

# Package: AlphaSimR (via r-universe)

September 19, 2024

**Type** Package

**Title** Breeding Program Simulations

**Version** 1.6.0

**Date** 2024-08-15

**Description** The successor to the 'AlphaSim' software for breeding program simulation [Faux et al. (2016) <[doi:10.3835/plantgenome2016.02.0013](https://doi.org/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](https://doi.org/10.1101/gr.083634.108)>].

**License** MIT + file LICENSE

**URL** <https://github.com/gaynorr/AlphaSimR>,  
<https://gaynorr.github.io/AlphaSimR/>,  
[https://www.edx.org/learn/animal-breeding/  
the-university-of-edinburgh-breeding-programme-modelling-with-alphasimr](https://www.edx.org/learn/animal-breeding/the-university-of-edinburgh-breeding-programme-modelling-with-alphasimr)

**Encoding** UTF-8

**Depends** R (>= 4.0.0), methods, R6

**Imports** Rcpp (>= 0.12.7), Rdpack

**RdMacros** Rdpack

**LinkingTo** Rcpp, RcppArmadillo (>= 0.7.500.0.0), BH

**RoxygenNote** 7.3.2

**Suggests** knitr, rmarkdown, testthat

**VignetteBuilder** knitr

**NeedsCompilation** yes

**Author** Chris Gaynor [aut, cre]

(<<https://orcid.org/0000-0003-0558-6656>>), Gregor Gorjanc [ctb]

(<<https://orcid.org/0000-0001-8008-2787>>), John Hickey [ctb]

(<<https://orcid.org/0000-0001-5675-3974>>), Daniel Money [ctb]

(<<https://orcid.org/0000-0001-5151-3648>>), David Wilson [ctb],

Thiago Oliveira [ctb]

(<<https://orcid.org/0000-0002-4555-2584>>), Audrey Martin [ctb]

(<<https://orcid.org/0000-0003-2235-0098>>), Philip Greenspoon

[ctb] (<<https://orcid.org/0000-0001-6284-7248>>)

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

**Repository** CRAN

**Date/Publication** 2024-08-17 22:20:13 UTC

## Contents

.newPop . . . . .	5
aa . . . . .	6
addSegSite . . . . .	6
attrition . . . . .	7
bv . . . . .	8
calcGCA . . . . .	9
cChr . . . . .	9
dd . . . . .	10
doubleGenome . . . . .	11
ebv . . . . .	11
editGenome . . . . .	12
editGenomeTopQtl . . . . .	13
fastRRBLUP . . . . .	14
genicVarA . . . . .	15
genicVarAA . . . . .	16
genicVarD . . . . .	17
genicVarG . . . . .	17
genParam . . . . .	18
getGenMap . . . . .	20
getNumThreads . . . . .	21
getPed . . . . .	21
getQtlMap . . . . .	22
getSnpMap . . . . .	23
gv . . . . .	24
hybridCross . . . . .	24
HybridPop-class . . . . .	25
importGenMap . . . . .	27
importHaplo . . . . .	27
importInbredGeno . . . . .	28

isFemale . . . . .	30
isPop . . . . .	31
LociMap-class . . . . .	31
makeCross . . . . .	32
makeCross2 . . . . .	33
makeDH . . . . .	34
MapPop-class . . . . .	35
meanEBV . . . . .	36
meanG . . . . .	36
meanP . . . . .	37
mergeGenome . . . . .	38
mergePops . . . . .	39
MultiPop-class . . . . .	39
mutate . . . . .	40
NamedMapPop-class . . . . .	41
newEmptyPop . . . . .	42
newMapPop . . . . .	43
newMultiPop . . . . .	44
newPop . . . . .	45
nInd . . . . .	46
pedigreeCross . . . . .	46
pheno . . . . .	48
Pop-class . . . . .	48
popVar . . . . .	50
pullIbdHaplo . . . . .	50
pullMarkerGeno . . . . .	51
pullMarkerHaplo . . . . .	52
pullQtlGeno . . . . .	53
pullQtlHaplo . . . . .	54
pullSegSiteGeno . . . . .	55
pullSegSiteHaplo . . . . .	56
pullSnpGeno . . . . .	57
pullSnpHaplo . . . . .	58
quickHaplo . . . . .	59
randCross . . . . .	59
randCross2 . . . . .	61
RawPop-class . . . . .	62
reduceGenome . . . . .	63
resetPop . . . . .	64
RRBLUP . . . . .	65
RRBLUP2 . . . . .	66
RRBLUPMemUse . . . . .	68
RRBLUP_D . . . . .	69
RRBLUP_D2 . . . . .	70
RRBLUP_GCA . . . . .	71
RRBLUP_GCA2 . . . . .	73
RRBLUP_SCA . . . . .	74
RRBLUP_SCA2 . . . . .	76

RRsol-class	77
runMacs	78
runMacs2	80
sampleHaplo	81
selectCross	82
selectFam	84
selectInd	85
selectOP	87
selectWithinFam	89
self	90
selIndex	91
selInt	92
setEBV	92
setMarkerHaplo	94
setPheno	95
setPhenoGCA	96
setPhenoProgTest	98
SimParam	99
smithHazel	125
solveMKM	126
solveMVM	126
solveRRBLUP	127
solveRRBLUPMK	127
solveRRBLUPMV	128
solveRRBLUP_EM	128
solveRRBLUP_EM2	129
solveRRBLUP_EM3	129
solveUVM	130
TraitA-class	130
TraitA2-class	131
TraitA2D-class	131
TraitAD-class	131
TraitADE-class	131
TraitADEG-class	132
TraitADG-class	132
TraitAE-class	132
TraitAEG-class	133
TraitAG-class	133
transMat	133
usefulness	134
varA	135
varAA	136
varD	137
varEBV	137
varG	138
varP	139
writePlink	139
writeRecords	141

---

.newPop                      *Create new population (internal)*

---

**Description**

Creates a new [Pop-class](#) from an object of of the Pop superclass.

**Usage**

```
.newPop(  
  rawPop,  
  id = NULL,  
  mother = NULL,  
  father = NULL,  
  iMother = NULL,  
  iFather = NULL,  
  isDH = NULL,  
  femaleParentPop = NULL,  
  maleParentPop = NULL,  
  hist = NULL,  
  simParam = NULL,  
  ...  
)
```

**Arguments**

- rawPop                      an object of the pop superclass
- id                            optional id for new individuals
- mother                      optional id for mothers
- father                       optional id for fathers
- iMother                     optional internal id for mothers
- iFather                      optional internal id for fathers
- isDH                         optional indicator for DH/inbred individuals
- femaleParentPop            optional population of female parents
- maleParentPop              optional population of male parents
- hist                         optional recombination history
- simParam                    an object of [SimParam](#)
- ...                          additional arguments passed to the finalizePop function in simParam

**Value**

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

---

aa *Additive-by-additive epistatic deviations*

---

### Description

Returns additive-by-additive epistatic deviations for all traits

### Usage

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

### Arguments

pop	an object of <a href="#">Pop-class</a>
simParam	an object of <a href="#">SimParam</a>

### 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)
```

---

addSegSite *Add segregating site to MapPop*

---

### Description

This function allows for adding a new segregating site with user supplied genotypes to a MapPop. The position of the site is set using a genetic map position.

### Usage

```
addSegSite(mapPop, siteName, chr, mapPos, haplo)
```

**Arguments**

mapPop	an object of <a href="#">MapPop-class</a>
siteName	name to give the segregating site
chr	which chromosome to add the site
mapPos	genetic map position of site in Morgans
haplo	haplotypes for the site

**Value**

an object of [MapPop-class](#)

**Examples**

```
# Creates a populations of 10 outbred individuals
# Their genome consists of 1 chromosome and 2 segregating sites
founderPop = quickHaplo(nInd=10,nChr=1,segSites=2)

# Add a locus a the 0.5 Morgan map position
haplo = matrix(sample(x=0:1, size=20, replace=TRUE), ncol=1)

founderPop2 = addSegSite(founderPop, siteName="x", chr=1, mapPos=0.5, haplo=haplo)

pullSegSiteHaplo(founderPop2)
```

---

attrition

*Lose individuals at random*


---

**Description**

Samples individuals at random to remove from the population. The user supplies a probability for the individuals to be removed from the population.

**Usage**

```
attrition(pop, p)
```

**Arguments**

pop	an object of <a href="#">Pop-class</a>
p	the expected proportion of individuals that will be lost to attrition.

**Value**

an object of [Pop-class](#)

**Examples**

```
#Create founder haplotypes
founderPop = quickHaplo(nInd=100, nChr=1, segSites=10)

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

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

#Lose an expected 5% of individuals
pop = attrition(pop, p=0.05)
```

---

bv	<i>Breeding value</i>
----	-----------------------

---

**Description**

Returns breeding values for all traits

**Usage**

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

**Arguments**

pop	an object of <a href="#">Pop-class</a>
simParam	an object of <a href="#">SimParam</a>

**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	<i>Calculate GCA</i>
---------	----------------------

---

**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 <a href="#">Pop-class</a> or <a href="#">HybridPop-class</a>
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	<i>Combine MapPop chromosomes</i>
------	-----------------------------------

---

**Description**

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

**Usage**

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

**Arguments**

... [MapPop-class](#) or [NamedMapPop-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 <a href="#">Pop-class</a>
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 <a href="#">SimParam</a>

**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 <a href="#">Pop-class</a> to serve as the training population
traits	an integer indicating the trait to model, a trait name, 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 <a href="#">SimParam</a>
...	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

*Additive genic variance*

---

### 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)
geneticVarAA(pop, simParam=SP)
```

---

geneticVarD                      *Dominance genetic variance*

---

**Description**

Returns dominance genetic variance for all traits

**Usage**

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

**Arguments**

pop	an object of <a href="#">Pop-class</a>
simParam	an object of <a href="#">SimParam</a>

**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)
geneticVarD(pop, simParam=SP)
```

---

geneticVarG                      *Total genetic variance*

---

**Description**

Returns total genetic variance for all traits

**Usage**

```
geneticVarG(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  
**alpha** a list of average allele substitution effects with length nTraits  
**alpha\_HW** a list of average allele substitution effects at Hardy-Weinberg equilibrium with length 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)
  
```

---

getGenMap	<i>Get genetic map</i>
-----------	------------------------

---

### Description

Retrieves the genetic map for all loci.

### Usage

```
getGenMap(object = NULL, sex = "A")
```

### Arguments

<b>object</b>	where to retrieve the genetic map. Can be an object of <a href="#">SimParam</a> or <a href="#">MapPop-class</a> . If NULL, the function will look for a SimParam object called "SP" in your global environment.
<b>sex</b>	determines which sex 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 sex specific maps or using pulling from a MapPop.

### Value

Returns a data.frame with:

**id** Unique identifier for locus  
**chr** Chromosome containing the locus  
**pos** Genetic map position

### Examples

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

#Set simulation parameters
getGenMap(founderPop)
```

---

getNumThreads	<i>Number of available threads</i>
---------------	------------------------------------

---

**Description**

Gets the number of available threads by calling the OpenMP function `omp_get_max_threads()`

**Usage**

```
getNumThreads()
```

**Value**

integer

**Examples**

```
getNumThreads()
```

---

getPed	<i>Get pedigree</i>
--------	---------------------

---

**Description**

Returns the population's pedigree as stored in the id, mother and father slots. NULL is returned if the input population lacks the required.

**Usage**

```
getPed(pop)
```

**Arguments**

pop	a population
-----	--------------

**Examples**

```
# Create a founder population
founderPop = quickHaplo(2,1,2)

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

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

```
# Get the pedigree
getPed(pop)

# Returns NULL when a population lacks a pedigree
getPed(founderPop)
```

---

getQtlMap

*Get QTL genetic map*


---

### Description

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

### Usage

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

### Arguments

trait	an integer for the
sex	determines which sex 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 sex specific maps.
simParam	an object of <a href="#">SimParam</a>

### Value

Returns a data.frame with:

**id** Unique identifier for the QTL  
**chr** Chromosome containing the QTL  
**site** Segregating site on the chromosome  
**pos** Genetic map position

### 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, sex = "A", simParam = NULL)
```

### Arguments

snpChip	an integer. Indicates which SNP chip's map to retrieve.
sex	determines which sex 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 sex specific maps.
simParam	an object of <a href="#">SimParam</a>

### Value

Returns a data.frame with:

**id** Unique identifier for the SNP  
**chr** Chromosome containing the SNP  
**site** Segregating site on the chromosome  
**pos** Genetic map position

### 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	<i>Genetic value</i>
----	----------------------

---

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

**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, ...)

isHybridPop(x)
```

**Arguments**

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

**Methods (by generic)**

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

**Functions**

- `isHybridPop()`: Test if object is of a HybridPop class

**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 p=0.5. Dimensions are nInd by nTraits.

pheno matrix of phenotypic values. Dimensions are nInd by nTraits.

gxe list containing GxE slopes for GxE traits

---

importGenMap	<i>Import genetic map</i>
--------------	---------------------------

---

### Description

Formats a genetic map stored in a data.frame to AlphaSimR's internal format. Map positions must be in Morgans.

### Usage

```
importGenMap(genMap)
```

### Arguments

genMap	genetic map as a data.frame. The first three columns must be: marker name, chromosome, and map position (Morgans). Marker name and chromosome are coerced using as.character.
--------	-------------------------------------------------------------------------------------------------------------------------------------------------------------------------------

### Value

a list of named vectors

### Examples

```
genMap = data.frame(markerName=letters[1:5],
                    chromosome=c(1,1,1,2,2),
                    position=c(0,0.5,1,0.15,0.4))

asrMap = importGenMap(genMap=genMap)

str(asrMap)
```

---

importHaplo	<i>Import haplotypes</i>
-------------	--------------------------

---

### Description

Formats haplotype in a matrix format to an AlphaSimR population that can be used to initialize a simulation. This function serves as wrapper for [newMapPop](#) that utilizes a more user friendly input format.

### Usage

```
importHaplo(haplo, genMap, ploidy = 2L, ped = NULL)
```

**Arguments**

haplo	a matrix of haplotypes
genMap	genetic map as a data.frame. The first three columns must be: marker name, chromosome, and map position (Morgans). Marker name and chromosome are coerced using as.character. See <a href="#">importGenMap</a>
ploidy	ploidy level of the organism
ped	an optional pedigree for the supplied genotypes. See details.

**Details**

The optional pedigree can be a data.frame, matrix or a vector. If the object is a data.frame or matrix, the first three columns must include information in the following order: id, mother, and father. All values are coerced using as.character. If the object is a vector, it is assumed to only include the id. In this case, the mother and father will be set to "0" for all individuals.

**Value**

a [MapPop-class](#) if ped is NULL, otherwise a [NamedMapPop-class](#)

**Examples**

```
haplo = rbind(c(1,1,0,1,0),
             c(1,1,0,1,0),
             c(0,1,1,0,0),
             c(0,1,1,0,0))
colnames(haplo) = letters[1:5]

genMap = data.frame(markerName=letters[1:5],
                   chromosome=c(1,1,1,2,2),
                   position=c(0,0.5,1,0.15,0.4))

ped = data.frame(id=c("a","b"),
                mother=c(0,0),
                father=c(0,0))

founderPop = importHaplo(haplo=haplo,
                        genMap=genMap,
                        ploidy=2L,
                        ped=ped)
```

**Description**

Formats the genotypes from inbred, diploid lines to an AlphaSimR population that can be used to initialize a simulation. An attempt is made to automatically detect 0,1,2 or -1,0,1 genotype coding. Heterozygotes or probabilistic genotypes are allowed, but will be coerced to the nearest homozygote. Pedigree information is optional and when provided will be passed to the population for easier identification in the simulation.

**Usage**

```
importInbredGeno(geno, genMap, ped = NULL)
```

**Arguments**

geno	a matrix of genotypes
genMap	genetic map as a data.frame. The first three columns must be: marker name, chromosome, and map position (Morgans). Marker name and chromosome are coerced using as.character. See <a href="#">importGenMap</a>
ped	an optional pedigree for the supplied genotypes. See details.

**Details**

The optional pedigree can be a data.frame, matrix or a vector. If the object is a data.frame or matrix, the first three columns must include information in the following order: id, mother, and father. All values are coerced using as.character. If the object is a vector, it is assumed to only include the id. In this case, the mother and father will be set to "0" for all individuals.

**Value**

a [MapPop-class](#) if ped is NULL, otherwise a [NamedMapPop-class](#)

**Examples**

```
geno = rbind(c(2,2,0,2,0),
             c(0,2,2,0,0))
colnames(geno) = letters[1:5]

genMap = data.frame(markerName=letters[1:5],
                   chromosome=c(1,1,1,2,2),
                   position=c(0,0.5,1,0.15,0.4))

ped = data.frame(id=c("a","b"),
                mother=c(0,0),
                father=c(0,0))

founderPop = importInbredGeno(geno=geno,
                              genMap=genMap,
                              ped=ped)
```

---

isFemale	<i>Test if individuals of a population are female or male</i>
----------	---------------------------------------------------------------

---

**Description**

Test if individuals of a population are female or male

**Usage**

```
isFemale(x)
```

```
isMale(x)
```

**Arguments**

x [Pop-class](#)

**Value**

logical

**Functions**

- `isMale()`: Test if individuals of a population are female or male

**Examples**

```
founderGenomes <- quickHaplo(nInd = 3, nChr = 1, segSites = 100)
SP <- SimParam$new(founderGenomes)
SP$setSexes(sexes = "yes_sys")
pop <- newPop(founderGenomes)

isFemale(pop)
isMale(pop)

pop[isFemale(pop)]
pop[isFemale(pop)]@sex
```

---

isPop	<i>Test if object is of a Population class</i>
-------	------------------------------------------------

---

**Description**

Utilify function to test if object is of a Population class

**Usage**

```
isPop(x)
```

**Arguments**

x                    [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)
isPop(pop)
isPop(SP)
```

---

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  
 name optional name for LociMap object

---

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 <a href="#">Pop-class</a>
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 <a href="#">SimParam</a>

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

---

### 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 <a href="#">Pop-class</a> for female parents.
males	an object of <a href="#">Pop-class</a> 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 <a href="#">SimParam</a>

### 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	<i>Generates DH lines</i>
--------	---------------------------

---

### 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

*Raw population with genetic map*

---

## 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, ...)

isMapPop(x)
```

## Arguments

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

## Methods (by generic)

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

## Functions

- `isMapPop()`: Test if object is of a MapPop class

## Slots

genMap list of chromosome genetic maps  
centromere vector of centromere positions  
inbred indicates whether the individuals are fully inbred

---

meanEBV                      *Mean estimated breeding values*

---

### Description

Returns the mean estimated breeding values for all traits

### Usage

```
meanEBV(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)
trtH2 = 0.5
SP$setVarE(h2=trtH2)

#Create population
pop = newPop(founderPop, simParam=SP)
pop@ebv = trtH2 * (pop@pheno - meanP(pop)) #ind performance based EBV
meanEBV(pop)
```

---

meanG                      *Mean genetic values*

---

### 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

*Mean phenotypic values*

---

**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 <a href="#">Pop-class</a> for female parents.
males	an object of <a href="#">Pop-class</a> 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 <a href="#">SimParam</a>

### 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 or a [MultiPop-class](#)

**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)
pop@misc$tmp = rnorm(n=10)
pop@misc$tmp2 = rnorm(n=10)

popList = list(pop, pop)
pop2 = mergePops(popList)
```

---

MultiPop-class	<i>Multi-Population</i>
----------------	-------------------------

---

**Description**

The mega-population represents a population of populations. It is designed to behave like a list of populations.

**Usage**

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

## S4 method for signature 'MultiPop'
x[[i]]

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

## S4 method for signature 'MultiPop'
length(x)

isMultiPop(x)
```

**Arguments**

x	a 'MultiPop' object
i	index of populations or mega-populations
...	additional 'MultiPop' or 'Pop' objects

**Methods (by generic)**

- `[]`: Extract MultiPop by index
- `[[`: Extract Pop by index
- `c(MultiPop)`: Combine multiple MultiPops
- `length(MultiPop)`: Number of pops in MultiPop

**Functions**

- `isMultiPop()`: Test if object is of a MultiPop class

**Slots**

pops list of [Pop-class](#) and/or [MultiPop-class](#)

---

mutate

*Add Random Mutations*


---

**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 <a href="#">Pop-class</a>
mutRate	rate of new mutations
returnPos	should the positions of mutations be returned
simParam	an object of <a href="#">SimParam</a>

**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)
```

---

NamedMapPop-class      *Raw population with genetic map and id*

---

**Description**

Extends [MapPop-class](#) to add id, mother and father.

**Usage**

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

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

isNamedMapPop(x)
```

**Arguments**

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

**Methods (by generic)**

- []: Extract NamedMapPop by index
- c(NamedMapPop): Combine multiple NamedMapPops

**Functions**

- isNamedMapPop(): Test if object is a NamedMapPop class

**Slots**

id	an individual's identifier
mother	the identifier of the individual's mother
father	the identifier of the individual's father

---

newEmptyPop	<i>Creates an empty population</i>
-------------	------------------------------------

---

**Description**

Creates an empty [Pop-class](#) object with user defined ploidy and other parameters taken from `simParam`.

**Usage**

```
newEmptyPop(ploidy = 2L, simParam = NULL)
```

**Arguments**

ploidy	the ploidy of the population
simParam	an object of <a href="#">SimParam</a>

**Value**

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

**Examples**

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

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

#Create empty population
pop = newEmptyPop(simParam=SP)
isPop(pop)
```

---

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 the 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. The number of rows must match the number of individuals times the ploidy if using inbred=FALSE. If using inbred=TRUE, the number of rows must equal the number of individuals. The haplotypes can be stored as numeric, integer or raw. The underlying C++ function will use raw.

**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)
```

---

newMultiPop

*Create new Multi Population*


---

**Description**

Creates a new [MultiPop-class](#) from one or more [Pop-class](#) and/or [MultiPop-class](#) objects.

**Usage**

```
newMultiPop(...)
```

**Arguments**

... one or more [Pop-class](#) and/or [MultiPop-class](#) objects.

**Value**

Returns an object of [MultiPop-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)
megaPop = newMultiPop(pop=pop)
isMultiPop(megaPop)
```

---

newPop	<i>Create new population</i>
--------	------------------------------

---

### Description

Creates an initial [Pop-class](#) from an object of [MapPop-class](#) or [NamedMapPop-class](#). The function is intended for use with output from functions such as [runMacs](#), [newMapPop](#), or [quickHaplo](#).

### Usage

```
newPop(rawPop, simParam = NULL, ...)
```

### Arguments

rawPop	an object of <a href="#">MapPop-class</a> or <a href="#">NamedMapPop-class</a>
simParam	an object of <a href="#">SimParam</a>
...	additional arguments used internally

### Details

Note that newPop takes genomes from the rawPop and uses them without recombination! Hence, if you call newPop(rawPop = founderGenomes) twice, you will get two sets of individuals with different id but the same genomes. To get genetically different sets of individuals you can subset the rawPop input, say first half for one set and the second half for the other set.

### 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)
isPop(pop)

#Misc
pop@misc$tmp1 = rnorm(n=2)
pop@misc$tmp2 = rnorm(n=2)

#MiscPop
pop@miscPop$tmp1 = sum(pop@misc$tmp1)
pop@miscPop$tmp2 = sum(pop@misc$tmp2)
```

---

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,
  maxCycle = 100,
  DH = NULL,
  nSelf = NULL,
  useFemale = TRUE,
  simParam = NULL
)
```

**Arguments**

founderPop	a <a href="#">Pop-class</a>
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.
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.
nSelf	an optional vector indicating how many generations an individual should be selfed.
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 sex, 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)

## S4 method for signature 'Pop'
length(x)
```

**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(Pop)`: Combine multiple Pops
- `show(Pop)`: Show population summary
- `length(Pop)`: Number of individuals in Pop (the same as `nInd()`)

**Slots**

`id` an individual's identifier

`iid` an individual's internal identifier

`mother` the identifier of the individual's mother

`father` the identifier of the individual's father

`sex` sex of individuals: "M" for males, "F" for females, and "H" for hermaphrodites

`nTraits` number of traits

`gv` matrix of genetic values. When using GxE traits, `gv` reflects `gv` when  $p=0.5$ . 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.

`misc` a list whose elements correspond to additional miscellaneous nodes with the items for individuals in the population (see example in `newPop`). This list is normally empty and exists solely as an open slot available for uses to store extra information about individuals.

`miscPop` a list of any length containing optional meta data for the population (see example in `newPop`). This list is empty unless information is supplied by the user. Note that the list is emptied every time the population is subsetted or combined because the meta data for old population might not be valid anymore.

### See Also

`newPop`, `newEmptyPop`, `resetPop`

---

`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 IBD haplotypes*

---

### Description

Retrieves IBD haplotype data

### Usage

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

**Arguments**

pop	an object of <a href="#">Pop-class</a>
chr	a vector of chromosomes to retrieve. If NULL, all chromosomes are retrieved.
snpChip	an integer indicating which SNP array loci are to be retrieved. If NULL, all sites are retrieved.
simParam	an object of <a href="#">SimParam</a>

**Value**

Returns a matrix of IBD 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)
SP$setTrackRec(TRUE)

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

---

pullMarkerGeno	<i>Pull marker genotypes</i>
----------------	------------------------------

---

**Description**

Retrieves genotype data for user specified loci

**Usage**

```
pullMarkerGeno(pop, markers, asRaw = FALSE, simParam = NULL)
```

**Arguments**

pop	an object of <a href="#">RawPop-class</a> or <a href="#">MapPop-class</a>
markers	a character vector. Indicates the names of the loci to be retrieved.
asRaw	return in raw (byte) format
simParam	an object of <a href="#">SimParam</a> , not used if pop is <a href="#">MapPop-class</a>

**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)

#Pull genotype data for first two markers on chromosome one.
#Marker name is consistent with default naming in AlphaSimR.
pullMarkerGeno(pop, markers=c("1_1", "1_2"), simParam=SP)
```

---

pullMarkerHaplo	<i>Pull marker haplotypes</i>
-----------------	-------------------------------

---

**Description**

Retrieves haplotype data for user specified loci

**Usage**

```
pullMarkerHaplo(pop, markers, haplo = "all", asRaw = FALSE, simParam = NULL)
```

**Arguments**

pop	an object of <a href="#">RawPop-class</a> or <a href="#">MapPop-class</a>
markers	a character vector. Indicates the names of the loci to be retrieved
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 in diploids.
asRaw	return in raw (byte) format
simParam	an object of <a href="#">SimParam</a> , not used if pop is <a href="#">MapPop-class</a>

**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)
SP$setTrackRec(TRUE)

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

#Pull haplotype data for first two markers on chromosome one.
#Marker name is consistent with default naming in AlphaSimR.
pullMarkerHaplo(pop, markers=c("1_1", "1_2"), simParam=SP)
```

---

pullQtlGeno

*Pull QTL genotypes*

---

## Description

Retrieves QTL genotype data

## Usage

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

## Arguments

pop	an object of <a href="#">Pop-class</a>
trait	an integer. Indicates which trait's QTL genotypes to retrieve.
chr	a vector of chromosomes to retrieve. If NULL, all chromosome are retrieved.
asRaw	return in raw (byte) format
simParam	an object of <a href="#">SimParam</a>

## 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

*Pull QTL haplotypes*

---

**Description**

Retrieves QTL haplotype data

**Usage**

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

**Arguments**

pop	an object of <a href="#">Pop-class</a>
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 in diploids.
chr	a vector of chromosomes to retrieve. If NULL, all chromosome are retrieved.
asRaw	return in raw (byte) format
simParam	an object of <a href="#">SimParam</a>

**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	<i>Pull segregating site genotypes</i>
-----------------	----------------------------------------

---

**Description**

Retrieves genotype data for all segregating sites

**Usage**

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

**Arguments**

pop	an object of <a href="#">RawPop-class</a> or <a href="#">MapPop-class</a>
chr	a vector of chromosomes to retrieve. If NULL, all chromosome are retrieved.
asRaw	return in raw (byte) format
simParam	an object of <a href="#">SimParam</a> , not used if pop is <a href="#">MapPop-class</a>

**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,
  asRaw = FALSE,
  simParam = NULL
)
```

### Arguments

pop	an object of <a href="#">RawPop-class</a> or <a href="#">MapPop-class</a>
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 in diploids.
chr	a vector of chromosomes to retrieve. If NULL, all chromosome are retrieved.
asRaw	return in raw (byte) format
simParam	an object of <a href="#">SimParam</a> , not used if pop is <a href="#">MapPop-class</a>

### 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	<i>Pull SNP genotypes</i>
-------------	---------------------------

---

### Description

Retrieves SNP genotype data

### Usage

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

### Arguments

pop	an object of <a href="#">Pop-class</a>
snpChip	an integer. Indicates which SNP chip's genotypes to retrieve.
chr	a vector of chromosomes to retrieve. If NULL, all chromosome are retrieved.
asRaw	return in raw (byte) format
simParam	an object of <a href="#">SimParam</a>

### 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                      *Pull SNP haplotypes*

---

### Description

Retrieves SNP haplotype data

### Usage

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

### Arguments

pop	an object of <a href="#">Pop-class</a>
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 in diploids.
chr	a vector of chromosomes to retrieve. If NULL, all chromosome are retrieved.
asRaw	return in raw (byte) format
simParam	an object of <a href="#">SimParam</a>

### 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,  
  ignoreSexes = FALSE,  
  simParam = NULL  
)
```

**Arguments**

pop	an object of <a href="#">Pop-class</a>
nCrosses	total number of crosses to make
nProgeny	number of progeny per cross
balance	if using sexes, this option will balance the number of progeny per parent
parents	an optional vector of indices for allowable parents
ignoreSexes	should sexes be ignored
simParam	an object of <a href="#">SimParam</a>

**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,
  ignoreSexes = FALSE,
  simParam = NULL
)
```

### Arguments

females	an object of <a href="#">Pop-class</a> for female parents.
males	an object of <a href="#">Pop-class</a> 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
ignoreSexes	should sex be ignored
simParam	an object of <a href="#">SimParam</a>

### 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)

isRawPop(x)
```

### Arguments

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

### Methods (by generic)

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

### Functions

- `isRawPop()`: Test if object is of a RawPop class

**Slots**

nInd number of individuals  
 nChr number of chromosomes  
 ploidy level of ploidy  
 nLoci number of loci per chromosome  
 geno list of nChr length containing chromosome genotypes. Each element is a three dimensional array of raw values. The array dimensions are nLoci by ploidy by nInd.

---

reduceGenome	<i>Create individuals with reduced ploidy</i>
--------------	-----------------------------------------------

---

**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 <a href="#">Pop-class</a>
simParam	an object of <a href="#">SimParam</a>

## 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,  
  simParam = NULL,  
  ...  
)
```

### Arguments

pop	a <a href="#">Pop-class</a> to serve as the training population
traits	an integer indicating the trait or traits to model, a vector of trait names, 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.
simParam	an object of <a href="#">SimParam</a>
...	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,
  simParam = NULL,
  ...
)

```

## Arguments

pop	a <a href="#">Pop-class</a> to serve as the training population
traits	an integer indicating the trait to model, a trait name, or a function of the traits returning a single value. Unlike <a href="#">RRBLUP</a> , 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
simParam	an object of <a href="#">SimParam</a>
...	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 algorithm is set by maxIter. The default value is typically too small for convergence. When the algorithm fails to converge 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 <a href="#">RRBLUP</a> , <a href="#">RRBLUP_GCA</a> and <a href="#">RRBLUP_SCA</a> 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,
  simParam = NULL,
  ...
)
```

**Arguments**

pop	a <a href="#">Pop-class</a> to serve as the training population
traits	an integer indicating the trait to model, a trait name, 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.
simParam	an object of <a href="#">SimParam</a>
...	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,
  simParam = NULL,
  ...
)

```

### Arguments

pop	a <a href="#">Pop-class</a> to serve as the training population
traits	an integer indicating the trait to model, a trait name, 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
simParam	an object of <a href="#">SimParam</a>
...	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))
```

**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,
  simParam = NULL,
  ...
)
```

**Arguments**

pop	a <a href="#">Pop-class</a> to serve as the training population
traits	an integer indicating the trait to model, a trait name, 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.
simParam	an object of <a href="#">SimParam</a>
...	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,
  simParam = NULL,
  ...
)
```

### Arguments

<code>pop</code>	a <a href="#">Pop-class</a> to serve as the training population
<code>traits</code>	an integer indicating the trait to model, a trait name, or a function of the traits returning a single value.
<code>use</code>	train model using phenotypes "pheno", genetic values "gv", estimated breeding values "ebv", breeding values "bv", or randomly "rand"
<code>snpChip</code>	an integer indicating which SNP chip genotype to use
<code>useQtl</code>	should QTL genotypes be used instead of a SNP chip. If TRUE, <code>snpChip</code> specifies which trait's QTL to use, and thus these QTL may not match the QTL underlying the phenotype supplied in <code>traits</code> .

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
simParam	an object of <a href="#">SimParam</a>
...	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,
  simParam = NULL,
  ...
)
```

**Arguments**

pop	a <a href="#">Pop-class</a> to serve as the training population
traits	an integer indicating the trait to model, a trait name, 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.
simParam	an object of <a href="#">SimParam</a>
...	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,
  simParam = NULL,
  ...
)
```

**Arguments**

pop	a <a href="#">Pop-class</a> to serve as the training population
traits	an integer indicating the trait to model, a trait name, 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
simParam	an object of <a href="#">SimParam</a>
...	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 (Chen et al. 2009).

**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

There are currently three species histories available: GENERIC, CATTLE, WHEAT, and MAIZE.

The GENERIC history is meant to be a reasonable all-purpose choice. It runs quickly and models a population with an effective populations size that has gone through several historic bottlenecks. This species history is used as the default arguments in the `runMacs2` function, so the user should examine this function for the details of how the species is modeled.

The CATTLE history is based off of real genome sequence data (MacLeod et al. 2013).

The WHEAT (Gaynor et al. 2017) and MAIZE (Hickey et al. 2014) histories have been included due to their use in previous simulations. However, it should be noted that neither faithfully simulates its respective species. This is apparent by the low number of segregating sites simulated by each history relative to their real-world analogs. Adjusting these histories to better represent their real-world analogs would result in a drastic increase to runtime.

## Value

an object of `MapPop-class`

## References

Chen GK, Marjoram P, Wall JD (2009). “Fast and Flexible Simulation of DNA Sequence Data.” *Genome Research*, **19**, 136-142. <https://genome.cshlp.org/content/19/1/136>.

Gaynor RC, Gorjanc G, Bentley AR, Ober ES, Howell P, Jackson R, Mackay IJ, Hickey JM (2017). “A Two-Part Strategy for Using Genomic Selection to Develop Inbred Lines.” *Crop Science*, **57**(5), 2372–2386. ISSN 0011-183X, [doi:10.2135/cropsci2016.09.0742](https://doi.org/10.2135/cropsci2016.09.0742), <https://access.onlinelibrary.wiley.com/doi/full/10.2135/cropsci2016.09.0742>.

Hickey JMDS, Crossa J, Hearne S, Babu R, Prasanna BM, Grondona M, Zambelli A, Windhausen VS, Mathews K, Gorjanc G (2014). “Evaluation of Genomic Selection Training Population Designs and Genotyping Strategies in Plant Breeding Programs Using Simulation.” *Crop Science*, **54**(4), 1476-1488. [doi:10.2135/cropsci2013.03.0195](https://doi.org/10.2135/cropsci2013.03.0195).

MacLeod IM, Larkin DM, Lewin HAHBJ, Goddard ME (2013). “Inferring Demography from Runs of Homozygosity in Whole-Genome Sequence, with Correction for Sequence Errors.” *Molecular Biology and Evolution*, **30**(9), 2209–2223. [doi:10.1093/molbev/mst125](https://doi.org/10.1093/molbev/mst125).

## Examples

```
# Creates a populations of 10 outbred individuals
# Their genome consists of 1 chromosome and 100 segregating sites
## Not run:
founderPop = runMacs(nInd=10,nChr=1,segSites=100)

## End(Not run)
```

runMacs2

*Alternative wrapper for MaCS***Description**

A wrapper function for `runMacs`. This wrapper is designed to provide a more intuitive interface for writing custom commands in MaCS (Chen et al. 2009). It effectively automates the creation of an appropriate line for the `manualCommand` argument in `runMacs` using user supplied variables, but only allows for a subset of the functionality offered by this argument. The default arguments of this function 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**

<code>nInd</code>	number of individuals to simulate
<code>nChr</code>	number of chromosomes to simulate
<code>segSites</code>	number of segregating sites to keep per chromosome
<code>Ne</code>	effective population size
<code>bp</code>	base pair length of chromosome
<code>genLen</code>	genetic length of chromosome in Morgans
<code>mutRate</code>	per base pair mutation rate
<code>histNe</code>	effective population size in previous generations
<code>histGen</code>	number of generations ago for effective population sizes given in <code>histNe</code>
<code>inbred</code>	should founder individuals be inbred
<code>split</code>	an optional historic population split in terms of generations ago
<code>ploidy</code>	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`.

### References

Chen GK, Marjoram P, Wall JD (2009). "Fast and Flexible Simulation of DNA Sequence Data." *Genome Research*, **19**, 136-142. <https://genome.cshlp.org/content/19/1/136>.

### 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
## Not run:
founderPop = runMacs2(nInd=10,nChr=1,segSites=100)

# runMacs() Implementation of the cattle demography following
# Macleod et al. (2013) https://doi.org/10.1093/molbev/mst125
cattleChrSum = 2.8e9 # https://www.ncbi.nlm.nih.gov/datasets/genome/GCF_002263795.3/
(cattleChrBp = cattleChrSum / 30)
recRate = 9.26e-09
(cattleGenLen = recRate * cattleChrBp)
mutRate = 1.20e-08
runMacs2(nInd = 10, nChr = 1, Ne = 90, bp = cattleChrBp,
         genLen = cattleGenLen, mutRate = 1.20e-08,
         histNe = c(120, 250, 350, 1000, 1500, 2000, 2500, 3500, 7000, 10000, 17000, 62000),
         histGen = c( 3,  6, 12,  18,  24, 154,  454,  654, 1754,  2354,  3354, 33154),
         returnCommand = TRUE)

## End(Not run)
```

---

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 <a href="#">MapPop-class</a> 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=1,segSites=11,inbred=TRUE)
founderPop = sampleHaplo(mapPop=founderPop,nInd=20)
```

---

selectCross

*Select and randomly cross*

---

**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**

pop	an object of <a href="#">Pop-class</a>
nInd	the number of individuals to select. These individuals are selected without regards to sex and it supercedes values for nFemale and nMale. Thus if the simulation uses sexes, it is likely better to leave this value as NULL and use nFemale and nMale instead.
nFemale	the number of females to select. This value is ignored if nInd is set.
nMale	the number of males to select. This value is ignored if nInd is set.
nCrosses	total number of crosses to make
nProgeny	number of progeny per cross
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.
simParam	an object of <a href="#">SimParam</a>
...	additional arguments if using a function for trait
balance	if using sexes, 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	<i>Select families</i>
-----------	------------------------

---

### Description

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

### Usage

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

### Arguments

pop	and object of <a href="#">Pop-class</a> , <a href="#">HybridPop-class</a> or <a href="#">MultiPop-class</a>
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. The function must work on a vector or matrix of use values as <code>trait(pop@use, ...)</code> - depending on what use is. See the examples and <a href="#">selIndex</a> .
use	the selection criterion. Either a character (genetic values "gv", estimated breeding values "ebv", breeding values "bv", phenotypes "pheno", or randomly "rand") or a function returning a vector of length nInd. The function must work on pop as <code>use(pop, trait, ...)</code> or as <code>trait(pop@use, ...)</code> depending on what trait is. See the examples.
sex	which sex to select. Use "B" for both, "F" for females and "M" for males. If the simulation is not using sexes, 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 <a href="#">Pop-class</a> . If FALSE, only the index of selected individuals is returned.
candidates	an optional vector of eligible selection candidates.
simParam	an object of <a href="#">SimParam</a>
...	additional arguments if using a function for trait and use

**Value**

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

*Select individuals*

---

**Description**

Selects a subset of nInd individuals from a population.

**Usage**

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

**Arguments**

pop	and object of <a href="#">Pop-class</a> , <a href="#">HybridPop-class</a> or <a href="#">MultiPop-class</a>
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. The function must work on a vector or matrix of use values as <code>trait(pop@use, ...)</code> - depending on what use is. See the examples and <a href="#">selIndex</a> .
use	the selection criterion. Either a character (genetic values "gv", estimated breeding values "ebv", breeding values "bv", phenotypes "pheno", or randomly "rand") or a function returning a vector of length nInd. The function must work on pop as <code>use(pop, trait, ...)</code> or as <code>trait(pop@use, ...)</code> depending on what trait is. See the examples.
sex	which sex to select. Use "B" for both, "F" for females and "M" for males. If the simulation is not using sexes, the argument is ignored.
selectTop	selects highest values if true. Selects lowest values if false.
returnPop	should results be returned as a <a href="#">Pop-class</a> . If FALSE, only the index of selected individuals is returned.
candidates	an optional vector of eligible selection candidates.
simParam	an object of <a href="#">SimParam</a>
...	additional arguments if using a function for trait or use

**Value**

Returns an object of [Pop-class](#), [HybridPop-class](#) or [MultiPop-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 top 5 (directional selection)
pop2 = selectInd(pop, 5, simParam=SP)
hist(pop@pheno); abline(v=pop@pheno, lwd=2)
abline(v=pop2@pheno, col="red", lwd=2)

#Select 5 most deviating from an optima (disruptive selection)
squaredDeviation = function(x, optima=0) (x - optima)^2
pop3 = selectInd(pop, 5, trait=squaredDeviation, selectTop=TRUE, simParam=SP)
```

```

hist(pop@pheno); abline(v=pop@pheno, lwd=2)
abline(v=pop3@pheno, col="red", lwd=2)

#Select 5 least deviating from an optima (stabilising selection)
pop4 = selectInd(pop, 5, trait=squaredDeviation, selectTop=FALSE, simParam=SP)
hist(pop@pheno); abline(v=pop@pheno, lwd=2)
abline(v=pop4@pheno, col="red", lwd=2)

#Select 5 individuals based on miscellaneous information with use function
pop@misc = list(smth=rnorm(10), smth2=rnorm(10))
useFunc = function(pop, trait=NULL) pop@misc$smth + pop@misc$smth2
pop5 = selectInd(pop, 5, use=useFunc, simParam=SP)
pop5@id

#... equivalent result with the use & trait function
useFunc2 = function(pop, trait=NULL) cbind(pop@misc$smth, pop@misc$smth2)
trtFunc = function(x) rowSums(x)
pop6 = selectInd(pop, 5, trait=trtFunc, use=useFunc2, simParam=SP)
pop6@id

```

---

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	and object of <a href="#">Pop-class</a> or <a href="#">MultiPop-class</a>
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. The function must work on a vector or matrix of use values as <code>trait(pop@use, ...)</code> - depending on what use is. See the examples and <a href="#">selIndex</a> .
use	the selection criterion. Either a character (genetic values "gv", estimated breeding values "ebv", breeding values "bv", phenotypes "pheno", or randomly "rand") or a function returning a vector of length nInd. The function must work on pop as <code>use(pop, trait, ...)</code> or as <code>trait(pop@use, ...)</code> depending on what trait is. See the examples.
selectTop	selects highest values if true. Selects lowest values if false.
candidates	an optional vector of eligible selection candidates.
simParam	an object of <a href="#">SimParam</a>
...	additional arguments if using a function for trait and use

**Value**

Returns an object of [Pop-class](#) or [MultiPop-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	<i>Select individuals within families</i>
-----------------	-------------------------------------------

---

### 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",
  sex = "B",
  famType = "B",
  selectTop = TRUE,
  returnPop = TRUE,
  candidates = NULL,
  simParam = NULL,
  ...
)
```

### Arguments

pop	and object of <a href="#">Pop-class</a> , <a href="#">HybridPop-class</a> or <a href="#">MultiPop-class</a>
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. The function must work on a vector or matrix of use values as trait(pop@use, ...) - depending on what use is. See the examples and <a href="#">selIndex</a> .
use	the selection criterion. Either a character (genetic values "gv", estimated breeding values "ebv", breeding values "bv", phenotypes "pheno", or randomly "rand") or a function returning a vector of length nInd. The function must work on pop as use(pop, trait, ...) or as trait(pop@use, ...) depending on what trait is. See the examples.
sex	which sex to select. Use "B" for both, "F" for females and "M" for males. If the simulation is not using sexes, 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 <a href="#">Pop-class</a> . 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 and use

### Value

Returns an object of [Pop-class](#), [HybridPop-class](#) or [MultiPop-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                      *Self individuals*

---

### Description

Creates selfed progeny from each individual in a population. Only works when sexes 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

*Selection index*


---

**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 estimated breeding values (EBV)</i>
--------	--------------------------------------------

---

### Description

Adds genomic estimated values to a populations's EBV slot using output from a genomic selection functions. 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 <a href="#">Pop-class</a>
solution	an object of <a href="#">RRsol-class</a>
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 <a href="#">SimParam</a>

**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))
```

---

setMarkerHaplo	<i>Set marker haplotypes</i>
----------------	------------------------------

---

### Description

Manually sets the haplotypes in a population for all individuals at one or more loci.

### Usage

```
setMarkerHaplo(pop, haplo, simParam = NULL)
```

### Arguments

pop	an object of <a href="#">RawPop-class</a> or <a href="#">MapPop-class</a>
haplo	a matrix of haplotypes, see details
simParam	an object of <a href="#">SimParam</a> , not used if pop is <a href="#">MapPop-class</a>

### Details

The format of the haplotype matrix should match the format of the output from [pullMarkerHaplo](#) with the option haplo="all". Thus, it is recommended that this function is first used to extract the haplotypes and that any desired changes be made to the output of [pullMarkerHaplo](#) before passing the matrix to [setMarkerHaplo](#). Any changes made to QTL may potentially result in changes to an individuals genetic value. These changes will be reflected in the gv and/or gxe slot. All other slots will remain unchanged, so the ebv and pheno slots will not reflect the new genotypes.

### Value

an object of the same class as the "pop" input

### Examples

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

# Extract haplotypes for marker "1_1"
H = pullMarkerHaplo(founderPop, markers="1_1")

# Set the first haplotype to 1
H[1,1] = 1L

# Set marker haplotypes
founderPop = setMarkerHaplo(founderPop, haplo=H)
```

---

setPheno	<i>Set phenotypes</i>
----------	-----------------------

---

### Description

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

### Usage

```
setPheno(
  pop,
  h2 = NULL,
  H2 = NULL,
  varE = NULL,
  corE = NULL,
  reps = 1,
  fixEff = 1L,
  p = NULL,
  onlyPheno = FALSE,
  traits = NULL,
  simParam = NULL
)
```

### Arguments

pop	an object of <a href="#">Pop-class</a> or <a href="#">HybridPop-class</a>
h2	a vector of desired narrow-sense heritabilities for each trait. See details.
H2	a vector of desired broad-sense heritabilities for each trait. See details.
varE	error (co)variances for traits. See details.
corE	an optional matrix for correlations between errors. See details.
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
traits	an integer vector indicate which traits to set. If NULL, all traits will be set.
simParam	an object of <a href="#">SimParam</a>

### Details

There are three arguments for setting the error variance of a phenotype: h2, H2, and varE. The user should only use one of these arguments. If the user supplies values for more than one, only one will be used according to order in which they are listed above.

The h2 argument allows the user to specify the error variance according to narrow-sense heritability. This calculation uses the additive genetic variance and total genetic variance in the founder population. Thus, the heritability relates to the founder population and not the current population.

The H2 argument allows the user to specify the error variance according to broad-sense heritability. This calculation uses the total genetic variance in the founder population. Thus, the heritability relates to the founder population and not the current population.

The varE argument allows the user to specify the error variance directly. The user may supply a vector describing the error variance for each trait or supply a matrix that specify the covariance of the errors.

The corE argument allows the user to specify correlations for the error covariance matrix. These correlations are be supplied in addition to the h2, H2, or varE arguments. These correlations will be used to construct a covariance matrix from a vector of variances. If the user supplied a covariance matrix to varE, these correlations will supercede values provided in that matrix.

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 the 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",
  h2 = NULL,
  H2 = NULL,
  varE = NULL,
  corE = NULL,
  reps = 1,
  fixEff = 1L,
  p = NULL,
  inbred = FALSE,
  onlyPheno = FALSE,
  simParam = NULL
)

```

**Arguments**

pop	an object of <a href="#">Pop-class</a>
testers	an object of <a href="#">Pop-class</a>
use	true genetic value (gv) or phenotypes (pheno, default)
h2	a vector of desired narrow-sense heritabilities for each trait. See details in <a href="#">setPheno</a> .
H2	a vector of desired broad-sense heritabilities for each trait. See details in <a href="#">setPheno</a> .
varE	error (co)variances for traits. See details in <a href="#">setPheno</a> .
corE	an optional matrix for correlations between errors. See details in <a href="#">setPheno</a> .
reps	number of replications for phenotype. See details in <a href="#">setPheno</a> .
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.
inbred	are both pop and testers fully inbred. They are only fully inbred if created by <a href="#">newPop</a> using inbred founders or by the <a href="#">makeDH</a> function
onlyPheno	should only the phenotype be returned, see return
simParam	an object of <a href="#">SimParam</a>

**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)
```

---

setPhenoProgTest      *Set progeny test as phenotype*

---

### Description

Models a progeny test of individuals in 'pop'. Returns 'pop' with a phenotype representing the average performance of their progeny. The phenotype is generated by mating individuals in 'pop' to randomly chosen individuals in testPop a number of times equal to 'nMatePerInd'.

### Usage

```
setPhenoProgTest(
  pop,
  testPop,
  nMatePerInd = 1L,
  use = "pheno",
  h2 = NULL,
  H2 = NULL,
  varE = NULL,
  corE = NULL,
  reps = 1,
  fixEff = 1L,
  p = NULL,
  onlyPheno = FALSE,
  simParam = NULL
)
```

### Arguments

pop	an object of <a href="#">Pop-class</a>
testPop	an object of <a href="#">Pop-class</a>
nMatePerInd	number of times an individual in 'pop' is mated to an individual in testPop
use	true genetic value (gv) or phenotypes (pheno, default)
h2	a vector of desired narrow-sense heritabilities for each trait. See details in <a href="#">setPheno</a> .

H2	a vector of desired broad-sense heritabilities for each trait. See details in <a href="#">setPheno</a> .
varE	error (co)variances for traits. See details in <a href="#">setPheno</a> .
corE	an optional matrix for correlations between errors. See details in <a href="#">setPheno</a> .
reps	number of replications for phenotype. See details in <a href="#">setPheno</a> .
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 <a href="#">SimParam</a>

### 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 two populations of 5 individuals
pop1 = newPop(founderPop[1:5], simParam=SP)
pop2 = newPop(founderPop[6:10], simParam=SP)

#Set phenotype according to a progeny test
pop3 = setPhenoProgTest(pop1, pop2, use="gv", 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  
 finalizePop function applied to newly created populations. Currently does nothing and should only be changed by expert users.  
 allowEmptyPop if true, population arguments with nInd=0 will return an empty population with a warning instead of an error.  
 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 2.6)  
 p the proportion of crossovers coming from a non-interfering pathway. (default is 0)  
 quadProb the probability of quadrivalent pairing in an autopolyploid. (default is 0)

**Active bindings**

traitNames vector of trait names  
 snpChipNames vector of chip names  
 traits list of traits  
 nChr number of chromosomes  
 nTraits number of traits  
 nSnpChips number of SNP chips  
 segSites segregating sites per chromosome  
 sexes sexes 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  
 haplotypes list of computed IBD haplotypes  
 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$restrSegSites()`
- `SimParam$setSexes()`
- `SimParam$setFounderHap()`
- `SimParam$addSnpChip()`
- `SimParam$addSnpChipByName()`
- `SimParam$addStructuredSnpChip()`
- `SimParam$addTraitA()`
- `SimParam$addTraitAD()`
- `SimParam$altAddTraitAD()`
- `SimParam$addTraitAG()`
- `SimParam$addTraitADG()`
- `SimParam$addTraitAE()`
- `SimParam$addTraitADE()`
- `SimParam$addTraitAEG()`
- `SimParam$addTraitADEG()`
- `SimParam$manAddTrait()`
- `SimParam$importTrait()`
- `SimParam$switchTrait()`
- `SimParam$removeTrait()`
- `SimParam$setVarE()`
- `SimParam$setCorE()`
- `SimParam$rescaleTraits()`
- `SimParam$setRecombRatio()`
- `SimParam$switchGenMap()`
- `SimParam$switchFemaleMap()`
- `SimParam$switchMaleMap()`
- `SimParam$addToRec()`
- `SimParam$ibdHaplo()`
- `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$new(founderPop)
\dontshow{SP$nThreads = 1L}
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$new(founderPop)
\dontshow{SP$nThreads = 1L}
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$new(founderPop)
\dontshow{SP$nThreads = 1L}
```

```
#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 a SNP and/or QTL.

*Usage:*

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

*Arguments:*

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

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

`excludeQtl` an optional vector of segregating site names to exclude from consideration as a viable QTL.

`excludeSnp` an optional vector of segregating site names to exclude from consideration as a viable SNP.

`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$new(founderPop)
\dontshow{SP$nThreads = 1L}
SP$restrSegSites(minQtlPerChr=5, minSnpPerChr=5)
```

**Method** `setSexes()`: Changes how sexes are determined in the simulation. The default sexes is "no", indicating all individuals are hermaphrodites. To add sexes to the simulation, run this function with "yes\_sys" or "yes\_rand". The value "yes\_sys" will systematically assign sexes to newly created individuals as first male and then female. Populations with an odd number of individuals will have one more male than female. The value "yes\_rand" will randomly assign a sex to each individual.

*Usage:*

```
SimParam$setSexes(sexes, force = FALSE)
```

*Arguments:*

`sexes` 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$new(founderPop)
\dontshow{SP$nThreads = 1L}
SP$setSexes("yes_sys")
```

**Method** `setFounderHap()`: Allows for the manual setting of founder haplotypes. This functionality is not fully documented, because it is still experimental.

*Usage:*

```
SimParam$setFounderHap(hapMap)
```

*Arguments:*

`hapMap` a list of founder haplotypes

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



*Usage:*

```
SimParam$addSnpChip(nSnpPerChr, minSnpFreq = NULL, refPop = NULL, name = 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.

name optional name for chip

*Examples:*

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

```
#Set simulation parameters
SP = SimParam$new(founderPop)
\dontshow{SP$nThreads = 1L}
SP$addSnpChip(10)
```

**Method** addSnpChipByName(): Assigns SNPs to a SNP chip by supplying marker names. This function does check against excluded SNPs and will not add the SNPs to the list of excluded QTL for the purpose of avoiding overlap between SNPs and QTL. Excluding these SNPs from being used as QTL can be accomplished using the excludeQtl argument in SimParam's restrSegSites function.

*Usage:*

```
SimParam$addSnpChipByName(markers, name = NULL)
```

*Arguments:*

markers a vector of names for the markers

name optional name for chip

*Examples:*

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

```
#Set simulation parameters
SP = SimParam$new(founderPop)
SP$addSnpChipByName(c("1_1", "1_3"))
```

**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 pleiotropic with correlated additive effects.

*Usage:*

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

*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 (the rate/scale parameter of the gamma distribution is accounted for via the desired level of genetic variance, the var argument)

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

name optional name for trait(s)

*Examples:*

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

```
#Set simulation parameters
SP = SimParam$new(founderPop)
\dontshow{SP$nThreads = 1L}
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,
    name = NULL
)

```

*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 (the rate/scale parameter of the gamma distribution is accounted for via the desired level of genetic variance, the *var* argument)

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

*name* optional name for trait(s)

*Examples:*

```

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

#Set simulation parameters
SP = SimParam$new(founderPop)
\dontshow{SP$nThreads = 1L}
SP$addTraitAD(10, meanDD=0.5)

```

**Method** `altAddTraitAD()`: An alternative method for adding a trait with additive and dominance effects to an AlphaSimR simulation. The function attempts to create a trait matching user defined values for number of QTL, inbreeding depression, additive genetic variance and dominance genetic variance.

*Usage:*

```

SimParam$altAddTraitAD(
  nQtlPerChr,
  mean = 0,
  varA = 1,
  varD = 0,
  inbrDepr = 0,
  limMeanDD = c(0, 1.5),
  limVarDD = c(0, 0.5),

```

```

    silent = FALSE,
    force = FALSE,
    name = NULL
)

```

*Arguments:*

nQtlPerChr number of QTLs per chromosome. Can be a single value or nChr values.  
 mean desired mean of the trait  
 varA desired additive variance  
 varD desired dominance variance  
 inbrDepr desired inbreeding depression, see details  
 limMeanDD limits for meanDD, see details  
 limVarDD limits for varDD, see details  
 silent should summary details be printed to the console  
 force should the check for a running simulation be ignored. Only set to TRUE if you know what you are doing.  
 name optional name for trait

*Details:* This function will always add a trait to 'SimParam', unless an error occurs with picking QTLs. The resulting trait will always have the desired mean and additive genetic variance. However, it may not have the desired values for inbreeding depression and dominance variance. Thus, it is strongly recommended to check the output printed to the console to determine how close the trait's parameters came to these desired values.

The mean and additive genetic variance will always be achieved exactly. The function attempts to achieve the desired dominance variance and inbreeding depression while staying within the user supplied constraints for the acceptable range of dominance degree mean and variance. If the desired values are not being achieved, the acceptable range need to be increased and/or the number of QTL may need to be increased. There are not limits to setting the range for dominance degree mean and variance, but care should be taken to with regards to the biological feasibility of the limits that are supplied. The default limits were somewhat arbitrarily set, so I make not claim to how reasonable these limits are for routine use.

Inbreeding depression in this function is defined as the difference in mean genetic value between a population with the same allele frequency as the reference population (population used to initialize SimParam) in Hardy-Weinberg equilibrium compared to a population with the same allele frequency that is fully inbred. This is equivalent to the amount the mean of a population increases when going from an inbreeding coefficient of 1 (fully inbred) to a population with an inbreeding coefficient of 0 (Hardy-Weinberg equilibrium). Note that the sign of the value should (usually) be positive. This corresponds to a detrimental effect of inbreeding when higher values of the trait are considered biologically beneficial.

Summary information on this trait is printed to the console when silent=FALSE. The summary information reports the inbreeding depression and dominance variance for the population as well as the dominance degree mean and variance applied to the trait.

*Examples:*

```

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

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

```

```
\dontshow{SP$nThreads = 1L}
SP$altAddTraitAD(nQt1PerChr=10, mean=0, varA=1, varD=0.05, inbrDepr=0.2)
```

**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(
  nQt1PerChr,
  mean = 0,
  var = 1,
  varGxE = 1e-06,
  varEnv = 0,
  corA = NULL,
  corGxE = NULL,
  gamma = FALSE,
  shape = 1,
  force = FALSE,
  name = NULL
)
```

*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  
 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 (the rate/scale parameter of the gamma distribution is accounted for via the desired level of genetic variance, the var argument)  
 force should the check for a running simulation be ignored. Only set to TRUE if you know what you are doing.  
 name optional name for trait(s)

*Examples:*

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

#Set simulation parameters
SP = SimParam$new(founderPop)
\dontshow{SP$nThreads = 1L}
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 = 0,
  varGxE = 1e-06,
  meanDD = 0,
  varDD = 0,
  corA = NULL,
  corDD = NULL,
  corGxE = NULL,
  useVarA = TRUE,
  gamma = FALSE,
  shape = 1,
  force = FALSE,
  name = NULL
)

```

*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  
*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 (the rate/scale parameter of the gamma distribution is accounted for via the desired level of genetic variance, the var argument)  
*force* should the check for a running simulation be ignored. Only set to TRUE if you know what you are doing.  
*name* optional name for trait(s)

*Examples:*

```

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

#Set simulation parameters
SP = SimParam$new(founderPop)
\dontshow{SP$nThreads = 1L}
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 pleiotropic with correlated additive effects.

*Usage:*

```

SimParam$addTraitAE(
  nQtlPerChr,
  mean = 0,
  var = 1,
  relAA = 0,
  corA = NULL,
  corAA = NULL,
  useVarA = TRUE,
  gamma = FALSE,
  shape = 1,
  force = FALSE,
  name = NULL
)

```

*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

`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 (the rate/scale parameter of the gamma distribution is accounted for via the desired level of genetic variance, the `var` argument)

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

`name` optional name for trait(s)

*Examples:*

```

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

#Set simulation parameters
SP = SimParam$new(founderPop)
\dontshow{SP$nThreads = 1L}
SP$addTraitAE(10, relAA=0.1)

```

**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 pleiotropic with correlated effects.

*Usage:*

```

SimParam$addTraitADE(
  nQtlPerChr,

```

```

mean = 0,
var = 1,
meanDD = 0,
varDD = 0,
relAA = 0,
corA = NULL,
corDD = NULL,
corAA = NULL,
useVarA = TRUE,
gamma = FALSE,
shape = 1,
force = FALSE,
name = NULL
)

```

*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 (the rate/scale parameter of the gamma distribution is accounted for via the desired level of genetic variance, the *var* argument)

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

*name* optional name for trait(s)

*Examples:*

```

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

#Set simulation parameters
SP = SimParam$new(founderPop)
\dontshow{SP$nThreads = 1L}
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 pleiotropic with correlated effects.

*Usage:*



```

SimParam$addTraitAEG(
  nQtlPerChr,
  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,
  name = NULL
)

```

*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  
*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 (the rate/scale parameter of the gamma distribution is accounted for via the desired level of genetic variance, the var argument)  
*force* should the check for a running simulation be ignored. Only set to TRUE if you know what you are doing.  
*name* optional name for trait(s)

*Examples:*

```

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

#Set simulation parameters
SP = SimParam$new(founderPop)
\dontshow{SP$nThreads = 1L}
SP$addTraitAEG(10, varGxE=2)

```

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

*Usage:*

```

SimParam$addTraitADEG(
  nQt1PerChr,
  mean = 0,
  var = 1,
  varEnv = 0,
  varGxE = 1e-06,
  meanDD = 0,
  varDD = 0,
  re1AA = 0,
  corA = NULL,
  corDD = NULL,
  corAA = NULL,
  corGxE = NULL,
  useVarA = TRUE,
  gamma = FALSE,
  shape = 1,
  force = FALSE,
  name = NULL
)

```

*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  
*re1AA* 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 (the rate/scale parameter of the gamma distribution is accounted for via the desired level of genetic variance, the var argument)  
*force* should the check for a running simulation be ignored. Only set to TRUE if you know what you are doing.  
*name* optional name for trait(s)

*Examples:*

```

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

```

```
#Set simulation parameters
SP = SimParam$new(founderPop)
\dontshow{SP$nThreads = 1L}
SP$addTraitADEG(10, meanDD=0.5, varGxE=2)
```

**Method** `manAddTrait()`: Manually add a new trait to the simulation. Trait must be formatted as a [LociMap-class](#). If the trait is not already formatted, consider using `importTrait`.

*Usage:*

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

*Arguments:*

`lociMap` a new object descended from [LociMap-class](#)

`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** `importTrait()`: Manually add a new trait(s) to the simulation. Unlike the `manAddTrait` function, this function does not require formatting the trait as a [LociMap-class](#). The formatting is performed automatically for the user, with more user friendly `data.frames` or matrices taken as inputs. This function only works for A and AD trait types.

*Usage:*

```
SimParam$importTrait(
  markerNames,
  addEff,
  domEff = NULL,
  intercept = NULL,
  name = NULL,
  varE = NULL,
  force = FALSE
)
```

*Arguments:*

`markerNames` a vector of names for the QTL

`addEff` a matrix of additive effects (nLoci x nTraits). Alternatively, a vector of length nLoci can be supplied for a single trait.

`domEff` optional dominance effects for each locus

`intercept` optional intercepts for each trait

`name` optional name(s) for the trait(s)

`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, varE = NA_real_, force = FALSE)
```

*Arguments:*

`traitPos` an integer indicate which trait to switch

lociMap a new object descended from [LociMap-class](#)  
 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 values for error variances used in [setPheno](#). These defaults will be used to automatically generate phenotypes when new populations are created. See the details section of [setPheno](#) for more information about each arguments and how they should be used.

*Usage:*

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

*Arguments:*

h2 a vector of desired narrow-sense heritabilities  
 H2 a vector of desired broad-sense heritabilities  
 varE a vector or matrix of error variances  
 corE an optional matrix of error correlations

*Examples:*

```

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

#Set simulation parameters
SP = SimParam$new(founderPop)
\dontshow{SP$nThreads = 1L}
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$new(founderPop)
\dontshow{SP$nThreads = 1L}
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$rescaleTraits(mean=1)
\dontshow{SP$nThreads = 1L}
#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 sex-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$new(founderPop)
\dontshow{SP$nThreads = 1L}
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(lastId, id, mother, father, isDH, hist, ploidy)
```

*Arguments:*

lastId ID of last individual  
id the name of each individual  
mother vector of mother iids  
father vector of father iids  
isDH indicator for DH lines  
hist new recombination history  
ploidy ploidy level

**Method** ibdHaplo(): For internal use only.

*Usage:*

```
SimParam$ibdHaplo(iid)
```

*Arguments:*

iid internal ID

**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, id, mother, father, isDH)
```

*Arguments:*

lastId ID of last individual  
id the name of each individual  
mother vector of mother iids  
father vector of father iids  
isDH indicator for DH lines

**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(minQtlPerChr=5, minSnpPerChr=5)

## -----
## Method `SimParam$setSexes`
## -----

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

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

SP$setSexes("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$addSnpChipByName`
## -----

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

#Set simulation parameters
SP = SimParam$new(founderPop)
SP$addSnpChipByName(c("1_1", "1_3"))

## -----
## 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$altAddTraitAD`
## -----

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

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

SP$altAddTraitAD(nQtlPerChr=10, mean=0, varA=1, varD=0.05, inbrDepr=0.2)

## -----
## 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)

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

SP$addTraitAE(10, relAA=0.1)

## -----
## 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)
```

---

 solveMKM

*Solve Multikernel Model*


---

**Description**

Solves a univariate mixed model with multiple random effects.

**Usage**

```
solveMKM(y, X, Zlist, Klist, maxIter = 40L, tol = 1e-04)
```

**Arguments**

y	a matrix with n rows and 1 column
X	a matrix with n rows and x columns
Zlist	a list of Z matrices
Klist	a list of K matrices
maxIter	maximum number of iteration
tol	tolerance for convergence

---

 solveMVM

*Solve Multivariate Model*


---

**Description**

Solves a multivariate mixed model of form  $Y = X\beta + Zu + e$

**Usage**

```
solveMVM(Y, X, Z, K, tol = 1e-06, maxIter = 1000L)
```

**Arguments**

Y	a matrix with n rows and q columns
X	a matrix with n rows and x columns
Z	a matrix with n rows and m columns
K	a matrix with m rows and m columns
tol	tolerance for convergence
maxIter	maximum number of iteration

---

solveRRBLUP	<i>Solve RR-BLUP</i>
-------------	----------------------

---

**Description**

Solves a univariate mixed model of form  $y = X\beta + Mu + e$

**Usage**

```
solveRRBLUP(y, X, M)
```

**Arguments**

y	a matrix with n rows and 1 column
X	a matrix with n rows and x columns
M	a matrix with n rows and m columns

---

solveRRBLUPMK	<i>Solve Multikernel RR-BLUP</i>
---------------	----------------------------------

---

**Description**

Solves a univariate mixed model with multiple random effects.

**Usage**

```
solveRRBLUPMK(y, X, Mlist, maxIter = 40L)
```

**Arguments**

y	a matrix with n rows and 1 column
X	a matrix with n rows and x columns
Mlist	a list of M matrices
maxIter	maximum number of iteration

---

solveRRBLUPMV	<i>Solve Multivariate RR-BLUP</i>
---------------	-----------------------------------

---

**Description**

Solves a multivariate mixed model of form  $Y = X\beta + Mu + e$

**Usage**

`solveRRBLUPMV(Y, X, M, maxIter = 1000L, tol = 1e-06)`

**Arguments**

Y	a matrix with n rows and q columns
X	a matrix with n rows and x columns
M	a matrix with n rows and m columns
maxIter	maximum number of iteration
tol	tolerance for convergence

---

solveRRBLUP_EM	<i>Solve RR-BLUP with EM</i>
----------------	------------------------------

---

**Description**

Solves a univariate mixed model of form  $y = X\beta + Mu + e$  using the Expectation-Maximization algorithm.

**Usage**

`solveRRBLUP_EM(Y, X, M, Vu, Ve, tol, maxIter, useEM)`

**Arguments**

Y	a matrix with n rows and 1 column
X	a matrix with n rows and x columns
M	a matrix with n rows and m columns
Vu	initial guess for variance of marker effects
Ve	initial guess for error variance
tol	tolerance for declaring convergence
maxIter	maximum iteration for attempting convergence
useEM	should EM algorithm be used. If false, no estimation of variance components is performed. The initial values are treated as true.



---

solveRRBLUP_EM2	<i>Solve RR-BLUP with EM and 2 random effects</i>
-----------------	---------------------------------------------------

---

**Description**

Solves a univariate mixed model of form  $y = X\beta + M_1u_1 + M_2u_2 + e$  using the Expectation-Maximization algorithm.

**Usage**

```
solveRRBLUP_EM2(Y, X, M1, M2, Vu1, Vu2, Ve, tol, maxIter, useEM)
```

**Arguments**

Y	a matrix with n rows and 1 column
X	a matrix with n rows and x columns
M1	a matrix with n rows and m1 columns
M2	a matrix with n rows and m2 columns
Vu1	initial guess for variance of the first marker effects
Vu2	initial guess for variance of the second marker effects
Ve	initial guess for error variance
tol	tolerance for declaring convergence
maxIter	maximum iteration for attempting convergence
useEM	should EM algorithm be used. If false, no estimation of variance components is performed. The initial values are treated as true.

---

solveRRBLUP_EM3	<i>Solve RR-BLUP with EM and 3 random effects</i>
-----------------	---------------------------------------------------

---

**Description**

Solves a univariate mixed model of form  $y = X\beta + M_1u_1 + M_2u_2 + M_3u_3 + e$  using the Expectation-Maximization algorithm.

**Usage**

```
solveRRBLUP_EM3(Y, X, M1, M2, M3, Vu1, Vu2, Vu3, Ve, tol, maxIter, useEM)
```

**Arguments**

Y	a matrix with n rows and 1 column
X	a matrix with n rows and x columns
M1	a matrix with n rows and m1 columns
M2	a matrix with n rows and m2 columns
M3	a matrix with n rows and m3 columns
Vu1	initial guess for variance of the first marker effects
Vu2	initial guess for variance of the second marker effects
Vu3	initial guess for variance of the second marker effects
Ve	initial guess for error variance
tol	tolerance for declaring convergence
maxIter	maximum iteration for attempting convergence
useEM	should EM algorithm be used. If false, no estimation of variance components is performed. The initial values are treated as true.

---

solveUVM	<i>Solve Univariate Model</i>
----------	-------------------------------

---

**Description**

Solves a univariate mixed model of form  $y = X\beta + Zu + e$

**Usage**

```
solveUVM(y, X, Z, K)
```

**Arguments**

y	a matrix with n rows and 1 column
X	a matrix with n rows and x columns
Z	a matrix with n rows and m columns
K	a matrix with m rows and m columns

---

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>Sex 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>Sex 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**

gxeEff 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**

gxeEff 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

---

transMat	<i>Linear transformation matrix</i>
----------	-------------------------------------

---

**Description**

Creates an m by m linear transformation matrix that can be applied to n by m uncorrelated deviates sampled from a standard normal distribution to produce correlated deviates with an arbitrary correlation of R. If R is not positive semi-definite, the function returns smoothing and returns a warning (see details).

**Usage**

```
transMat(R)
```

**Arguments**

R a correlation matrix

**Details**

An eigendecomposition is applied to the correlation matrix and used to test if it is positive semi-definite. If the matrix is not positive semi-definite, it is not a valid correlation matrix. In this case, smoothing is applied to the matrix (as described in the 'cor.smooth' of the 'psych' library) to obtain a valid correlation matrix. The resulting deviates will thus not exactly match the desired correlation, but will hopefully be close if the input matrix wasn't too far removed from a valid correlation matrix.

**Examples**

```
# Create an 2x2 correlation matrix
R = 0.5*diag(2) + 0.5

# Sample 1000 uncorrelated deviates from a
# bivariate standard normal distribution
X = matrix(rnorm(2*1000), ncol=2)

# Compute the transformation matrix
T = transMat(R)

# Apply the transformation to the deviates
Y = X%*%T

# Measure the sample correlation
cor(Y)
```

---

usefulness

*Usefulness criterion*


---

**Description**

Calculates the usefulness criterion

**Usage**

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

**Arguments**

pop	and object of <a href="#">Pop-class</a> or <a href="#">HybridPop-class</a>
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 <a href="#">SimParam</a>
...	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

*Additive-by-additive epistatic variance*


---

**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                                      *Dominance variance*

---

### 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)
```

---

varEBV                                    *Variance of estimated breeding values*

---

### Description

Returns variance of estimated breeding values for all traits

### Usage

```
varEBV(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)
trtH2 = 0.5
SP$setVarE(h2=trtH2)

#Create population
pop = newPop(founderPop, simParam=SP)
pop@ebv = trtH2 * (pop@pheno - meanP(pop)) #ind performance based EBV
varA(pop)
varEBV(pop)
```

---

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

*Writes a Pop-class as PLINK files*


---

### Description

Writes a Pop-class to PLINK PED and MAP files. The arguments for this function were chosen for consistency with [RRBLUP2](#). The base pair coordinate will be the locus position as stored in AlphaSimR and not an actual base pair position. This is because AlphaSimR doesn't track base pair positions, only relative positions for the loci used in the simulation.

**Usage**

```
writePlink(
  pop,
  baseName,
  traits = 1,
  use = "pheno",
  snpChip = 1,
  useQtl = FALSE,
  simParam = NULL,
  ...
)
```

**Arguments**

pop	an object of <a href="#">Pop-class</a>
baseName	basename for PED and MAP files.
traits	an integer indicating the trait to write, a trait name, or a function of the traits returning a single value.
use	what to use for PLINK's phenotype field. Either phenotypes "pheno", genetic values "gv", estimated breeding values "ebv", breeding values "bv", or random values "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.
simParam	an object of <a href="#">SimParam</a>
...	additional arguments if using a function for traits

**Examples**

```
## Not run:
#Create founder haplotypes
founderPop = quickHaplo(nInd=10, nChr=1, segSites=15)

#Set simulation parameters
SP = SimParam$new(founderPop)
\dontshow{SP$nThreads = 1L}
SP$setSexes(sex="yes_rand")
SP$addTraitA(nQtlPerChr=10)
SP$addSnpChip(nSnpPerChr=5)
SP$setVarE(h2=0.5)

#Create population
pop = newPop(rawPop = founderPop)

# Write out PLINK files
writePlink(pop, baseName="test")

## End(Not run)
```

---

writeRecords	<i>Write data records</i>
--------------	---------------------------

---

**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 <a href="#">Pop-class</a>
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 <a href="#">SimParam</a>

# Index

- .newPop, 5
- [,HybridPop-method (HybridPop-class), 25
- [,MapPop-method (MapPop-class), 35
- [,MultiPop-method (MultiPop-class), 39
- [,NamedMapPop-method (NamedMapPop-class), 41
- [,Pop-method (Pop-class), 48
- [,RawPop-method (RawPop-class), 62
- [[,MultiPop-method (MultiPop-class), 39
  
- aa, 6
- addSegSite, 6
- attrition, 7
  
- bv, 8
  
- c,HybridPop-method (HybridPop-class), 25
- c,MapPop-method (MapPop-class), 35
- c,MultiPop-method (MultiPop-class), 39
- c,NamedMapPop-method (NamedMapPop-class), 41
- c,Pop-method (Pop-class), 48
- c,RawPop-method (RawPop-class), 62
- calcGCA, 9
- cChr, 9
  
- dd, 10
- doubleGenome, 11, 34
  
- ebv, 11
- editGenome, 12
- editGenomeTopQtl, 13
  
- fastRRBLUP, 14
  
- genicVarA, 15
- genicVarAA, 16
- genicVarD, 17
- genicVarG, 17
- genParam, 18
- getGenMap, 20
  
- getNumThreads, 21
- getPed, 21
- getQtlMap, 22
- getSnpMap, 23
- gv, 24
  
- hybridCross, 24, 102
- HybridPop-class, 25
  
- importGenMap, 27, 28, 29
- importHaplo, 27
- importInbredGeno, 28
- isFemale, 30
- isHybridPop (HybridPop-class), 25
- isMale (isFemale), 30
- isMapPop (MapPop-class), 35
- isMultiPop (MultiPop-class), 39
- isNamedMapPop (NamedMapPop-class), 41
- isPop, 31
- isRawPop (RawPop-class), 62
  
- length,MultiPop-method (MultiPop-class), 39
- length,Pop-method (Pop-class), 48
- LociMap-class, 31
  
- makeCross, 32, 59
- makeCross2, 33, 61
- makeDH, 24, 34, 97
- MapPop-class, 35
- meanEBV, 36
- meanG, 36
- meanP, 37
- mergeGenome, 38
- mergePops, 39
- MultiPop-class, 39
- mutate, 40
  
- NamedMapPop-class, 41
- newEmptyPop, 42, 50
- newMapPop, 27, 43, 45

- newMultiPop, 44
- newPop, 24, 45, 50, 97
- nInd, 46
  
- pedigreeCross, 46
- pheno, 48
- Pop-class, 48
- popVar, 50
- pullIbdHaplo, 50
- pullMarkerGeno, 51
- pullMarkerHaplo, 52, 94
- pullQtlGeno, 53
- pullQtlHaplo, 54
- pullSegSiteGeno, 55
- pullSegSiteHaplo, 56
- pullSnpgeno, 57
- pullSnpgenoHaplo, 58
  
- quickHaplo, 45, 59
  
- randCross, 59, 82
- randCross2, 61
- RawPop-class, 62
- reduceGenome, 34, 63
- resetPop, 50, 64, 119
- RRBLUP, 14, 65, 66–68
- RRBLUP2, 14, 66, 139
- RRBLUP\_D, 69, 70
- RRBLUP\_D2, 70
- RRBLUP\_GCA, 68, 71, 73, 74
- RRBLUP\_GCA2, 73
- RRBLUP\_SCA, 68, 74, 76
- RRBLUP\_SCA2, 76
- RRBLUPMemUse, 68
- RRsol-class, 77
- runMacs, 45, 78, 80, 81
- runMacs2, 79, 80
  
- sampleHaplo, 81
- selectCross, 82
- selectFam, 84
- selectInd, 82, 85, 91
- selectOP, 87
- selectWithinFam, 89
- self, 90
- selIndex, 84, 86, 88, 89, 91
- selInt, 92
- setEBV, 40, 92
- setMarkerHaplo, 94
- setPheno, 40, 95, 97–99, 116
- setPhenoGCA, 96
- setPhenoProgTest, 98
- show, Pop-method (Pop-class), 48
- show, RawPop-method (RawPop-class), 62
- SimParam, 5, 6, 8, 10, 12, 13, 15–18, 20, 22, 23, 25, 32, 33, 35, 38, 41, 42, 45, 51–58, 60, 61, 64, 65, 67, 69, 71, 72, 74, 75, 77, 83, 84, 86, 88, 90, 93–95, 97, 99, 99, 135–137, 140, 141
- smithHazel, 125
- solveMKM, 126
- solveMVM, 126
- solveRRBLUP, 127
- solveRRBLUP\_EM, 128
- solveRRBLUP\_EM2, 129
- solveRRBLUP\_EM3, 129
- solveRRBLUPMK, 127
- solveRRBLUPMV, 128
- solveUVM, 130
  
- TraitA-class, 130
- TraitA2-class, 131
- TraitA2D-class, 131
- TraitAD-class, 131
- TraitADE-class, 131
- TraitADEG-class, 132
- TraitADG-class, 132
- TraitAE-class, 132
- TraitAEG-class, 133
- TraitAG-class, 133
- transMat, 133
  
- usefulness, 134
  
- var, 50
- varA, 135
- varAA, 136
- varD, 137
- varEBV, 137
- varG, 138
- varP, 139
  
- writePlink, 139
- writeRecords, 141