

Today is December 11, 2012.

```
> library("vcf2R")
> library("trioClasses")
> library("trio")
> data("BMP4-european-all.geno")
> data(ped, package = "trioClasses")
```

After loading the necessary packages and data we first make sure that the pedigree data frame contains fields F, M and O for father, mother and offspring ids. Note that these ids should match those in the vcf file.

```
> pedigreeInfo <- within(ped.df, {
  F <- as.character(fid)
  M <- as.character(mid)
  O <- as.character(id)
})
> tg.ped <- Pedigree(pedigreeInfo = pedigreeInfo)
> tg.ped
```

This pedigree object contains 1812 complete trios.

For access to the data frame use the `trios()` accessor function.

First we do our best to retrieve ids from the vcf/geno data and manipulate them to match the pedigree file.

```
> id.vec <- colnames(geno.mat)
> head(id.vec)

[1] "H_ME-DS11103_02-DS11103_02" "H_ME-DS11103_03-DS11103_03"
[3] "H_ME-DS11107_02-DS11107_02" "H_ME-DS11107_03-DS11107_03"
[5] "H_ME-DS11107_01-DS11107_01" "H_ME-DS11108_02-DS11108_02"

> foo <- strsplit(x = id.vec, split = "-")
> id.vec <- as.character(data.frame((do.call("rbind", foo)))[,
  3])
> length(id.vec)

[1] 218

> head(id.vec)

[1] "DS11103_02" "DS11103_03" "DS11107_02" "DS11107_03" "DS11107_01"
[6] "DS11108_02"
```

There are 218 subjects in the genotype matrix. Now with ids formatted properly we create the genotype object needed for the `gTrio` class, as well as the accompanying `ped` object such that all trio members have data in the genotype matrix. These values may all be NA, as NA is not precluded from the genotype matrix.

```
> genomat <- t(ifelse(geno.mat == "0/0", 0L, ifelse(geno.mat ==
  "0/1", 1L, ifelse(geno.mat == "1/1", 2L, NA))))
> colnames(genomat) <- rownames(geno.mat)
> rownames(genomat) <- id.vec
> geno.trio <- genoMat(tg.ped, genomat)
> (tg.ped.complete <- completeTrios(tg.ped, colnames(geno.trio)))
```

This pedigree object contains 65 complete trios.  
For access to the data frame use the `trios()` accessor function.

Now we create the `gTrio` object.

```
> (gTrio.obj <- gTrio(tg.ped.complete, geno = geno.trio))
```

This object has 65 trios and 1559 markers (likely SNPs).

```
> missing.snp <- rowSums(is.na(geno(gTrio.obj)))/dim(geno(gTrio.obj))[2]/dim(geno(gTrio.obj))[3]
> missing.subject <- colSums(is.na(geno(gTrio.obj)))/dim(geno(gTrio.obj))[1]
> length(missing.subject)
```

```
[1] 195
```

```
> length(missing.snp)
```

```
[1] 1559
```

```
> geno <- geno(gTrio.obj)
> maf <- rowSums(geno[, , 1:2], dims = 1, na.rm = TRUE)/rowSums(!is.na(geno[,
, 1:2]), dims = 1)/2
> maf <- ifelse(maf >= 0.5, 1 - maf, maf)
> summary(maf)
```

```
   Min. 1st Qu.  Median    Mean 3rd Qu.    Max.
0.00000 0.00000 0.00000 0.01857 0.00000 0.50000
```

```
> geno.trio.2 <- geno.trio[-c(which(missing.snp > 0.1),
which(maf < 0.05)), ]
> (gTrio.obj <- gTrio(tg.ped.complete, geno = geno.trio.2))
```

This object has 65 trios and 111 markers (likely SNPs).

```
> geno <- getGeno(gTrio.obj, type = "holger")
> geno <- geno(gTrio.obj)
> maf2 <- rowSums(geno[, , 1:2], dims = 1, na.rm = TRUE)/rowSums(!is.na(geno[,
, 1:2]), dims = 1)/2
> maf2 <- ifelse(maf2 >= 0.5, 1 - maf2, maf2)
```

Now, it's easy to perform any method in Holger's trio package, such as `allelicTDT`.

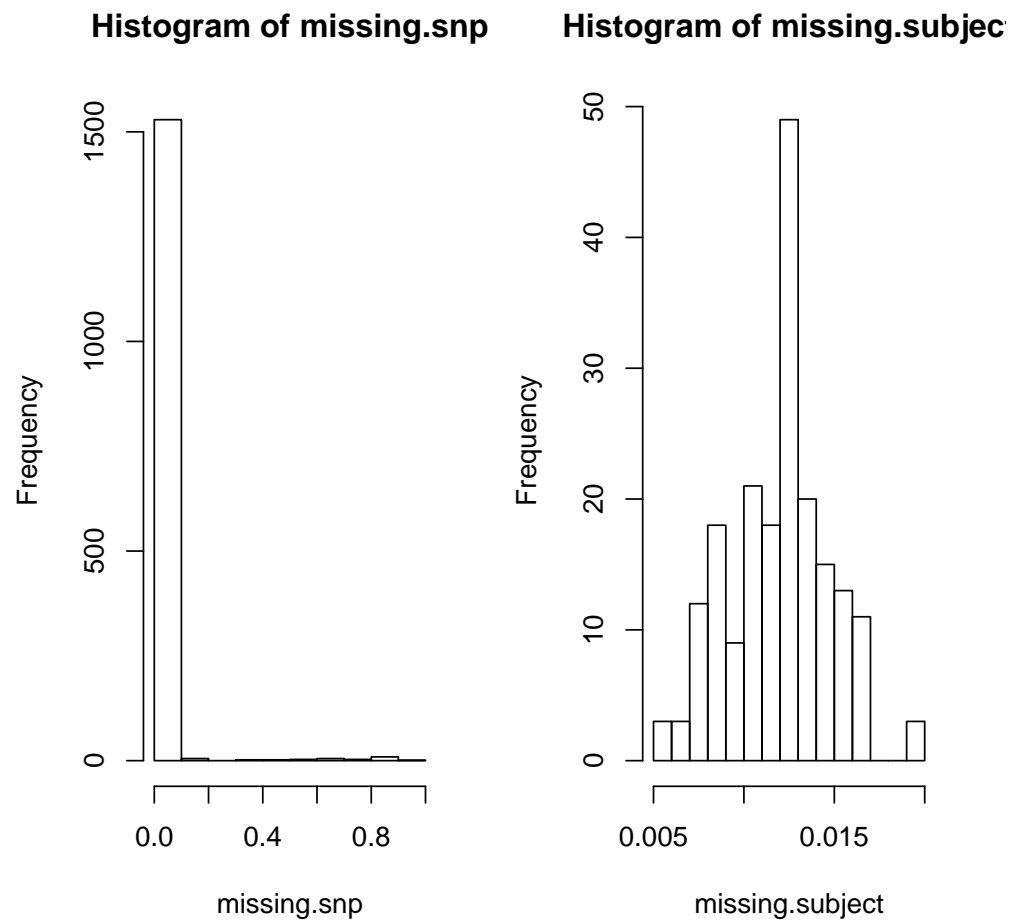
```
> geno <- getGeno(gTrio.obj, type = "holger")
> (aTDT <- allelicTDT(mat.snp = geno, size = 10000))
```

Allelic TDT

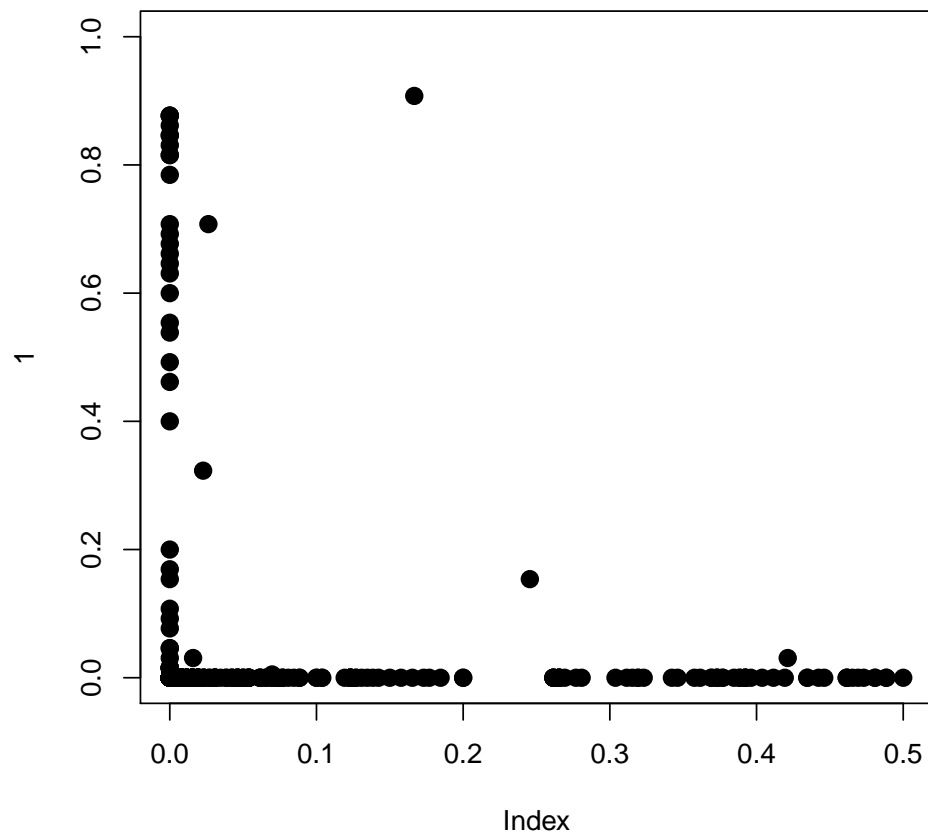
Top 5 SNPs:

	Statistic	p-value
14:54427602	7.143	0.007526
14:54429292	7.143	0.007526
14:54429785	7.143	0.007526
14:54430408	7.143	0.007526
14:54431127	7.143	0.007526

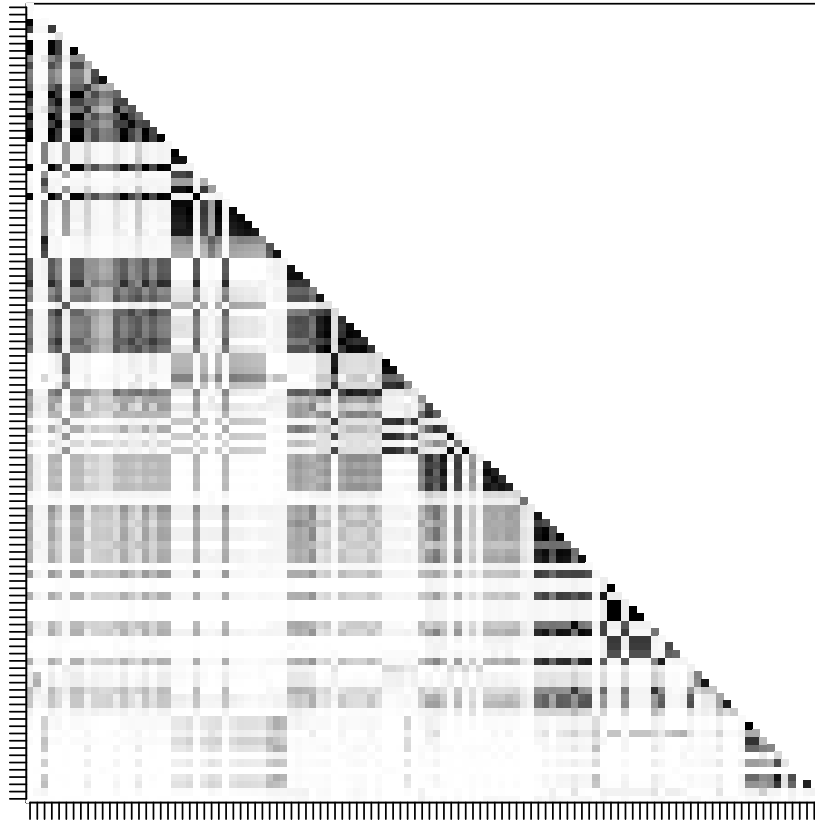
```
> layout(matrix(1:2, ncol = 2, nrow = 1))
> hist(missing.snp, breaks = 10)
> hist(missing.subject, breaks = 10)
```



```
> plot(1, type = "n", ylim = c(0, 1), xlim = c(0, 0.5))
> points(x = maf, y = missing.snp, pch = 20, cex = 2)
```



```
> ld <- getLD(x = geno, which = "both", parentsOnly = TRUE)
> plot(ld, y = "rSquare", xlab = "", ylab = "", cexAxis = 0.01)
```



```
> (gTDT <- tdt(snp = geno))
```

Genotypic TDT Based on 3 Pseudo Controls

Model Type: Additive

Coef	OR	Lower	Upper	SE	Statistic	p-Value
-0.0836	0.9198	0.8686	0.974	0.02922	8.183	0.004229

```
> hist(maf, breaks = 10)
```

