

Classes *gTrio* and *iTrio*: Derivatives of *TrioSet* for use with genotype and intensity data in package *trioClasses*

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I found it difficult to construct an extension of *gSet* that was flexible with the *AssayData*, e.g., *geno*, *lrr* and *baf*. I think it makes sense to begin with the individual class scheme, and to strip the *TrioSet* class down to bare-bones so we can build it up into something that suits our present needs. Ultimately, I think a one class scheme is best, but I think it will be challenging to implement. So, what I've done is define two classes *gTrio*, for genotype data, and *iTrio* for probe intensity data (*lrr* and *baf*). Each of these classes is the *TrioSet* class defined in MD gutted to have only the bare essentials. I would like to add *eSet/gSet*-type objects in as we go, so that I understand exactly what they do. The class *gTrio* has some methods defined for it to as demonstrated in this vignette. The class *iTrio* has no methods defined for it yet. Presumably we can simply port in methods from MD.

```
> rm(list = ls())
> library("trioClasses")
> library("trio")
```

First we load the sample pedigree data frame included in local versions of the *trioClasses* package.

```
> data(ped)
> head(ped.df)
```

	id	mid	fid	Population	PI	Ethnicity
578_01	578_01	578_03	578_02	PHILIPPINES	Murray	filipino
578_02	578_02	<NA>	<NA>	PHILIPPINES	Murray	filipino
578_03	578_03	<NA>	<NA>	PHILIPPINES	Murray	filipino
1539_01	1539_01	1539_03	1539_02	IOWA	Murray	european
1539_02	1539_02	<NA>	<NA>	IOWA	Murray	european
1539_03	1539_03	<NA>	<NA>	IOWA	Murray	european

```
> 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.

After we ensure that F, M and O exist in the data frame we create a Pedigree object. Note the terse show method for the Pedigree object.

1 *gTrio* class

Next we load the genotype matrix with well-named rows and columns, with rows for subjects and columns for SNPs.

```
> data(geno)
> head(geno.mat[, 1:6])
```

	snp1	snp2	snp3	snp4	snp5	snp6
578_01	2	2	0	2	2	1
578_02	1	0	1	0	2	2
578_03	0	2	2	0	2	2
1539_01	2	0	2	2	2	0
1539_02	1	1	2	0	1	0
1539_03	1	0	2	2	0	2

Now we format the genotype matrix for input into `gTrio()` and use the `completeTrios` method to remove trios that do not have genotype information for all members.

```
> geno.trio <- genoMat(tg.ped, geno.mat)
> (tg.ped.comp <- completeTrios(tg.ped, colnames(geno.trio)))
```

This pedigree object contains 33 complete trios.
For access to the data frame use the `trios()` accessor function.

Now we create the `gTrio` object from a complete pedigree and properly formatted, well-named, genotype matrix.

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

```
gTrio (storageMode: lockedEnvironment)
assayData: 10 features, 33 samples
  element names: geno
protocolData: none
phenoData: none
featureData
  featureNames: snp1 snp2 ... snp10 (10 total)
  fvarLabels: position chromosome isSnp
  fvarMetadata: labelDescription
experimentData: use 'experimentData(object)'
Annotation:
genome: hg19
```

```
> class(gTrio.obj)
```

```
[1] "gTrio"
attr(,"package")
[1] "trioClasses"
```

Now we use the `getGeno` method to retrieve a genotype matrix formatted, in this case, for use in Holger's trio package.

```
> geno <- getGeno(gTrio.obj, type = "holger")
> dim(geno)
```

```
[1] 99 10
```

```
> (aTDT <- allelicTDT(mat.snp = geno, size = 10000))
```

```
Allelic TDT
```

```
Top 5 SNPs:
```

	Statistic	p-value
snp10	3.8571	0.04953
snp4	2.5714	0.10881
snp5	1.1429	0.28505
snp8	1.0000	0.31731
snp6	0.6923	0.40538

2 *iTrio* class

```
> (iTrio.obj <- iTrio(tg.ped.comp, lrr = geno.trio, baf = geno.trio))
```

```
iTrio (storageMode: lockedEnvironment)
assayData: 10 features, 33 samples
  element names: baf, lrr
protocolData: none
phenoData: none
featureData
  featureNames: snp1 snp2 ... snp10 (10 total)
  fvarLabels: position chromosome isSnp
  fvarMetadata: labelDescription
experimentData: use 'experimentData(object)'
Annotation:
genome: hg19
```