

# Package ‘EFGLmh’

January 28, 2022

**Type** Package

**Title** Functions For Working With Microhaps for EFGL

**Version** 0.1.0

**Description** Written to work with microhaps and SNPs (which are just short microhaps).  
More generally, will function with codominant, diploid genotypes.  
Uses ``Progeny-style" (and FishGen-style) inputs.  
Performs basic manipulations, data summaries, and exporting data in  
formats for other packages/programs.

**Imports** tibble,  
dplyr,  
readr,  
tidyr

**License** MIT + file LICENSE

**Encoding** UTF-8

**LazyData** true

**RoxygenNote** 7.1.2

**Depends** R (>= 2.10)

**Suggests** knitr,  
rmarkdown,  
tidyverse,  
rubias

**VignetteBuilder** knitr

## R topics documented:

aRich . . . . .	2
calcAF . . . . .	3
calcHet . . . . .	3
cleanGrandma . . . . .	4
combineEFGLdata . . . . .	4
construct_EFGLdata . . . . .	5
createF1Hybrids . . . . .	5
dumpTable . . . . .	6
exampleData . . . . .	6
exportCKMRsimAF . . . . .	7
exportCKMRsimLG . . . . .	7

exportGenAIEx . . . . .	8
exportGenePop . . . . .	8
exportGrandma . . . . .	9
exportHierFstat . . . . .	10
exportPlink . . . . .	10
exportProgenyStyle . . . . .	11
exportRubias_baseline . . . . .	12
exportRubias_mixture . . . . .	12
exportSNPPIT . . . . .	13
exportStructure . . . . .	14
genoSuccess . . . . .	15
getInds . . . . .	15
getLoci . . . . .	16
getMeta . . . . .	16
getPops . . . . .	16
lociSuccess . . . . .	17
moveInds . . . . .	17
movePops . . . . .	18
numInds . . . . .	18
print.EFGLdata . . . . .	19
readInData . . . . .	19
removeInds . . . . .	20
removeLoci . . . . .	21
removePops . . . . .	21
whichLower . . . . .	22

<b>Index</b>	<b>23</b>
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aRich	<i>calculate allelic richness of loci</i>
-------	---

---

## Description

calculate allelic richness of loci

## Usage

aRich(x)

## Arguments

x                      an EFGLdata object

## Value

a tibble giving the allelic richness of each locus

---

calcAF	<i>calculate allele frequencies within populations Loci that are all missing genotypes within a pop will be (silently) excluded from the output.</i>
--------	--

---

**Description**

calculate allele frequencies within populations Loci that are all missing genotypes within a pop will be (silently) excluded from the output.

**Usage**

```
calcAF(x)
```

**Arguments**

x                      an EFGLdata object

**Value**

a tibble

---

calcHet	<i>calculate expected and observed heterozygosity within populations</i>
---------	--

---

**Description**

calculate expected and observed heterozygosity within populations

**Usage**

```
calcHet(x)
```

**Arguments**

x                      an EFGLdata object

**Value**

a tibble

---

cleanGrandma	<i>convenience function to remove loci with all fails or no variation from gRandma input</i>
--------------	--

---

### Description

convenience function to remove loci with all fails or no variation from gRandma input

### Usage

```
cleanGrandma(baseline, mixture = NULL)
```

### Arguments

baseline	a gRandma baseline input
mixture	a gRandma mixture input

### Value

a list with two components, one is the baseline, one is the mixture

---

combineEFGLdata	<i>combine multiple EFGL objects into one</i>
-----------------	---

---

### Description

combine multiple EFGL objects into one

### Usage

```
combineEFGLdata(
  ...,
  genoComb = c("intersect", "union"),
  metaComb = c("intersect", "union")
)
```

### Arguments

...	multiple EFGLdata objects separated by commas
genoComb	if the objects have different loci, whether to create a new object with the intersection or union of loci. If union, genotypes for missing loci are all NA.
metaComb	if the objects have different metadata fields, whether to create a new object with the intersection or union of the fields. If union, missing fields are all NA.

---

construct_EFGLdata	<i>some basic checks on EFGLdata objects</i>
--------------------	--

---

**Description**

some basic checks on EFGLdata objects

**Usage**

```
construct_EFGLdata(x)
```

**Arguments**

x	an EFGLdata object
---	--------------------

---

createF1Hybrids	<i>Create F1 hybrids from two populations</i>
-----------------	---

---

**Description**

Creates hybrids by sampling one allele from one pop and the other allele from the other pop based on observed allele frequencies. Missing genotypes are sampled based on the combined rate of missing genotypes or data can be simulated with no missing genotypes. Alleles at different loci are sampled independently. Note that if a sex marker exists in the data set, it is treated like any other marker, resulting in XX, XY, and YY genotypes in the hybrids. You may want to remove it prior to simulation using the 'removeLoci' function. Loci with all missing genotypes in one population are simulated to have all missing genotypes in the hybrids REGARDLESS of the input value for 'missingGenos'.

**Usage**

```
createF1Hybrids(x, pop1, pop2, newName, n = 50, missingGenos = TRUE)
```

**Arguments**

x	an EFGLdata object
pop1	one parent population (must be in 'x')
pop2	the other parent population (must be in 'x')
newName	a string giving the name of the population to put hybrids into. This MUST be a new pop.
n	the number of hybrids to simulate
missingGenos	TRUE to simulate missing genotypes, FALSE to not

**Value**

an EFGLdata object with all the populations in 'x', plus the new simulated hybrid population

---

dumpTable	<i>wrapper for write table with commonly used options - carried over from IDFGEN</i>
-----------	--

---

**Description**

wrapper for write table with commonly used options - carried over from IDFGEN

**Usage**

```
dumpTable(x, filename, row.names = FALSE, sep = "\t")
```

**Arguments**

x	object to write out
filename	filename to write out as
row.names	passed to write.table
sep	passed to write.table

**Value**

nothing, just writes a file

---

exampleData	<i>An example input dataset used in the vignette</i>
-------------	--

---

**Description**

An example input dataset used in the vignette

**Usage**

```
exampleData
```

**Format**

a tibble

---

exportCKMRsimAF	<i>return a CKMRsim allele frequency input tibble.</i>
-----------------	--

---

**Description**

No chromosome and position information is used. All populations specified are combined into one allele frequency output. Any loci with only missing genotypes are omitted from the output. The output should then be run through the CKMRsim function `reindex_markers()`.

**Usage**

```
exportCKMRsimAF(x, pops = NULL, loci = NULL)
```

**Arguments**

x	an EFGLdata object
pops	a vector of pops to include. If not specified, all pops are used.
loci	a vector of loci to include. If not specified, all loci are used.

**Value**

a tibble

---

exportCKMRsimLG	<i>Return a long genotype tibble for input to CKMRsim for relationship inference.</i>
-----------------	---

---

**Description**

Return a long genotype tibble for input to CKMRsim for relationship inference.

**Usage**

```
exportCKMRsimLG(x, pops = NULL, loci = NULL)
```

**Arguments**

x	an EFGLdata object
pops	a vector of pops to include. If not specified, all pops are used.
loci	a vector of loci to include. If not specified, all loci are used.

**Value**

a tibble

---

exportGenAlEx	<i>write a GenAlEx input file</i>
---------------	-----------------------------------

---

**Description**

write a GenAlEx input file

**Usage**

```
exportGenAlEx(
  x,
  filename,
  pops = NULL,
  loci = NULL,
  title = "",
  useNames = TRUE
)
```

**Arguments**

x	an EFGLdata object
filename	the name of the file to write
pops	a vector of pops to include. If not specified, all pops are used.
loci	a vector of loci to include. If not specified, all loci are used.
title	a string to use as the "title" row
useNames	TRUE to use samples names, FALSE to replace with unique numerical identifiers

**Value**

nothing, just writes a file

---

exportGenePop	<i>write a genepop input file</i>
---------------	-----------------------------------

---

**Description**

write a genepop input file

**Usage**

```
exportGenePop(
  x,
  filename,
  header = "genePop file",
  pops = NULL,
  loci = NULL,
  useIndNames = FALSE
)
```



**Arguments**

x	an EFGLdata object
filename	the name of the file to write
header	a string to use as the header line of the genepop file
pops	a vector of pops to include. If not specified, all pops are used.
loci	a vector of loci to include. If not specified, all loci are used.
useIndNames	TRUE to use individual names as sample identifiers. Otherwise, population names are used

**Value**

nothing, just writes a file

---

exportGrandma	<i>export a gRandma baseline or mixture</i>
---------------	---

---

**Description**

export a gRandma baseline or mixture

**Usage**

```
exportGrandma(x, pops = NULL, loci = NULL, baseline = TRUE)
```

**Arguments**

x	an EFGLdata object
pops	a vector of pops to include in the baseline. If not specified, all pops are used.
loci	a vector of loci to use. If not specified, all loci are used.
baseline	TRUE to make a baseline input, FALSE to make a mixture input.

**Value**

a tibble

---

exportHierFstat	<i>export a hierfstat input dataframe</i>
-----------------	---

---

### Description

export a hierfstat input dataframe

### Usage

```
exportHierFstat(x, pops = NULL, loci = NULL)
```

### Arguments

x	an EFGldata object
pops	a vector of pops to include. If not specified, all pops are used.
loci	a vector of loci to use. If not specified, all loci are used.

### Value

a dataframe coded to be used as input for hierfstat

---

exportPlink	<i>export a Plink PED file and optionally a (not very useful except as a template) MAP file</i>
-------------	---

---

### Description

Only biallelic markers are used, exports as two column per call. This format can be used as the input for "Admixture".

### Usage

```
exportPlink(
  x,
  filename,
  pops = NULL,
  loci = NULL,
  FID = "Ind",
  IID = "Ind",
  pa = NULL,
  ma = NULL,
  sex = NULL,
  pheno = NULL,
  map = NULL
)
```

**Arguments**

x	an EFGldata object
filename	the name of the file to write
pops	a vector of pops to include. If not specified, all pops are used.
loci	a vector of loci to use. If not specified, all loci are used.
FID	Metadata column name to use as the family ID
IID	Metadata column name to use as the within-family ID
pa	Metadata column name to use as the within-family ID of the father. If NULL, all are listed as unknown (0).
ma	Metadata column name to use as the within-family ID of the mother. If NULL, all are listed as unknown (0).
sex	Metadata column to use as the sex. This should be coded as "M" for male, "F" for female, and any other values are treated as Unknown. If NULL, all are listed as unknown (0).
pheno	Metadata column to use as the phenotype. This should be coded as "1" for control, "2" for case, and "-9" or "0" for Unknown. If NULL, all are listed as unknown (0).
map	filename to write a MAP file. If NULL, no MAP file is written. This just writes a dummy MAP file with the loci names in order and all given the same chromosome code and positions of "0". If you need a valid MAP file, you will need to edit this.

**Value**

nothing, just writes a file

---

exportProgenyStyle	<i>export a "Progeny-style" export file for later reading into EFGlmh</i>
--------------------	---

---

**Description**

Columns in order are Pop, Ind, metadata, genotypes (2-column per call) Missing genotypes are "0" for SNPs (biallelic or nonvariable with alleles represented by 1 character) and "000" for others. If a locus is all missing, it is treated as a SNP. Pop column is called "Pedigree" and Ind column is called "Individual.Name".

**Usage**

```
exportProgenyStyle(x, filename, pops = NULL, loci = NULL, metadata = NULL)
```

**Arguments**

x	an EFGldata object
filename	the name of the file to write
pops	a vector of pops to include. If not specified, all pops are used.
loci	a vector of loci to use. If not specified, all loci are used.
metadata	a vector of metadata fields to include. If not specified, all fields are used.

**Value**

nothing, just writes a file

---

exportRubias\_baseline *export a rubias baseline*

---

**Description**

export a rubias baseline

**Usage**

```
exportRubias_baseline(
  x,
  pops = NULL,
  repunit = NULL,
  collection = NULL,
  loci = NULL
)
```

**Arguments**

x	an EFGLdata object
pops	a vector of pops to include in the baseline. If not specified, all pops are used.
repunit	the column name of the metadata variable designating repunit. This can be Pop to use the population name. If not specified, NA is used for all samples.
collection	the column name of the metadata variable designating collection. This can be Pop to use the population name. If not specified, NA is used for all samples.
loci	a vector of loci to use. If not specified, all loci are used.

**Value**

a tibble

---

exportRubias\_mixture *export a rubias mixture*

---

**Description**

export a rubias mixture

**Usage**

```
exportRubias_mixture(x, pops = NULL, collection = NULL, loci = NULL)
```

**Arguments**

x	an EFGLdata object
pops	a vector of pops to include in the baseline. If not specified, all pops are used.
collection	the column name of the metadata variable designating collection. This can be Pop to use the population name. If not specified, NA is used for all samples. For mixtures, this variable indicates what samples come from the same "stratum" - to be analyzed together.
loci	a vector of loci to use. If not specified, all loci are used.

**Value**

a tibble

---

exportSNPPIT	<i>write a SNPPIT input file. Will warn about skipping loci with &gt; 2 alleles.</i>
--------------	--

---

**Description**

write a SNPPIT input file. Will warn about skipping loci with > 2 alleles.

**Usage**

```
exportSNPPIT(
  x,
  filename,
  baseline,
  mixture,
  loci = NULL,
  errorRate = 0.005,
  POPCOLUMN_SEX = NULL,
  POPCOLUMN_REPRO_YEARS = NULL,
  POPCOLUMN_SPAWN_GROUP = NULL,
  OFFSPRINGCOLUMN_BORN_YEAR = NULL,
  OFFSRPINGCOLUMN_SAMPLE_YEAR = NULL,
  OFFSPRINGCOLUMN_AGE_AT_SAMPLING = NULL
)
```

**Arguments**

x	an EFGLdata object
filename	the name of the file to write
baseline	a vector of pops to use as the baseline (potential parents).
mixture	a vector of pops to use as the mixture (potential offspring).
loci	a vector of loci to include. If not specified, all loci are used.
errorRate	per allele error rate for all loci
POPCOLUMN_SEX	metadata column with sex info (coded as M, F, and ?)

POPCOLUMN\_REPRO\_YEARS  
                             metadata column with repro years

POPCOLUMN\_SPAWN\_GROUP  
                             metadata column with spawn group

OFFSPRINGCOLUMN\_BORN\_YEAR  
                             metadata column with birth year

OFFSRPINGCOLUMN\_SAMPLE\_YEAR  
                             metadata column with sample year

OFFSPRINGCOLUMN\_AGE\_AT\_SAMPLING  
                             metadata column with age at sampling

### Value

nothing, just writes a file

---

exportStructure	<i>write a Structure input file.</i>
-----------------	--------------------------------------

---

### Description

Columns are Ind, Pop (converted to integers), then genotypes. Missing alleles are coded as -9

### Usage

```
exportStructure(x, filename, pops = NULL, loci = NULL)
```

### Arguments

x	an EFGldata object
filename	the name of the file to write
pops	a vector of pops to include. If not specified, all pops are used.
loci	a vector of loci to include. If not specified, all loci are used.

### Value

nothing, just writes a file

---

genoSuccess	<i>calculate genotyping success of individuals (uses only allele 1 for each genotype - assumes if allele 1 is (is not) NA, so is (is not) allele 2)</i>
-------------	---

---

**Description**

calculate genotyping success of individuals (uses only allele 1 for each genotype - assumes if allele 1 is (is not) NA, so is (is not) allele 2)

**Usage**

```
genoSuccess(x, loci = NULL)
```

**Arguments**

x	an EFGldata object
loci	a vector of loci to include. If not specified, all loci are used.

**Value**

a tibble giving the genotyping success of each individual as a proportion and number of missing genotypes

---

getInds	<i>get a vector of individuals present</i>
---------	--

---

**Description**

get a vector of individuals present

**Usage**

```
getInds(x, pops = NULL)
```

**Arguments**

x	an EFGldata object
pops	a vector of pops that you want individual names for. If not specified, names for all pops are returned

**Value**

a vector of the Individual names present

---

getLoci	<i>get a vector of loci names present</i>
---------	---

---

**Description**

get a vector of loci names present

**Usage**

```
getLoci(x)
```

**Arguments**

x	an EFGLdata object
---	--------------------

---

getMeta	<i>get a vector of metadata column names present</i>
---------	--

---

**Description**

get a vector of metadata column names present

**Usage**

```
getMeta(x)
```

**Arguments**

x	an EFGLdata object
---	--------------------

---

getPops	<i>get a vector of populations (pedigrees) present</i>
---------	--

---

**Description**

get a vector of populations (pedigrees) present

**Usage**

```
getPops(x)
```

**Arguments**

x	an EFGLdata object
---	--------------------

**Value**

a vector of the unique population names present



---

lociSuccess	<i>calculate genotyping success of loci (uses only allele 1 for each genotype - assumes if allele 1 is (is not) NA, so is (is not) allele 2)</i>
-------------	--

---

**Description**

calculate genotyping success of loci (uses only allele 1 for each genotype - assumes if allele 1 is (is not) NA, so is (is not) allele 2)

**Usage**

```
lociSuccess(x)
```

**Arguments**

x	an EFGLdata object
---	--------------------

**Value**

a tibble giving the genotyping success of each locus as a proportion

---

moveInds	<i>combine individuals into one population AND REMOVE the previous entry for those individuals</i>
----------	--

---

**Description**

combine individuals into one population AND REMOVE the previous entry for those individuals

**Usage**

```
moveInds(x, inds, newName)
```

**Arguments**

x	an EFGLdata object
inds	a vector of individuals to put in the new pop
newName	a string giving the name of population to add the individuals too. This can be a new pop or an existing pop (a warning is issued if existing).

**Value**

an EFGLdata object

---

movePops	<i>combine populations into one AND REMOVE the old populations</i>
----------	--

---

**Description**

combine populations into one AND REMOVE the old populations

**Usage**

```
movePops(x, pops, newName)
```

**Arguments**

x	an EFGLdata object
pops	a vector of populations to combine
newName	a string giving the name of the population to combine pops into. This can be a new pop or an existing pop (a warning is issued if existing).

**Value**

an EFGLdata object

---

numInds	<i>get the number of individuals present in each pop</i>
---------	--

---

**Description**

get the number of individuals present in each pop

**Usage**

```
numInds(x, pops = NULL)
```

**Arguments**

x	an EFGLdata object
pops	a vector of pops that you want individual names for. If not specified, numbers for all pops are returned

**Value**

a named vector with the number of individuals in each pop

---

print.EFGLdata	<i>print method for EFGLdata</i>
----------------	----------------------------------

---

**Description**

print method for EFGLdata

**Usage**

```
## S3 method for class 'EFGLdata'
print(x, ...)
```

**Arguments**

x	an EFGLdata object
...	ignored

---

readInData	<i>read in data from a Progeny-style output file or matrix/dataframe/tibble and create an EFGLdata object</i>
------------	---

---

**Description**

read in data from a Progeny-style output file or matrix/dataframe/tibble and create an EFGLdata object

**Usage**

```
readInData(
  input,
  genotypeStart = NULL,
  pedigreeColumn = 1,
  nameColumn = 2,
  convertNames = TRUE,
  convertMetaDataNames = TRUE,
  missingAlleles = c("0", "00", "000"),
  guess_max = 10000
)
```

**Arguments**

input	Either a character string to the tab-separated input file with a header row or a matrix/dataframe/tibble. Structure of the input: one row per individual. One column giving pedigree (population) names, one column giving individual names, optional additional metadata columns, genotype columns. Pedigree and individual name columns can be anywhere, if specified. Genotype columns <b>MUST</b> be consecutive and be the right most columns. Genotypes are given as two columns per call (diploidy assumed).
-------	---

genotypeStart	The column number that genotypes start at. If not specified, the first column with a column name ending in ".A1", ".a1", "-A1", or "-a1" is chosen.
pedigreeColumn	The column number that contains pedigree (population) names.
nameColumn	The column number that contains individual names. These MUST be unique.
convertNames	TRUE to convert genotype and pedigree names in the same way that IDF-GEN does (remove special characters from both and remove "." from genotype names).
convertMetaDataNames	TRUE to remove special characters and spaces from metadata column names. This makes accessing them easier.
missingAlleles	a vector of values (not NA) to treat as missing alleles. They will be converted to NA.
guess_max	If input is a character string, this is the maximum number of lines to use when guessing input data types. Making this smaller results in quicker loading, making it larger can fix some parsing errors

### Value

An EFGLdata object, which is just a list with two elements. The first element is a tibble with genotype data, the second is a tibble with metadata

---

removeInds	<i>remove individuals from an EFGLdata object</i>
------------	---

---

### Description

remove individuals from an EFGLdata object

### Usage

```
removeInds(x, inds)
```

### Arguments

x	an EFGLdata object
inds	a vector of individuals to remove

### Value

an EFGLdata object

---

removeLoci	<i>remove loci from an EFGLdata object</i>
------------	--

---

**Description**

remove loci from an EFGLdata object

**Usage**

```
removeLoci(x, lociRemove)
```

**Arguments**

x	an EFGLdata object
lociRemove	a vector of loci names to remove

---

removePops	<i>remove pops from an EFGLdata object</i>
------------	--

---

**Description**

remove pops from an EFGLdata object

**Usage**

```
removePops(x, pops)
```

**Arguments**

x	an EFGLdata object
pops	a vector of pops to remove

**Value**

an EFGLdata object

---

whichLower	<i>Identify which individual out of duplicate pairs has the lower genotyping success</i>
------------	--

---

**Description**

Identify which individual out of duplicate pairs has the lower genotyping success

**Usage**

```
whichLower(dupTable, geno_success)
```

**Arguments**

dupTable	the output of close_matching_samples (from rubias)
geno_success	the output of genoSuccess

**Value**

a vector of unique individual names representing the individuals with lower genotyping success from each pair

# Index

\* **datasets**  
    exampleData, [6](#)

aRich, [2](#)

calcAF, [3](#)  
calcHet, [3](#)  
cleanGrandma, [4](#)  
combineEFGLdata, [4](#)  
construct\_EFGLdata, [5](#)  
createF1Hybrids, [5](#)

dumpTable, [6](#)

exampleData, [6](#)  
exportCKMRsimAF, [7](#)  
exportCKMRsimLG, [7](#)  
exportGenAlEx, [8](#)  
exportGenePop, [8](#)  
exportGrandma, [9](#)  
exportHierFstat, [10](#)  
exportPlink, [10](#)  
exportProgenyStyle, [11](#)  
exportRubias\_baseline, [12](#)  
exportRubias\_mixture, [12](#)  
exportSNPPIT, [13](#)  
exportStructure, [14](#)

genoSuccess, [15](#)  
getInds, [15](#)  
getLoci, [16](#)  
getMeta, [16](#)  
getPops, [16](#)

lociSuccess, [17](#)

moveInds, [17](#)  
movePops, [18](#)

numInds, [18](#)

print.EFGLdata, [19](#)

readInData, [19](#)  
removeInds, [20](#)  
removeLoci, [21](#)  
removePops, [21](#)  
whichLower, [22](#)