

# MultiGWAS report for Naive GWAS model

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This report, created by the MultiGWAS tool, is a summary of the input parameters used to run the tool and its main outputs. The outputs include the best-ranked and significant SNPs found for each tool. For each one, there is a score table, Venn diagram, and an SNP profile.

## 1 Input Parameters

MultiGWAS uses as input a configuration file where the user specifies the input genomic data (genotype and phenotype files) along with different values for parameters used by the MultiGWAS tool and by the other four GWAS software (GWASpoly, SHEsis, PLINK, and TASSEL).

The following table shows the current input parameters specified by the user in the configuration file:

PARAMETER	VALUE
Genotype filename	example-genotype.tbl
Phenotype filename	example-phenotype.tbl
Significance level (Genome-wide significance level)	0.05
Correction method (Bonferroni or FDR)	Bonferroni
GWAS model (Full or Naive)	Naive
NBest (Number of best-ranked SNPs to be reported)	7
Filtering (TRUE or FALSE)	TRUE
MIND Filter (Individual with missing genotype)	0.1
GENO Filter (SNPs with missing genotype)	0.1
MAF Filter (Minor allele frequency)	0.01
HWE Filter (Hardy-Weinberg test)	1e-10

## 2 Best-ranked SNPs

This section shows a table and a Venn diagram for the best N ranked SNPs (For this report, N=7). The configuration file allows defining the parameter N.

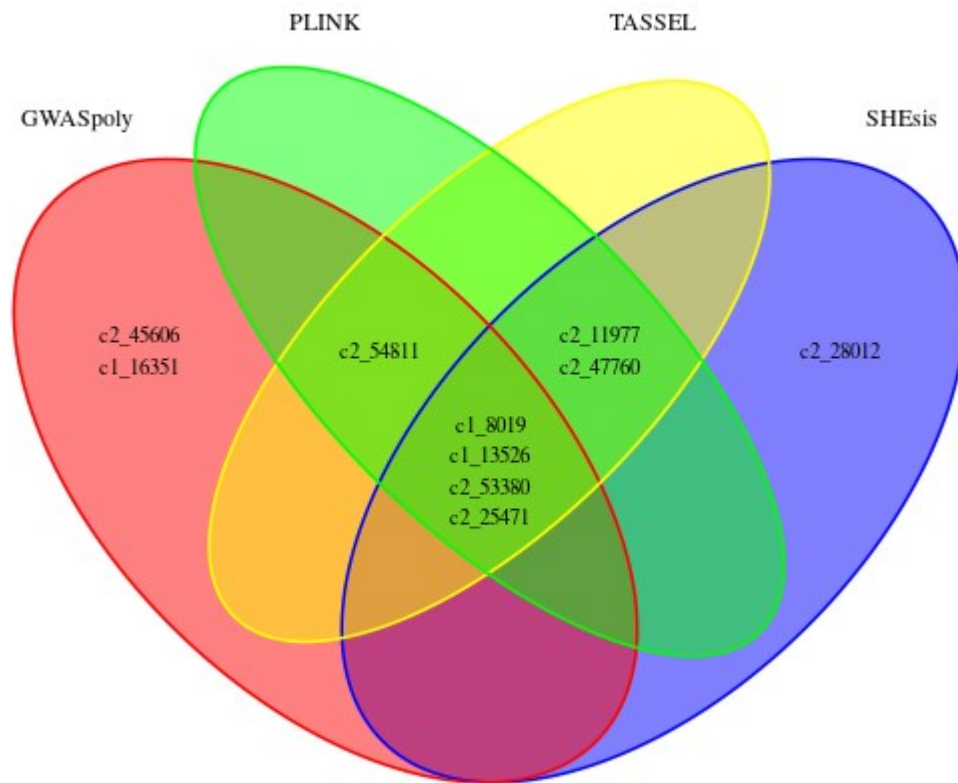
### 2.1 Table of best-ranked SNPs

The following table shows the associations between SNPs and the traits as transformed p-values to scores as  $-\log_{10}$  (p-value) sorted from the highest to the lowest scores based on each tool analysis and the model selected (Full or Naive). Each SNP has its associated information, such as the chromosome, position in the genome, and name. Also, based on the threshold calculated by a multiple testing correction (i.e., Bonferroni or FDR), the table marks the significance of each SNP as a TRUE or FALSE.

TOOL	MODEL	CHROM	POSITION	SNP	PVALUE	SCORE	THRESHOLD	SIGNIFICANCE
GWASpoly	Naive	10	48863165	c1_8019	0	11.560000	4.500000	TRUE
GWASpoly	Naive	10	48020996	c1_13526	0	10.610000	4.500000	TRUE
GWASpoly	Naive	1	70371898	c2_53380	0	10.370000	4.500000	TRUE
GWASpoly	Naive	10	48808404	c2_25471	0	9.890000	4.500000	TRUE
GWASpoly	Naive	1	46270954	c2_54811	0	8.540000	4.500000	TRUE
GWASpoly	Naive	10	48218826	c2_45606	0	7.970000	4.500000	TRUE
GWASpoly	Naive	10	48761642	c1_16351	0	7.970000	4.500000	TRUE
PLINK	Naive	5	32820618	c2_11977	0	6.848054	3.283301	TRUE
PLINK	Naive	10	13697423	c1_8019	0	5.040577	3.301030	TRUE
PLINK	Naive	1	46046095	c2_53380	0	4.967240	3.260071	TRUE
PLINK	Naive	2	72026885	c2_47760	0	4.771482	3.292256	TRUE
PLINK	Naive	1	49804489	c2_54811	0	4.691867	3.269513	TRUE
PLINK	Naive	10	30837971	c1_13526	0	4.560996	3.292256	TRUE
PLINK	Naive	10	39255236	c2_25471	0	4.536164	3.292256	TRUE
SHEsis	Naive	5	46046095	c2_53380	0	9.497204	3.313867	TRUE
SHEsis	Naive	2	13697423	c1_8019	0	8.031405	3.357935	TRUE
SHEsis	Naive	1	30837971	c1_13526	0	7.933190	3.330414	TRUE
SHEsis	Naive	2	32820618	c2_11977	0	6.565176	3.334454	TRUE
SHEsis	Naive	3	39255236	c2_25471	0	6.264880	3.342423	TRUE
SHEsis	Naive	4	72026885	c2_47760	0	5.941574	3.342423	TRUE
SHEsis	Naive	3	46475259	c2_28012	0	5.781570	3.318063	TRUE
TASSEL	Naive	5	3515956	c2_11977	0	8.830326	4.549984	TRUE
TASSEL	Naive	10	48863165	c1_8019	0	7.040577	4.549984	TRUE
TASSEL	Naive	1	70371898	c2_53380	0	6.926245	4.549984	TRUE
TASSEL	Naive	1	46270954	c2_54811	0	6.859964	4.549984	TRUE
TASSEL	Naive	2	20034673	c2_47760	0	6.762783	4.549984	TRUE
TASSEL	Naive	10	48020996	c1_13526	0	6.552175	4.549984	TRUE
TASSEL	Naive	10	48808404	c2_25471	0	6.527434	4.549984	TRUE

## 2.2 Venn diagram of best-ranked SNPs

The Venn diagram shows the best-ranked SNPs with high scores that are either unique or shared by the four software. Shared SNPs appear in the intersection areas of the diagram.



# 3 Significant SNPs

This section shows a table and a Venn diagram for the significant SNPs (score is above the significance threshold for each tool).

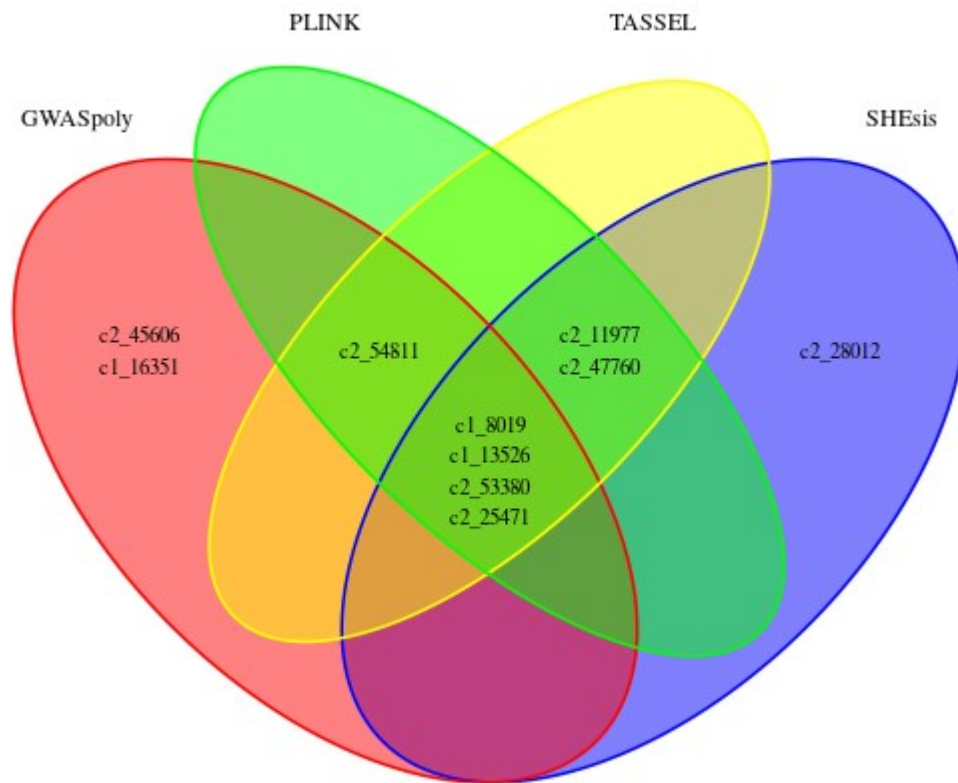
## 3.1 Table of significant SNPs

The following table shows the associations between SNPs and the traits as transformed p-values to scores as  $-\log_{10}(\text{p-value})$  sorted from the highest to the lowest scores based on each tool analysis and the model selected (Full or Naive). Each SNP has its associated information, such as the chromosome, position in the genome, and name.

TOOL	MODEL	CHROM	POSITION	SNP	PVALUE	SCORE	THRESHOLD	SIGNIFICANCE
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PLINK	Naive	5	32820618	c2_11977	0	6.848054	3.283301	TRUE
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TASSEL	Naive	10	48808404	c2_25471	0	6.527434	4.549984	TRUE

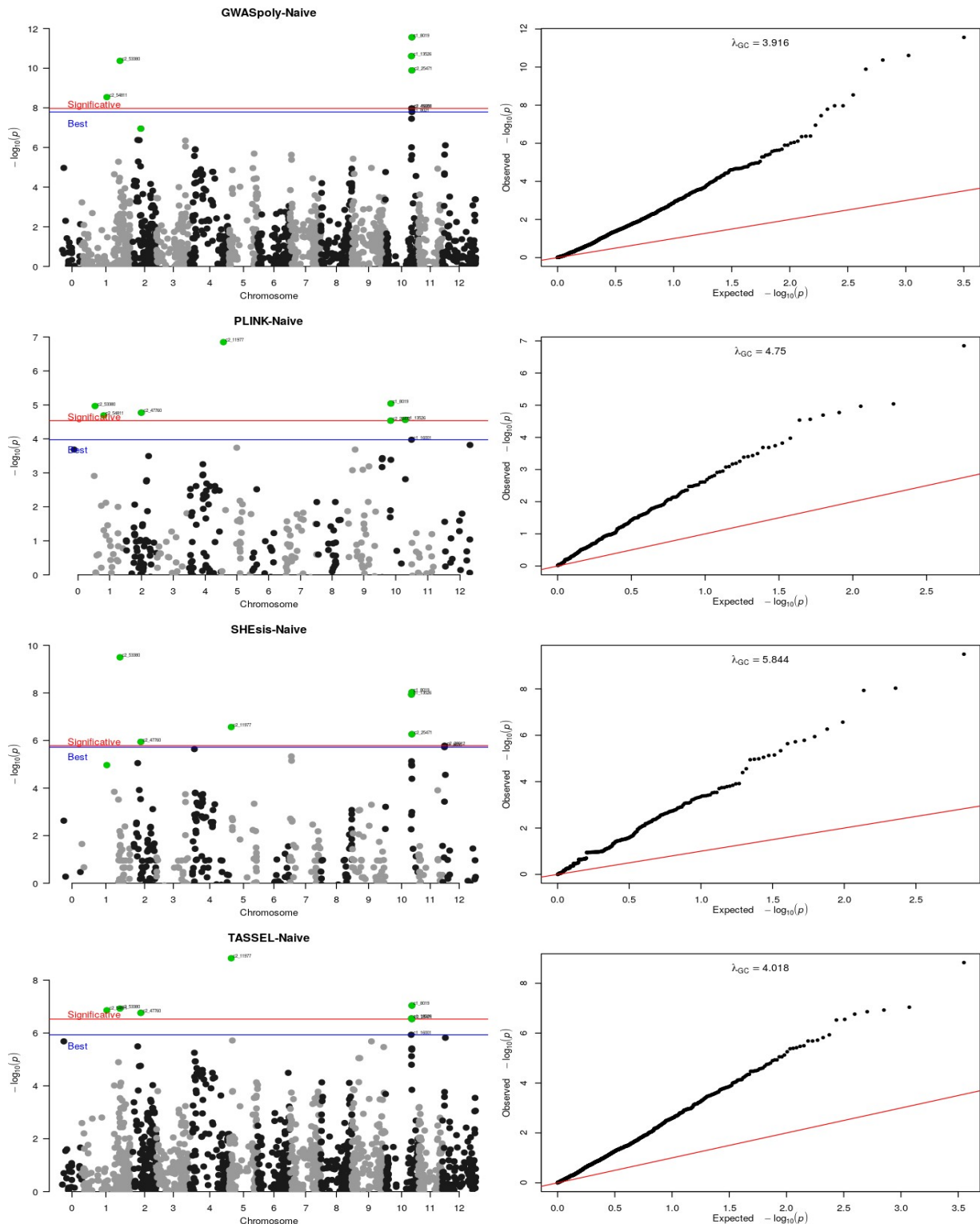
## 3.2 Venn diagram of significant SNPs

The Venn diagram shows the significant SNPs that are either unique or shared by the four software. Shared SNPs appear in the intersection areas of the diagram.



# 4 Manhattan and QQ plots

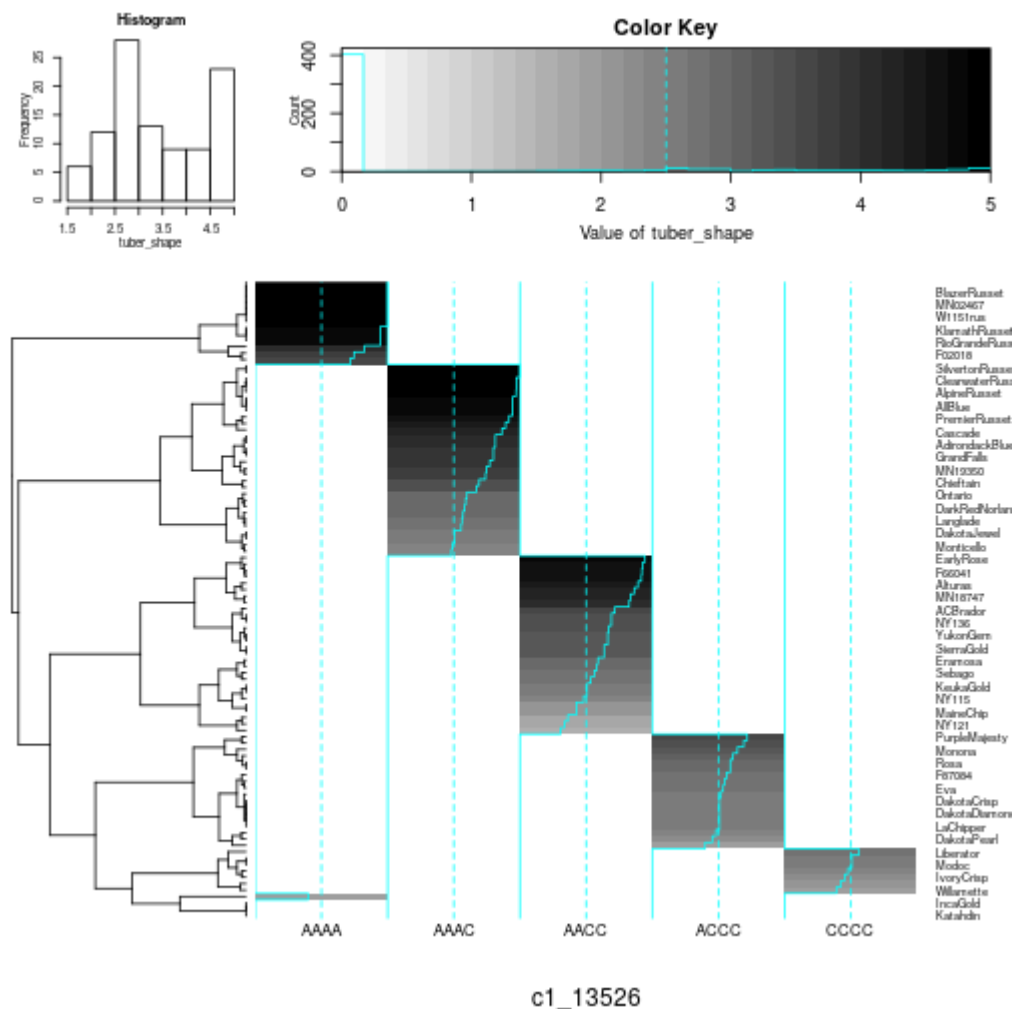
MultiGWAS uses classical Manhattan and Quantile–Quantile (QQ) plots to visually summarize GWAS results and identify both best-ranked and significant SNPs for each GWAS software. The Manhattan plot shows each SNPs distributed in the genetic location (x-axis) vs. the p-value (y-axis). The best-ranked SNPs are above the blue line and the significant SNPs above the blue line. In green are the SNPs that got high scores in more than one software. The QQ plot plots the observed (black dot lines) vs. expected  $-\log_{10}(p\text{-value})$  (red line).

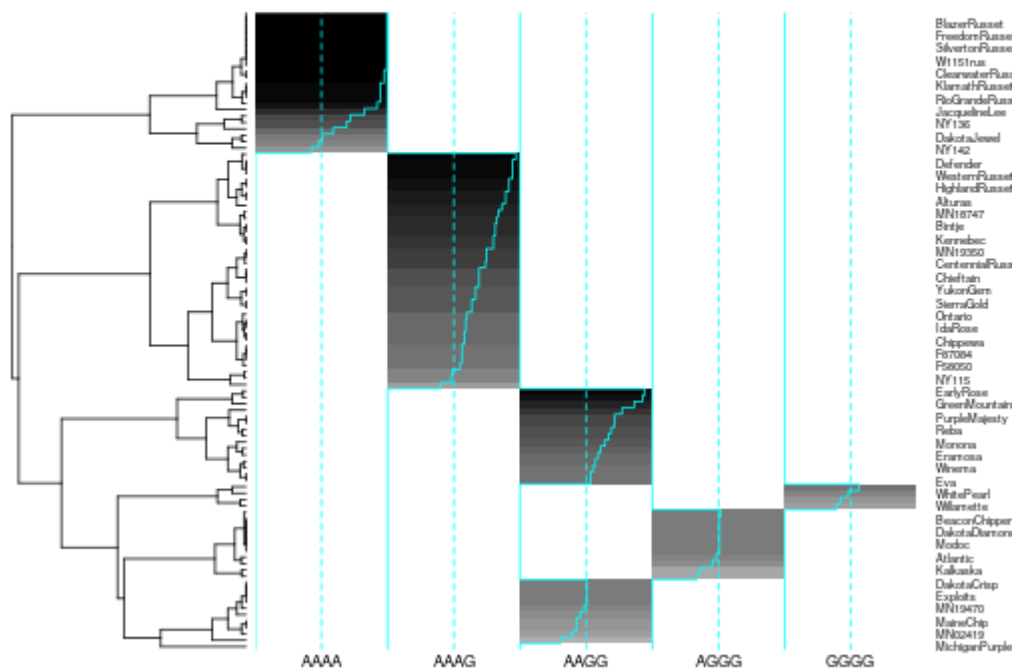
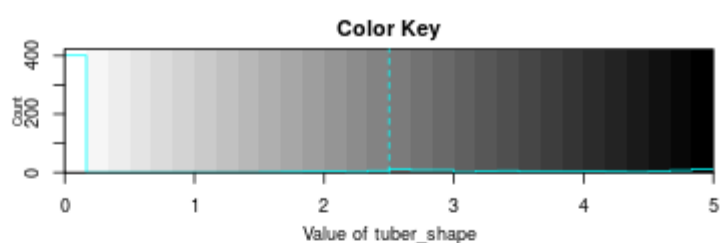


# 5 Profiles for common significant SNPs

For the common significant SNPs, MultiGWAS provides a figure to visualize each trait by individuals (rows) and genotypes (columns). At the left, the individuals are grouped in a dendrogram by their genotype. At the right, there is the name or ID of each individual. At the bottom, the genotypes are ordered from left to right, starting from the major to the minor allele (i.e., AAAA, AAAC, AACC, ACCC, CCCC). At the top, there is a description of the trait based on a histogram of frequency (top left) and by an assigned color for each numerical phenotype value using a grayscale (top right). Thus, each individual appears as a colored line by its phenotype value on its genotype column. For each column, there is a solid cyan line with the mean of each column and a broken cyan line that indicates how far the cell deviates from the mean.

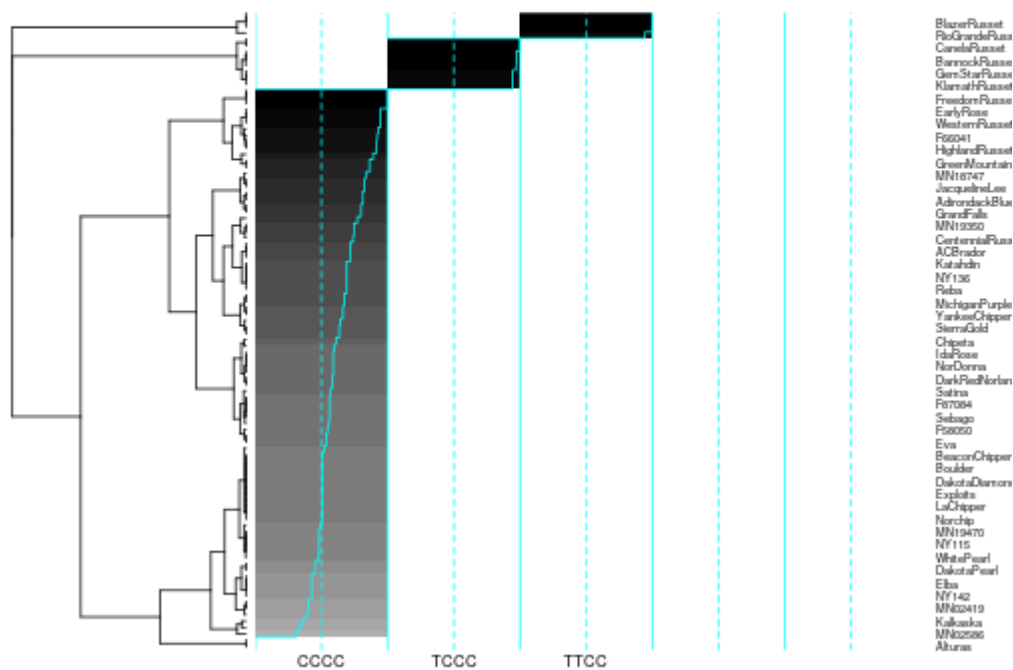
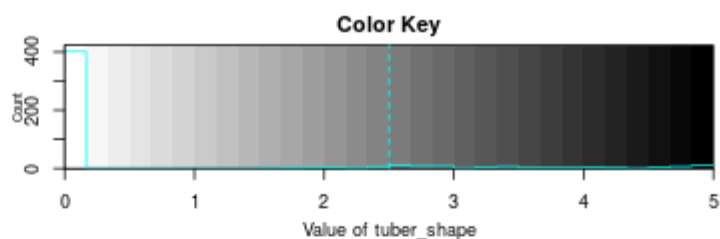
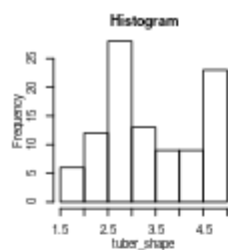
Because each multiGWAS report shows one specific trait at a time, the histogram and color key will remain the same for all the best-ranked SNPs.



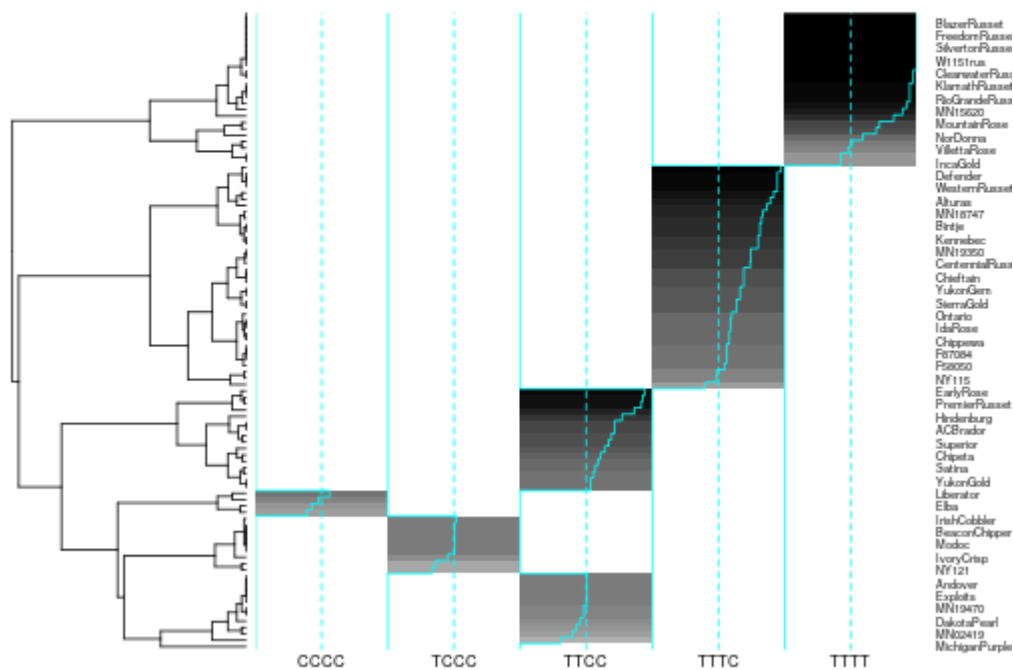
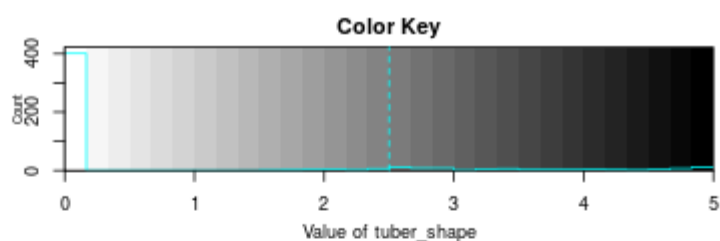
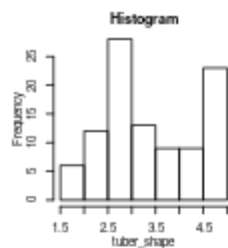


c1\_8019

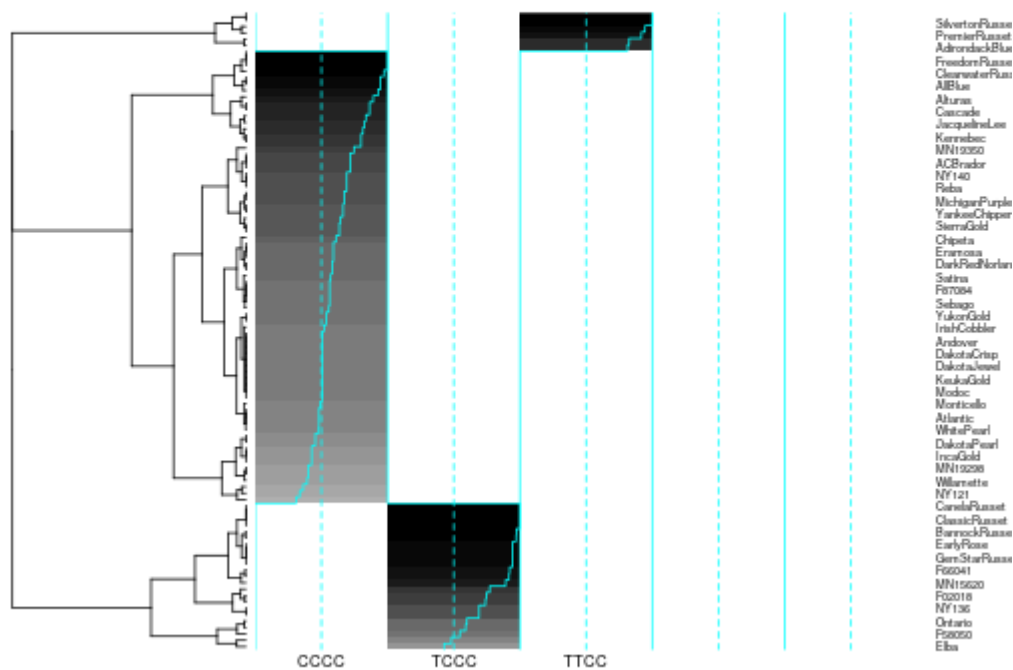
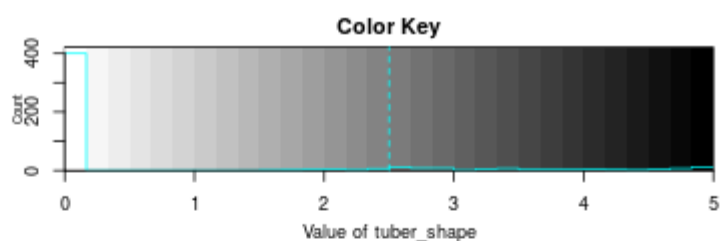
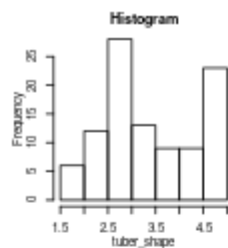




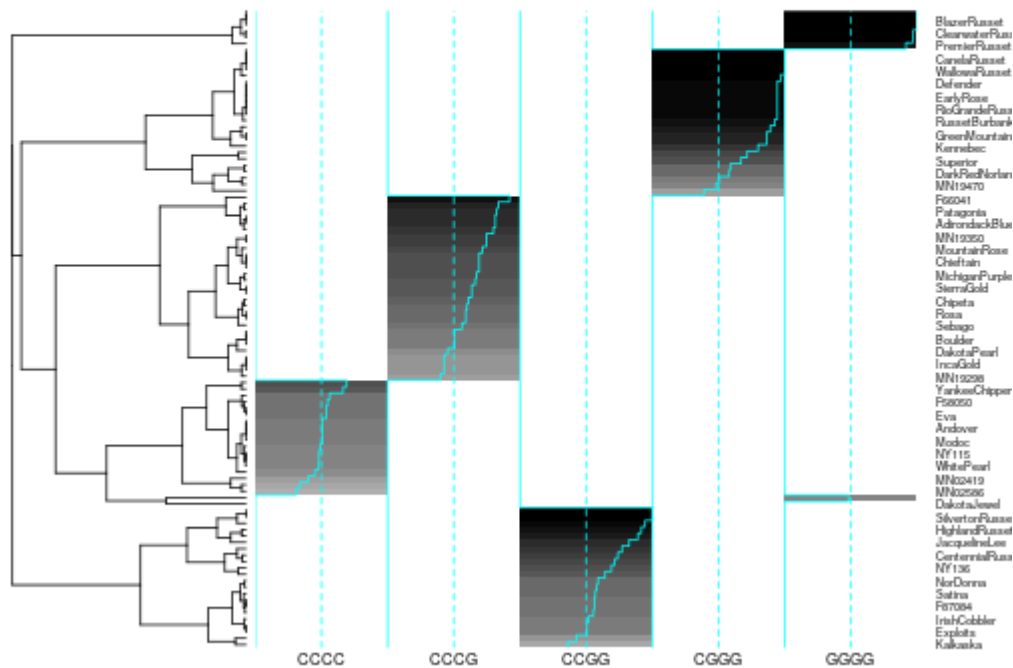
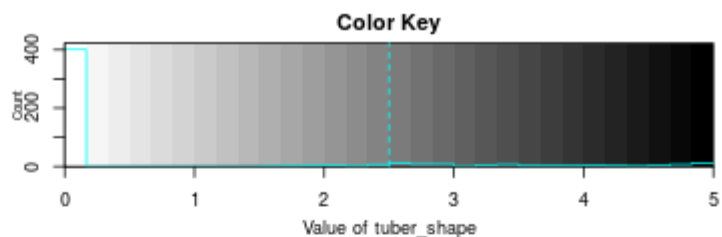
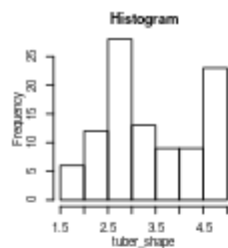
c2\_11977



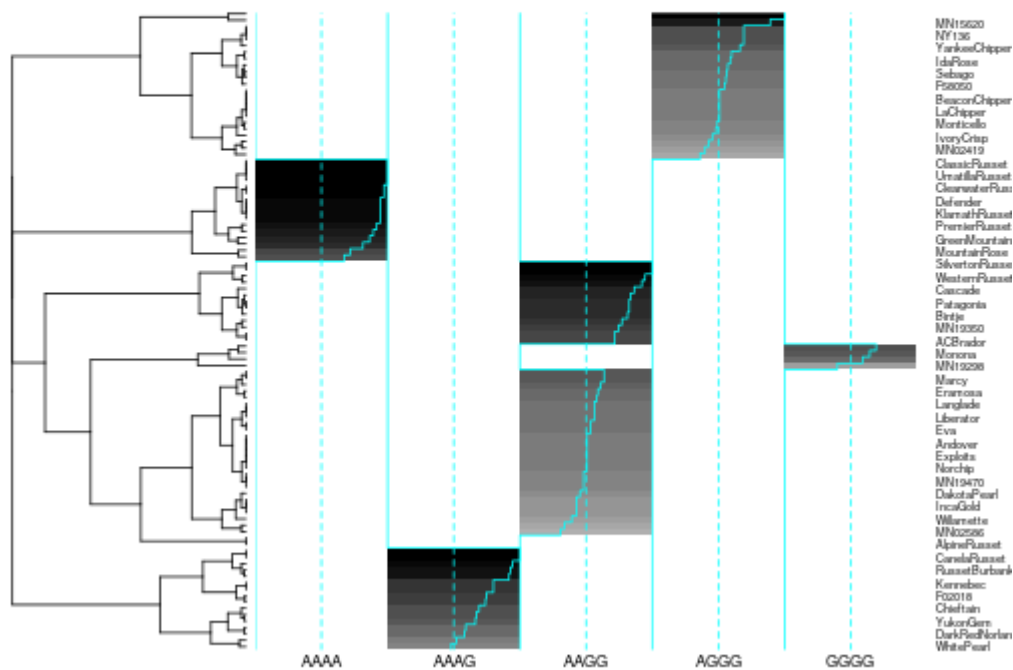
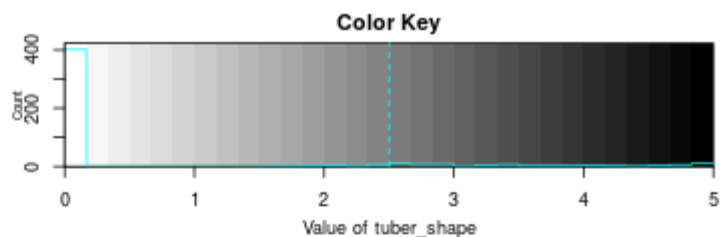
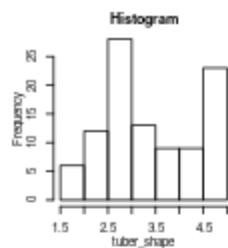
c2\_25471



c2\_47760



c2\_53380



c2\_54811