

Package ‘AlphaSimR’

May 19, 2020

Type Package

Title Breeding Program Simulations

Version 0.12.2

Date 2020-05-19

Description The successor to the 'AlphaSim' software for breeding program simulation [Faux et al. (2016) <doi:10.3835/plantgenome2016.02.0013>]. Used for stochastic simulations of breeding programs to the level of DNA sequence for every individual. Contained is a wide range of functions for modeling common tasks in a breeding program, such as selection and crossing. These functions allow for constructing simulations of highly complex plant and animal breeding programs via scripting in the R software environment. Such simulations can be used to evaluate overall breeding program performance and conduct research into breeding program design, such as implementation of genomic selection. Included is the 'Markovian Coalescent Simulator' ('MaCS') for fast simulation of biallelic sequences according to a population demographic history [Chen et al. (2009) <doi:10.1101/gr.083634.108>].

License MIT + file LICENSE

URL <https://alphagenes.roslin.ed.ac.uk/wp/software-2/alphasimr/>,
<https://bitbucket.org/hickeyjohnnteam/alphasimr>

Encoding UTF-8

LazyData true

Depends R (>= 3.3.0), methods, R6

Imports Rcpp (>= 0.12.7)

LinkingTo Rcpp, RcppArmadillo (>= 0.7.500.0.0), BH

RoxygenNote 7.1.0

Suggests knitr, rmarkdown, testthat

VignetteBuilder knitr

NeedsCompilation yes

Author Chris Gaynor [aut, cre] (<<https://orcid.org/0000-0003-0558-6656>>),
Gregor Gorjanc [aut] (<<https://orcid.org/0000-0001-8008-2787>>),

David Wilson [aut],
 John Hickey [aut] (<<https://orcid.org/0000-0001-5675-3974>>),
 Daniel Money [ctb] (<<https://orcid.org/0000-0001-5151-3648>>)

Maintainer Chris Gaynor <gaynor.robert@hotmail.com>

Repository CRAN

Date/Publication 2020-05-19 09:30:02 UTC

R topics documented:

aa	4
AlphaSimR	5
bv	5
calcGCA	6
cChr	6
dd	7
doubleGenome	8
ebv	8
editGenome	9
editGenomeTopQtl	10
fastRRBLUP	11
genicVarA	12
genicVarAA	13
genicVarD	14
genicVarG	14
genParam	15
getQtlMap	16
getSnpMap	17
gv	18
hybridCross	19
HybridPop-class	20
LociMap-class	21
makeCross	21
makeCross2	22
makeDH	23
MapPop-class	24
meanG	24
meanP	25
mergeGenome	26
mergePops	27
mutate	27
newMapPop	28
newPop	29
nInd	30
pedigreeCross	31
pheno	32
Pop-class	33
popVar	34

pullIbdHaplo	35
pullQtlGeno	36
pullQtlHaplo	37
pullSegSiteGeno	38
pullSegSiteHaplo	39
pullSnpGeno	40
pullSnpHaplo	41
quickHaplo	42
randCross	42
randCross2	44
RawPop-class	45
reduceGenome	46
resetPop	47
RRBLUP	48
RRBLUP2	49
RRBLUPMemUse	51
RRBLUP_D	52
RRBLUP_D2	53
RRBLUP_GCA	55
RRBLUP_GCA2	56
RRBLUP_SCA	58
RRBLUP_SCA2	59
RRsol-class	61
runMacs	61
runMacs2	62
sampleHaplo	64
selectCross	65
selectFam	66
selectInd	68
selectOP	69
selectWithinFam	70
self	72
selIndex	73
selInt	74
setEBV	74
setPheno	75
setPhenoGCA	77
SimParam	78
smithHazel	99
TraitA-class	99
TraitA2-class	100
TraitA2D-class	100
TraitAD-class	100
TraitADE-class	100
TraitADEG-class	101
TraitADG-class	101
TraitAE-class	101
TraitAEG-class	102

TraitAG-class	102
usefulness	102
varA	103
varAA	104
varD	105
varG	105
varP	106
writePlink	107
writeRecords	108

Index	109
--------------	------------

aa	<i>Additive-by-additive epistatic deviations</i>
----	--------------------------------------------------

Description

Returns additive-by-additive epistatic deviations for all traits

Usage

```
aa(pop, simParam = NULL)
```

Arguments

pop	an object of Pop-class
simParam	an object of SimParam

Examples

```
#Create founder haplotypes
founderPop = quickHaplo(nInd=10, nChr=1, segSites=10)

#Set simulation parameters
SP = SimParam$new(founderPop)
SP$addTraitAD(10, meanDD=0.5)
SP$setVarE(h2=0.5)

#Create population
pop = newPop(founderPop, simParam=SP)
aa(pop, simParam=SP)
```

Description

The successor to the 'AlphaSim' software for breeding program simulation [Faux et al. (2016) <doi:10.3835/plantgenome2016.02.0013>]. Used for stochastic simulations of breeding programs to the level of DNA sequence for every individual. Contained is a wide range of functions for modeling common tasks in a breeding program, such as selection and crossing. These functions allow for constructing simulations of highly complex plant and animal breeding programs via scripting in the R software environment. Such simulations can be used to evaluate overall breeding program performance and conduct research into breeding program design, such as implementation of genomic selection. Included is the 'Markovian Coalescent Simulator' ('MaCS') for fast simulation of biallelic sequences according to a population demographic history [Chen et al. (2009) <doi:10.1101/gr.083634.108>].

Please see the introductory vignette for instructions for using this package. The vignette can be viewed using the following command: `vignette("intro", package="AlphaSimR")`

bv

Breeding value

Description

Returns breeding values for all traits

Usage

```
bv(pop, simParam = NULL)
```

Arguments

pop	an object of Pop-class
simParam	an object of SimParam

Examples

```
#Create founder haplotypes
founderPop = quickHaplo(nInd=10, nChr=1, segSites=10)

#Set simulation parameters
SP = SimParam$new(founderPop)
SP$addTraitAD(10, meanDD=0.5)
SP$setVarE(h2=0.5)

#Create population
pop = newPop(founderPop, simParam=SP)
bv(pop, simParam=SP)
```

 calcGCA

Calculate GCA

Description

Calculate general combining ability of test crosses. Intended for output from hybridCross using the "testcross" option, but will work for any population.

Usage

```
calcGCA(pop, use = "pheno")
```

Arguments

pop	an object of Pop-class or HybridPop-class
use	tabulate either genetic values "gv", estimated breeding values "ebv", or phenotypes "pheno"

Examples

```
#Create founder haplotypes
founderPop = quickHaplo(nInd=10, nChr=1, segSites=10, inbred=TRUE)

#Set simulation parameters
SP = SimParam$new(founderPop)
SP$addTraitA(10)

#Create population
pop = newPop(founderPop, simParam=SP)

#Make crosses for full diallele
pop2 = hybridCross(pop, pop, simParam=SP)
GCA = calcGCA(pop2, use="gv")
```

 cChr

Combine MapPop chromosomes

Description

Merges the chromosomes of multiple [MapPop-class](#) objects. Each MapPop must have the same number of chromosomes

Usage

```
cChr(...)
```

Arguments

... [MapPop-class](#) objects to be combined

Value

Returns an object of [MapPop-class](#)

Examples

```
pop1 = quickHaplo(nInd=10, nChr=1, segSites=10)
pop2 = quickHaplo(nInd=10, nChr=1, segSites=10)

combinedPop = cChr(pop1, pop2)
```

dd

Dominance deviations

Description

Returns dominance deviations for all traits

Usage

```
dd(pop, simParam = NULL)
```

Arguments

pop an object of [Pop-class](#)
simParam an object of [SimParam](#)

Examples

```
#Create founder haplotypes
founderPop = quickHaplo(nInd=10, nChr=1, segSites=10)

#Set simulation parameters
SP = SimParam$new(founderPop)
SP$addTraitAD(10, meanDD=0.5)
SP$setVarE(h2=0.5)

#Create population
pop = newPop(founderPop, simParam=SP)
dd(pop, simParam=SP)
```

doubleGenome	<i>Double the ploidy of individuals</i>
--------------	-----------------------------------------

Description

Creates new individuals with twice the ploidy. This function was created to model the formation of tetraploid potatoes from diploid potatoes. This function will work on any population.

Usage

```
doubleGenome(pop, keepParents = TRUE, simParam = NULL)
```

Arguments

pop	an object of 'Pop' superclass
keepParents	should previous parents be used for mother and father.
simParam	an object of 'SimParam' class

Value

Returns an object of [Pop-class](#)

Examples

```
#Create founder haplotypes
founderPop = quickHaplo(nInd=2, nChr=1, segSites=10)

#Set simulation parameters
SP = SimParam$new(founderPop)

#Create population
pop = newPop(founderPop, simParam=SP)

#Create individuals with doubled ploidy
pop2 = doubleGenome(pop, simParam=SP)
```

ebv	<i>Estimated breeding value</i>
-----	---------------------------------

Description

A wrapper for accessing the ebv slot

Usage

```
ebv(pop)
```


Arguments

pop a [Pop-class](#) or similar object

Examples

```
#Create founder haplotypes
founderPop = quickHaplo(nInd=10, nChr=1, segSites=10)

#Set simulation parameters
SP = SimParam$new(founderPop)
SP$addTraitAD(10, meanDD=0.5)
SP$setVarE(h2=0.5)

#Create population
pop = newPop(founderPop, simParam=SP)
pop@ebv = matrix(rnorm(pop@nInd), nrow=pop@nInd, ncol=1)
ebv(pop)
```

editGenome

Edit genome

Description

Edits selected loci of selected individuals to a homozygous state for either the 1 or 0 allele. The gv slot is recalculated to reflect the any changes due to editing, but other slots remain the same.

Usage

```
editGenome(pop, ind, chr, segSites, allele, simParam = NULL)
```

Arguments

pop an object of [Pop-class](#)

ind a vector of individuals to edit

chr a vector of chromosomes to edit. Length must match length of segSites.

segSites a vector of segregating sites to edit. Length must match length of chr.

allele either 0 or 1 for desired allele

simParam an object of [SimParam](#)

Value

Returns an object of [Pop-class](#)

Examples

```
#Create founder haplotypes
founderPop = quickHaplo(nInd=2, nChr=1, segSites=10)

#Set simulation parameters
SP = SimParam$new(founderPop)
SP$addTraitA(10)

#Create population
pop = newPop(founderPop, simParam=SP)

#Change individual 1 to homozygous for the 1 allele
#at locus 1, chromosome 1
pop2 = editGenome(pop, ind=1, chr=1, segSites=1,
                  allele=1, simParam=SP)
```

editGenomeTopQtl	<i>Edit genome - the top QTL</i>
------------------	----------------------------------

Description

Edits the top QTL (with the largest additive effect) to a homozygous state for the allele increasing. Only nonfixed QTL are edited. The gv slot is recalculated to reflect the any changes due to editing, but other slots remain the same.

Usage

```
editGenomeTopQtl(pop, ind, nQtl, trait = 1, increase = TRUE, simParam = NULL)
```

Arguments

pop	an object of Pop-class
ind	a vector of individuals to edit
nQtl	number of QTL to edit
trait	which trait effects should guide selection of the top QTL
increase	should the trait value be increased or decreased
simParam	an object of SimParam

Value

Returns an object of [Pop-class](#)

Examples

```
#Create founder haplotypes
founderPop = quickHaplo(nInd=2, nChr=1, segSites=10)

#Set simulation parameters
SP = SimParam$new(founderPop)
SP$addTraitA(10)

#Create population
pop = newPop(founderPop, simParam=SP)

#Change up to 10 loci for individual 1
pop2 = editGenomeTopQtl(pop, ind=1, nQtl=10, simParam=SP)
```

fastRRBLUP

Fast RR-BLUP

Description

Solves an RR-BLUP model for genomic predictions given known variance components. This implementation is meant as a fast and low memory alternative to [RRBLUP](#) or [RRBLUP2](#). Unlike the those functions, the fastRRBLUP does not fit fixed effects (other than the intercept) or account for unequal replication.

Usage

```
fastRRBLUP(
  pop,
  traits = 1,
  use = "pheno",
  snpChip = 1,
  useQtl = FALSE,
  maxIter = 1000,
  Vu = NULL,
  Ve = NULL,
  simParam = NULL,
  ...
)
```

Arguments

pop	a Pop-class to serve as the training population
traits	an integer indicating the trait to model or a function of the traits returning a single value. Only univariate models are supported.
use	train model using phenotypes "pheno", genetic values "gv", estimated breeding values "ebv", breeding values "bv", or randomly "rand"

snpChip	an integer indicating which SNP chip genotype to use
useQtl	should QTL genotypes be used instead of a SNP chip. If TRUE, snpChip specifies which trait's QTL to use, and thus these QTL may not match the QTL underlying the phenotype supplied in traits.
maxIter	maximum number of iterations.
Vu	marker effect variance. If value is NULL, a reasonable value is chosen automatically.
Ve	error variance. If value is NULL, a reasonable value is chosen automatically.
simParam	an object of SimParam
...	additional arguments if using a function for traits

Examples

```
#Create founder haplotypes
founderPop = quickHaplo(nInd=10, nChr=1, segSites=20)

#Set simulation parameters
SP = SimParam$new(founderPop)
SP$addTraitA(10)
SP$setVarE(h2=0.5)
SP$addSnpChip(10)

#Create population
pop = newPop(founderPop, simParam=SP)

#Run GS model and set EBV
ans = fastRRBLUP(pop, simParam=SP)
pop = setEBV(pop, ans, simParam=SP)

#Evaluate accuracy
cor(gv(pop), ebv(pop))
```

genicVarA	<i>Additive genic variance</i>
-----------	--------------------------------

Description

Returns additive genic variance for all traits

Usage

```
genicVarA(pop, simParam = NULL)
```

Arguments

pop	an object of Pop-class
simParam	an object of SimParam

Examples

```
#Create founder haplotypes
founderPop = quickHaplo(nInd=10, nChr=1, segSites=10)

#Set simulation parameters
SP = SimParam$new(founderPop)
SP$addTraitAD(10, meanDD=0.5)
SP$setVarE(h2=0.5)

#Create population
pop = newPop(founderPop, simParam=SP)
genicVarA(pop, simParam=SP)
```

genicVarAA

Additive-by-additive genic variance

Description

Returns additive-by-additive epistatic genic variance for all traits

Usage

```
genicVarAA(pop, simParam = NULL)
```

Arguments

pop	an object of Pop-class
simParam	an object of SimParam

Examples

```
#Create founder haplotypes
founderPop = quickHaplo(nInd=10, nChr=1, segSites=10)

#Set simulation parameters
SP = SimParam$new(founderPop)
SP$addTraitAD(10, meanDD=0.5)
SP$setVarE(h2=0.5)

#Create population
pop = newPop(founderPop, simParam=SP)
genicVarAA(pop, simParam=SP)
```

genicVarD *Dominance genic variance*

Description

Returns dominance genic variance for all traits

Usage

```
genicVarD(pop, simParam = NULL)
```

Arguments

pop an object of [Pop-class](#)
simParam an object of [SimParam](#)

Examples

```
#Create founder haplotypes  
founderPop = quickHaplo(nInd=10, nChr=1, segSites=10)  
  
#Set simulation parameters  
SP = SimParam$new(founderPop)  
SP$addTraitAD(10, meanDD=0.5)  
SP$setVarE(h2=0.5)  
  
#Create population  
pop = newPop(founderPop, simParam=SP)  
genicVarD(pop, simParam=SP)
```

genicVarG *Total genic variance*

Description

Returns total genic variance for all traits

Usage

```
genicVarG(pop, simParam = NULL)
```

Arguments

pop an object of [Pop-class](#)
simParam an object of [SimParam](#)

Examples

```
#Create founder haplotypes
founderPop = quickHaplo(nInd=10, nChr=1, segSites=10)

#Set simulation parameters
SP = SimParam$new(founderPop)
SP$addTraitAD(10, meanDD=0.5)
SP$setVarE(h2=0.5)

#Create population
pop = newPop(founderPop, simParam=SP)
genicVarG(pop, simParam=SP)
```

genParam

Sumarize genetic parameters

Description

Calculates genetic and genic additive and dominance variances for an object of [Pop-class](#)

Usage

```
genParam(pop, simParam = NULL)
```

Arguments

pop an object of [Pop-class](#)
simParam an object of [SimParam](#)

Value

varA an nTrait by nTrait matrix of additive genetic variances
varD an nTrait by nTrait matrix of dominance genetic variances
varAA an nTrait by nTrait matrix of additive-by-additive genetic variances
varG an nTrait by nTrait matrix of total genetic variances
genicVarA an nTrait vector of additive genic variances
genicVarD an nTrait vector of dominance genic variances
genicVarAA an nTrait vector of additive-by-additive genic variances
genicVarG an nTrait vector of total genic variances
covA_HW an nTrait vector of additive covariances due to non-random mating
covD_HW an nTrait vector of dominance covariances due to non-random mating
covAA_HW an nTrait vector of additive-by-additive covariances due to non-random mating
covG_HW an nTrait vector of total genic covariances due to non-random mating

covA_L an nTrait vector of additive covariances due to linkage disequilibrium
covD_L an nTrait vector of dominance covariances due to linkage disequilibrium
covAA_L an nTrait vector of additive-by-additive covariances due to linkage disequilibrium
covAD_L an nTrait vector of additive by dominance covariances due to linkage disequilibrium
covAAA_L an nTrait vector of additive by additive-by-additive covariances due to linkage disequilibrium
covDAA_L an nTrait vector of dominance by additive-by-additive covariances due to linkage disequilibrium
covG_L an nTrait vector of total genic covariances due to linkage disequilibrium
mu an nTrait vector of trait means
mu_HW an nTrait vector of expected trait means under random mating
gv a matrix of genetic values with dimensions nInd by nTraits
bv a matrix of breeding values with dimensions nInd by nTraits
dd a matrix of dominance deviations with dimensions nInd by nTraits
aa a matrix of additive-by-additive epistatic deviations with dimensions nInd by nTraits
gv_mu an nTrait vector of intercepts with dimensions nInd by nTraits
gv_a a matrix of additive genetic values with dimensions nInd by nTraits
gv_d a matrix of dominance genetic values with dimensions nInd by nTraits
gv_aa a matrix of additive-by-additive genetic values with dimensions nInd by nTraits

Examples

```

#Create founder haplotypes
founderPop = quickHaplo(nInd=10, nChr=1, segSites=10)

#Set simulation parameters
SP = SimParam$new(founderPop)
SP$addTraitAD(10, meanDD=0.5)
SP$setVarE(h2=0.5)

#Create population
pop = newPop(founderPop, simParam=SP)
ans = genParam(pop, simParam=SP)

```

getQtlMap

Get QTL genetic map

Description

Retrieves the genetic map for the QTL of a given trait.

Usage

```
getQtlMap(trait = 1, gender = "A", simParam = NULL)
```

Arguments

trait	an integer for the
gender	determines which gender specific map is returned. Options are "A" for average map, "F" for female map, and "M" for male map. All options are equivalent if not using gender specific maps.
simParam	an object of SimParam

Value

Returns a data.frame for the SNP map.

Examples

```
#Create founder haplotypes
founderPop = quickHaplo(nInd=10, nChr=1, segSites=10)

#Set simulation parameters
SP = SimParam$new(founderPop)
SP$addTraitA(5)

#Pull SNP map
getQtlMap(trait=1, simParam=SP)
```

getSnpMap	<i>Get SNP genetic map</i>
-----------	----------------------------

Description

Retrieves the genetic map for a given SNP chip.

Usage

```
getSnpMap(snpChip = 1, gender = "A", simParam = NULL)
```

Arguments

snpChip	an integer. Indicates which SNP chip's map to retrieve.
gender	determines which gender specific map is returned. Options are "A" for average map, "F" for female map, and "M" for male map. All options are equivalent if not using gender specific maps.
simParam	an object of SimParam

Value

Returns a data.frame for the SNP map.

Examples

```
#Create founder haplotypes
founderPop = quickHaplo(nInd=10, nChr=1, segSites=10)

#Set simulation parameters
SP = SimParam$new(founderPop)
SP$addSnpChip(5)

#Pull SNP map
getSnpMap(snpChip=1, simParam=SP)
```

gv

Genetic value

Description

A wrapper for accessing the gv slot

Usage

```
gv(pop)
```

Arguments

pop a [Pop-class](#) or similar object

Examples

```
#Create founder haplotypes
founderPop = quickHaplo(nInd=10, nChr=1, segSites=10)

#Set simulation parameters
SP = SimParam$new(founderPop)
SP$addTraitAD(10, meanDD=0.5)
SP$setVarE(h2=0.5)

#Create population
pop = newPop(founderPop, simParam=SP)
gv(pop)
```

hybridCross	<i>Hybrid crossing</i>
-------------	------------------------

Description

A convenience function for hybrid plant breeding simulations. Allows for easy specification of a test cross scheme and/or creation of an object of [HybridPop-class](#). Note that the [HybridPop-class](#) should only be used if the parents were created using the [makeDH](#) function or [newPop](#) using inbred founders. The id for new individuals is [mother_id]_[father_id]

Usage

```
hybridCross(
  females,
  males,
  crossPlan = "testcross",
  returnHybridPop = FALSE,
  simParam = NULL
)
```

Arguments

females	female population, an object of Pop-class
males	male population, an object of Pop-class
crossPlan	either "testcross" for all possible combinations or a matrix with two columns for designed crosses
returnHybridPop	should results be returned as HybridPop-class . If false returns results as Pop-class . Population must be fully inbred if TRUE.
simParam	an object of SimParam

Examples

```
#Create founder haplotypes
founderPop = quickHaplo(nInd=2, nChr=1, segSites=10)

#Set simulation parameters
SP = SimParam$new(founderPop)

#Create population
pop = newPop(founderPop, simParam=SP)

#Make crosses for full diallele
pop2 = hybridCross(pop, pop, simParam=SP)
```

HybridPop-class	<i>Hybrid population</i>
-----------------	--------------------------

Description

A lightweight version of [Pop-class](#) for hybrid lines. Memory is saved by not storing genotypic data.

Usage

```
## S4 method for signature 'HybridPop'
x[i]

## S4 method for signature 'HybridPop'
c(x, ...)
```

Arguments

x	a 'HybridPop'
i	index of individuals
...	additional 'HybridPop' objects

Methods (by generic)

- `[]`: Extract HybridPop using index or id
- `c`: Combine multiple HybridPops

Slots

nInd number of individuals
 id an individual's identifier
 mother the identifier of the individual's mother
 father the identifier of the individual's father
 nTraits number of traits
 gv matrix of genetic values. When using GxE traits, gv reflects gv when w=0. Dimensions are nInd by nTraits.
 pheno matrix of phenotypic values. Dimensions are nInd by nTraits.
 gxe list containing GxE slopes for GxE traits

LociMap-class	<i>Loci metadata</i>
---------------	----------------------

Description

used for both SNPs and QTLs

Slots

nLoci total number of loci

lociPerChr number of loci per chromosome

lociLoc physical position of loci

makeCross	<i>Make designed crosses</i>
-----------	------------------------------

Description

Makes crosses within a population using a user supplied crossing plan.

Usage

```
makeCross(pop, crossPlan, nProgeny = 1, simParam = NULL)
```

Arguments

pop an object of [Pop-class](#)

crossPlan a matrix with two column representing female and male parents. Either integers for the position in population or character strings for the IDs.

nProgeny number of progeny per cross

simParam an object of [SimParam](#)

Value

Returns an object of [Pop-class](#)

Examples

```
#Create founder haplotypes
founderPop = quickHaplo(nInd=10, nChr=1, segSites=10)

#Set simulation parameters
SP = SimParam$new(founderPop)

#Create population
pop = newPop(founderPop, simParam=SP)

#Cross individual 1 with individual 10
crossPlan = matrix(c(1,10), nrow=1, ncol=2)
pop2 = makeCross(pop, crossPlan, simParam=SP)
```

makeCross2

Make designed crosses

Description

Makes crosses between two populations using a user supplied crossing plan.

Usage

```
makeCross2(females, males, crossPlan, nProgeny = 1, simParam = NULL)
```

Arguments

females	an object of Pop-class for female parents.
males	an object of Pop-class for male parents.
crossPlan	a matrix with two column representing female and male parents. Either integers for the position in population or character strings for the IDs.
nProgeny	number of progeny per cross
simParam	an object of SimParam

Value

Returns an object of [Pop-class](#)

Examples

```
#Create founder haplotypes
founderPop = quickHaplo(nInd=10, nChr=1, segSites=10)

#Set simulation parameters
SP = SimParam$new(founderPop)
```

```
#Create population
pop = newPop(founderPop, simParam=SP)

#Cross individual 1 with individual 10
crossPlan = matrix(c(1,10), nrow=1, ncol=2)
pop2 = makeCross2(pop, pop, crossPlan, simParam=SP)
```

makeDH

Generates DH lines

Description

Creates DH lines from each individual in a population. Only works with diploid individuals. For polyploids, use [reduceGenome](#) and [doubleGenome](#).

Usage

```
makeDH(pop, nDH = 1, useFemale = TRUE, keepParents = TRUE, simParam = NULL)
```

Arguments

pop	an object of 'Pop' superclass
nDH	total number of DH lines per individual
useFemale	should female recombination rates be used.
keepParents	should previous parents be used for mother and father.
simParam	an object of 'SimParam' class

Value

Returns an object of [Pop-class](#)

Examples

```
#Create founder haplotypes
founderPop = quickHaplo(nInd=2, nChr=1, segSites=10)

#Set simulation parameters
SP = SimParam$new(founderPop)

#Create population
pop = newPop(founderPop, simParam=SP)

#Create 1 DH for each individual
pop2 = makeDH(pop, simParam=SP)
```

MapPop-class	<i>Raw population with genetic map</i>
--------------	----------------------------------------

Description

Extends [RawPop-class](#) to add a genetic map. This is the first object created in a simulation. It is used for creating initial populations and setting traits in the [SimParam](#).

Usage

```
## S4 method for signature 'MapPop'
x[i]
```

```
## S4 method for signature 'MapPop'
c(x, ...)
```

Arguments

x	a 'MapPop' object
i	index of chromosomes
...	additional 'MapPop' objects

Methods (by generic)

- `[]`: Extract MapPop by index
- `c`: Combine multiple MapPops

Slots

genMap	"matrix" of chromosome genetic maps
centromere	vector of centromere positions

meanG	<i>Mean genetic values</i>
-------	----------------------------

Description

Returns the mean genetic values for all traits

Usage

```
meanG(pop)
```


Arguments

pop an object of [Pop-class](#) or [HybridPop-class](#)

Examples

```
#Create founder haplotypes
founderPop = quickHaplo(nInd=10, nChr=1, segSites=10)

#Set simulation parameters
SP = SimParam$new(founderPop)
SP$addTraitA(10)
SP$setVarE(h2=0.5)

#Create population
pop = newPop(founderPop, simParam=SP)
meanG(pop)
```

meanP	<i>Mean phenotypic values</i>
-------	-------------------------------

Description

Returns the mean phenotypic values for all traits

Usage

```
meanP(pop)
```

Arguments

pop an object of [Pop-class](#) or [HybridPop-class](#)

Examples

```
#Create founder haplotypes
founderPop = quickHaplo(nInd=10, nChr=1, segSites=10)

#Set simulation parameters
SP = SimParam$new(founderPop)
SP$addTraitA(10)
SP$setVarE(h2=0.5)

#Create population
pop = newPop(founderPop, simParam=SP)
meanP(pop)
```

mergeGenome	<i>Combine genomes of individuals</i>
-------------	---------------------------------------

Description

This function is designed to model the pairing of gametes. The male and female individuals are treated as gametes, so the ploidy of newly created individuals will be the sum of it parents.

Usage

```
mergeGenome(females, males, crossPlan, simParam = NULL)
```

Arguments

females	an object of Pop-class for female parents.
males	an object of Pop-class for male parents.
crossPlan	a matrix with two column representing female and male parents. Either integers for the position in population or character strings for the IDs.
simParam	an object of SimParam

Value

Returns an object of [Pop-class](#)

Examples

```
#Create founder haplotypes
founderPop = quickHaplo(nInd=10, nChr=1, segSites=10)

#Set simulation parameters
SP = SimParam$new(founderPop)

#Create population
pop = newPop(founderPop, simParam=SP)

#Cross individual 1 with individual 10
crossPlan = matrix(c(1,10), nrow=1, ncol=2)
pop2 = mergeGenome(pop, pop, crossPlan, simParam=SP)
```

mergePops	<i>Merge list of populations</i>
-----------	----------------------------------

Description

Rapidly merges a list of populations into a single population

Usage

```
mergePops(popList)
```

Arguments

popList a list containing [Pop-class](#) elements

Value

Returns a [Pop-class](#)

Examples

```
#Create founder haplotypes
founderPop = quickHaplo(nInd=10, nChr=1, segSites=10)

#Set simulation parameters
SP = SimParam$new(founderPop)

#Create a list of populations and merge list
pop = newPop(founderPop, simParam=SP)
popList = list(pop, pop)
pop2 = mergePops(popList)
```

mutate	<i>Add Random Mutations</i>
--------	-----------------------------

Description

Adds random mutations to individuals in a population. Note that any existing phenotypes or EBVs are kept. Thus, the user will need to run [setPheno](#) and/or [setEBV](#) to generate new phenotypes or EBVs that reflect changes introduced by the new mutations.

Usage

```
mutate(pop, mutRate = 2.5e-08, returnPos = FALSE, simParam = NULL)
```

Arguments

pop	an object of Pop-class
mutRate	rate of new mutations
returnPos	should the positions of mutations be returned
simParam	an object of SimParam

Value

an object of [Pop-class](#) if returnPos=FALSE or a list containing a [Pop-class](#) and a data.frame containing the positions of mutations if returnPos=TRUE

Examples

```
#Create founder haplotypes
founderPop = quickHaplo(nInd=2, nChr=1, segSites=10)

#Set simulation parameters
SP = SimParam$new(founderPop)
SP$addTraitA(10)

#Create population
pop = newPop(founderPop, simParam=SP)

#Introduce mutations
pop = mutate(pop, simParam=SP)
```

newMapPop

New MapPop

Description

Creates a new [MapPop-class](#) from user supplied genetic maps and haplotypes.

Usage

```
newMapPop(genMap, haplotypes, inbred = FALSE, ploidy = 2L)
```

Arguments

genMap	a list of genetic maps
haplotypes	a list of matrices or data.frames that can be coerced to matrices. See details.
inbred	are individuals fully inbred
ploidy	ploidy level of organism

Details

Each item of `genMap` must be a vector of ordered genetic lengths in Morgans. The first value must be zero. The length of the vector determines the number of segregating sites on the chromosome.

Each item of `haplotypes` must be coercible to a matrix. The columns of this matrix correspond to segregating sites and their number must match

Value

an object of `MapPop-class`

Examples

```
# Create genetic map for two chromosomes, each 1 Morgan long
# Each chromosome contains 11 equally spaced segregating sites
genMap = list(seq(0,1,length.out=11),
              seq(0,1,length.out=11))

# Create haplotypes for 10 outbred individuals
chr1 = sample(x=0:1,size=20*11,replace=TRUE)
chr1 = matrix(chr1,nrow=20,ncol=11)
chr2 = sample(x=0:1,size=20*11,replace=TRUE)
chr2 = matrix(chr2,nrow=20,ncol=11)
haplotypes = list(chr1,chr2)

founderPop = newMapPop(genMap=genMap,haplotypes=haplotypes)
```

newPop

Create new Population

Description

Creates a new `Pop-class` from an object of `MapPop-class` or `RawPop-class`. The function is intended for creating initial populations from 'FOUNDERPOP' created by `runMacs`.

Usage

```
newPop(
  rawPop,
  mother = NULL,
  father = NULL,
  origM = NULL,
  origF = NULL,
  isDH = FALSE,
  simParam = NULL
)
```

Arguments

rawPop	an object of MapPop-class or RawPop-class
mother	optional id for mothers. Must match id in pedigree if using track pedigree.
father	optional id for fathers. Must match id in pedigree if using track pedigree.
origM	optional alternative id for mothers
origF	optional alternative id for fathers
isDH	optional value indicating if the individuals are doubled haploids and/or inbred founders
simParam	an object of SimParam

Value

Returns an object of [Pop-class](#)

Examples

```
#Create founder haplotypes
founderPop = quickHaplo(nInd=2, nChr=1, segSites=10)

#Set simulation parameters
SP = SimParam$new(founderPop)
SP$addTraitA(10)

#Create population
pop = newPop(founderPop, simParam=SP)
```

nInd	<i>Number of individuals</i>
------	------------------------------

Description

A wrapper for accessing the nInd slot

Usage

```
nInd(pop)
```

Arguments

pop	a Pop-class or similar object
-----	-----------------------------------------------

Examples

```
#Create founder haplotypes
founderPop = quickHaplo(nInd=10, nChr=1, segSites=10)

#Set simulation parameters
SP = SimParam$new(founderPop)
SP$addTraitAD(10, meanDD=0.5)
SP$setVarE(h2=0.5)

#Create population
pop = newPop(founderPop, simParam=SP)
nInd(pop)
```

pedigreeCross	<i>Pedigree cross</i>
---------------	-----------------------

Description

Creates a [Pop-class](#) from a generic pedigree and a set of founder individuals.

The way in which the user supplied pedigree is used depends on the value of matchID. If matchID is TRUE, the IDs in the user supplied pedigree are matched against founderNames. If matchID is FALSE, founder individuals in the user supplied pedigree are randomly sampled from founderPop.

Usage

```
pedigreeCross(
  founderPop,
  id,
  mother,
  father,
  matchID = FALSE,
  founderNames = NULL,
  maxCycle = 100,
  DH = NULL,
  useFemale = TRUE,
  simParam = NULL
)
```

Arguments

founderPop	a Pop-class
id	a vector of unique identifiers for individuals in the pedigree. The values of these IDs are separate from the IDs in the founderPop if matchID=FALSE.
mother	a vector of identifiers for the mothers of individuals in the pedigree. Must match one of the elements in the id vector or they will be treated as unknown.

father	a vector of identifiers for the fathers of individuals in the pedigree. Must match one of the elements in the id vector or they will be treated as unknown.
matchID	indicates if the IDs in founderPop should be matched to the id argument. See details.
founderNames	names for individuals in the founder population. Must be provided when matchID=TRUE.
maxCycle	the maximum number of loops to make over the pedigree to sort it.
DH	an optional vector indicating if an individual should be made a doubled haploid.
useFemale	If creating DH lines, should female recombination rates be used. This parameter has no effect if, recombRatio=1.
simParam	an object of 'SimParam' class

Examples

```
#Create founder haplotypes
founderPop = quickHaplo(nInd=2, nChr=1, segSites=10)

#Set simulation parameters
SP = SimParam$new(founderPop)

#Create population
pop = newPop(founderPop, simParam=SP)

#Pedigree for a biparental cross with 7 generations of selfing
id = 1:10
mother = c(0,0,1,3:9)
father = c(0,0,2,3:9)
pop2 = pedigreeCross(pop, id, mother, father, simParam=SP)
```

pheno

Phenotype

Description

A wrapper for accessing the pheno slot

Usage

```
pheno(pop)
```

Arguments

pop a [Pop-class](#) or similar object

Examples

```
#Create founder haplotypes
founderPop = quickHaplo(nInd=10, nChr=1, segSites=10)

#Set simulation parameters
SP = SimParam$new(founderPop)
SP$addTraitAD(10, meanDD=0.5)
SP$setVarE(h2=0.5)

#Create population
pop = newPop(founderPop, simParam=SP)
pheno(pop)
```

 Pop-class

Population

Description

Extends [RawPop-class](#) to add gender, genetic values, phenotypes, and pedigrees.

Usage

```
## S4 method for signature 'Pop'
x[i]

## S4 method for signature 'Pop'
c(x, ...)

## S4 method for signature 'Pop'
show(object)
```

Arguments

x	a 'Pop' object
i	index of individuals
...	additional 'Pop' objects
object	a 'Pop' object

Methods (by generic)

- [: Extract Pop by index or id
- c: Combine multiple Pops
- show: Show population summary

Slots

- id an individual's identifier
- mother the identifier of the individual's mother
- father the identifier of the individual's father
- gender gender of individuals
- nTraits number of traits
- gv matrix of genetic values. When using GxE traits, gv reflects gv when w=0. Dimensions are nInd by nTraits.
- pheno matrix of phenotypic values. Dimensions are nInd by nTraits.
- ebv matrix of estimated breeding values. Dimensions are nInd rows and a variable number of columns.
- gxe list containing GxE slopes for GxE traits
- fixEff a fixed effect relating to the phenotype. Used by genomic selection models but otherwise ignored.
- reps the number of replications used to measure the phenotype. Used by genomic selection models, but otherwise ignored.

 popVar

Population variance

Description

Calculates the population variance matrix as opposed to the sample variance matrix calculated by [var](#). i.e. divides by n instead of n-1

Usage

```
popVar(X)
```

Arguments

X an n by m matrix

Value

an m by m variance-covariance matrix

pullIbdHaplo *Pull Identity By Descent (IBD) haplotypes*

Description

Retrieves Identity By Descent (IBD) haplotype data

Usage

```
pullIbdHaplo(
  pop = NULL,
  chr = NULL,
  snpChip = NULL,
  pedigree = NULL,
  simParam = NULL
)
```

Arguments

pop	an object of Pop-class or RawPop-class . If NULL, haplotypes for the whole ancestral pedigree are retrieved. Otherwise, haplotypes just for the pop individuals are retrieved. In both cases the base population is controlled by pedigree.
chr	a vector of chromosomes to retrieve. If NULL, all chromosomes are retrieved.
snpChip	an integer. Indicates which SNP array loci are retrieved. If NULL, all sites are retrieved.
pedigree	a matrix with ancestral pedigree to set a base population. It should be of the same form as <code>simParam\$pedigree</code> (see <code>setTrackPed</code> in SimParam), i.e., two columns (mother and father) and the same number of rows as <code>simParam\$pedigree.Base</code> . Base population can be set by setting parents as 0. If NULL, pedigree from SimParam is taken.
simParam	an object of SimParam

Value

Returns a matrix of haplotypes with Identity By Descent (IBD) coding of locus alleles. The matrix colnames reflect whether all segregating loci (sites) are retrieved or only SNP array loci.

Examples

```
#Create founder haplotypes
founderPop = quickHaplo(nInd=10, nChr=1, segSites=15)

#Set simulation parameters
SP = SimParam$new(founderPop)
SP$addTraitA(10)
SP$addSnpChip(5)
SP$setTrackRec(TRUE)
```

```
#Create population
pop = newPop(founderPop, simParam=SP)
pullIbdHaplo(pop, simParam=SP)
```

pullQtlGeno

Pull QTL genotype

Description

Retrieves QTL genotype data

Usage

```
pullQtlGeno(pop, trait = 1, chr = NULL, simParam = NULL)
```

Arguments

pop	an object of Pop-class
trait	an integer. Indicates which trait's QTL genotypes to retrieve.
chr	a vector of chromosomes to retrieve. If NULL, all chromosome are retrieved.
simParam	an object of SimParam

Details

```
#' @title Pull SNP genotype for multiple snp chips #' #' @description Retrieves SNP genotype
data for multiple snp chips #' #' @param pop an object of Pop-class #' @param chips a vec-
tor. For each animal indicates what snp #' chip to use #' @param missing What value to use for
missing #' @param simParam an object of SimParam #' #' @return Returns a matrix of SNP geno-
types. #' #' @export pullMultipleSnpGeno = function(pop, chips, missing=9, simParam=NULL)
if(is.null(simParam)) simParam = get("SP",envir=.GlobalEnv)
```

```
# I feel like the next line shouldn't be needed but I don't know # enough R! (dmoney) missing
= as.integer(missing) allSnps = numeric(0) uniqueChips = unique(chips) for (c in uniqueChips)
allSnps = sort(union(allSnps,simParam$snpChips[[c]]@lociLoc))
```

```
output = matrix(pop@nInd,length(allSnps),data=missing) if(class(pop)=="Pop") rownames(output)
= pop@id else rownames(output) = as.character(1:pop@nInd)
```

```
for (snpChip in uniqueChips) mask = allSnps one = getGeno(pop@geno, simParam$snpChips[[snpChip]]@lociPerChr,
simParam$snpChips[[snpChip]]@lociLoc, simParam$nThreads) one = convToImat(one) for (i in
1:pop@nInd) if (chips[i] == snpChip) output[i,mask] = one[i,] output[i,mask] = one[i,]
```

```
colnames(output) = paste("SNP",1:ncol(output),sep="_")
```

```
return(output)
```

Value

Returns a matrix of QTL genotypes.

Examples

```
#Create founder haplotypes
founderPop = quickHaplo(nInd=10, nChr=1, segSites=15)

#Set simulation parameters
SP = SimParam$new(founderPop)
SP$addTraitA(10)
SP$addSnpChip(5)

#Create population
pop = newPop(founderPop, simParam=SP)
pullQtlGeno(pop, simParam=SP)
```

pullQtlHaplo	<i>Pull QTL haplotypes</i>
--------------	----------------------------

Description

Retrieves QTL haplotype data

Usage

```
pullQtlHaplo(pop, trait = 1, haplo = "all", chr = NULL, simParam = NULL)
```

Arguments

pop	an object of Pop-class
trait	an integer. Indicates which trait's QTL haplotypes to retrieve.
haplo	either "all" for all haplotypes or an integer for a single set of haplotypes. Use a value of 1 for female haplotypes and a value of 2 for male haplotypes.
chr	a vector of chromosomes to retrieve. If NULL, all chromosome are retrieved.
simParam	an object of SimParam

Details

```
#' @title Pull SNP haplotypes for multiple chips #' #' @description Retrieves SNP haplotype data
for multiple snp #' #' @param pop an object of Pop-class #' @param chips a vector. For each
animal indicates what snp #' chip to use #' @param haplo either "all" for all haplotypes or an integer
#' for a single set of haplotypes. Use a value of 1 for female #' haplotyes and a value of 2 for male
haplotypes. #' @param missing What value to use for missing #' @param simParam an object of
SimParam #' #' @return Returns a matrix of SNP haplotypes. #' #' @export pullMultipleSnpHaplo
= function(pop, chips, haplo="all", missing=9, simParam=NULL) if(is.null(simParam)) simParam
= get("SP",envir=.GlobalEnv)
```

```
# I feel like the next line shouldn't be needed but I don't know # enough R! (dmoney) missing
= as.integer(missing) allSnps = numeric(0) uniqueChips = unique(chips) for (c in uniqueChips)
allSnps = sort(union(allSnps,simParam$snpChips[[c]]@lociLoc))
```

```

if (haplo == "all") output = matrix(pop@nInd*2,length(allSnps),data=missing) if(class(pop)=="Pop")
rownames(output) = paste(rep(pop@id,each=pop@ploidy), rep(1:pop@ploidy,pop@nInd),sep="_")
else rownames(output) = paste(rep(1:pop@nInd,each=pop@ploidy), rep(1:pop@ploidy,pop@nInd),sep="_")
else output = matrix(pop@nInd,length(allSnps),data=missing) if(class(pop)=="Pop") rownames(output)
= paste(pop@id,rep(haplo,pop@nInd),sep="_") else rownames(output) = paste(1:pop@nInd,rep(haplo,pop@nInd),sep="_")
for (snpChip in uniqueChips) mask = allSnps if (haplo == "all") one = getHaplo(pop@geno, sim-
Param$snpChips[[snpChip]]@lociPerChr, simParam$snpChips[[snpChip]]@lociLoc, simParam$nThreads)
one = convToImat(one) for (i in 1:pop@nInd) if (chips[i] == snpChip) output[i*2-1,mask] = one[i*2-
1,] output[i*2,mask] = one[i*2,]
else one = getOneHaplo(pop@geno, simParam$snpChips[[snpChip]]@lociPerChr, simParam$snpChips[[snpChip]]@lociLo
as.integer(haplo), simParam$nThreads) one = convToImat(one) for (i in 1:pop@nInd) if (chips[i]
== snpChip) output[i,mask] = one[i,] output[i,mask] = one[i,]
colnames(output) = paste("SNP",1:ncol(output),sep="_")
return(output)

```

Value

Returns a matrix of QTL haplotypes.

Examples

```

#Create founder haplotypes
founderPop = quickHaplo(nInd=10, nChr=1, segSites=15)

#Set simulation parameters
SP = SimParam$new(founderPop)
SP$addTraitA(10)
SP$addSnpChip(5)

#Create population
pop = newPop(founderPop, simParam=SP)
pullQtlHaplo(pop, simParam=SP)

```

pullSegSiteGeno

Pull seg site genotypes

Description

Retrieves genotype data for all segregating sites

Usage

```
pullSegSiteGeno(pop, chr = NULL, simParam = NULL)
```

Arguments

pop an object of [Pop-class](#) or [RawPop-class](#)
 chr a vector of chromosomes to retrieve. If NULL, all chromosome are retrieved.
 simParam an object of [SimParam](#)

Value

Returns a matrix of genotypes

Examples

```
#Create founder haplotypes
founderPop = quickHaplo(nInd=10, nChr=1, segSites=15)

#Set simulation parameters
SP = SimParam$new(founderPop)
SP$addTraitA(10)
SP$addSnpChip(5)

#Create population
pop = newPop(founderPop, simParam=SP)
pullSegSiteGeno(pop, simParam=SP)
```

pullSegSiteHaplo *Pull seg site haplotypes*

Description

Retrieves haplotype data for all segregating sites

Usage

```
pullSegSiteHaplo(pop, haplo = "all", chr = NULL, simParam = NULL)
```

Arguments

pop an object of [Pop-class](#) or [RawPop-class](#)
 haplo either "all" for all haplotypes or an integer for a single set of haplotypes. Use a value of 1 for female haplotypes and a value of 2 for male haplotypes.
 chr a vector of chromosomes to retrieve. If NULL, all chromosome are retrieved.
 simParam an object of [SimParam](#)

Value

Returns a matrix of haplotypes

Examples

```
#Create founder haplotypes
founderPop = quickHaplo(nInd=10, nChr=1, segSites=15)

#Set simulation parameters
SP = SimParam$new(founderPop)
SP$addTraitA(10)
SP$addSnpChip(5)

#Create population
pop = newPop(founderPop, simParam=SP)
pullSegSiteHaplo(pop, simParam=SP)
```

pullSnpGeno

Pull SNP genotype

Description

Retrieves SNP genotype data

Usage

```
pullSnpGeno(pop, snpChip = 1, chr = NULL, simParam = NULL)
```

Arguments

pop	an object of Pop-class
snpChip	an integer. Indicates which SNP chip's genotypes to retrieve.
chr	a vector of chromosomes to retrieve. If NULL, all chromosome are retrieved.
simParam	an object of SimParam

Value

Returns a matrix of SNP genotypes.

Examples

```
#Create founder haplotypes
founderPop = quickHaplo(nInd=10, nChr=1, segSites=15)

#Set simulation parameters
SP = SimParam$new(founderPop)
SP$addTraitA(10)
SP$addSnpChip(5)

#Create population
pop = newPop(founderPop, simParam=SP)
pullSnpGeno(pop, simParam=SP)
```

pullSnpHaplo	<i>Pull SNP haplotypes</i>
--------------	----------------------------

Description

Retrieves SNP haplotype data

Usage

```
pullSnpHaplo(pop, snpChip = 1, haplo = "all", chr = NULL, simParam = NULL)
```

Arguments

pop	an object of Pop-class
snpChip	an integer. Indicates which SNP chip's haplotypes to retrieve.
haplo	either "all" for all haplotypes or an integer for a single set of haplotypes. Use a value of 1 for female haplotypes and a value of 2 for male haplotypes.
chr	a vector of chromosomes to retrieve. If NULL, all chromosome are retrieved.
simParam	an object of SimParam

Value

Returns a matrix of SNP haplotypes.

Examples

```
#Create founder haplotypes
founderPop = quickHaplo(nInd=10, nChr=1, segSites=15)

#Set simulation parameters
SP = SimParam$new(founderPop)
SP$addTraitA(10)
SP$addSnpChip(5)

#Create population
pop = newPop(founderPop, simParam=SP)
pullSnpHaplo(pop, simParam=SP)
```

quickHaplo	<i>Quick founder haplotype simulation</i>
------------	-------------------------------------------

Description

Rapidly simulates founder haplotypes by randomly sampling 0s and 1s. This is equivalent to having all loci with allele frequency 0.5 and being in linkage equilibrium.

Usage

```
quickHaplo(nInd, nChr, segSites, genLen = 1, ploidy = 2L, inbred = FALSE)
```

Arguments

nInd	number of individuals to simulate
nChr	number of chromosomes to simulate
segSites	number of segregating sites per chromosome
genLen	genetic length of chromosomes
ploidy	ploidy level of organism
inbred	should founder individuals be inbred

Value

an object of [MapPop-class](#)

Examples

```
# Creates a populations of 10 outbred individuals
# Their genome consists of 1 chromosome and 100 segregating sites
founderPop = quickHaplo(nInd=10,nChr=1,segSites=100)
```

randCross	<i>Make random crosses</i>
-----------	----------------------------

Description

A wrapper for [makeCross](#) that randomly selects parental combinations for all possible combinations.

Usage

```
randCross(  
  pop,  
  nCrosses,  
  nProgeny = 1,  
  balance = TRUE,  
  parents = NULL,  
  ignoreGender = FALSE,  
  simParam = NULL  
)
```

Arguments

pop	an object of Pop-class
nCrosses	total number of crosses to make
nProgeny	number of progeny per cross
balance	if using gender, this option will balance the number of progeny per parent
parents	an optional vector of indices for allowable parents
ignoreGender	should gender be ignored
simParam	an object of SimParam

Value

Returns an object of [Pop-class](#)

Examples

```
#Create founder haplotypes  
founderPop = quickHaplo(nInd=10, nChr=1, segSites=10)  
  
#Set simulation parameters  
SP = SimParam$new(founderPop)  
  
#Create population  
pop = newPop(founderPop, simParam=SP)  
  
#Make 10 crosses  
pop2 = randCross(pop, 10, simParam=SP)
```

randCross2	<i>Make random crosses</i>
------------	----------------------------

Description

A wrapper for [makeCross2](#) that randomly selects parental combinations for all possible combinations between two populations.

Usage

```
randCross2(  
  females,  
  males,  
  nCrosses,  
  nProgeny = 1,  
  balance = TRUE,  
  femaleParents = NULL,  
  maleParents = NULL,  
  ignoreGender = FALSE,  
  simParam = NULL  
)
```

Arguments

females	an object of Pop-class for female parents.
males	an object of Pop-class for male parents.
nCrosses	total number of crosses to make
nProgeny	number of progeny per cross
balance	this option will balance the number of progeny per parent
femaleParents	an optional vector of indices for allowable female parents
maleParents	an optional vector of indices for allowable male parents
ignoreGender	should gender be ignored
simParam	an object of SimParam

Value

Returns an object of [Pop-class](#)

Examples

```
#Create founder haplotypes  
founderPop = quickHaplo(nInd=10, nChr=1, segSites=10)  
  
#Set simulation parameters  
SP = SimParam$new(founderPop)
```

```
#Create population
pop = newPop(founderPop, simParam=SP)

#Make 10 crosses
pop2 = randCross2(pop, pop, 10, simParam=SP)
```

RawPop-class

Raw Population

Description

The raw population class contains only genotype data.

Usage

```
## S4 method for signature 'RawPop'
x[i]

## S4 method for signature 'RawPop'
c(x, ...)
```

```
## S4 method for signature 'RawPop'
show(object)
```

Arguments

x	a 'RawPop' object
i	index of individuals
...	additional 'RawPop' objects
object	a 'RawPop' object

Methods (by generic)

- `[]`: Extract RawPop by index
- `c`: Combine multiple RawPops
- `show`: Show population summary

Slots

nInd number of individuals
nChr number of chromosomes
ploidy level of ploidy
nLoci number of loci per chromosome

geno "matrix" containing chromosome genotypes. The "matrix" has dimensions nChr by 1 and each element is a three dimensional array of raw values. The array dimensions are nLoci by ploidy by nInd.

reduceGenome *Create individuals with reduced ploidy*

Description

Creates new individuals from gametes. This function was created to model the creation of diploid potatoes from tetraploid potatoes. It can be used on any population with an even ploidy level. The newly created individuals will have half the ploidy level of the originals. The reduction can occur with or without genetic recombination.

Usage

```
reduceGenome(
  pop,
  nProgeny = 1,
  useFemale = TRUE,
  keepParents = TRUE,
  simRecomb = TRUE,
  simParam = NULL
)
```

Arguments

pop	an object of 'Pop' superclass
nProgeny	total number of progeny per individual
useFemale	should female recombination rates be used.
keepParents	should previous parents be used for mother and father.
simRecomb	should genetic recombination be modeled.
simParam	an object of 'SimParam' class

Value

Returns an object of [Pop-class](#)

Examples

```
#Create founder haplotypes
founderPop = quickHaplo(nInd=2, nChr=1, segSites=10)

#Set simulation parameters
SP = SimParam$new(founderPop)

#Create population
```

```
pop = newPop(founderPop, simParam=SP)

#Create individuals with reduced ploidy
pop2 = reduceGenome(pop, simParam=SP)
```

resetPop

Reset population

Description

Recalculates a population's genetic values and resets phenotypes and EBVs.

Usage

```
resetPop(pop, simParam = NULL)
```

Arguments

pop	an object of Pop-class
simParam	an object of SimParam

Value

an object of [Pop-class](#)

Examples

```
#Create founder haplotypes
founderPop = quickHaplo(nInd=2, nChr=1, segSites=10)

#Set simulation parameters
SP = SimParam$new(founderPop)
SP$addTraitA(10)

#Create population
pop = newPop(founderPop, simParam=SP)

#Rescale to set mean to 1
SP$rescaleTraits(mean=1)
pop = resetPop(pop, simParam=SP)
```

RRBLUP

*RR-BLUP Model***Description**

Fits an RR-BLUP model for genomic predictions.

Usage

```
RRBLUP(
  pop,
  traits = 1,
  use = "pheno",
  snpChip = 1,
  useQtl = FALSE,
  maxIter = 1000L,
  useReps = FALSE,
  simParam = NULL,
  ...
)
```

Arguments

pop	a Pop-class to serve as the training population
traits	an integer indicating the trait or traits to model, or a function of the traits returning a single value.
use	train model using phenotypes "pheno", genetic values "gv", estimated breeding values "ebv", breeding values "bv", or randomly "rand"
snpChip	an integer indicating which SNP chip genotype to use
useQtl	should QTL genotypes be used instead of a SNP chip. If TRUE, snpChip specifies which trait's QTL to use, and thus these QTL may not match the QTL underlying the phenotype supplied in traits.
maxIter	maximum number of iterations. Only used when number of traits is greater than 1.
useReps	should population's reps slot be used to model heterogeneous error variance
simParam	an object of SimParam
...	additional arguments if using a function for traits

Examples

```
#Create founder haplotypes
founderPop = quickHaplo(nInd=10, nChr=1, segSites=20)

#Set simulation parameters
SP = SimParam$new(founderPop)
```



```

SP$addTraitA(10)
SP$setVarE(h2=0.5)
SP$addSnpChip(10)

#Create population
pop = newPop(founderPop, simParam=SP)

#Run GS model and set EBV
ans = RRBLUP(pop, simParam=SP)
pop = setEBV(pop, ans, simParam=SP)

#Evaluate accuracy
cor(gv(pop), ebv(pop))

```

RRBLUP2

RR-BLUP Model 2

Description

Fits an RR-BLUP model for genomic predictions. This implementation is meant for situations where [RRBLUP](#) is too slow. Note that RRBLUP2 is only faster in certain situations, see details below. Most users should use [RRBLUP](#).

Usage

```

RRBLUP2(
  pop,
  traits = 1,
  use = "pheno",
  snpChip = 1,
  useQtl = FALSE,
  maxIter = 10,
  Vu = NULL,
  Ve = NULL,
  useEM = TRUE,
  tol = 1e-06,
  useReps = FALSE,
  simParam = NULL,
  ...
)

```

Arguments

pop	a Pop-class to serve as the training population
traits	an integer indicating the trait to model or a function of the traits returning a single value. Unlike RRBLUP , only univariate models are supported.

use	train model using phenotypes "pheno", genetic values "gv", estimated breeding values "ebv", breeding values "bv", or randomly "rand"
snpChip	an integer indicating which SNP chip genotype to use
useQtl	should QTL genotypes be used instead of a SNP chip. If TRUE, snpChip specifies which trait's QTL to use, and thus these QTL may not match the QTL underlying the phenotype supplied in traits.
maxIter	maximum number of iterations.
Vu	marker effect variance. If value is NULL, a reasonable starting point is chosen automatically.
Ve	error variance. If value is NULL, a reasonable starting point is chosen automatically.
useEM	use EM to solve variance components. If false, the initial values are considered true.
tol	tolerance for EM algorithm convergence
useReps	should population's reps slot be used to model heterogeneous error variance
simParam	an object of SimParam
...	additional arguments if using a function for traits

Details

The RRBLUP2 function works best when the number of markers is not too large. This is because it solves the RR-BLUP problem by setting up and solving Henderson's mixed model equations. Solving these equations involves a square matrix with dimensions equal to the number of fixed effects plus the number of random effects (markers). Whereas the [RRBLUP](#) function solves the RR-BLUP problem using the EMMA approach. This approach involves a square matrix with dimensions equal to the number of phenotypic records. This means that the RRBLUP2 function uses less memory than RRBLUP when the number of markers is approximately equal to or smaller than the number of phenotypic records.

The RRBLUP2 function is not recommend for cases where the variance components are unknown. This is uses the EM algorithm to solve for unknown variance components, which is generally considerably slower than the EMMA approach of [RRBLUP](#). The number of iterations for the EM algorith is set by maxIter. The default value is typically too small for convergence. When the algorithm fails to converage a warning is displayed, but results are given for the last iteration. These results may be "good enough". However we make no claim to this effect, because we can not generalize to all possible use cases.

The RRBLUP2 function can quickly solve the mixed model equations without estimating variance components. The variance components are set by defining Vu and Ve. Estimation of components is suppressed by setting useEM to false. This may be useful if the model is being retrained multiple times during the simulation. You could run [RRBLUP](#) function the first time the model is trained, and then use the variance components from this output for all future runs with the RRBLUP2 functions. Again, we can make no claim to the general robustness of this approach.

Examples

```
#Create founder haplotypes
```

```

founderPop = quickHaplo(nInd=10, nChr=1, segSites=20)

#Set simulation parameters
SP = SimParam$new(founderPop)
SP$addTraitA(10)
SP$setVarE(h2=0.5)
SP$addSnpChip(10)

#Create population
pop = newPop(founderPop, simParam=SP)

#Run GS model and set EBV
ans = RRBLUP2(pop, simParam=SP)
pop = setEBV(pop, ans, simParam=SP)

#Evaluate accuracy
cor(gv(pop), ebv(pop))

```

RRBLUPMemUse

RRBLUP Memory Usage

Description

Estimates the amount of RAM needed to run the [RRBLUP](#) and its related functions for a given training population size. Note that this function may underestimate total usage.

Usage

```
RRBLUPMemUse(nInd, nMarker, model = "REG")
```

Arguments

nInd	the number of individuals in the training population
nMarker	the number of markers per individual
model	either "REG", "GCA", or "SCA" for RRBLUP , RRBLUP_GCA and RRBLUP_SCA respectively.

Value

Returns an estimate for the required gigabytes of RAM

Examples

```
RRBLUPMemUse(nInd=1000, nMarker=5000)
```

RRBLUP_D

*RR-BLUP Model with Dominance***Description**

Fits an RR-BLUP model for genomic predictions that includes dominance effects.

Usage

```
RRBLUP_D(
  pop,
  traits = 1,
  use = "pheno",
  snpChip = 1,
  useQtl = FALSE,
  maxIter = 40L,
  useReps = FALSE,
  simParam = NULL,
  ...
)
```

Arguments

pop	a Pop-class to serve as the training population
traits	an integer indicating the trait to model, or a function of the traits returning a single value.
use	train model using phenotypes "pheno", genetic values "gv", estimated breeding values "ebv", breeding values "bv", or randomly "rand"
snpChip	an integer indicating which SNP chip genotype to use
useQtl	should QTL genotypes be used instead of a SNP chip. If TRUE, snpChip specifies which trait's QTL to use, and thus these QTL may not match the QTL underlying the phenotype supplied in traits.
maxIter	maximum number of iterations. Only used when number of traits is greater than 1.
useReps	should population's reps slot be used to model heterogeneous error variance
simParam	an object of SimParam
...	additional arguments if using a function for traits

Examples

```
#Create founder haplotypes
founderPop = quickHaplo(nInd=10, nChr=1, segSites=20)

#Set simulation parameters
SP = SimParam$new(founderPop)
```

```

SP$addTraitAD(10, meanDD=0.5)
SP$setVarE(h2=0.5)
SP$addSnpChip(10)

#Create population
pop = newPop(founderPop, simParam=SP)

#Run GS model and set EBV
ans = RRBLUP_D(pop, simParam=SP)
pop = setEBV(pop, ans, simParam=SP)

#Evaluate accuracy
cor(gv(pop), ebv(pop))

```

RRBLUP_D2

RR-BLUP with Dominance Model 2

Description

Fits an RR-BLUP model for genomic predictions that includes dominance effects. This implementation is meant for situations where [RRBLUP_D](#) is too slow. Note that [RRBLUP_D2](#) is only faster in certain situations. Most users should use [RRBLUP_D](#).

Usage

```

RRBLUP_D2(
  pop,
  traits = 1,
  use = "pheno",
  snpChip = 1,
  useQtl = FALSE,
  maxIter = 10,
  Va = NULL,
  Vd = NULL,
  Ve = NULL,
  useEM = TRUE,
  tol = 1e-06,
  useReps = FALSE,
  simParam = NULL,
  ...
)

```

Arguments

pop	a Pop-class to serve as the training population
traits	an integer indicating the trait to model, or a function of the traits returning a single value.

use	train model using phenotypes "pheno", genetic values "gv", estimated breeding values "ebv", breeding values "bv", or randomly "rand"
snpChip	an integer indicating which SNP chip genotype to use
useQtl	should QTL genotypes be used instead of a SNP chip. If TRUE, snpChip specifies which trait's QTL to use, and thus these QTL may not match the QTL underlying the phenotype supplied in traits.
maxIter	maximum number of iterations. Only used when number of traits is greater than 1.
Va	marker effect variance for additive effects. If value is NULL, a reasonable starting point is chosen automatically.
Vd	marker effect variance for dominance effects. If value is NULL, a reasonable starting point is chosen automatically.
Ve	error variance. If value is NULL, a reasonable starting point is chosen automatically.
useEM	use EM to solve variance components. If false, the initial values are considered true.
tol	tolerance for EM algorithm convergence
useReps	should population's reps slot be used to model heterogeneous error variance
simParam	an object of SimParam
...	additional arguments if using a function for traits

Examples

```
#Create founder haplotypes
founderPop = quickHaplo(nInd=10, nChr=1, segSites=20)

#Set simulation parameters
SP = SimParam$new(founderPop)
SP$addTraitAD(10, meanDD=0.5)
SP$setVarE(h2=0.5)
SP$addSnpChip(10)

#Create population
pop = newPop(founderPop, simParam=SP)

#Run GS model and set EBV
ans = RRBLUP_D2(pop, simParam=SP)
pop = setEBV(pop, ans, simParam=SP)

#Evaluate accuracy
cor(gv(pop), ebv(pop))
```

RRBLUP_GCA

*RR-BLUP GCA Model***Description**

Fits an RR-BLUP model that estimates separate marker effects for females and males. Useful for predicting GCA of parents in single cross hybrids. Can also predict performance of specific single cross hybrids.

Usage

```
RRBLUP_GCA(
  pop,
  traits = 1,
  use = "pheno",
  snpChip = 1,
  useQtl = FALSE,
  maxIter = 40L,
  useReps = FALSE,
  simParam = NULL,
  ...
)
```

Arguments

pop	a Pop-class to serve as the training population
traits	an integer indicating the trait to model, or a function of the traits returning a single value.
use	train model using phenotypes "pheno", genetic values "gv", estimated breeding values "ebv", breeding values "bv", or randomly "rand"
snpChip	an integer indicating which SNP chip genotype to use
useQtl	should QTL genotypes be used instead of a SNP chip. If TRUE, snpChip specifies which trait's QTL to use, and thus these QTL may not match the QTL underlying the phenotype supplied in traits.
maxIter	maximum number of iterations for convergence.
useReps	should population's reps slot be used to model heterogeneous error variance
simParam	an object of SimParam
...	additional arguments if using a function for traits

Examples

```
#Create founder haplotypes
founderPop = quickHaplo(nInd=10, nChr=1, segSites=20)

#Set simulation parameters
```

```

SP = SimParam$new(founderPop)
SP$addTraitA(10)
SP$setVarE(h2=0.5)
SP$addSnpChip(10)

#Create population
pop = newPop(founderPop, simParam=SP)

#Run GS model and set EBV
ans = RRBLUP_GCA(pop, simParam=SP)
pop = setEBV(pop, ans, simParam=SP)

#Evaluate accuracy
cor(gv(pop), ebv(pop))

```

RRBLUP_GCA2

RR-BLUP GCA Model 2

Description

Fits an RR-BLUP model that estimates separate marker effects for females and males. This implementation is meant for situations where [RRBLUP_GCA](#) is too slow. Note that [RRBLUP_GCA2](#) is only faster in certain situations. Most users should use [RRBLUP_GCA](#).

Usage

```

RRBLUP_GCA2(
  pop,
  traits = 1,
  use = "pheno",
  snpChip = 1,
  useQtl = FALSE,
  maxIter = 10,
  VuF = NULL,
  VuM = NULL,
  Ve = NULL,
  useEM = TRUE,
  tol = 1e-06,
  useReps = FALSE,
  simParam = NULL,
  ...
)

```

Arguments

pop	a Pop-class to serve as the training population
traits	an integer indicating the trait to model, or a function of the traits returning a single value.

use	train model using phenotypes "pheno", genetic values "gv", estimated breeding values "ebv", breeding values "bv", or randomly "rand"
snpChip	an integer indicating which SNP chip genotype to use
useQtl	should QTL genotypes be used instead of a SNP chip. If TRUE, snpChip specifies which trait's QTL to use, and thus these QTL may not match the QTL underlying the phenotype supplied in traits.
maxIter	maximum number of iterations for convergence.
VuF	marker effect variance for females. If value is NULL, a reasonable starting point is chosen automatically.
VuM	marker effect variance for males. If value is NULL, a reasonable starting point is chosen automatically.
Ve	error variance. If value is NULL, a reasonable starting point is chosen automatically.
useEM	use EM to solve variance components. If false, the initial values are considered true.
tol	tolerance for EM algorithm convergence
useReps	should population's reps slot be used to model heterogeneous error variance
simParam	an object of SimParam
...	additional arguments if using a function for traits

Examples

```
#Create founder haplotypes
founderPop = quickHaplo(nInd=10, nChr=1, segSites=20)

#Set simulation parameters
SP = SimParam$new(founderPop)
SP$addTraitA(10)
SP$setVarE(h2=0.5)
SP$addSnpChip(10)

#Create population
pop = newPop(founderPop, simParam=SP)

#Run GS model and set EBV
ans = RRBLUP_GCA2(pop, simParam=SP)
pop = setEBV(pop, ans, simParam=SP)

#Evaluate accuracy
cor(gv(pop), ebv(pop))
```

RRBLUP_SCA

*RR-BLUP SCA Model***Description**

An extension of [RRBLUP_GCA](#) that adds dominance effects. Note that we have not seen any consistent benefit of this model over [RRBLUP_GCA](#).

Usage

```
RRBLUP_SCA(
  pop,
  traits = 1,
  use = "pheno",
  snpChip = 1,
  useQtl = FALSE,
  maxIter = 40L,
  useReps = FALSE,
  simParam = NULL,
  ...
)
```

Arguments

pop	a Pop-class to serve as the training population
traits	an integer indicating the trait to model, or a function of the traits returning a single value.
use	train model using phenotypes "pheno", genetic values "gv", estimated breeding values "ebv", breeding values "bv", or randomly "rand"
snpChip	an integer indicating which SNP chip genotype to use
useQtl	should QTL genotypes be used instead of a SNP chip. If TRUE, snpChip specifies which trait's QTL to use, and thus these QTL may not match the QTL underlying the phenotype supplied in traits.
maxIter	maximum number of iterations for convergence.
useReps	should population's reps slot be used to model heterogeneous error variance
simParam	an object of SimParam
...	additional arguments if using a function for traits

Examples

```
#Create founder haplotypes
founderPop = quickHaplo(nInd=2, nChr=1, segSites=20)

#Set simulation parameters
SP = SimParam$new(founderPop)
```

```

SP$addTraitA(10)
SP$setVarE(h2=0.5)
SP$addSnpChip(10)

#Create population
pop = newPop(founderPop, simParam=SP)

#Run GS model and set EBV
ans = RRBLUP_SCA(pop, simParam=SP)
pop = setEBV(pop, ans, simParam=SP)

#Evaluate accuracy
cor(gv(pop), ebv(pop))

```

RRBLUP_SCA2

RR-BLUP SCA Model 2

Description

Fits an RR-BLUP model that estimates separate additive effects for females and males and a dominance effect. This implementation is meant for situations where [RRBLUP_SCA](#) is too slow. Note that [RRBLUP_SCA2](#) is only faster in certain situations. Most users should use [RRBLUP_SCA](#).

Usage

```

RRBLUP_SCA2(
  pop,
  traits = 1,
  use = "pheno",
  snpChip = 1,
  useQtl = FALSE,
  maxIter = 10,
  VuF = NULL,
  VuM = NULL,
  VuD = NULL,
  Ve = NULL,
  useEM = TRUE,
  tol = 1e-06,
  useReps = FALSE,
  simParam = NULL,
  ...
)

```

Arguments

`pop` a [Pop-class](#) to serve as the training population

traits	an integer indicating the trait to model, or a function of the traits returning a single value.
use	train model using phenotypes "pheno", genetic values "gv", estimated breeding values "ebv", breeding values "bv", or randomly "rand"
snpChip	an integer indicating which SNP chip genotype to use
useQtl	should QTL genotypes be used instead of a SNP chip. If TRUE, snpChip specifies which trait's QTL to use, and thus these QTL may not match the QTL underlying the phenotype supplied in traits.
maxIter	maximum number of iterations for convergence.
VuF	marker effect variance for females. If value is NULL, a reasonable starting point is chosen automatically.
VuM	marker effect variance for males. If value is NULL, a reasonable starting point is chosen automatically.
VuD	marker effect variance for dominance. If value is NULL, a reasonable starting point is chosen automatically.
Ve	error variance. If value is NULL, a reasonable starting point is chosen automatically.
useEM	use EM to solve variance components. If false, the initial values are considered true.
tol	tolerance for EM algorithm convergence
useReps	should population's reps slot be used to model heterogeneous error variance
simParam	an object of SimParam
...	additional arguments if using a function for traits

Examples

```
#Create founder haplotypes
founderPop = quickHaplo(nInd=10, nChr=1, segSites=20)

#Set simulation parameters
SP = SimParam$new(founderPop)
SP$addTraitA(10)
SP$setVarE(h2=0.5)
SP$addSnpChip(10)

#Create population
pop = newPop(founderPop, simParam=SP)

#Run GS model and set EBV
ans = RRBLUP_SCA2(pop, simParam=SP)
pop = setEBV(pop, ans, simParam=SP)

#Evaluate accuracy
cor(gv(pop), ebv(pop))
```

`RRsol-class`*RR-BLUP Solution*

Description

Contains output from AlphaSimR's genomic selection functions.

Slots

`gv` Trait(s) for estimating genetic values
`bv` Trait(s) for estimating breeding values
`female` Trait(s) for estimating GCA in the female pool
`male` Trait(s) for estimating GCA in the male pool
`Vu` Estimated marker variance(s)
`Ve` Estimated error variance

`runMacs`*Create founder haplotypes using MaCS*

Description

Uses the MaCS software to produce founder haplotypes.

Usage

```
runMacs(  
  nInd,  
  nChr = 1,  
  segSites = NULL,  
  inbred = FALSE,  
  species = "GENERIC",  
  split = NULL,  
  ploidy = 2L,  
  manualCommand = NULL,  
  manualGenLen = NULL,  
  nThreads = NULL  
)
```

Arguments

nInd	number of individuals to simulate
nChr	number of chromosomes to simulate
segSites	number of segregating sites to keep per chromosome. A value of NULL results in all sites being retained.
inbred	should founder individuals be inbred
species	species history to simulate. See details.
split	an optional historic population split in terms of generations ago.
ploidy	ploidy level of organism
manualCommand	user provided MaCS options. For advanced users only.
manualGenLen	user provided genetic length. This must be supplied if using manualCommand. If not using manualCommand, this value will replace the predefined genetic length for the species. However, this the genetic length is only used by AlphaSimR and is not passed to MaCS, so MaCS still uses the predefined genetic length. For advanced users only.
nThreads	if OpenMP is available, this will allow for simulating chromosomes in parallel. If the value is NULL, the number of threads is automatically detected.

Details

The current species histories are included: GENERIC, CATTLE, WHEAT, MAIZE, and EUROPEAN.

Value

an object of [MapPop-class](#)

Examples

```
# Creates a populations of 10 outbred individuals
# Their genome consists of 1 chromosome and 100 segregating sites
founderPop = runMacs(nInd=10,nChr=1,segSites=100)
```

runMacs2

Alternative wrapper for MaCS

Description

A wrapper function for [runMacs](#). This wrapper is designed to be easier to use than supply custom comands to manualCommand in [runMacs](#). It effectively automates the creation of an appropriate manualCommand using user supplied variables, but only deals with a subset of the possibilities. The defaults were chosen to match species="GENERIC" in [runMacs](#).

Usage

```
runMacs2(
  nInd,
  nChr = 1,
  segSites = NULL,
  Ne = 100,
  bp = 1e+08,
  genLen = 1,
  mutRate = 2.5e-08,
  histNe = c(500, 1500, 6000, 12000, 1e+05),
  histGen = c(100, 1000, 10000, 1e+05, 1e+06),
  inbred = FALSE,
  split = NULL,
  ploidy = 2L,
  returnCommand = FALSE,
  nThreads = NULL
)
```

Arguments

nInd	number of individuals to simulate
nChr	number of chromosomes to simulate
segSites	number of segregating sites to keep per chromosome
Ne	effective population size
bp	base pair length of chromosome
genLen	genetic length of chromosome in Morgans
mutRate	per base pair mutation rate
histNe	effective population size in previous generations
histGen	number of generations ago for effective population sizes given in histNe
inbred	should founder individuals be inbred
split	an optional historic population split in terms of generations ago
ploidy	ploidy level of organism
returnCommand	should the command passed to manualCommand in runMacs be returned. If TRUE, MaCS will not be called and the command is returned instead.
nThreads	if OpenMP is available, this will allow for simulating chromosomes in parallel. If the value is NULL, the number of threads is automatically detected.

Value

an object of [MapPop-class](#) or if returnCommand is true a string giving the MaCS command passed to the manualCommand argument of [runMacs](#).

Examples

```
# Creates a populations of 10 outbred individuals
# Their genome consists of 1 chromosome and 100 segregating sites
# The command is equivalent to using species="GENERIC" in runMacs
founderPop = runMacs2(nInd=10,nChr=1,segSites=100)
```

sampleHaplo

Sample haplotypes from a MapPop

Description

Creates a new [MapPop-class](#) from an existing [MapPop-class](#) by randomly sampling haplotypes.

Usage

```
sampleHaplo(mapPop, nInd, inbred = FALSE, ploidy = NULL, replace = TRUE)
```

Arguments

mapPop	the MapPop-class used to sample haplotypes
nInd	the number of individuals to create
inbred	should new individuals be fully inbred
ploidy	new ploidy level for organism. If NULL, the ploidy level of the mapPop is used.
replace	should haplotypes be sampled with replacement

Value

an object of [MapPop-class](#)

Examples

```
founderPop = quickHaplo(nInd=2,nChr=2,segSites=11,inbred=TRUE)
founderPop = sampleHaplo(mapPop=founderPop,nInd=20)
```

selectCross	<i>Select and randomly cross</i>
-------------	----------------------------------

Description

This is a wrapper that combines the functionalities of [randCross](#) and [selectInd](#). The purpose of this wrapper is to combine both selection and crossing in one function call that minimized the amount of intermediate populations created. This reduces RAM usage and simplifies code writing. Note that this wrapper does not provide the full functionality of either function.

Usage

```
selectCross(
  pop,
  nInd = NULL,
  nFemale = NULL,
  nMale = NULL,
  nCrosses,
  nProgeny = 1,
  trait = 1,
  use = "pheno",
  selectTop = TRUE,
  simParam = NULL,
  ...,
  balance = TRUE
)
```

Arguments

<code>pop</code>	an object of Pop-class
<code>nInd</code>	the number of individuals to select. These individuals are selected without regards to gender and it supercedes values for <code>nFemale</code> and <code>nMale</code> . Thus if the simulation uses gender, it is likely better to leave this value as <code>NULL</code> and use <code>nFemale</code> and <code>nMale</code> instead.
<code>nFemale</code>	the number of females to select. This value is ignored if <code>nInd</code> is set.
<code>nMale</code>	the number of males to select. This value is ignored if <code>nInd</code> is set.
<code>nCrosses</code>	total number of crosses to make
<code>nProgeny</code>	number of progeny per cross
<code>trait</code>	the trait for selection. Either a number indicating a single trait or a function returning a vector of length <code>nInd</code> .
<code>use</code>	select on genetic values "gv", estimated breeding values "ebv", breeding values "bv", phenotypes "pheno", or randomly "rand"
<code>selectTop</code>	selects highest values if true. Selects lowest values if false.
<code>simParam</code>	an object of SimParam

... additional arguments if using a function for trait
 balance if using gender, this option will balance the number of progeny per parent. This argument occurs after ..., so the argument name must be matched exactly.

Value

Returns an object of [Pop-class](#)

Examples

```
#Create founder haplotypes
founderPop = quickHaplo(nInd=10, nChr=1, segSites=10)

#Set simulation parameters
SP = SimParam$new(founderPop)
SP$addTraitA(10)
SP$setVarE(h2=0.5)

#Create population
pop = newPop(founderPop, simParam=SP)

#Select 4 individuals and make 8 crosses
pop2 = selectCross(pop, nInd=4, nCrosses=8, simParam=SP)
```

selectFam

Select families

Description

Selects a subset of full-sib families from a population.

Usage

```
selectFam(
  pop,
  nFam,
  trait = 1,
  use = "pheno",
  gender = "B",
  famType = "B",
  selectTop = TRUE,
  returnPop = TRUE,
  candidates = NULL,
  simParam = NULL,
  ...
)
```

Arguments

pop	and object of Pop-class or HybridPop-class
nFam	the number of families to select
trait	the trait for selection. Either a number indicating a single trait or a function returning a vector of length nInd.
use	select on genetic values "gv", estimated breeding values "ebv", breeding values "bv", phenotypes "pheno", or randomly "rand"
gender	which gender to select. Use "B" for both, "F" for females and "M" for males. If the simulation is not using gender, the argument is ignored.
famType	which type of family to select. Use "B" for full-sib families, "F" for half-sib families on female side and "M" for half-sib families on the male side.
selectTop	selects highest values if true. Selects lowest values if false.
returnPop	should results be returned as a Pop-class . If FALSE, only the index of selected individuals is returned.
candidates	an optional vector of eligible selection candidates.
simParam	an object of SimParam
...	additional arguments if using a function for trait

Value

Returns an object of [Pop-class](#) or [HybridPop-class](#)

Examples

```
#Create founder haplotypes
founderPop = quickHaplo(nInd=10, nChr=1, segSites=10)

#Set simulation parameters
SP = SimParam$new(founderPop)
SP$addTraitA(10)
SP$setVarE(h2=0.5)

#Create population
pop = newPop(founderPop, simParam=SP)

#Create 3 biparental families with 10 progeny
pop2 = randCross(pop, nCrosses=3, nProgeny=10, simParam=SP)

#Select best 2 families
pop3 = selectFam(pop2, 2, simParam=SP)
```

selectInd	<i>Select individuals</i>
-----------	---------------------------

Description

Selects a subset of nInd individuals from a population.

Usage

```
selectInd(
  pop,
  nInd,
  trait = 1,
  use = "pheno",
  gender = "B",
  selectTop = TRUE,
  returnPop = TRUE,
  candidates = NULL,
  simParam = NULL,
  ...
)
```

Arguments

pop	and object of Pop-class or HybridPop-class
nInd	the number of individuals to select
trait	the trait for selection. Either a number indicating a single trait or a function returning a vector of length nInd.
use	select on genetic values "gv", estimated breeding values "ebv", breeding values "bv", phenotypes "pheno", or randomly "rand"
gender	which gender to select. Use "B" for both, "F" for females and "M" for males. If the simulation is not using gender, the argument is ignored.
selectTop	selects highest values if true. Selects lowest values if false.
returnPop	should results be returned as a Pop-class . If FALSE, only the index of selected individuals is returned.
candidates	an optional vector of eligible selection candidates.
simParam	an object of SimParam
...	additional arguments if using a function for trait

Value

Returns an object of [Pop-class](#) or [HybridPop-class](#)

Examples

```

#Create founder haplotypes
founderPop = quickHaplo(nInd=10, nChr=1, segSites=10)

#Set simulation parameters
SP = SimParam$new(founderPop)
SP$addTraitA(10)
SP$setVarE(h2=0.5)

#Create population
pop = newPop(founderPop, simParam=SP)

#Select best 5
pop2 = selectInd(pop, 5, simParam=SP)

```

selectOP

*Select open pollinating plants***Description**

This function models selection in an open pollinating plant population. It allows for varying the percentage of selfing. The function also provides an option for modeling selection as occurring before or after pollination.

Usage

```

selectOP(
  pop,
  nInd,
  nSeeds,
  probSelf = 0,
  pollenControl = FALSE,
  trait = 1,
  use = "pheno",
  selectTop = TRUE,
  candidates = NULL,
  simParam = NULL,
  ...
)

```

Arguments

pop	an object of Pop-class
nInd	the number of plants to select
nSeeds	number of seeds per plant
probSelf	percentage of seeds expected from selfing. Value ranges from 0 to 1.

pollenControl	are plants selected before pollination
trait	the trait for selection. Either a number indicating a single trait or a function returning a vector of length nInd.
use	select on genetic values "gv", estimated breeding values "ebv", breeding values "bv", phenotypes "pheno", or randomly "rand"
selectTop	selects highest values if true. Selects lowest values if false.
candidates	an optional vector of eligible selection candidates.
simParam	an object of SimParam
...	additional arguments if using a function for trait

Value

Returns an object of [Pop-class](#)

Examples

```
#Create founder haplotypes
founderPop = quickHaplo(nInd=10, nChr=1, segSites=10)

#Set simulation parameters
SP = SimParam$new(founderPop)
SP$addTraitA(10)
SP$setVarE(h2=0.5)

#Create population
pop = newPop(founderPop, simParam=SP)

#Create new population by selecting the best 3 plant
#Assuming 50% selfing in plants and 10 seeds per plant
pop2 = selectOP(pop, nInd=3, nSeeds=10, probSelf=0.5, simParam=SP)
```

selectWithinFam *Select individuals within families*

Description

Selects a subset of nInd individuals from each full-sib family within a population. Will return all individuals from a full-sib family if it has less than or equal to nInd individuals.

Usage

```
selectWithinFam(
  pop,
  nInd,
  trait = 1,
  use = "pheno",
```

```

    gender = "B",
    famType = "B",
    selectTop = TRUE,
    returnPop = TRUE,
    candidates = NULL,
    simParam = NULL,
    ...
)

```

Arguments

pop	and object of Pop-class or HybridPop-class
nInd	the number of individuals to select within a family
trait	the trait for selection. Either a number indicating a single trait or a function returning a vector of length nInd.
use	select on genetic values "gv", estimated breeding values "ebv", breeding values "bv", phenotypes "pheno", or randomly "rand"
gender	which gender to select. Use "B" for both, "F" for females and "M" for males. If the simulation is not using gender, the argument is ignored.
famType	which type of family to select. Use "B" for full-sib families, "F" for half-sib families on female side and "M" for half-sib families on the male side.
selectTop	selects highest values if true. Selects lowest values if false.
returnPop	should results be returned as a Pop-class . If FALSE, only the index of selected individuals is returned.
candidates	an optional vector of eligible selection candidates.
simParam	an object of SimParam
...	additional arguments if using a function for trait

Value

Returns an object of [Pop-class](#) or [HybridPop-class](#)

Examples

```

#Create founder haplotypes
founderPop = quickHaplo(nInd=10, nChr=1, segSites=10)

#Set simulation parameters
SP = SimParam$new(founderPop)
SP$addTraitA(10)
SP$setVarE(h2=0.5)

#Create population
pop = newPop(founderPop, simParam=SP)

#Create 3 biparental families with 10 progeny
pop2 = randCross(pop, nCrosses=3, nProgeny=10, simParam=SP)

```

```
#Select best individual per family
pop3 = selectWithinFam(pop2, 1, simParam=SP)
```

self	<i>Self individuals</i>
------	-------------------------

Description

Creates selfed progeny from each individual in a population. Only works when gender is "no".

Usage

```
self(pop, nProgeny = 1, parents = NULL, keepParents = TRUE, simParam = NULL)
```

Arguments

pop	an object of Pop-class
nProgeny	total number of selfed progeny per individual
parents	an optional vector of indices for allowable parents
keepParents	should previous parents be used for mother and father.
simParam	an object of SimParam

Value

Returns an object of [Pop-class](#)

Examples

```
#Create founder haplotypes
founderPop = quickHaplo(nInd=2, nChr=1, segSites=10)

#Set simulation parameters
SP = SimParam$new(founderPop)

#Create population
pop = newPop(founderPop, simParam=SP)

#Self pollinate each individual
pop2 = self(pop, simParam=SP)
```

selIndex	<i>Selection index</i>
----------	------------------------

Description

Calculates values of a selection index given trait values and weights. This function is intended to be used in combination with selection functions working on populations such as [selectInd](#).

Usage

```
selIndex(Y, b, scale = FALSE)
```

Arguments

Y	a matrix of trait values
b	a vector of weights
scale	should Y be scaled and centered

Examples

```
#Create founder haplotypes
founderPop = quickHaplo(nInd=10, nChr=1, segSites=10)

#Set simulation parameters
SP = SimParam$new(founderPop)
#Model two genetically correlated traits
G = 1.5*diag(2)-0.5 #Genetic correlation matrix
SP$addTraitA(10, mean=c(0,0), var=c(1,1), corA=G)
SP$setVarE(h2=c(0.5,0.5))

#Create population
pop = newPop(founderPop, simParam=SP)

#Calculate Smith-Hazel weights
econWt = c(1, 1)
b = smithHazel(econWt, varG(pop), varP(pop))

#Selection 2 best individuals using Smith-Hazel index
#selIndex is used as a trait
pop2 = selectInd(pop, nInd=2, trait=selIndex,
                 simParam=SP, b=b)
```

selInt	<i>Selection intensity</i>
--------	----------------------------

Description

Calculates the standardized selection intensity

Usage

```
selInt(p)
```

Arguments

p the proportion of individuals selected

Examples

```
selInt(0.1)
```

setEBV	<i>Set EBV</i>
--------	----------------

Description

Adds genomic estimated values to a population's EBV slot using output from a genomic selection function. The genomic estimated values can be either estimated breeding values, estimated genetic values, or estimated general combining values.

Usage

```
setEBV(  
  pop,  
  solution,  
  value = "gv",  
  targetPop = NULL,  
  append = FALSE,  
  simParam = NULL  
)
```

Arguments

pop	an object of Pop-class
solution	an object of RRsol-class
value	the genomic value to be estimated. Can be either "gv", "bv", "female", or "male".
targetPop	an optional target population that can be used when value is "bv", "female", or "male". When supplied, the allele frequency in the targetPop is used to set these values.
append	should estimated values be appended to existing data in the EBV slot. If TRUE, a new column is added. If FALSE, existing data is replaced with the new estimates.
simParam	an object of SimParam

Value

Returns an object of [Pop-class](#)

Examples

```
#Create founder haplotypes
founderPop = quickHaplo(nInd=10, nChr=1, segSites=20)

#Set simulation parameters
SP = SimParam$new(founderPop)
SP$addTraitA(10)
SP$setVarE(h2=0.5)
SP$addSnpChip(10)

#Create population
pop = newPop(founderPop, simParam=SP)

#Run GS model and set EBV
ans = RRBLUP(pop, simParam=SP)
pop = setEBV(pop, ans, simParam=SP)

#Evaluate accuracy
cor(gv(pop), ebv(pop))
```

setPheno

Set phenotypes

Description

Sets phenotypes for all traits by adding random error from a multivariate normal distribution.

Usage

```

setPheno(
  pop,
  varE = NULL,
  reps = 1,
  fixEff = 1L,
  p = NULL,
  onlyPheno = FALSE,
  simParam = NULL
)

```

Arguments

pop	an object of Pop-class or HybridPop-class
varE	error variances for phenotype. A vector of length nTraits for independent error or a square matrix of dimensions nTraits for correlated errors. If NULL, value in simParam is used.
reps	number of replications for phenotype. See details.
fixEff	fixed effect to assign to the population. Used by genomic selection models only.
p	the p-value for the environmental covariate used by GxE traits. If NULL, a value is sampled at random.
onlyPheno	should only the phenotype be returned, see return
simParam	an object of SimParam

Details

The reps parameter is for convenient representation of replicated data. It is intended to represent replicated yield trials in plant breeding programs. In this case, varE is set to the plot error and reps is set to the number of plots per entry. The resulting phenotype represents entry means.

Value

Returns an object of [Pop-class](#) or [HybridPop-class](#) if onlyPheno=FALSE, if onlyPheno=TRUE a matrix is returned

Examples

```

#Create founder haplotypes
founderPop = quickHaplo(nInd=10, nChr=1, segSites=10)

#Set simulation parameters
SP = SimParam$new(founderPop)
SP$addTraitA(10)

#Create population
pop = newPop(founderPop, simParam=SP)

#Add phenotype with error variance of 1

```

```
pop = setPheno(pop, varE=1)
```

setPhenoGCA *Set GCA as phenotype*

Description

Calculates general combining ability from a set of testers and returns these values as phenotypes for a population.

Usage

```
setPhenoGCA(
  pop,
  testers,
  use = "pheno",
  varE = NULL,
  reps = 1,
  fixEff = 1L,
  p = 0.5,
  inbred = FALSE,
  onlyPheno = FALSE,
  simParam = NULL
)
```

Arguments

pop	an object of Pop-class
testers	an object of Pop-class
use	true genetic value (gv) or phenotypes (pheno, default)
varE	error variances for phenotype if use="pheno". A vector of length nTraits for independent error or a square matrix of dimensions nTraits for correlated errors.
reps	number of replications for phenotype. See details.
fixEff	fixed effect to assign to the population. Used by genomic selection models only.
p	the p-value for the environmental covariate
inbred	are both pop and testers fully inbred. They are only fully inbred if created by newPop using inbred founders or by the makeDH function
onlyPheno	should only the phenotype be returned, see return
simParam	an object of SimParam

Details

The reps parameter is for convenient representation of replicated data. It was intended for representation of replicated yield trials in plant breeding programs. In this case, varE is set to the plot error and reps is set to the number plots per entry. The resulting phenotype would reflect the mean of all replications.

Value

Returns an object of `Pop-class` or a matrix if `onlyPheno=TRUE`

Examples

```
#Create founder haplotypes
founderPop = quickHaplo(nInd=10, nChr=1, segSites=10, inbred=TRUE)

#Set simulation parameters
SP = SimParam$new(founderPop)
SP$addTraitA(10)

#Create population
pop = newPop(founderPop, simParam=SP)

#Set phenotype to average per
pop2 = setPhenoGCA(pop, pop, use="gv", inbred=TRUE, simParam=SP)
```

SimParam

Simulation parameters

Description

Container for global simulation parameters. Saving this object as SP will allow it to be accessed by function defaults.

Public fields

`nThreads` number of threads used on platforms with OpenMP support
`snpChips` list of SNP chips
`invalidQtl` list of segregating sites that aren't valid QTL
`invalidSnp` list of segregating sites that aren't valid SNP
`founderPop` founder population used for variance scaling
`v` the crossover interference parameter for a gamma model of recombination. A value of 1 indicates no crossover interference (e.g. Haldane mapping function). A value of 2.6 approximates the degree of crossover interference implied by the Kosambi mapping function. (default is 1)
`quadProb` the probability of quadrivalent pairing in an autopolyploid. (default is 0)

Active bindings

`traits` list of traits
`nChr` number of chromosomes
`nTraits` number of traits
`nSnpChips` number of SNP chips

segSites segregating sites per chromosome
gender is gender used for mating
sepMap are there separate genetic maps for males and females
genMap "matrix" of chromosome genetic maps
femaleMap "matrix" of chromosome genetic maps for females
maleMap "matrix" of chromosome genetic maps for males
centromere position of centromeres genetic map
femaleCentromere position of centromeres on female genetic map
maleCentromere position of centromeres on male genetic map
lastId last ID number assigned
isTrackPed is pedigree being tracked
pedigree pedigree matrix for all individuals
isTrackRec is recombination being tracked
rechist list of historic recombination events
varA additive genetic variance in founderPop
varG total genetic variance in founderPop
varE default error variance
version the version of AlphaSimR used to generate this object

Methods

Public methods:

- [SimParam\\$new\(\)](#)
- [SimParam\\$setTrackPed\(\)](#)
- [SimParam\\$setTrackRec\(\)](#)
- [SimParam\\$resetPed\(\)](#)
- [SimParam\\$restrSegSites\(\)](#)
- [SimParam\\$setGender\(\)](#)
- [SimParam\\$addSnpChip\(\)](#)
- [SimParam\\$addStructuredSnpChip\(\)](#)
- [SimParam\\$addTraitA\(\)](#)
- [SimParam\\$addTraitAD\(\)](#)
- [SimParam\\$addTraitAG\(\)](#)
- [SimParam\\$addTraitADG\(\)](#)
- [SimParam\\$addTraitAE\(\)](#)
- [SimParam\\$addTraitADE\(\)](#)
- [SimParam\\$addTraitAEG\(\)](#)
- [SimParam\\$addTraitADEG\(\)](#)
- [SimParam\\$manAddTrait\(\)](#)
- [SimParam\\$switchTrait\(\)](#)

- `SimParam$removeTrait()`
- `SimParam$setVarE()`
- `SimParam$setCorE()`
- `SimParam$rescaleTraits()`
- `SimParam$setRecombRatio()`
- `SimParam$switchGenMap()`
- `SimParam$switchFemaleMap()`
- `SimParam$switchMaleMap()`
- `SimParam$addToRec()`
- `SimParam$updateLastId()`
- `SimParam$addToPed()`
- `SimParam$clone()`

Method `new()`: Starts the process of building a new simulation by creating a new `SimParam` object and assigning a founder population to the class. It is recommended that you save the object with the name "SP", because subsequent functions will check your global environment for an object of this name if their `simParam` arguments are `NULL`. This allows you to call these functions without explicitly supplying a `simParam` argument with every call.

Usage:

```
SimParam$new(founderPop)
```

Arguments:

`founderPop` an object of `MapPop-class`

Examples:

```
#Create founder haplotypes
founderPop = quickHaplo(nInd=10, nChr=1, segSites=10)

#Set simulation parameters
SP = SimParam$new(founderPop)
```

Method `setTrackPed()`: Sets pedigree tracking for the simulation. By default pedigree tracking is turned off. When turned on, the pedigree of all individuals created will be tracked, except those created by `hybridCross`. Turning off pedigree tracking will turn off recombination tracking if it is turned on.

Usage:

```
SimParam$setTrackPed(isTrackPed, force = FALSE)
```

Arguments:

`isTrackPed` should pedigree tracking be on.

`force` should the check for a running simulation be ignored. Only set to `TRUE` if you know what you are doing.

Examples:

```
#Create founder haplotypes
founderPop = quickHaplo(nInd=10, nChr=1, segSites=10)

#Set simulation parameters
SP = SimParam$new(founderPop)
SP$setTrackPed(TRUE)
```


Method setTrackRec(): Sets recombination tracking for the simulation. By default recombination tracking is turned off. When turned on recombination tracking will also turn on pedigree tracking. Recombination tracking keeps records of all individuals created, except those created by `hybridCross`, because their pedigree is not tracked.

Usage:

```
SimParam$setTrackRec(isTrackRec, force = FALSE)
```

Arguments:

`isTrackRec` should recombination tracking be on.

`force` should the check for a running simulation be ignored. Only set to TRUE if you know what you are doing.

Examples:

```
#Create founder haplotypes
founderPop = quickHaplo(nInd=10, nChr=1, segSites=10)

#Set simulation parameters
SP = SimParam$new(founderPop)
SP$setTrackRec(TRUE)
```

Method resetPed(): Resets the internal `lastId`, the pedigree and recombination tracking (if in use) to the supplied `lastId`. Be careful using this function because it may introduce a bug if you use individuals from the deleted portion of the pedigree.

Usage:

```
SimParam$resetPed(lastId = 0L)
```

Arguments:

`lastId` last ID to include in pedigree

Examples:

```
#Create founder haplotypes
founderPop = quickHaplo(nInd=10, nChr=1, segSites=10)

#Set simulation parameters
SP = SimParam$new(founderPop)

#Create population
pop = newPop(founderPop, simParam=SP)
pop@id # 1:10

#Create another population after resetting pedigree
SP$resetPed()
pop2 = newPop(founderPop, simParam=SP)
pop2@id # 1:10
```

Method restrSegSites(): Sets restrictions on which segregating sites can serve as SNP and/or QTL.

Usage:

```

SimParam$restrSegSites(
  minQtlPerChr = NULL,
  minSnpPerChr = NULL,
  overlap = FALSE,
  minSnpFreq = NULL
)

```

Arguments:

`minQtlPerChr` the minimum number of `segSites` for QTLs. Can be a single value or a vector values for each chromosome.

`minSnpPerChr` the minimum number of `segSites` for SNPs. Can be a single value or a vector values for each chromosome.

`overlap` should SNP and QTL sites be allowed to overlap.

`minSnpFreq` minimum allowable frequency for SNP loci. No minimum SNP frequency is used if value is NULL.

Examples:

```

#Create founder haplotypes
founderPop = quickHaplo(nInd=10, nChr=1, segSites=10)

#Set simulation parameters
SP = SimParam$new(founderPop)
SP$restrSegSites(minQtlPerChr=5, minSnpPerChr=5)

```

Method `setGender()`: Changes how gender is used in the simulation. The default gender of a simulation is "no". To add gender to the simulation, run this function with "yes_sys" or "yes_rand". The value "yes_sys" will systematically assign gender to newly created individuals as first male, then female. Thus, odd numbers of individuals will have one more male than female. The value "yes_rand" will randomly assign gender to individuals.

Usage:

```

SimParam$setGender(gender, force = FALSE)

```

Arguments:

`gender` acceptable value are "no", "yes_sys", or "yes_rand"

`force` should the check for a running simulation be ignored. Only set to TRUE if you know what you are doing.

Examples:

```

#Create founder haplotypes
founderPop = quickHaplo(nInd=10, nChr=1, segSites=10)

#Set simulation parameters
SP = SimParam$new(founderPop)
SP$setGender("yes_sys")

```

Method `addSnpChip()`: Randomly assigns eligible SNPs to a SNP chip

Usage:

```

SimParam$addSnpChip(nSnpPerChr, minSnpFreq = NULL, refPop = NULL)

```

Arguments:

nSnPPerChr number of SNPs per chromosome. Can be a single value or nChr values.
 minSnPFreq minimum allowable frequency for SNP loci. If NULL, no minimum frequency is used.
 refPop reference population for calculating SNP frequency. If NULL, the founder population is used.

Examples:

```
#Create founder haplotypes
founderPop = quickHaplo(nInd=10, nChr=1, segSites=10)

#Set simulation parameters
SP = SimParam$new(founderPop)
SP$addSnPChip(10)
```

Method addStructuredSnPChip(): Randomly selects the number of snps in structure and then assigns them to chips based on structure

Usage:

```
SimParam$addStructuredSnPChip(nSnPPerChr, structure, force = FALSE)
```

Arguments:

nSnPPerChr number of SNPs per chromosome. Can be a single value or nChr values.
 structure a matrix. Rows are snp chips, columns are chips. If value is true then that snp is on that chip.
 force should the check for a running simulation be ignored. Only set to TRUE if you know what you are doing.

Method addTraitA(): Randomly assigns eligible QTLs for one or more additive traits. If simulating more than one trait, all traits will be pleiotrophic with correlated additive effects.

Usage:

```
SimParam$addTraitA(
  nQtlPerChr,
  mean = 0,
  var = 1,
  corA = NULL,
  gamma = FALSE,
  shape = 1,
  force = FALSE
)
```

Arguments:

nQtlPerChr number of QTLs per chromosome. Can be a single value or nChr values.
 mean a vector of desired mean genetic values for one or more traits
 var a vector of desired genetic variances for one or more traits
 corA a matrix of correlations between additive effects
 gamma should a gamma distribution be used instead of normal
 shape the shape parameter for the gamma distribution
 force should the check for a running simulation be ignored. Only set to TRUE if you know what you are doing.

Examples:

```
#Create founder haplotypes
founderPop = quickHaplo(nInd=10, nChr=1, segSites=10)

#Set simulation parameters
SP = SimParam$new(founderPop)
SP$addTraitA(10)
```

Method `addTraitAD()`: Randomly assigns eligible QTLs for one or more traits with dominance. If simulating more than one trait, all traits will be pleiotropic with correlated effects.

Usage:

```
SimParam$addTraitAD(
  nQtlPerChr,
  mean = 0,
  var = 1,
  meanDD = 0,
  varDD = 0,
  corA = NULL,
  corDD = NULL,
  useVarA = TRUE,
  gamma = FALSE,
  shape = 1,
  force = FALSE
)
```

Arguments:

`nQtlPerChr` number of QTLs per chromosome. Can be a single value or `nChr` values.
`mean` a vector of desired mean genetic values for one or more traits
`var` a vector of desired genetic variances for one or more traits
`meanDD` mean dominance degree
`varDD` variance of dominance degree
`corA` a matrix of correlations between additive effects
`corDD` a matrix of correlations between dominance degrees
`useVarA` tune according to additive genetic variance if true. If FALSE, tuning is performed according to total genetic variance.
`gamma` should a gamma distribution be used instead of normal
`shape` the shape parameter for the gamma distribution
`force` should the check for a running simulation be ignored. Only set to TRUE if you know what you are doing.

Examples:

```
#Create founder haplotypes
founderPop = quickHaplo(nInd=10, nChr=1, segSites=10)

#Set simulation parameters
SP = SimParam$new(founderPop)
SP$addTraitAD(10, meanDD=0.5)
```

Method addTraitAG(): Randomly assigns eligible QTLs for one or more additive GxE traits. If simulating more than one trait, all traits will be pleiotropic with correlated effects.

Usage:

```
SimParam$addTraitAG(
  nQtlPerChr,
  mean = 0,
  var = 1,
  varGxE = 1e-06,
  varEnv = 0,
  corA = NULL,
  corGxE = NULL,
  gamma = FALSE,
  shape = 1,
  force = FALSE
)
```

Arguments:

`nQtlPerChr` number of QTLs per chromosome. Can be a single value or `nChr` values.
`mean` a vector of desired mean genetic values for one or more traits
`var` a vector of desired genetic variances for one or more traits
`varGxE` a vector of total genotype-by-environment variances for the traits
`varEnv` a vector of environmental variances for one or more traits
`corA` a matrix of correlations between additive effects
`corGxE` a matrix of correlations between GxE effects
`gamma` should a gamma distribution be used instead of normal
`shape` the shape parameter for the gamma distribution
`force` should the check for a running simulation be ignored. Only set to TRUE if you know what you are doing.

Examples:

```
#Create founder haplotypes
founderPop = quickHaplo(nInd=10, nChr=1, segSites=10)

#Set simulation parameters
SP = SimParam$new(founderPop)
SP$addTraitAG(10, varGxE=2)
```

Method addTraitADG(): Randomly assigns eligible QTLs for a trait with dominance and GxE.

Usage:

```
SimParam$addTraitADG(
  nQtlPerChr,
  mean = 0,
  var = 1,
  varEnv = 1e-06,
  varGxE = 1e-06,
  meanDD = 0,
  varDD = 0,
```

```

corA = NULL,
corDD = NULL,
corGxE = NULL,
useVarA = TRUE,
gamma = FALSE,
shape = 1,
force = FALSE
)

```

Arguments:

nQt1PerChr number of QTLs per chromosome. Can be a single value or nChr values.
 mean a vector of desired mean genetic values for one or more traits
 var a vector of desired genetic variances for one or more traits
 varEnv a vector of environmental variances for one or more traits
 varGxE a vector of total genotype-by-environment variances for the traits
 meanDD mean dominance degree
 varDD variance of dominance degree
 corA a matrix of correlations between additive effects
 corDD a matrix of correlations between dominance degrees
 corGxE a matrix of correlations between GxE effects
 useVarA tune according to additive genetic variance if true
 gamma should a gamma distribution be used instead of normal
 shape the shape parameter for the gamma distribution
 force should the check for a running simulation be ignored. Only set to TRUE if you know what you are doing.

Examples:

```

#Create founder haplotypes
founderPop = quickHaplo(nInd=10, nChr=1, segSites=10)

#Set simulation parameters
SP = SimParam$new(founderPop)
SP$addTraitADG(10, meanDD=0.5, varGxE=2)

```

Method addTraitAE(): Randomly assigns eligible QTLs for one or more additive and epistasis traits. If simulating more than one trait, all traits will be pleiotrophic with correlated additive effects.

Usage:

```

SimParam$addTraitAE(
  nQt1PerChr,
  mean = 0,
  var = 1,
  relAA = 0,
  corA = NULL,
  corAA = NULL,
  useVarA = TRUE,
  gamma = FALSE,
)

```

```

    shape = 1,
    force = FALSE
)

```

Arguments:

nQt1PerChr number of QTLs per chromosome. Can be a single value or nChr values.
 mean a vector of desired mean genetic values for one or more traits
 var a vector of desired genetic variances for one or more traits
 relAA the relative value of additive-by-additive variance compared to additive variance in a diploid organism with allele frequency 0.5
 corA a matrix of correlations between additive effects
 corAA a matrix of correlations between additive-by-additive effects
 useVarA tune according to additive genetic variance if true. If FALSE, tuning is performed according to total genetic variance.
 gamma should a gamma distribution be used instead of normal
 shape the shape parameter for the gamma distribution
 force should the check for a running simulation be ignored. Only set to TRUE if you know what you are doing.

Examples:

```

#Create founder haplotypes
founderPop = quickHaplo(nInd=10, nChr=1, segSites=10)

```

Method addTraitADE(): Randomly assigns eligible QTLs for one or more traits with dominance and epistasis. If simulating more than one trait, all traits will be pleiotrophic with correlated effects.

Usage:

```

SimParam$addTraitADE(
  nQt1PerChr,
  mean = 0,
  var = 1,
  meanDD = 0,
  varDD = 0,
  relAA = 0,
  corA = NULL,
  corDD = NULL,
  corAA = NULL,
  useVarA = TRUE,
  gamma = FALSE,
  shape = 1,
  force = FALSE
)

```

Arguments:

nQt1PerChr number of QTLs per chromosome. Can be a single value or nChr values.
 mean a vector of desired mean genetic values for one or more traits
 var a vector of desired genetic variances for one or more traits
 meanDD mean dominance degree

varDD variance of dominance degree
 relAA the relative value of additive-by-additive variance compared to additive variance in a diploid organism with allele frequency 0.5
 corA a matrix of correlations between additive effects
 corDD a matrix of correlations between dominance degrees
 corAA a matrix of correlations between additive-by-additive effects
 useVarA tune according to additive genetic variance if true. If FALSE, tuning is performed according to total genetic variance.
 gamma should a gamma distribution be used instead of normal
 shape the shape parameter for the gamma distribution
 force should the check for a running simulation be ignored. Only set to TRUE if you know what you are doing.

Examples:

```

#Create founder haplotypes
founderPop = quickHaplo(nInd=10, nChr=1, segSites=10)

#Set simulation parameters
SP = SimParam$new(founderPop)
SP$addTraitADE(10)
  
```

Method addTraitAEG(): Randomly assigns eligible QTLs for one or more additive and epistasis GxE traits. If simulating more than one trait, all traits will be pleiotrophic with correlated effects.

Usage:

```

SimParam$addTraitAEG(
  nQt1PerChr,
  mean = 0,
  var = 1,
  relAA = 0,
  varGxE = 1e-06,
  varEnv = 0,
  corA = NULL,
  corAA = NULL,
  corGxE = NULL,
  useVarA = TRUE,
  gamma = FALSE,
  shape = 1,
  force = FALSE
)
  
```

Arguments:

nQt1PerChr number of QTLs per chromosome. Can be a single value or nChr values.
 mean a vector of desired mean genetic values for one or more traits
 var a vector of desired genetic variances for one or more traits
 relAA the relative value of additive-by-additive variance compared to additive variance in a diploid organism with allele frequency 0.5
 varGxE a vector of total genotype-by-environment variances for the traits

varEnv a vector of environmental variances for one or more traits
 corA a matrix of correlations between additive effects
 corAA a matrix of correlations between additive-by-additive effects
 corGxE a matrix of correlations between GxE effects
 useVarA tune according to additive genetic variance if true. If FALSE, tuning is performed according to total genetic variance.
 gamma should a gamma distribution be used instead of normal
 shape the shape parameter for the gamma distribution
 force should the check for a running simulation be ignored. Only set to TRUE if you know what you are doing.

Examples:

```

#Create founder haplotypes
founderPop = quickHaplo(nInd=10, nChr=1, segSites=10)

#Set simulation parameters
SP = SimParam$new(founderPop)
SP$addTraitAEG(10, varGxE=2)

```

Method addTraitAEG(): Randomly assigns eligible QTLs for a trait with dominance, epistasis and GxE.

Usage:

```

SimParam$addTraitAEG(
  nQtlPerChr,
  mean = 0,
  var = 1,
  varEnv = 1e-06,
  varGxE = 1e-06,
  meanDD = 0,
  varDD = 0,
  relAA = 0,
  corA = NULL,
  corDD = NULL,
  corAA = NULL,
  corGxE = NULL,
  useVarA = TRUE,
  gamma = FALSE,
  shape = 1,
  force = FALSE
)

```

Arguments:

nQtlPerChr number of QTLs per chromosome. Can be a single value or nChr values.
 mean a vector of desired mean genetic values for one or more traits
 var a vector of desired genetic variances for one or more traits
 varEnv a vector of environmental variances for one or more traits
 varGxE a vector of total genotype-by-environment variances for the traits

meanDD mean dominance degree
 varDD variance of dominance degree
 relAA the relative value of additive-by-additive variance compared to additive variance in a
 diploid organism with allele frequency 0.5
 corA a matrix of correlations between additive effects
 corDD a matrix of correlations between dominance degrees
 corAA a matrix of correlations between additive-by-additive effects
 corGxE a matrix of correlations between GxE effects
 useVarA tune according to additive genetic variance if true
 gamma should a gamma distribution be used instead of normal
 shape the shape parameter for the gamma distribution
 force should the check for a running simulation be ignored. Only set to TRUE if you know
 what you are doing.

Examples:

```

#Create founder haplotypes
founderPop = quickHaplo(nInd=10, nChr=1, segSites=10)

#Set simulation parameters
SP = SimParam$new(founderPop)
SP$addTraitADEG(10, meanDD=0.5, varGxE=2)
  
```

Method `manAddTrait()`: Manually add a new trait to the simulation.

Usage:

```

SimParam$manAddTrait(
  lociMap,
  varA = NA_real_,
  varG = NA_real_,
  varE = NA_real_,
  force = FALSE
)
  
```

Arguments:

lociMap a new object descended from [LociMap-class](#)
 varA the value for varA in the base population, optional
 varG the value for varG in the base population, optional
 varE default error variance for phenotype, optional
 force should the check for a running simulation be ignored. Only set to TRUE if you know
 what you are doing

Method `switchTrait()`: Switch a trait in the simulation.

Usage:

```

SimParam$switchTrait(
  traitPos,
  lociMap,
  varA = NA_real_,
  varG = NA_real_,
  )
  
```

```

    varE = NA_real_,
    force = FALSE
)

```

Arguments:

traitPos an integer indicate which trait to switch
 lociMap a new object descended from [LociMap-class](#)
 varA the value for varA in the base population, optional
 varG the value for varG in the base population, optional
 varE default error variance for phenotype, optional
 force should the check for a running simulation be ignored. Only set to TRUE if you know what you are doing

Method `removeTrait()`: Remove a trait from the simulation

Usage:

```
SimParam$removeTrait(traits, force = FALSE)
```

Arguments:

traits an integer vector indicating which traits to remove
 force should the check for a running simulation be ignored. Only set to TRUE if you know what you are doing

Method `setVarE()`: Defines a default value for error variances in the simulation.

Usage:

```
SimParam$setVarE(h2 = NULL, H2 = NULL, varE = NULL)
```

Arguments:

h2 a vector of desired narrow-sense heritabilities
 H2 a vector of desired broad-sense heritabilities
 varE a vector of error variances

Examples:

```

#Create founder haplotypes
founderPop = quickHaplo(nInd=10, nChr=1, segSites=10)

#Set simulation parameters
SP = SimParam$new(founderPop)
SP$addTraitA(10)
SP$setVarE(h2=0.5)

```

Method `setCorE()`: Defines a correlation structure for default error variances. You must call `setVarE` first to define the default error variances.

Usage:

```
SimParam$setCorE(corE)
```

Arguments:

corE a correlation matrix for the error variances

Examples:

```
#Create founder haplotypes
founderPop = quickHaplo(nInd=10, nChr=1, segSites=10)

#Set simulation parameters
SP = SimParam$new(founderPop)
SP$addTraitA(10, mean=c(0,0), var=c(1,1), corA=diag(2))
SP$setVarE(varE=c(1,1))
E = 0.5*diag(2)+0.5 #Positively correlated error
SP$setCorE(E)
```

Method `rescaleTraits()`: Linearly scales all traits to achieve desired values of means and variances in the founder population.

Usage:

```
SimParam$rescaleTraits(
  mean = 0,
  var = 1,
  varEnv = 0,
  varGxE = 1e-06,
  useVarA = TRUE
)
```

Arguments:

`mean` a vector of new trait means
`var` a vector of new trait variances
`varEnv` a vector of new environmental variances
`varGxE` a vector of new GxE variances
`useVarA` tune according to additive genetic variance if true

Examples:

```
#Create founder haplotypes
founderPop = quickHaplo(nInd=10, nChr=1, segSites=10)

#Set simulation parameters
SP = SimParam$new(founderPop)
SP$addTraitA(10)

#Create population
pop = newPop(founderPop, simParam=SP)
meanG(pop)

#Change mean to 1
SP$rescaleTraits(mean=1)
#Run resetPop for change to take effect
pop = resetPop(pop, simParam=SP)
meanG(pop)
```

Method `setRecombRatio()`: Set the relative recombination rates between males and females. This allows for gender specific recombination rates, under the assumption of equivalent recombination landscapes.

Usage:

```
SimParam$setRecombRatio(femaleRatio)
```

Arguments:

femaleRatio relative ratio of recombination in females compared to males. A value of 2 indicate twice as much recombination in females. The value must be greater than 0. (default is 1)

Examples:

```
#Create founder haplotypes
founderPop = quickHaplo(nInd=10, nChr=1, segSites=10)

#Set simulation parameters
SP = SimParam$new(founderPop)
SP$setRecombRatio(2) #Twice as much recombination in females
```

Method `switchGenMap()`: Replaces existing genetic map.

Usage:

```
SimParam$switchGenMap(genMap, centromere = NULL)
```

Arguments:

genMap a list of length *nChr* containing numeric vectors for the position of each segregating site on a chromosome.

centromere a numeric vector of centromere positions. If NULL, the centromere are assumed to be metacentric.

Method `switchFemaleMap()`: Replaces existing female genetic map.

Usage:

```
SimParam$switchFemaleMap(genMap, centromere = NULL)
```

Arguments:

genMap a list of length *nChr* containing numeric vectors for the position of each segregating site on a chromosome.

centromere a numeric vector of centromere positions. If NULL, the centromere are assumed to be metacentric.

Method `switchMaleMap()`: Replaces existing male genetic map.

Usage:

```
SimParam$switchMaleMap(genMap, centromere = NULL)
```

Arguments:

genMap a list of length *nChr* containing numeric vectors for the position of each segregating site on a chromosome.

centromere a numeric vector of centromere positions. If NULL, the centromere are assumed to be metacentric.

Method `addToRec()`: For internal use only.

Usage:

```
SimParam$addToRec(hist)
```

Arguments:

hist new recombination history

Method updateLastId(): For internal use only.

Usage:

SimParam\$updateLastId(lastId)

Arguments:

lastId last ID assigned

Method addToPed(): For internal use only.

Usage:

SimParam\$addToPed(lastId, mother, father, isDH)

Arguments:

lastId ID of last individual
 mother vector of mother IDs
 father vector of father IDs
 isDH vector of DH indicators

Method clone(): The objects of this class are cloneable with this method.

Usage:

SimParam\$clone(deep = FALSE)

Arguments:

deep Whether to make a deep clone.

Note

By default the founder population is the population used to initialize the SimParam object. This population can be changed by replacing the population in the founderPop slot. You must run [resetPop](#) on any existing populations to obtain the new trait values.

Examples

```
## -----
## Method `SimParam$new`
## -----

#Create founder haplotypes
founderPop = quickHaplo(nInd=10, nChr=1, segSites=10)

#Set simulation parameters
SP = SimParam$new(founderPop)

## -----
## Method `SimParam$setTrackPed`
## -----
```

```

#Create founder haplotypes
founderPop = quickHaplo(nInd=10, nChr=1, segSites=10)

#Set simulation parameters
SP = SimParam$new(founderPop)
SP$setTrackPed(TRUE)

## -----
## Method `SimParam$setTrackRec`
## -----

#Create founder haplotypes
founderPop = quickHaplo(nInd=10, nChr=1, segSites=10)

#Set simulation parameters
SP = SimParam$new(founderPop)
SP$setTrackRec(TRUE)

## -----
## Method `SimParam$resetPed`
## -----

#Create founder haplotypes
founderPop = quickHaplo(nInd=10, nChr=1, segSites=10)

#Set simulation parameters
SP = SimParam$new(founderPop)

#Create population
pop = newPop(founderPop, simParam=SP)
pop@id # 1:10

#Create another population after resetting pedigree
SP$resetPed()
pop2 = newPop(founderPop, simParam=SP)
pop2@id # 1:10

## -----
## Method `SimParam$restrSegSites`
## -----

#Create founder haplotypes
founderPop = quickHaplo(nInd=10, nChr=1, segSites=10)

#Set simulation parameters
SP = SimParam$new(founderPop)
SP$restrSegSites(minQt1PerChr=5, minSnpPerChr=5)

## -----
## Method `SimParam$setGender`
## -----

#Create founder haplotypes

```

```

founderPop = quickHaplo(nInd=10, nChr=1, segSites=10)

#Set simulation parameters
SP = SimParam$new(founderPop)
SP$setGender("yes_sys")

## -----
## Method `SimParam$addSnpChip`
## -----

#Create founder haplotypes
founderPop = quickHaplo(nInd=10, nChr=1, segSites=10)

#Set simulation parameters
SP = SimParam$new(founderPop)
SP$addSnpChip(10)

## -----
## Method `SimParam$addTraitA`
## -----

#Create founder haplotypes
founderPop = quickHaplo(nInd=10, nChr=1, segSites=10)

#Set simulation parameters
SP = SimParam$new(founderPop)
SP$addTraitA(10)

## -----
## Method `SimParam$addTraitAD`
## -----

#Create founder haplotypes
founderPop = quickHaplo(nInd=10, nChr=1, segSites=10)

#Set simulation parameters
SP = SimParam$new(founderPop)
SP$addTraitAD(10, meanDD=0.5)

## -----
## Method `SimParam$addTraitAG`
## -----

#Create founder haplotypes
founderPop = quickHaplo(nInd=10, nChr=1, segSites=10)

#Set simulation parameters
SP = SimParam$new(founderPop)
SP$addTraitAG(10, varGxE=2)

## -----
## Method `SimParam$addTraitADG`
## -----

```



```

#Create founder haplotypes
founderPop = quickHaplo(nInd=10, nChr=1, segSites=10)

#Set simulation parameters
SP = SimParam$new(founderPop)
SP$addTraitADG(10, meanDD=0.5, varGxE=2)

## -----
## Method `SimParam$addTraitAE`
## -----

#Create founder haplotypes
founderPop = quickHaplo(nInd=10, nChr=1, segSites=10)

## -----
## Method `SimParam$addTraitADE`
## -----

#Create founder haplotypes
founderPop = quickHaplo(nInd=10, nChr=1, segSites=10)

#Set simulation parameters
SP = SimParam$new(founderPop)
SP$addTraitADE(10)

## -----
## Method `SimParam$addTraitAEG`
## -----

#Create founder haplotypes
founderPop = quickHaplo(nInd=10, nChr=1, segSites=10)

#Set simulation parameters
SP = SimParam$new(founderPop)
SP$addTraitAEG(10, varGxE=2)

## -----
## Method `SimParam$addTraitADEG`
## -----

#Create founder haplotypes
founderPop = quickHaplo(nInd=10, nChr=1, segSites=10)

#Set simulation parameters
SP = SimParam$new(founderPop)
SP$addTraitADEG(10, meanDD=0.5, varGxE=2)

## -----
## Method `SimParam$setVarE`
## -----

#Create founder haplotypes

```

```

founderPop = quickHaplo(nInd=10, nChr=1, segSites=10)

#Set simulation parameters
SP = SimParam$new(founderPop)
SP$addTraitA(10)
SP$setVarE(h2=0.5)

## -----
## Method `SimParam$setCorE`
## -----

#Create founder haplotypes
founderPop = quickHaplo(nInd=10, nChr=1, segSites=10)

#Set simulation parameters
SP = SimParam$new(founderPop)
SP$addTraitA(10, mean=c(0,0), var=c(1,1), corA=diag(2))
SP$setVarE(varE=c(1,1))
E = 0.5*diag(2)+0.5 #Positively correlated error
SP$setCorE(E)

## -----
## Method `SimParam$rescaleTraits`
## -----

#Create founder haplotypes
founderPop = quickHaplo(nInd=10, nChr=1, segSites=10)

#Set simulation parameters
SP = SimParam$new(founderPop)
SP$addTraitA(10)

#Create population
pop = newPop(founderPop, simParam=SP)
meanG(pop)

#Change mean to 1
SP$rescaleTraits(mean=1)
#Run resetPop for change to take effect
pop = resetPop(pop, simParam=SP)
meanG(pop)

## -----
## Method `SimParam$setRecombRatio`
## -----

#Create founder haplotypes
founderPop = quickHaplo(nInd=10, nChr=1, segSites=10)

#Set simulation parameters
SP = SimParam$new(founderPop)
SP$setRecombRatio(2) #Twice as much recombination in females

```

smithHazel	<i>Calculate Smith-Hazel weights</i>
------------	--------------------------------------

Description

Calculates weights for Smith-Hazel index given economic weights and phenotypic and genotypic variance-covariance matrices.

Usage

```
smithHazel(econWt, varG, varP)
```

Arguments

econWt	vector of economic weights
varG	the genetic variance-covariance matrix
varP	the phenotypic variance-covariance matrix

Value

a vector of weight for calculating index values

Examples

```
G = 1.5*diag(2)-0.5
E = diag(2)
P = G+E
wt = c(1,1)
smithHazel(wt, G, P)
```

TraitA-class	<i>Additive trait</i>
--------------	-----------------------

Description

Extends [LociMap-class](#) to model additive traits

Slots

addEff	additive effects
intercept	adjustment factor for gv

TraitA2-class	<i>Gender specific additive trait</i>
---------------	---------------------------------------

Description

Extends [TraitA-class](#) to model separate additive effects for parent of origin. Used exclusively for genomic selection.

Slots

addEffMale additive effects

TraitA2D-class	<i>Gender specific additive and dominance trait</i>
----------------	-----------------------------------------------------

Description

Extends [TraitA2-class](#) to add dominance

Slots

domEff dominance effects

TraitAD-class	<i>Additive and dominance trait</i>
---------------	-------------------------------------

Description

Extends [TraitA-class](#) to add dominance

Slots

domEff dominance effects

TraitADE-class	<i>Additive, dominance, and epistatic trait</i>
----------------	-------------------------------------------------

Description

Extends [TraitAD-class](#) to add epistasis

Slots

epiEff epistatic effects

TraitADEG-class	<i>Additive, dominance, epistasis, and GxE trait</i>
-----------------	------------------------------------------------------

Description

Extends [TraitADE-class](#) to add GxE effects

Slots

gxEff GxE effects
gxEInt GxE intercept
envVar Environmental variance

TraitADG-class	<i>Additive, dominance and GxE trait</i>
----------------	------------------------------------------

Description

Extends [TraitAD-class](#) to add GxE effects

Slots

gxEff GxE effects
gxEInt GxE intercept
envVar Environmental variance

TraitAE-class	<i>Additive and epistatic trait</i>
---------------	-------------------------------------

Description

Extends [TraitA-class](#) to add epistasis

Slots

epiEff epistatic effects

TraitAEG-class	<i>Additive, epistasis and GxE trait</i>
----------------	------------------------------------------

Description

Extends [TraitAE-class](#) to add GxE effects

Slots

gxeEff GxE effects
 gxeInt GxE intercept
 envVar Environmental variance

TraitAG-class	<i>Additive and GxE trait</i>
---------------	-------------------------------

Description

Extends [TraitA-class](#) to add GxE effects

Slots

gxeEff GxE effects
 gxeInt GxE intercept
 envVar Environmental variance

usefulness	<i>Usefulness criterion</i>
------------	-----------------------------

Description

Calculates the usefulness criterion

Usage

```
usefulness(
  pop,
  trait = 1,
  use = "gv",
  p = 0.1,
  selectTop = TRUE,
  simParam = NULL,
  ...
)
```

Arguments

pop	and object of Pop-class or HybridPop-class
trait	the trait for selection. Either a number indicating a single trait or a function returning a vector of length nInd.
use	select on genetic values (gv, default), estimated breeding values (ebv), breeding values (bv), or phenotypes (pheno)
p	the proportion of individuals selected
selectTop	selects highest values if true. Selects lowest values if false.
simParam	an object of SimParam
...	additional arguments if using a function for trait

Value

Returns a numeric value

Examples

```
#Create founder haplotypes
founderPop = quickHaplo(nInd=2, nChr=1, segSites=10)

#Set simulation parameters
SP = SimParam$new(founderPop)
SP$addTraitA(10)

#Create population
pop = newPop(founderPop, simParam=SP)

#Determine usefulness of population
usefulness(pop, simParam=SP)

#Should be equivalent to GV of best individual
max(gv(pop))
```

varA

Additive variance

Description

Returns additive variance for all traits

Usage

```
varA(pop, simParam = NULL)
```

Arguments

pop an object of [Pop-class](#)
 simParam an object of [SimParam](#)

Examples

```
#Create founder haplotypes
founderPop = quickHaplo(nInd=10, nChr=1, segSites=10)

#Set simulation parameters
SP = SimParam$new(founderPop)
SP$addTraitAD(10, meanDD=0.5)
SP$setVarE(h2=0.5)

#Create population
pop = newPop(founderPop, simParam=SP)
varA(pop, simParam=SP)
```

varAA	<i>Additive-by-additive epistatic variance</i>
-------	------------------------------------------------

Description

Returns additive-by-additive epistatic variance for all traits

Usage

```
varAA(pop, simParam = NULL)
```

Arguments

pop an object of [Pop-class](#)
 simParam an object of [SimParam](#)

Examples

```
#Create founder haplotypes
founderPop = quickHaplo(nInd=10, nChr=1, segSites=10)

#Set simulation parameters
SP = SimParam$new(founderPop)
SP$addTraitAD(10, meanDD=0.5)
SP$setVarE(h2=0.5)

#Create population
pop = newPop(founderPop, simParam=SP)
varAA(pop, simParam=SP)
```

varD	<i>Dominance variance</i>
------	---------------------------

Description

Returns dominance variance for all traits

Usage

```
varD(pop, simParam = NULL)
```

Arguments

pop	an object of Pop-class
simParam	an object of SimParam

Examples

```
#Create founder haplotypes
founderPop = quickHaplo(nInd=10, nChr=1, segSites=10)

#Set simulation parameters
SP = SimParam$new(founderPop)
SP$addTraitAD(10, meanDD=0.5)
SP$setVarE(h2=0.5)

#Create population
pop = newPop(founderPop, simParam=SP)
varD(pop, simParam=SP)
```

varG	<i>Total genetic variance</i>
------	-------------------------------

Description

Returns total genetic variance for all traits

Usage

```
varG(pop)
```

Arguments

pop	an object of Pop-class or HybridPop-class
-----	---------------------------------------------------------------------------

Examples

```
#Create founder haplotypes
founderPop = quickHaplo(nInd=10, nChr=1, segSites=10)

#Set simulation parameters
SP = SimParam$new(founderPop)
SP$addTraitA(10)
SP$setVarE(h2=0.5)

#Create population
pop = newPop(founderPop, simParam=SP)
varG(pop)
```

varP

Phenotypic variance

Description

Returns phenotypic variance for all traits

Usage

```
varP(pop)
```

Arguments

pop an object of [Pop-class](#) or [HybridPop-class](#)

Examples

```
#Create founder haplotypes
founderPop = quickHaplo(nInd=10, nChr=1, segSites=10)

#Set simulation parameters
SP = SimParam$new(founderPop)
SP$addTraitA(10)
SP$setVarE(h2=0.5)

#Create population
pop = newPop(founderPop, simParam=SP)
varP(pop)
```

writePlink	<i>Writes a Pop-class as PLINK files</i>
------------	------------------------------------------

Description

Writes a Pop-class as PLINK PED and MAP files

Usage

```
writePlink(
  pop,
  baseName,
  trait = 1L,
  snpChip = 1L,
  simParam = NULL,
  chromLength = 10L^8
)
```

Arguments

pop	an object of Pop-class
baseName	a character. Basename of PED and MAP files.
trait	an integer. Which phenotype trait should be used.
snpChip	an integer. Which SNP array should be used.
simParam	an object of SimParam
chromLength	an integer. The size of chromosomes in base pairs; assuming all chromosomes are of the same size.

Examples

```
## Not run:
#Create founder haplotypes
founderPop = quickHaplo(nInd=10, nChr=1, segSites=10)

#Set simulation parameters
SP = SimParam$new(founderPop)
SP$setGender(gender = "yes_rand")
SP$addTraitA(nQt1PerChr = 10)
SP$addSnpChip(nSnpPerChr = 5)

#Create population
pop = newPop(rawPop = founderPop)
pop = setPheno(pop, varE = SP$varA)
writePlink(pop, baseName="test")

#Test
test = read.table(file = "test.ped")
```

```

#...gender
if (!identical(x = c("M", "F")[test[[5]]], y = pop@gender)) { stop() }
#...pheno (issues with rounding)
# if (!identical(x = test[[6]], y = pop@pheno[, 1])) { stop() }
#...genotypes
x = test[, -(1:6)] - 1
x[, 1] = x[, 1] + x[, 2]
x[, 2] = x[, 3] + x[, 4]
x[, 3] = x[, 5] + x[, 6]
x[, 4] = x[, 7] + x[, 8]
x[, 5] = x[, 9] + x[, 10]
y = pullSnpGeno(pop)
if (sum(x[, 1:5] - y) != 0) { stop() }

## End(Not run)

```

writeRecords

Write data records

Description

Saves a population's phenotypic and marker data to a directory.

Usage

```

writeRecords(
  pop,
  dir,
  snpChip = 1,
  useQtl = FALSE,
  includeHaplo = FALSE,
  append = TRUE,
  simParam = NULL
)

```

Arguments

pop	an object of Pop-class
dir	path to a directory for saving output
snpChip	which SNP chip genotype to save. If useQtl=TRUE, this value will indicate which trait's QTL genotype to save. A value of 0 will skip writing a snpChip.
useQtl	should QTL genotype be written instead of SNP chip genotypes.
includeHaplo	should markers be separated by female and male haplotypes.
append	if true, new records are added to any existing records. If false, any existing records are deleted before writing new records. Note that this will delete all files in the 'dir' directory.
simParam	an object of SimParam

Index

[,HybridPop-method (HybridPop-class), 20
[,MapPop-method (MapPop-class), 24
[,Pop-method (Pop-class), 33
[,RawPop-method (RawPop-class), 45

aa, 4
AlphaSimR, 5

bv, 5

c,HybridPop-method (HybridPop-class), 20
c,MapPop-method (MapPop-class), 24
c,Pop-method (Pop-class), 33
c,RawPop-method (RawPop-class), 45
calcGCA, 6
cChr, 6

dd, 7
doubleGenome, 8, 23

ebv, 8
editGenome, 9
editGenomeTopQtl, 10

fastRRBLUP, 11

genicVarA, 12
genicVarAA, 13
genicVarD, 14
genicVarG, 14
genParam, 15
getQtlMap, 16
getSnpMap, 17
gv, 18

hybridCross, 19, 80, 81
HybridPop-class, 20

LociMap-class, 21

makeCross, 21, 42

makeCross2, 22, 44
makeDH, 19, 23, 77
MapPop-class, 24
meanG, 24
meanP, 25
mergeGenome, 26
mergePops, 27
mutate, 27

newMapPop, 28
newPop, 19, 29, 77
nInd, 30

pedigreeCross, 31
pheno, 32
Pop-class, 33
popVar, 34
pullIbdHaplo, 35
pullQtlGeno, 36
pullQtlHaplo, 37
pullSegSiteGeno, 38
pullSegSiteHaplo, 39
pullSnpGeno, 40
pullSnpHaplo, 41

quickHaplo, 42

randCross, 42, 65
randCross2, 44
RawPop-class, 45
reduceGenome, 23, 46
resetPop, 47, 94
RRBLUP, 11, 48, 49–51
RRBLUP2, 11, 49
RRBLUP_D, 52, 53
RRBLUP_D2, 53
RRBLUP_GCA, 51, 55, 56, 58
RRBLUP_GCA2, 56
RRBLUP_SCA, 51, 58, 59
RRBLUP_SCA2, 59

RRBLUPMemUse, 51
RRsol-class, 61
runMacs, 29, 61, 62, 63
runMacs2, 62

sampleHaplo, 64
selectCross, 65
selectFam, 66
selectInd, 65, 68, 73
selectOP, 69
selectWithinFam, 70
self, 72
selIndex, 73
selInt, 74
setEBV, 27, 74
setPheno, 27, 75
setPhenoGCA, 77
show, Pop-method (Pop-class), 33
show, RawPop-method (RawPop-class), 45
SimParam, 4, 5, 7, 9, 10, 12–15, 17, 19, 21, 22,
24, 26, 28, 30, 35–37, 39–41, 43, 44,
47, 48, 50, 52, 54, 55, 57, 58, 60, 65,
67, 68, 70–72, 75–77, 78, 103–105,
107, 108
smithHazel, 99

TraitA-class, 99
TraitA2-class, 100
TraitA2D-class, 100
TraitAD-class, 100
TraitADE-class, 100
TraitADEG-class, 101
TraitADG-class, 101
TraitAE-class, 101
TraitAEG-class, 102
TraitAG-class, 102

usefulness, 102

var, 34
varA, 103
varAA, 104
varD, 105
varG, 105
varP, 106

writePlink, 107
writeRecords, 108