

fbat

April 19, 2009

checkMarkers *Basic data quality checks for markers*

Description

Basic data quality checks for markers.

Usage

```
checkMarkers(geneSetObj, founderOnly=TRUE, thrsh=0.05, quiet=TRUE)
checkMarkers.default(pedObj, founderOnly=TRUE, thrsh=0.05, quiet=TRUE)
```

Arguments

`geneSetObj` a `geneSet` object.

`pedObj` a list with five elements: `ped`, `columns`, `markerNames`, `Position`, and `filename`. `ped` is a pedigree data frame whose first 6 columns are family (pedigree id), `pid` (patient id), father (father id), mother (mother id), sex, affected (affection status). The remaining columns are pairs of marker alleles. Each row corresponds to an individual; `columns` are the names of the first 5 (or 6) columns of `ped` file. It should be either equal to `c("family","pid","father","mother","sex","affected")` or equal to `c("family","pid","father","mother","sex")`; `founderOnly` indicates if using only founder info; `markerNames` is a vector of marker names; `Position` is a vector of marker positions; `fileName` is the pedigree file name

`founderOnly` indicates if using only founder info

`thrsh` threshold for Hardy-Weinberg test. If the pvalue of the HW test for a marker is greater than `thrsh`, then the marker is a good marker.

`quiet` print intermediate results if `quiet=FALSE`.

Value

a data frame contains components:

`Name` marker names.

`Position` marker positions.

`ObsHET` marker's observed heterozygosity (i.e., proportion of heterozygotes at markes). Missing alleles are excluded in the calculation.

PredHET	marker's predicted heterozygosity (i.e., $2 * MAF * (1 - MAF)$). Missing alleles are excluded in the calculation.
HWpval	pvalues for Hardy-Weinberg test
pGeno	percentage of non-missing genotypes for markers
MAF	minor allele frequencies. missing allele are excluded from calculation
Rating	Rating[i]=1 means that the <i>i</i> -th marker passes HW test (do not reject H0 that HW equilibrium holds). Rating[i]=0 means HW equilibrium does hold for the <i>i</i> -th marker.

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Examples

```
data(CAMP)
res<-checkMarkers(CAMP)
print(res)
```

checkMendelian *Check Mendelian Errors*

Description

Check Mendelian errors.

Usage

```
checkMendelian(geneSetObj, quiet = TRUE)

checkMendelian.default(pedObj, quiet=TRUE)
```

Arguments

geneSetObj	a geneSet object.
pedObj	a list with five elements: ped, columns, markerNames, Position, and filename. ped is a pedigree data frame whose first 6 columns are family (pedigree id), pid (patient id), father (father id), mother (mother id), sex, affected (affection status). The remaining columns are pairs of marker alleles. Each row corresponds to an individual; columns are the names of the first 5 (or 6) columns of ped file. It should be either equal to c("family","pid","father","mother","sex","affected") or equal to c("family","pid","father","mother","sex"); founderOnly indicates if using only founder info; markerNames is a vector of marker names; Position is a vector of marker positions; fileName is the pedigree file name
quiet	print intermediate results if quiet=FALSE.

Details

check the following errors:

- 1 father id = subject id
- 2 mother id = subject id
- 3 could not determine if an individual is a parent or a child in a family
- 4 inconsistent parental sex in a family
- 5 parental genotypes are not compatible with childrens' genotypes in a family
- 6 all childrens' genotypes are missing in a family
- 7 inconsistent sib genotypes in a family

Value

A list with following elements:

<code>errorFlag</code>	<code>errorFlag=1</code> indicates the occurrence of errors; <code>errorFlag=0</code> indicates no error.
<code>compatibleFlag</code>	<code>compatibleFlag=0</code> indicates the occurrence of non-compatibility; <code>compatibleFlag=1</code> indicates compatibility.
<code>nMerrMarker</code>	A $nMarkers \times 1$ vector records the numbers of families with non-compatible genotypes, where $nMarkers$ is the number of markers.
<code>nMerrFamily</code>	A $nFamily \times 1$ vector records the numbers of markers with non-compatible genotypes, where $nFamily$ is the number of families.
<code>nErrFamilySample</code>	A $nFamily \times 1$ vector records the numbers of times that father id is equal to subject id or mother id is equal to subject id in a family.

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Examples

```
data(CAMP)
checkMendelian(CAMP, quiet = TRUE)
```

fbat

*Family-Based Association Tests***Description**

Family-Based Association Tests for biallelic markers.

Usage

```
fbat (geneSetObj,
      model="a",
      traitMethod=3,
      traitOffset=0,
      quiet=TRUE)

fbat.default (pedObj,
              model="a",
              traitMethod=3,
              traitOffset=0,
              quiet=TRUE)
```

Arguments

<code>geneSetObj</code>	an object of <code>geneSet</code> .
<code>pedObj</code>	a list with five elements: <code>ped</code> , <code>columns</code> , <code>markerNames</code> , <code>Position</code> , and <code>filename</code> . <code>ped</code> is a pedigree data frame whose first 6 columns are family (pedigree id), <code>pid</code> (patient id), father (father id), mother (mother id), sex, affected (affection status). The remaining columns are pairs of marker alleles. Each row corresponds to an individual; <code>columns</code> are the names of the first 5 (or 6) columns of <code>ped</code> file. It should be either equal to <code>c("family", "pid", "father", "mother", "sex", "affected")</code> or equal to <code>c("family", "pid", "father", "mother", "sex")</code> ; <code>founderOnly</code> indicates if using only founder info; <code>markerNames</code> is a vector of marker names; <code>Position</code> is a vector of marker positions; <code>fileName</code> is the pedigree file name
<code>model</code>	Genotype coding method. <code>model="d"</code> means GDOM (dominant) coding; <code>model="r"</code> means GREC (recessive) coding; <code>model="g"</code> means GEN (genotype) coding; <code>model="a"</code> or otherwise means GTDT (additive) coding.
<code>traitMethod</code>	Trait coding method. <code>traitMethod=1</code> means $T=y-\text{offset}$, where y is the trait and <code>offset</code> is an offset. In a <code>.ped</code> file, $y=2$ if affected; $y=1$ if unaffected; and $y=0$ if unknown. <code>traitMethod=2</code> means $T=1$ if affected, $T=0$ otherwise.
<code>traitOffset</code>	Offset if <code>traitMethod=1</code> .
<code>quiet</code>	Print some intermediate results if <code>quiet=FALSE</code> .

Value

<code>statPvalue</code>	A m by 3 matrix with the 3 columns: test statistics, degree of freedom and pvalues, where m is the number of markers.
<code>S.list</code>	A list of S scores for markers.
<code>ES.list</code>	A list of expected S scores for markers.

Arguments

`pedObj` a list with five elements: `ped`, `columns`, `markerNames`, `Position`, and `filename`. `ped` is a pedigree data frame whose first 6 columns are family (pedigree id), `pid` (patient id), father (father id), mother (mother id), sex, affected (affection status). The remaining columns are pairs of marker alleles. Each row corresponds to an individual; `columns` are the names of the first 5 (or 6) columns of ped file. It should be either equal to `c("family", "pid", "father", "mother", "sex", "affected")` or equal to `c("family", "pid", "father", "mother", "sex")`; `founderOnly` indicates if using only founder info; `markerNames` is a vector of marker names; `Position` is a vector of marker positions; `fileName` is the pedigree file name

Value

An pedigree object contains only founders' information.

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References

~put references to the literature/web site here ~

<code>missGFreq</code>	<i>Count frequencies of missing genotypes</i>
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Description

Count frequencies of missing genotypes

Usage

```
missGFreq(geneSetObj, founderOnly = TRUE, quiet = FALSE)
```

```
missGFreq.default(pedObj, founderOnly=TRUE)
```

Arguments

`geneSetObj` a `geneSet` object.

`pedObj` a list with five elements: `ped`, `columns`, `markerNames`, `Position`, and `filename`. `ped` is a pedigree data frame whose first 6 columns are family (pedigree id), `pid` (patient id), father (father id), mother (mother id), sex, affected (affection status). The remaining columns are pairs of marker alleles. Each row corresponds to an individual; `columns` are the names of the first 5 (or 6) columns of ped file. It should be either equal to `c("family", "pid", "father", "mother", "sex", "affected")` or equal to `c("family", "pid", "father", "mother", "sex")`; `founderOnly` indicates if using only founder info; `markerNames` is a vector of marker names; `Position` is a vector of marker positions; `fileName` is the pedigree file name

`founderOnly` indicates if using only founder info

`quiet` print intermediate results if `quiet=FALSE`.

Value

A matrix with the following three columns:

column 1	counts of genotypes, of which both alleles are missing.
column 2	counts of genotypes, of which the first allele is missing and the second allele is not missing.
column 3	counts of genotypes, of which the first allele is not missing and the second allele is missing.

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Examples

```
data(CAMP)
res<-missGFreq(CAMP, founderOnly=FALSE)
# number of missing genotypes per marker
print(res$MissMarkers)
# number of missing genotypes per subject
print(res$MissSubjects[1:10,])
```

pedAFreq	<i>get allele frequencies</i>
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Description

Get allele frequencies (missing alleles allowed).

Usage

```
pedAFreq(geneSetObj, founderOnly=TRUE, missingOutput=FALSE, quiet=FALSE)

pedAFreq.default(pedObj,
                 founderOnly=TRUE,
                 missingOutput=FALSE,
                 quiet=FALSE)
```

Arguments

geneSetObj	a <code>geneSet</code> object.
pedObj	a list with five elements: <code>ped</code> , <code>columns</code> , <code>markerNames</code> , <code>Position</code> , and <code>filename</code> . <code>ped</code> is a pedigree data frame whose first 6 columns are family (pedigree id), <code>pid</code> (patient id), father (father id), mother (mother id), sex, affected (affection status). The remaining columns are pairs of marker alleles. Each row corresponds to an individual; <code>columns</code> are the names of the first 5 (or 6) columns of <code>ped</code> file. It should be either equal to <code>c("family", "pid", "father", "mother", "sex", "affected")</code> or equal to <code>c("family", "pid", "father", "mother", "sex")</code> ; <code>founderOnly</code> indicates if using only founder info; <code>markerNames</code> is a vector of marker names; <code>Position</code> is a vector of marker positions; <code>fileName</code> is the pedigree file name

founderOnly indicates if using only founder info.
 missingOutput indicates if missing allele frequency should be output.
 quiet print intermediate results if quiet=FALSE.

Value

aFreqMat allele frequencies.
 aPercMat allele percentages.
 missingOutput indicates if missing allele frequency should be output.

Author(s)

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Examples

```
data(CAMP)
res<-pedAFreq(CAMP)
res$aFreqMat
res$aPercMat
res$missingOutput
```

pedFlagHomo *flag homo/heterozygotes*

Description

Flag homo/heterozygotes.

Usage

```
pedFlagHomo(geneSetObj, founderOnly=TRUE, quiet=FALSE)

pedFlagHomo.default(pedObj,
                    founderOnly=TRUE,
                    quiet=FALSE)
```

Arguments

geneSetObj a geneSet object.
 pedObj a list with five elements: ped, columns, markerNames, Position, and filename. ped is a pedigree data frame whose first 6 columns are family (pedigree id), pid (patient id), father (father id), mother (mother id), sex, affected (affection status). The remaining columns are pairs of marker alleles. Each row corresponds to an individual; columns are the names of the first 5 (or 6) columns of ped file. It should be either equal to c("family","pid","father","mother","sex","affected") or equal to c("family","pid","father","mother","sex"); founderOnly indicates if using only founder info; markerNames is a vector of marker names; Position is a vector of marker positions; fileName is the pedigree file name

founderOnly indicates if using only founder info
 quiet print intermediate results if quiet=FALSE.

Value

countMat Count the number of homo/heterozygotes.
 flagHomoMat Flag homo/heterozygotes. 1 – homozygotes; 0 – heterozygotes; -1 – genotype contains one missing allele; -2 – genotype contains two missing alleles.
 markerNames marker names.

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Examples

```
data (CAMP)
res<-pedFlagHomo (CAMP)
res$countMat
res$flagHomoMat
res$markerNames
```

pedGFreq *get genotype frequencies*

Description

Get genotype frequencies (missing alleles allowed).

Usage

```
pedGFreq(geneSetObj, founderOnly=TRUE, missingOutput=FALSE, quiet=FALSE)

pedGFreq.default (pedObj,
  founderOnly=TRUE,
  missingOutput=FALSE,
  quiet=FALSE)
```

Arguments

geneSetObj a geneSet object.
 pedObj a list with five elements: ped, columns, markerNames, Position, and filename. ped is a pedigree data frame whose first 6 columns are family (pedigree id), pid (patient id), father (father id), mother (mother id), sex, affected (affection status). The remaining columns are pairs of marker alleles. Each row corresponds to an individual; columns are the names of the first 5 (or 6) columns of ped file. It should be either equal to c("family","pid","father","mother","sex","affected") or equal to c("family","pid","father","mother","sex"); founderOnly indicates if using only founder info; markerNames is a vector of marker names; Position is a vector of marker positions; fileName is the pedigree file name

founderOnly indicates if using only founder info.
 missingOutput indicates if missing allele frequency should be output.
 quiet print intermediate results if quiet=FALSE.

Value

gFreqMat genotype frequencies.
 gPercMat genotype percentages.
 missingOutput indicates if missing allele frequency should be output.

Author(s)

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Examples

```
data(CAMP)
res<-pedGFreq(CAMP)
res$gFreqMat
res$gPercMat
```

pedHardyWeinberg *Test Hardy-Weinberg equilibrium for each marker based on parental data*

Description

Test Hardy-Weinberg equilibrium for each marker based on parental data.

Usage

```
pedHardyWeinberg(geneSetObj, threshold=3, founderOnly=TRUE, quiet=FALSE)

pedHardyWeinberg.default(pedObj,
  threshold=3, founderOnly=TRUE, quiet=FALSE)
```

Arguments

geneSetObj a geneSet object.
 pedObj a list with five elements: ped, columns, markerNames, Position, and filename. ped is a pedigree data frame whose first 6 columns are family (pedigree id), pid (patient id), father (father id), mother (mother id), sex, affected (affection status). The remaining columns are pairs of marker alleles. Each row corresponds to an individual; columns are the names of the first 5 (or 6) columns of ped file. It should be either equal to c("family", "pid", "father", "mother", "sex", "affected") or equal to c("family", "pid", "father", "mother", "sex"); founderOnly indicates if using only founder info; markerNames is a vector of marker names; Position is a vector of marker positions; fileName is the pedigree file name

threshold a threshold to check if expected frequencies of genotypes are too small.
 founderOnly indicates if using only founder info
 quiet print intermediate results if quiet=FALSE.

Value

resMat A matrix records the following quantities for all markers (rows correspond to markers): nInfoInd (number of informative individuals, i.e. individuals whose genotypes contain no missing alleles for the specified marker), nGenotype (number of possible genotypes), nHET (number of heterozygous genotypes), nHOM (number of homozygous genotypes), nAllele (number of alleles), nMissing (number of missing alleles), chi2 (chi square test statistic), df (degree of freedom of the chi square test statistic under H0), p-value (pvalue of the test).

genotype A list of possible genotypes and their frequencies for all markers.
 nGenotype.vec A vector of numbers of possible genotypes for all markers.
 piVec Allele frequencies for all markers.

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Examples

```
data(CAMP)
res<-pedHardyWeinberg(CAMP)
viewHW(res, "m709")
viewHW(res, "m654")
viewHW(res, "m47")
viewHW(res, "p46")
viewHW(res, "p79")
viewHW(res, "p252")
viewHW(res, "p491")
viewHW(res, "p523")
```

readHapMap *Import HapMap data*

Description

Import HapMap data and convert it to pedigree format.

Usage

```
readHapMap(hapmapfile, race="CEU", skip = 2, comment.char = "&", quiet = FALSE)
```

Arguments

hapmapfile	the hapmap file name
race	can take values CEU, YRI, CHB, and JPT
skip	first skip lines in the file hapmapfile will be skipped.
comment.char	hapmapfile snp names contain the symbol # which is the comment command of R. So by default, we set comment.char as "&".
quiet	print intermediate results if quiet=FALSE.

Details

HapMap files are those snp files output by HapMap browsers.

Value

A list with five elements: ped, columns, markerNames, Position, and fileName. ped is a pedigree data frame whose first 6 columns are family (pedigree id), pid (patient id), father (father id), mother (mother id), sex, affected (affection status). The remaining columns are pairs of marker alleles. Each row corresponds to an individual; columns are the names of the first 5 (or 6) columns of ped file. It should be either equal to c("family","pid","father","mother","sex","affected") or equal to c("family","pid","father","mother","sex"); founderOnly indicates if using only founder info; markerNames is a vector of marker names; Position is a vector of marker positions; fileName is the pedigree file name.

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readLink

Import file with PLINK data format

Description

Import file with PLINK data format and convert it to a pedigree object.

Usage

```
readLink(pedfile, gmfile, columns = c("family", "pid", "father", "mother", "sex"))
```

Arguments

pedfile	pedigree data file with no marker info
gmfile	marker info files. It contains three columns: marker IDs, marker names, and marker positions
columns	By default, the first five columns of pedfile are sample information: family id, patient id, father id, mother id, patient sex.
quiet	print intermediate results if quiet=FALSE.

Details

The data format is used by the software PLINK.

Value

A list with five elements: `ped`, `columns`, `markerNames`, `Position`, and `fileName`. `ped` is a pedigree data frame whose first 6 columns are family (pedigree id), pid (patient id), father (father id), mother (mother id), sex, affected (affection status). The remaining columns are pairs of marker alleles. Each row corresponds to an individual; `columns` are the names of the first 5 (or 6) columns of `ped` file. It should be either equal to `c("family","pid","father","mother","sex","affected")` or equal to `c("family","pid","father","mother","sex")`; `founderOnly` indicates if using only founder info; `markerNames` is a vector of marker names; `Position` is a vector of marker positions; `fileName` is the pedigree file name.

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 readPed

Import pedigree file from standard pedigree file format

Description

Import pedigree file from standard pedigree file format.

Usage

```
readPed (
  filename,
  columns=c("family", "pid", "father", "mother",
            "sex", "affected"),
  quiet=FALSE
)
```

Arguments

<code>filename</code>	File containing genotype data
<code>columns</code>	column names for sample info
<code>quiet</code>	indicates if intermediate output should be printed

Value

A list with five elements: `ped`, `columns`, `markerNames`, `Position`, and `fileName`. `ped` is a pedigree data frame whose first 6 columns are family (pedigree id), pid (patient id), father (father id), mother (mother id), sex, affected (affection status). The remaining columns are pairs of marker alleles. Each row corresponds to an individual; `columns` are the names of the first 5 (or 6) columns of `ped` file. It should be either equal to `c("family","pid","father","mother","sex","affected")` or equal to `c("family","pid","father","mother","sex")`; `founderOnly` indicates if using only founder info; `markerNames` is a vector of marker names; `Position` is a vector of marker positions; `fileName` is the pedigree file name.

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See Also

[read.table](#), etc

sampleInfoCEU *Information about HapMap CEU subjects*

Description

Information about HapMap CEU subjects.

Usage

```
data(sampleInfoCEU)
```

Format

A data frame with 90 observations on the following 6 variables.

pedID a numeric vector

id a numeric vector

fid a numeric vector

mid a numeric vector

sex a numeric vector

sampleID a factor with levels NA06985 NA06991 NA06993 NA06994 NA07000 NA07019

```
NA07022 NA07029 NA07034 NA07048 NA07055 NA07056 NA07345 NA07348 NA07357
NA10830 NA10831 NA10835 NA10838 NA10839 NA10846 NA10847 NA10851 NA10854
NA10855 NA10856 NA10857 NA10859 NA10860 NA10861 NA10863 NA11829 NA11830
NA11831 NA11832 NA11839 NA11840 NA11881 NA11882 NA11992 NA11993 NA11994
NA11995 NA12003 NA12004 NA12005 NA12006 NA12043 NA12044 NA12056 NA12057
NA12144 NA12145 NA12146 NA12154 NA12155 NA12156 NA12234 NA12236 NA12239
NA12248 NA12249 NA12264 NA12707 NA12716 NA12717 NA12740 NA12750 NA12751
NA12752 NA12753 NA12760 NA12761 NA12762 NA12763 NA12801 NA12802 NA12812
NA12813 NA12814 NA12815 NA12864 NA12865 NA12872 NA12873 NA12874 NA12875
NA12878 NA12891 NA12892
```

Examples

```
data(sampleInfoCEU)
```

sampleInfoCHB *Information about HapMap CHB subjects*

Description

Information about HapMap CHB subjects

Usage

```
data(sampleInfoCHB)
```

Format

A data frame with 45 observations on the following 6 variables.

pedid a numeric vector

id a numeric vector

fid a numeric vector

mid a numeric vector

sex a numeric vector

sampleID a factor with levels NA18524 NA18526 NA18529 NA18532 NA18537 NA18540
NA18542 NA18545 NA18547 NA18550 NA18552 NA18555 NA18558 NA18561 NA18562
NA18563 NA18564 NA18566 NA18570 NA18571 NA18572 NA18573 NA18576 NA18577
NA18579 NA18582 NA18592 NA18593 NA18594 NA18603 NA18605 NA18608 NA18609
NA18611 NA18612 NA18620 NA18621 NA18622 NA18623 NA18624 NA18632 NA18633
NA18635 NA18636 NA18637

Examples

```
data(sampleInfoCHB)
```

sampleInfoJPT *Information about HapMap JPT subjects*

Description

Information about HapMap JPT subjects

Usage

```
data(sampleInfoJPT)
```

Format

A data frame with 45 observations on the following 6 variables.

pedid a numeric vector

id a numeric vector

fid a numeric vector

mid a numeric vector

sex a numeric vector

sampleID a factor with levels NA18940 NA18942 NA18943 NA18944 NA18945 NA18947
 NA18948 NA18949 NA18951 NA18952 NA18953 NA18956 NA18959 NA18960 NA18961
 NA18964 NA18965 NA18966 NA18967 NA18968 NA18969 NA18970 NA18971 NA18972
 NA18973 NA18974 NA18975 NA18976 NA18978 NA18980 NA18981 NA18987 NA18990
 NA18991 NA18992 NA18994 NA18995 NA18997 NA18998 NA18999 NA19000 NA19003
 NA19005 NA19007 NA19012

Examples

```
data(sampleInfoJPT)
```

```
sampleInfoYRI
```

Information about HapMap YRI subjects

Description

Information about HapMap YRI subjects

Usage

```
data(sampleInfoYRI)
```

Format

A data frame with 90 observations on the following 6 variables.

pedID a numeric vector

id a numeric vector

fid a numeric vector

mid a numeric vector

sex a numeric vector

sampleID a factor with levels NA18500 NA18501 NA18502 NA18503 NA18504 NA18505
 NA18506 NA18507 NA18508 NA18515 NA18516 NA18517 NA18521 NA18522 NA18523
 NA18852 NA18853 NA18854 NA18855 NA18856 NA18857 NA18858 NA18859 NA18860
 NA18861 NA18862 NA18863 NA18870 NA18871 NA18872 NA18912 NA18913 NA18914
 NA19092 NA19093 NA19094 NA19098 NA19099 NA19100 NA19101 NA19102 NA19103
 NA19116 NA19119 NA19120 NA19127 NA19128 NA19129 NA19130 NA19131 NA19132
 NA19137 NA19138 NA19139 NA19140 NA19141 NA19142 NA19143 NA19144 NA19145
 NA19152 NA19153 NA19154 NA19159 NA19160 NA19161 NA19171 NA19172 NA19173
 NA19192 NA19193 NA19194 NA19200 NA19201 NA19202 NA19203 NA19204 NA19205
 NA19206 NA19207 NA19208 NA19209 NA19210 NA19211 NA19221 NA19222 NA19223
 NA19238 NA19239 NA19240

Examples

```
data(sampleInfoYRI)
```

summaryPvalue	<i>summarize the test statistics and p-values</i>
---------------	---

Description

summarize the test statistics and p-values

Usage

```
summaryPvalue(fbatObject)
```

Arguments

fbatObject Object for Family Based Association Tests. See references.

Details

Print summary of test statistics and p-value.

Author(s)

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References

<http://www.biostat.harvard.edu/~fbat/fbat.htm>

Examples

```
data(CAMP)
tmp<-fbat(CAMP)
summaryPvalue(tmp)
```

viewFlagHomo	<i>flag homo/heterozygotes for specified marker</i>
--------------	---

Description

Flag homo/heterozygotes for specified marker.

Usage

```
viewFlagHomo(flagHomo.object, markerName)
```

Arguments

`flagHomo.object` object returned by the function `pedFlagHomo()`.
`markerName` name of the specified marker.

Value

`countMatM` Count the number of homo/heterozygotes for the specified marker.
`flagHomoMatM` Flag homo/heterozygotes for the specified marker. 1 – homozygotes; 0 – heterozygotes; -1 – genotype contains one missing allele; -2 – genotype contains two missing alleles.

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Examples

```
data (CAMP)
res<-pedFlagHomo (CAMP)
viewFlagHomo (res, "p79")
```

<code>viewHW</code>	<i>View allele frequencies, Hardy-Weinberg equilibrium test statistics for specified marker</i>
---------------------	---

Description

View allele frequencies, Hardy-Weinberg equilibrium test statistics for specified marker.

Usage

```
viewHW(HW.object, markerName)
```

Arguments

`HW.object` object returned by the function `'pedHardyWeinberg'`.
`markerName` a character string indicating the name of marker whose statistics are to be viewed

Value

`resM` A vector records the following quantities for the specified marker: `nInfoInd` (number of informative individuals, i.e. individuals whose genotypes contain no missing alleles for the specified marker), `nGenotype` (number of possible genotypes), `nHET` (number of heterozygous genotypes), `nHOM` (number of homozygous genotypes), `nAllele` (number of alleles), `nMissing` (number of missing alleles), `chi2` (chi square test statistic), `df` (degree of freedom of the chi square test statistic under H0), `p-value` (pvalue of the test).
`nGenotypeM` number of possible genotypes for the specified marker.
`genotypeM` possible genotypes and their frequencies.
`piVecM` allele frequencies.

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Examples

```
data(CAMP)
res<-pedHardyWeinberg(CAMP)
viewHW(res, "m709")
viewHW(res, "m654")
viewHW(res, "m47")
viewHW(res, "p46")
viewHW(res, "p79")
viewHW(res, "p252")
viewHW(res, "p491")
viewHW(res, "p523")
```

viewstat

view statistics for a marker

Description

view statistics for a marker

Usage

```
viewstat(fbatObject, markerName)
```

Arguments

fbatObject Object for Family Based Association Tests. See references.
markerName name(s) of the marker(s) for which statistics is needed

Details

Print various stats for a marker, such as: family size, number of people in the family, number of informative families in the marker, the alleles of marker, scores for marker, expected score for marker, covariance matrix of the score for the marker, Moore-Penrose generalized inverse of covariance matrix and P-value.

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References

<http://www.biostat.harvard.edu/~fbat/fbat.htm>

Examples

```
data(CAMP)
res<-fbat(CAMP)
viewstat(res, "m709")
viewstat(res, "m654")
viewstat(res, "m47")
viewstat(res, "p46")
viewstat(res, "p79")
viewstat(res, "p252")
viewstat(res, "p491")
viewstat(res, "p523")
```

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