

Joint/conditional analysis and fine-mapping

The INTERVAL data is used as reference panel. The logic of this specific directory is a simple solution of the dilemma that the reference data, possibly like others, uses reference sequence ID (rsid) whenever possible. However, during meta-analysis the practice of using rsid is undesirable so SNPID, i.e., chr:pos_A1_A2, (A1<=A2) is necessary.

After a rather long and laborious process involving many software, it turned out a simple way out is to obtain sentinels using SNPID but return to rsid at this stage and forward. The implementation here reflects this. The file INTERVAL.rsid contains SNPID-rsid mapping and could be generated from programs such as `qctool/bgenix/plink`.

A note on regions is ready. It is attractive to use the last genomic region from iterative merging for analysis and perhaps a flanking version. This is more appropriate than genomewide hard and fast 10MB windows or approximately independent LD blocks. For the latter, we found that the boundaries from the distributed 1000Genomes project were often inappropriate and one may not attempt to compute them for specific reference panel. Nevertheless, the iterative procedure actually just does empirically. Again the HLA region is condensed.

The last point regards software `finemap`, which uses summary statistics associated with the reference panel rather than that from meta-analysis.

Main analyses

| File specification | Function |
|------------------------------------|---------------------------------|
| NLRP2.sh | the exclusion list |
| cs.sh | Credible sets |
| ma.sh | INF1 sumstats |
| INTERVAL-ma.sh | INTERVAL sumstats |
| prune.sh | pruning |
| slct.sh | GCTA --cojo-slct analysis |
| finemap.sh | <code>finemap</code> analysis |
| jam.sh | <code>JAM</code> analysis |
| coloc.sb | coloc analysis -- clumsy verion |
| coloc.R | coloc analysis via pQTLtools |
| fastenloc.sb | fastenloc analysis |
| garfield.sh | GARFIELD analysis |
| hyprcoloc.sh | hyprcoloc analysis |
| st.sh | batch command file |
| cs/, finemap/, jam/, prune/, work/ | working directories |

Miscellaneous analyses and utilities

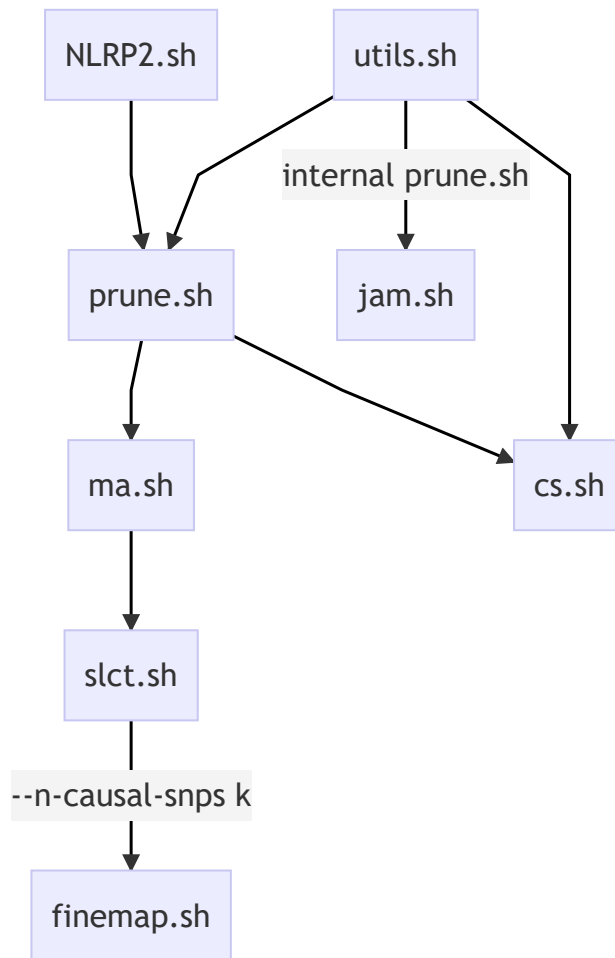
| File | Description |
|--------------|---|
| chembl.sh | toy ChEMBL queries |
| CRP.sh | an inflammation score |
| cvd1.sh | SCALLOP-CVDI supplementary tables |
| efo.R | experimental factor ontology |
| eQTL.R | cis-pQTL eQTL lookup (PhenoScanner) |
| GTEEx.R | cis-pQTL GTEEx eQTL lookup (PhenoScanner) |
| GTEEx.sh | cis-pQTL GTEEx eQTL lookup (GTEEx v8) |
| gdigt.R | GDI and gene-trials |
| gwasvcf.sh | GWAS to VCF conversion |
| gwas2vcf.sb | gwas2vcf + VCF operations |
| HGI.sh | HGI data analysis |
| latex.tex | LaTeX formulas embedded in Markdown |
| latex.docx | LaTeX formulas embedded in MicroSoft Word |
| LTBR.sh | LTBR LocusZoom plots |
| magma.sh | MAGMA for IL12B |
| neale.sh | signal selection for Neale UKB data (HbA1c) |
| pleiotropy.R | horizontal and vertical pleiotropy |
| ppi.R | PPI using EpiGraphDB |
| pqtIGWAS.R | pQTL-GWAS lookup |
| pQTL.R | pQTL lookup |
| pqtIMR.sh | pQTL MR analysis |
| rentrez.sh | reuse of rentrez |
| rGREAT.R | GREAT analysis |
| stringdb.sh | STRINGdb |
| tables.R | code to create Excel Tables |
| uniprot.R | UniProt IDs to others |
| utils.sh | utilities |
| vep.sh | VEP annotation |
| wgcna.sh | experiment on modules |

Stacked associaiton plots

1. IL.18-rs385076.sh
2. rs12075.sh
3. TNFB-rs2364485.sh
4. TNFB-rs2364485-MR.sh (a two-sample MR)
5. OPG-TRANCE.sh

Steps

`st.sh` conceptually executes the following elements,



Note that the `GCTA` `.ma`, `jma.cojo`, `.ldr.cojo` become `-rsid.ma`, `-rsid.jma.cojo`, `-rsid.ldr.cojo`, respectively; the same are true for files related to `finemap`.

Date last changed: **18/1/2021**