

This code is for transforming genotype of plink file to BLUPF90 form

笔记本: R—code

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This code is for transforming genotype of plink file to BLUPF90 form

plink file form is as the following

- ped file: 6+2*N columns and individual for each row

1	11103832	0	0	0	0	G	G	G	G	C
2	11103863	0	0	0	0	G	G	A	G	C
3	11103458	0	0	0	0	G	G	G	G	C
4	11109536	0	0	0	0	G	G	G	G	T
5	11102017	0	0	0	0	G	G	G	G	T
6	11199000	0	0	0	0	G	G	G	G	C
7	11105007	0	0	0	0	G	G	G	G	C
8	11105687	0	0	0	0	G	G	A	G	T
9	11104882	0	0	0	0	G	G	G	G	C
10	11102935	0	0	0	0	G	G	G	G	C
11	11104849	0	0	0	0	G	G	G	G	T
12	11103090	0	0	0	0	G	G	A	G	C
13	11101680	0	0	0	0	G	G	G	G	T

- map file: 4 columns for chromosome, SNP name, Mendel distance and position

14	ARS-BFGL-BAC-10172	0	6371334
14	ARS-BFGL-BAC-1020	0	7928189
14	ARS-BFGL-BAC-10245	28.23	31819743
14	ARS-BFGL-BAC-10345	0	6133529
14	ARS-BFGL-BAC-10375	0	6616434
14	ARS-BFGL-BAC-10591	0	17544926
14	ARS-BFGL-BAC-10793	24.56396	29259114
14	ARS-BFGL-BAC-10867	36.53289	34639444
14	ARS-BFGL-BAC-10919	26.84388	31267746
10	ARS-BFGL-BAC-10952	43.86	18882288
10	ARS-BFGL-BAC-10960	45.20664	20609250
10	ARS-BFGL-BAC-10972	45.3343	20792754

using plink and the following step to transform A T G C into 0 1
2 (example for cattle)

```
plink --cow --file test --recodeA --out test_qc
```

test.raw file will be exist

FID	IID	PAT	MAT	SEX	PHENOTYPE	ARS-BFGL-BAC-11718_C	ARS-BFGL-BAC-13111_G																									
1	z11103832	0	0	0	-9	0	0	1	0	1	0	0	2	0	1	0	2	0	0	1	0	1	0	1	0	0	0	2	0			
2	11103863	0	0	0	-9	0	0	2	0	0	0	0	2	1	0	0	1	1	0	0	0	2	1	1	0	2	1	0	1	0	2	1
3	11103458	0	0	0	-9	2	1	2	2	1	0	0	1	1	0	0	1	1	0	2	1	0	0	1	0	0	1	0	1	0	1	0
4	11109536	0	0	0	-9	0	0	1	0	1	1	0	0	0	2	0	0	1	1	0	2	1	0	0	0	0	1	2	0	1	1	1
5	11102017	0	0	0	-9	0	0	1	0	0	1	0	0	1	0	0	1	0	2	0	0	0	2	0	1	1	0	0	1	0	1	1
6	11199000	0	0	0	-9	0	0	1	0	0	1	0	0	0	0	1	0	2	0	1	1	1	0	0	2	0	0	1	0	1	0	1

Now we can use R to transform the file

```
#set path
setwd("D:\\2-test\\20190807SNP\\")
genotype_file_name<-"test_qc.raw"
map_file_name<-"test.map"
out_name<-"genotype"
```

```

plink_blupf90<-function(genotype_file_name,map_file_name,out_name){
  if(!require(data.table)) install.packages("data.table")
  ped<-fread(genotype_file_name,header = F)
  map<-fread("test_qc.map",header = F)
  #genotype for blupf90
  ped<-ped[-1,-c(1,3:6)]
  ped[is.na(ped)]=5
  ped_blupf90<-matrix(nrow = nrow(ped),ncol = 2)
  n<-max(nchar(ped[1,]))
  for(i in 1:nrow(ped)){
    id<-as.character(ped[i,1])
    ped_blupf90[i,1]<-sprintf(paste("%",n,"s",sep = ""),id)
    geno<-as.numeric(ped[i,-1])
    ped_blupf90[i,2]<-paste(geno,collapse = "")
  }
  fwrite(as.data.frame(ped_blupf90),paste(out_name,"_blupf90",sep
= ""),row.names = F,col.names = F,sep = " ",quote = F)
  #map infomation for blupf90
  map$snp_order<-seq(1:nrow(map))
  fwrite(map[,c(5,1,4)],paste(out_name,"_map_blupf90",sep = ""),row.names =
F,col.names = F,sep = " ",quote = F)
}
plink_blupf90(genotype_file_name = genotype_file_name,map_file_name
= map_file_name,out_name = out_name)

```