A)

Step 1: one file per protein per study: merge multi-chr, split two-study combinations, split one-big-file Ţ Step 2: have a standardized file naming scheme, unify column structure and naming schemes, re-map the a1/a2 columns for indels (I/D->A/ATC) T Step 3: standardize markernames to <chr>:<position hg19>:<A1 A2>, with A1_A2 as alphabetical sort and compare frequencies with 1kgenomes T first change in row-count: filtering SNPs according to imputation quality, as well as genotyped/imputed column ("IMP") П Step 5: run METAL analysis on prepared data

Meta analysis

Sumstats data freeze