

GWAS Protocol

Notebook: GWAS

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1. Download data
2. Pylmm (get it from https://github.com/michaelbilow/pylmm_zarlab)
 1. python pylmmKinship.py --bfile (base of the binary plink files) outfile.kinship
 1. INPUT - Binary PLINK - *.bed *.bim *.fam all in same base - i.e.
 2. OUTPUT - Kinship matrix -- I prefer *.kinship with same base as the binary plink files
 2. Extract pheno file from fam file
 1. cat base.fam | awk '{print \$1, \$2, \$6}' >> base.pheno
 3. python pylmmGWAS.py -- bfile (base of the binary plink file) --kfile (the kinship including the extension) --phenofile (pheno file generated in step 2) outfile (*.gwas)
 1. INPUT - Plink Binary files, Kinship, Pheno
 2. OUTPUT - *.gwas file
 3. GWAS file
 1. SNP ID
 2. Beta value
 3. Stdev of Beta
 4. Z-score
 5. P-value
3. Manhattan Plot from GWAS
 1. Get map file from the Binary PLINK files
 1. plink --bfile (Base of the binary plink) --out (Base of the binary plink file)
 1. INPUT - Binary Plink
 2. OUTPUT - Regular Plink files (*.ped, *.map) -- the map file is required
 2. Extract the coordinates and p value for manhattan plot
 1. extract_data_for_manhattan_plot.py
 1. INPUT
 1. GWAS file from pylmm
 2. MAP file from plink
 3. OUTPUT file name
 - 2.
 3. R script for Manhattan Plot
 1. library(qqman)
f <- read.table('THE MANHATTAN PLOT DATA FILE FROM STEP 2')
colnames(f)[1]='CHR'
colnames(f)[2]='BP'
colnames(f)[3]='SNP'
colnames(f)[4]='P'
manhattan(f, main = "THE NAME OF THE STUDY", col = c("blue4", "orange3"))
4. Metasoft Analysis
 1. GET SNP_list and GET metasoft_input
 1. meta_analysis_prep.py
 1. INPUT == list of *.gwas files from pylmmGWAS
 2. metasoft - <http://genetics.cs.ucla.edu/meta/>
 1. java -jar (Metasoft.jar) -input (OUTPUT from meta_analysis_prep) -output

(DESIRED OUTPUT NAME) -mvalue

3. ForestPMPLOT

1. #python pmplot.py u4c/new_metasoft_input.txt
u4c/meta_result_with_m.txt u4c/study_names.txt u4c/study_order.txt
rs3013451 eh test_three.pdf
2. python pmplot.py (INPUT FOR METASOFT) (OUTPUT FROM METASOFT)
(list of study names) (list of order - 1 \n 2 \n 3\n 4\n ... is fine) (GENE
NAME of the SNP) (OUTPUT.pdf)

5. CAVIAR

1. ### SPECIFY THIS PART ###

6. Data Handling - if there are discrepancies in data

1. If the indexing is different
 1. It is in chr:coordinate, rather than rsid
 1. change_coordinate_gwas_file.py
 1. GWAS FILE
 2. INDEX FILE
 3. OUTPUT FILE
 1. outputs the gwas file with rsid
2. if map file does not have coordinates
 1. extract_data_for_manhattan_plot_without_map.py
 1. GWAS FILE
 2. INDEX FILE
 3. OUTPUT FILE
 1. outputs the manhattan plot data