Overview of GGtools for expression genetics

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1 Introduction

The *GGtools* package contains infrastructure and demonstration data for joint analysis of transcriptome and genome through combination of DNA expression microarray and high-density SNP genotyping data. For Bioconductor 2.2 we adopted a representation of genotypes due to Clayton (in package *snpMatrix*) allowing reasonably convenient storage and manipulation of 4 megaSNP phase II HapMap genotypes on all the CEPH CEU samples. This contrasts with the previous version of *GGtools* which was limited to 550 kiloSNP and 58 CEU founders.

To give an immediate taste of the capabilities, we attach the package and load some test data.

```
> library(GGtools)
> data(hmceuB36.2021)
> hmceuB36.2021
snp.matrix-based genotype set:
number of samples: 90
number of chromosomes present:
annotation: illuminaHumanv1.db
Expression data dims: 47293 x 90
Phenodata: An object of class "AnnotatedDataFrame"
  sampleNames: NA06985, NA06991, ..., NA12892
                                               (90 total)
 varLabels and varMetadata description:
    famid: hapmap family id
    persid: hapmap person id
    male: logical TRUE if male
    (7 total)
```

Expression data are recoverable in a familiar way:

> exprs(hmceuB36.2021)[1:5, 1:5]

```
NA06985 NA06991 NA06993 NA06994 NA07000 GI_10047089-S 5.983962 5.939529 5.912270 5.891347 5.906675 GI_10047091-S 6.544493 6.286516 6.244446 6.277397 6.330893 GI_10047093-S 9.905235 10.353804 10.380972 9.889223 10.155686 GI_10047099-S 7.993935 7.593970 8.261215 6.598430 6.728085 GI_10047103-S 11.882265 12.204753 12.249708 11.798415 12.015252
```

Genotype data have more complex representation.

> smList(hmceuB36.2021)

\$`20`

A snp.matrix with 90 rows and 119921 columns

Row names: NA06985 ... NA12892 Col names: rs4814683 ... rs6090120

\$`21`

A snp.matrix with 90 rows and 50165 columns

Row names: NA06985 ... NA12892 Col names: rs885550 ... rs10483083

> class(smList(hmceuB36.2021)[["20"]])

[1] "snp.matrix"

This shows that we use a named list to hold items of the *snp.matrix* class from *snpMatrix*. It will generally be unnecessary to probe to this level, but it is instructive to check the underlying representation:

- > schunk = smList(hmceuB36.2021)[["20"]]
- > schunk@.Data[1:4, 1:4]

	rs4814683	rs6076506	rs6139074	rs1418258
NA06985	03	03	03	03
NA06991	02	03	02	02
NA06993	01	03	01	01
NA06994	01	03	01	01

The leading zeroes show that a raw byte representation is used. We can convert to allele codes as follows:

> as(schunk[1:4, 1:4], "character")

```
rs4814683 rs6076506 rs6139074 rs1418258
NA06985 "B/B"
                   "B/B"
                              "B/B"
                                         "B/B"
                                         "A/B"
NA06991 "A/B"
                   "B/B"
                              "A/B"
                                         "A/A"
NA06993 "A/A"
                   "B/B"
                              "A/A"
NA06994 "A/A"
                   "B/B"
                              "A/A"
                                         "A/A"
```

The primary method of interest is the genome-wide association study, here applied with expression as the phenotype. Here we execute a founders-only analysis, adjusting for gender, confining attention to chromosome 20:

```
> pd = pData(hmceuB36.2021)
> hmFou = hmceuB36.2021[, which(pd$mothid == 0 & pd$fathid == 0)]
> f1 = gwSnpTests(genesym("CPNE1") ~ male, hmFou, chrnum(20))
```

2 Conversion to nucleotide codes

This is currently somewhat cumbersome. Suppose we want to know the specific nucleotide assignments for a given genotype call. For example, rs4814683 for subject NA06985.

```
> schunk["NA06985", "rs4814683"]
Autosomal snp(s):
[1] "B/B"
```

We need to know a) that the A/B tokens map in lexical order to the nucleotides (A will be the alphabetically first nucleotide for the diallelic call).

Using the SNPlocs. Hsapiens. dbSNP. 20071016 package, we can get the nucleotides:

```
> library(SNPlocs.Hsapiens.dbSNP.20071016)
> s20 = getSNPlocs("chr20")
> s20[s20[, 1] == 4814683, ]

RefSNP_id alleles_as_ambig loc
4 4814683 K 9795
```

Now we need to translate the IUPAC code to the nucleotides:

```
> library(Biostrings)
```

> IUPAC_CODE_MAP

```
Α
                    G
                           Τ
                                   М
                                           R
                                                           S
                                                                   Y
                                                                          K
  " A "
          "C"
                                "AC"
                                                "AT"
                                                        "CG"
                                                                "CT"
                                        "AG"
                                                                        "GT"
                                                                              "ACG"
            D
                    В
    Η
"ACT" "AGT" "CGT" "ACGT"
```