1:100292476     Patient1        0.0000          -0.3793
1:101064936     Patien1         0.0171          -0.1651
1:103380393     Patient1        0.9595          -0.0225
1:104303716     Patient1        0.9974          -0.0944
1:104864464     Patient1        1.0000          -0.0837
```

Name	ID.Log R Ratio	ID.B Allele Freq
1:100292476	-0.3793	0.0000
1:101064936	-0.1651	0.0171
1:103380393	-0.0225	0.9595
1:104303716	-0.0944	0.9974
1:104864464	-0.0837	1.0000

Figure 7. Individual signal intensity obtained by PennCNV

Rule CREATE_PennCNV_INPUT_FILES

Description: Generate the Population frequency of B allele (PFB) file and the GCModel file used by PennCNV in the CNV detection.

Input: a file with the list of names for each intensity file created, and the SNPs list file (name, Chr, Position).

Output: the pfb file and the gc-model file.

A log file is generated inside the logs directory with the module name (logs/data_conversion.log) showing useful information for future reports.

2. Module Data Calling

Rule DETECT_CNVs

Description: Run the PennCNV for CNVs detection

Input: the pfb file and the gc-model file.

Output: list of calls per individual in PennCNV *.rawcn (Figure 7) and log files. This file

```
chr1:149095346-149143879      numsnp=4      length=48,534      state2,cn=1 split1 startsnp=GSA-1:149095346 endsnp=GSA-rs79760750 conf=18.072
chr4:156963692-156966696      numsnp=3      length=3,005       state2,cn=1 split1 startsnp=rs4691238 endsnp=GSA-rs112975832 conf=12.029
chr5:97066021-97096042        numsnp=5      length=30,022      state2,cn=1 split1 startsnp=GSA-rs76728709 endsnp=rs4380692 conf=20.359
chr6:76261562-76265642        numsnp=3      length=4,081       state2,cn=1 split1 startsnp=rs61234544 endsnp=rs9360921 conf=16.661
chr11:95979334-95980389       numsnp=3      length=1,056       state2,cn=1 split1 startsnp=rs79686966 endsnp=rs535912 conf=14.694
chr14:106208082-106232585     numsnp=3      length=24,504      state2,cn=1 split1 startsnp=GSA-rs11621259 endsnp=BOT2-rs10136766 conf=8.561
chr22:42527793-42528976       numsnp=4      length=1,184       state2,cn=1 split1 startsnp=rs1080989 endsnp=rs28360521 conf=17.310
chr3:155481492-155505991      numsnp=6      length=24,500      state5,cn=3 split1 startsnp=chr3-155481492 endsnp=rs112074828 conf=16.732
chr7:141763387-141775383      numsnp=10     length=11,997      state5,cn=3 split1 startsnp=chr7-141763387 endsnp=GSA-rs77848363 conf=28.198
chr8:12009572-12009597        numsnp=3      length=26          state5,cn=3 split1 startsnp=GSA-rs9773610 endsnp=GSA-rs75619199 conf=10.326
```

shows CNV chromosome coordinates, number of SNPs contained in the CNV, length, type of variation (CN=0 or 1 means there is a deletion and CN>=3 means there is a duplication), the starting marker identifier and the ending marker identifier in the CNV, respectively.

A log file is generated inside the logs directory with the module name (logs/data_calling.log) showing useful information for future reports.

Figure 7. PennCNV calling file rawcn format.

3. Module Data Clean

Rule FILTER_LOW_QUALITY_SAMPLES

Description: filter low quality samples based mainly on LRR and BAF statistics.

Input: rawcn and log file from PennCNV.

Output: rawcn file with good quality samples and their CNVs, file with the list of samples passed the QC, and summary file with all samples and their calls (passed and not passed the QC) include LRR and BAF statistics.

A plot showing the statistics LRR, BAF and WF distribution is generated. Also a log file is created inside the logs directory with the module name (logs/data_clean.log) showing useful information for future reports.

Rule REMOVE_SPURIOUS_CNVs

Description: Remove CNVs from spurious regions such as HLA, centromere and telomere.

Input: rawcn file with good quality samples and their CNVs.

Output: rawcn file with good quality samples and their CNVs not located on spurious regions

A log file is generated inside the logs directory with the module name (logs/data_clean.log) showing useful information for future reports.

Rule MERGING_ADJACENT_CNV

Description: Merge adjacent CNVcalls, in one single one, that could have been split from a large CNVs (>500kb) into smaller parts.

Input: rawcn file with good quality samples and their CNVs not located on spurious regions.

Output: merged CNVs in rawcn format. Violin plot showing the calls number distribution before and after the merging process is also generated.

A log file is generated inside the logs directory with the module name (logs/data_clean.log) showing useful information for future reports.

Rare copy number variants (CNV) pipeline

1. Module Data Conversion

Rule CONVERT_PENNCNV_TO_PLINK_FORMAT

Description: convert rawcn files from PennCNV to Plink cnv, fam and map files. File with **all** CNVs previously detected at Module Data Calling in QC pipeline is converted in plink format. Information about number of calls, number of cases, controls, males and females can be extracted from these files. External function (function.sh) is referenced in this rule to execute external bash code.

Input: file with all calls (rawcn format) previously detected.

Output: cnv, fam and map files for all calls (Plink format).

A log file is created inside the logs directory with the module name (logs/data_conversion.log) showing useful information for future reports.

Rule EXTRACT_CORE_SAMPLES

Description: extract CORE homogeneous unrelated samples, and their CNVs in Plink format files. Users should provide the ID list of unrelated samples (e.g. European descendant individuals). This rule could be modified according to the project requirements. External bash code, included in the function.sh file, is called in this rule.

Input: rawcn file with clean CNVs (passed QC and merged CNVs) and the list of unrelated European samples (IDs in a column).

Output: core samples and clean CNVs in cnv, fam and map plink files format.

A log file is created inside the logs directory with the module name (logs/data_conversion.log) showing useful information for future reports.

Rule EXTRACT_FINAL_CNVs

Description: A set of clean big calls covered by a minimum of markers is extracted in this step. Both, length and number of markers, can be set at variables.py file (by default we use > 50 kb and > 5 markers). This rule could be adapted or removed according to the project requirements.

Input: rawcn file with core samples and clean calls (passed QC and merged CNVs)

Output: core samples and clean CNVs (passed QC and merged CNVs) bigger than 50 kb and covered by more than 5 markers, in cnv, fam and map plink files format. Individuals with 0 CNVs are also included in this final set.

A log file is created inside the logs directory with the module name (logs/data_conversion.log) showing useful information for future reports.

2. Module Burden Analysis

Rule BURDEN_CNVs

Description: Perform the global burden test for CNVS in the core samples, cases vs. controls.

```
*.cnv.grp.summary
TEST      GRP          AFF        UNAFF
N         ALL         2270         7728
RATE      ALL         1.92        1.927
PROP      ALL       0.8477       0.8536
TOTKB     ALL        362.6        356.8
AVGKB     ALL        160.5        157.2

*.cnv.summary.mperm
CHR             SNP         EMP1
  S            RATE     0.562744
  S            PROP     0.706829
  S           KBTOT     0.341366
  S           KBAVG     0.284572
  S           GRATE            1
  S           GPROP            1
  S           GRICH            1
```

Input: CNVs for core samples in cnv, fam and cnv.map files.

Output: burden test result for CNVs in the core samples, showed at *.summary.mperm and grp.summary files (Figure 8).

A log file is created inside the logs directory with the module name (logs/burden_analysis.log) showing useful information for future reports.

Figure 8. Burden test output. This test reports four tests in both cases and controls, RATE (Number of segments), PROP (Proportion of sample with one or more segments), TOTKB (Total kb length spanned), and AVGKB (Average segment size). Tests are based (1-sided) on comparing these metrics in cases versus. For more detail see https://zzz.bwh.harvard.edu/plink/cnv.shtml#write_cnvlist. In this example there in not apparent significant difference in any of the four test (EMP1 > 0.05)

Rule SPLIT_CNVs_DEL_AND_DUP

Description: Split CNVs in deletions and duplications and then obtain the frequency of deletions and duplications in cases vs. controls.

Input: CNVs for core samples in cnv, fam and cnv.map files.

Output: *.cnv.indiv files for deletions and duplications in core samples. Also generate a tsv file containing the deletions and duplications ratio, split by length (by default 50kb, 100kb, 200kb, 500kb and 1Mb) which can be configured in variable.py file. Moreover, generate a tsv file containing the total mean length (in KB) for deletions and duplications in cases and controls (Figure 9); and a plot showing the CNVs length-distribution in cases vs. controls

```
Deletion frequency TSV file
pheno   length  numCNV  ratio
1       50KB    3502    0.873317
1       100KB   1667    0.415711
1       200KB   424     0.105736
1       500KB   83      0.0206983
1       1000KB  12      0.00299252
2       50KB    1027    0.868866
2       100KB   460     0.389171
2       200KB   132     0.111675
2       500KB   30      0.0253807
2       1000KB  13      0.0109983

Total mean length (in KB) TSV file
Deletions
pheno   tot_length_CNVs(Kb)     tot_CNVs        tot_mean_length_CNVs(Kb)
1       463500                  3502            132.353
2       145182                  1027            141.365

Duplications
pheno   tot_length_CNVs(Kb)     tot_CNVs        tot_mean_length_CNVs(Kb)
1       757743                  4226            179.305
2       218126                  1243            175.484
```

(Figure 10).

A log file is created inside the logs directory with the module name (logs/burden_analysis.log) showing useful information for future reports.

Figure 9. TSV files generated in the module Burden Analysis. The first example shows the deletions frequency classified by length and phenotype (1:controls, 2:cases); and the second example shows the mean length for all deletions and duplications in controls and cases, 1 and 2 for pheno column respectively.

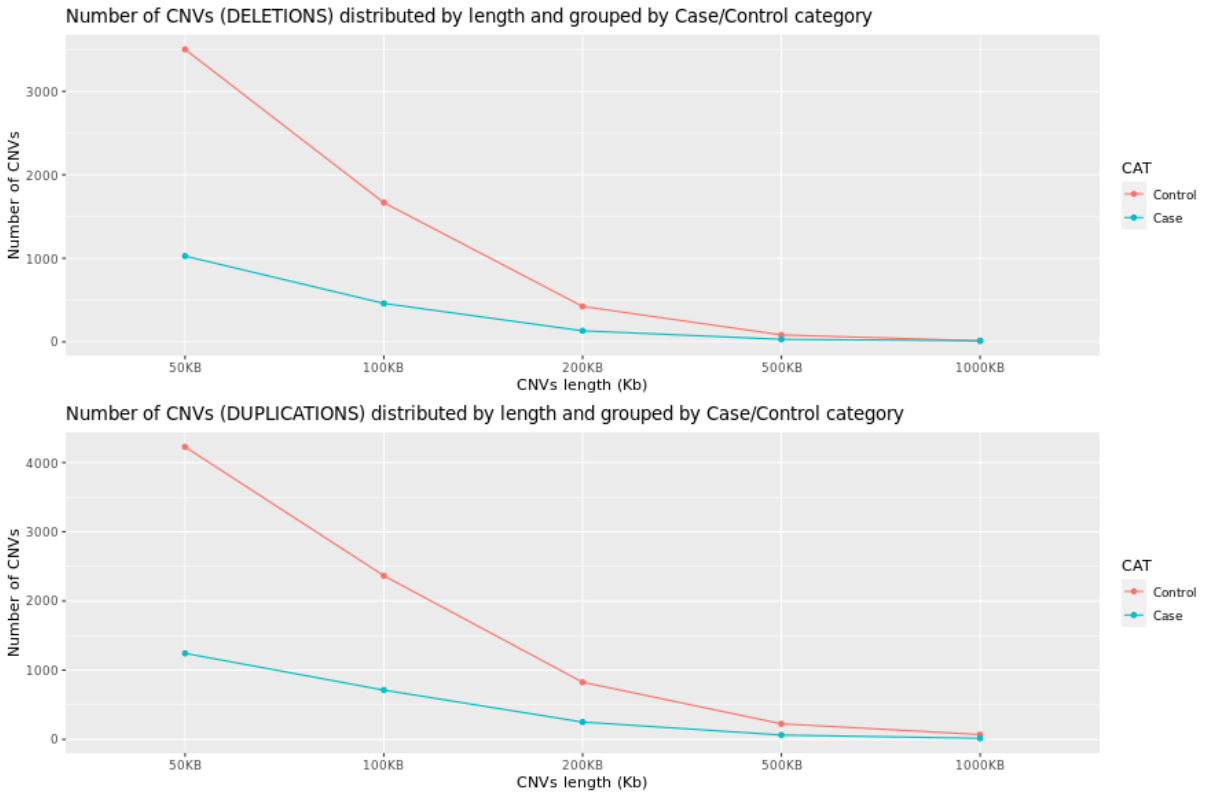

Figure 10. Length distribution in deletions and duplications. In this example CNVs are split in deletions and duplications and classified by cases and controls. Then they are distributed in 5 length intervals [50kb, 100kb, 200kb, 500kb, 1000kb]. This plot shows that bigger is the CNV length less is the number of CNVs in that interval, in cases and controls.

3. Module Rare CNVs Analysis

Rule GET_REF_CONTROLS_COMMON_CNVs

Description: Based on a subset of control samples, it obtains the **common** CNVs (deletions and duplications). The number of controls included in this subset can be defined in the config file variables.py. Common CNVs are defined as CNVs with high frequency, and this frequency also can be defined in variables.py.

Input: The ID list for the set of reference controls. These samples should be obtained randomly and including the same number of males and females (and any other population variable as nationality).

Output: *.cnv, *.fam and *.cnv.map files for reference controls and **common variants** (high frequency) split by type: deletions and duplications

A log file is created inside the logs directory with the module name (logs/rare_cnvs.log) showing useful information for future reports.

Rule FILTER_REFERENCE_CONTROLS

Description: Remove the reference control samples and their CNVs, from the original control cohort, since this dataset has been used to extract the common CNVs.

Input: Reference controls ID list, *.cnv, *.fam and *.cnv.map for common variants and reference controls.

Output: *.cnv, *.fam and *.cnv.map files for the new set of samples and CNVs which exclude the reference controls.

A log file is created inside the logs directory with the module name (logs/rare_cnvs.log) showing useful information for future reports.

Rule GET_RARE_CNVs

Description: Extract rare CNVs, deletions and duplications. All CNVs overlapping at least 50% with common variants will be removed to retain just low frequency CNVs.

Input: *.cnv, *.fam, *.map files for common variants and new dataset excluding reference controls.

Output: *.cnv, *.fam and *.cnv.map files for rare deletions and duplications. Also BED files were created for a graphic visualization in the Genome Browser and any genome viewer as IGV (Figure 11)

A log file is created inside the logs directory with the module name (logs/rare_cnvs.log) showing useful information for future reports.

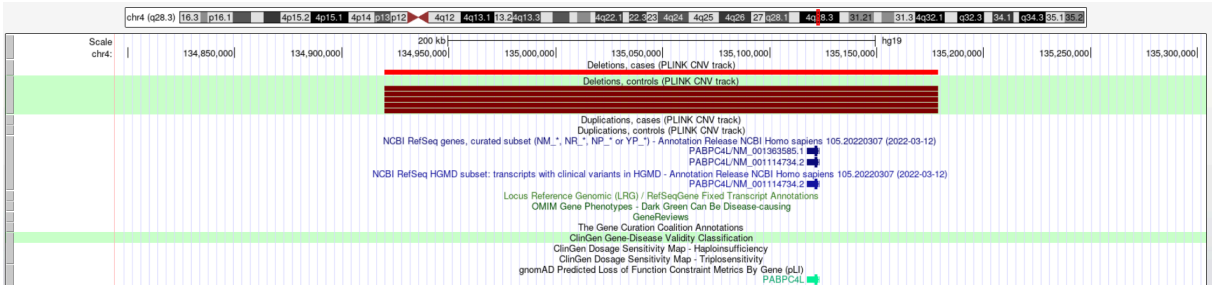

Figure 11. CNVs visualized in the Genome Browser. Deletions in cases and controls (light red versus dark red) visualized using the BED files.

Rule FREQUENCY_CNVs_INSPECTION

Description: Inspect the frequency of rare CNVs, deletions and duplications. CNVs regions (CNVR) are defined to calculate the CNVs frequencies. After that, frequency is plotted to verify that frequencies are approximately below the frequency threshold set in the variable.py file (Figure 12).

Input: *.cnv, *.fam, *.map files for rare deletions and duplications, and the fam file for all CNVs included in the study to calculate the frequency in each CNVR.

Output: *.cnv, *.fam and *.cnv.map files for CNVR. Bar plot showing the CNVs frequency distribution.

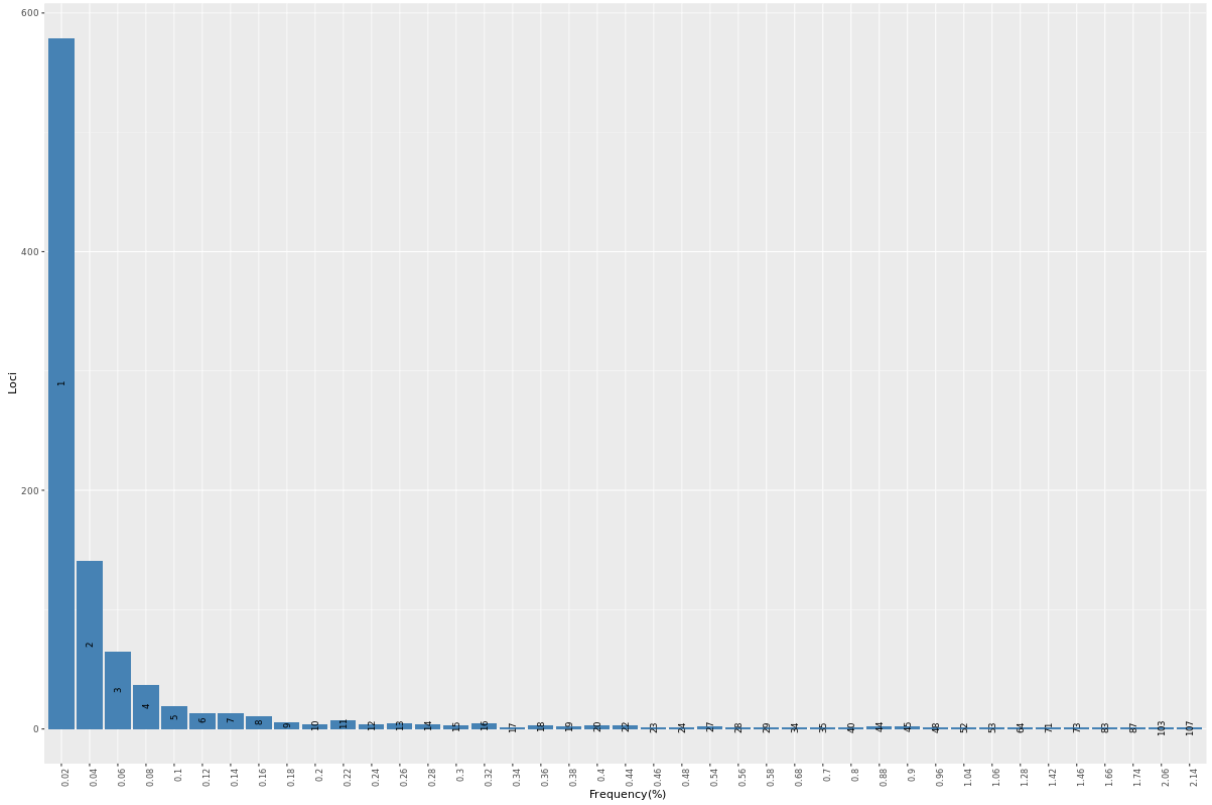

Figure 12. Deletions frequency distribution. After removing common variants, rare CNVs frequency are inspected showing the bar plot with the CNVs frequency which should be approximately below the threshold defined in the config file (in this example frequency should be approximately less than 2%) . The Y axis shows the number of CNVs in a specific locus (region), the X axis shows the CNVs frequency, and the number in the bar shows the amount of CNVs which represent this frequency. E.g for the second bar, there are 141 regions with 2 deletions in each, and the frequency of these deletions is equal to 0.04%, in a total of 4992 samples (2/4992).

Rule FREQUENCY_CNVs_ASSESSMENT

Description: assess the CNVs frequencies. A frequencies summary file for all deletions and duplications in cases and controls is created. This file includes the two proportion test evaluation and the odds ratio (OR) estimation. Similarly, a summary file containing the deletions and duplications frequencies split by CNVs interval sizes, and the statistics tests is created in this step. These results were graphically plotted in two forest plots, showing the frequencies confidence interval in each CNVs interval size, and the p-value associated (Figure 13). Also, bars plots are generated showing the proportion of individuals with specific number of CNVs, and the proportion of individuals with CNVs in each size interval.

Input: rare CNVs (deletions and duplications) *.cnv, *.fam and *.cnv.map files.

Output: summary frequencies *.csv files, forest plot and samples distribution bar plots. A log file is created inside the logs directory with the module name (logs/rare_cnvs.log) showing useful information for future reports.

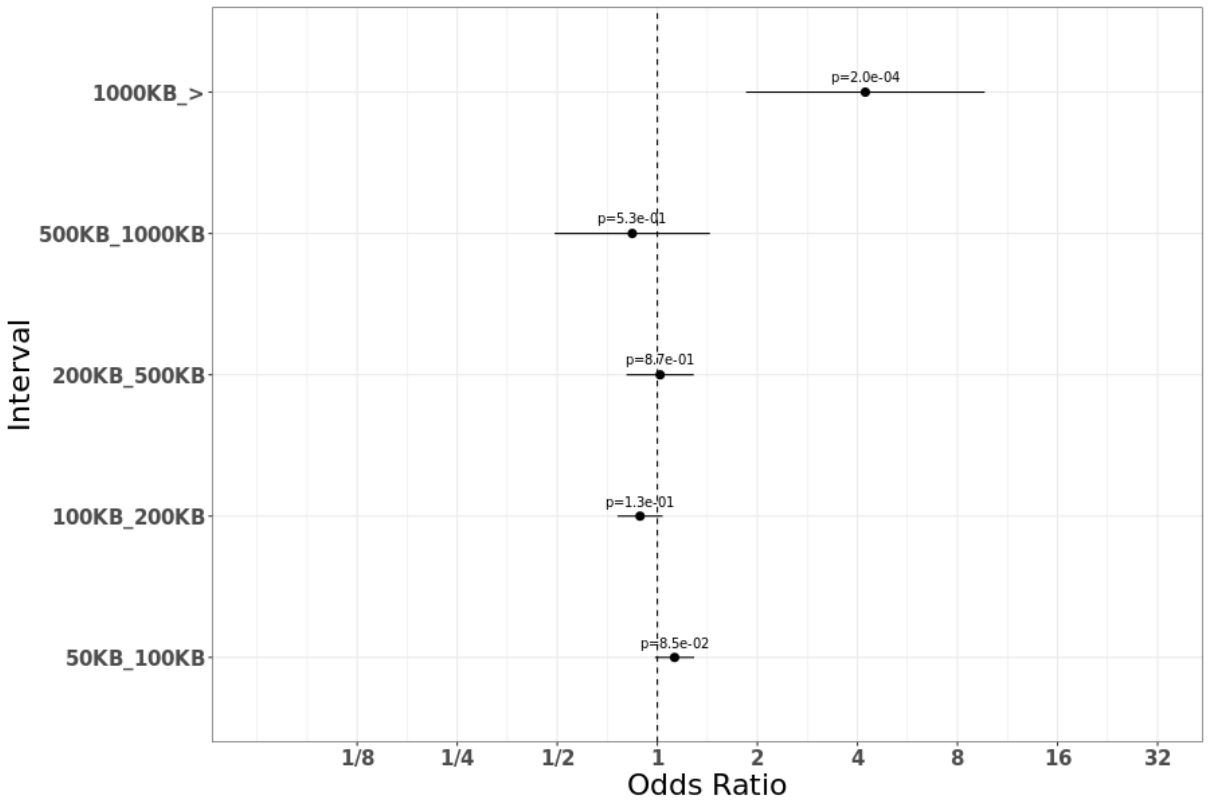

Figure 13. Deletions frequencies forest plot.

4. Module Rare CNVs Enrichment

Rule GENIC_ENRICHMENT_ANALYSIS

Description: Perform the basic geneset-enrichment test to evaluate which genes are enriched for CNVs. This test is a permutation-based test with 10000 null permutations to generate empirical p-values. Also, Gene regions borders are set at 20 kb around each region. Both, permutations and regions border, can be adjusted inside the rule.

Input: rare deletions and duplications *.cnv, *.fam and *.cnv.map files, and the gene annotation for the entire genome.

Output: *.cnv.burden files with the Performing GLM-based CNV burden test results. Also, a text file containing the list of genes (chromosome, coordinates, number of CNVs in cases and controls) overlapped by any CNV is generated.

A log file is created inside the logs directory with the module name (logs/enrichment_rare_cnvs.log) showing useful information for future reports.

Rule PATHWAY_ENRICHMENT_ANALYSIS

Description: Perform the basic geneset-enrichment test to evaluate whether a subset of genes are enriched, relative to the whole genome. This test is a permutation-based test with 10000 null permutations to generate empirical p-values. Also, Gene regions borders are set at 20 kb around each region. Both, permutations and regions border, can be adjusted inside the rule.

Input: rare deletions and duplications *.cnv, *.fam and *.cnv.map files, and the gene annotation for the entire genome and file of gene names forming the pathway to be tested.

Output: *.cnv.burden files with the Performing GLM-based CNV burden test results. Also, a text file containing the list of genes (chromosome, coordinates, number of CNVs in cases and controls) overlapped by any CNV is generated.

A log file is created inside the logs directory with the module name (logs/enrichment_rare_cnvs.log) showing useful information for future reports.