

Package: MoBPS (via r-universe)

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Type Package

Title Modular Breeding Program Simulator

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Description Framework for the simulation framework for the simulation of complex breeding programs and compare their economic and genetic impact. The package is also used as the background simulator for our a web-based interface <<http://www.mobps.de>>. Associated publication: Pook et al. (2020) <[doi:10.1534/g3.120.401193](https://doi.org/10.1534/g3.120.401193)>.

Depends R (>= 3.0),

Imports graphics, stats, utils

License GPL (>= 3)

LazyData TRUE

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Enhances miraculix (>= 0.9.10), RandomFieldsUtils (>= 0.5.9), MoBPSmaps

Additional_repositories <https://tpook92.github.io/drat/>

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add.array	<i>Add a genotyping array</i>
-----------	-------------------------------

Description

Function to add a genotyping array for the population

Usage

```
add.array(population, marker.included = TRUE, array.name = NULL)
```

Arguments

population	population list
marker.included	Vector with number of SNP entries coding if each marker is on the array (TRUE/FALSE)
array.name	Name of the added array

Value

Population list

Examples

```
data(ex_pop)
population <- add.array(ex_pop, marker.included = c(TRUE, FALSE), array.name="Half-density")
```

add.combi	<i>Add a trait as a linear combination of other traits</i>
-----------	--

Description

Function to create an additional trait that is the results of a linear combination of the other traits

Usage

```
add.combi(population, trait, combi.weights, trait.name = NULL)
```

Arguments

population	population list
trait	trait nr. for which to implement a combination of other traits
combi.weights	Weights (only linear combinations of other traits are allowed!)
trait.name	Name of the trait generated

Value

Population list

Population list

Examples

```
data(ex_pop)
population <- creating.trait(ex_pop, n.additive = 100)
population <- add.combi(population, trait = 3, combi.weights = c(1,5))
```

add.diag	<i>Add something to the diagonal</i>
----------	--------------------------------------

Description

Function to add numeric to the diagonal of a matrix

Usage

```
add.diag(M, d)
```

Arguments

M	Matrix
d	Vector to add to the diagonal of the matrix

Value

Matrix with increased diagonal elements

Matrix with modified diagonal entries

Examples

```
A <- matrix(c(1,2,3,4), ncol=2)
B <- add.diag(A, 5)
```

add.founder.kinship *Add a relationship matrix for founder individuals*

Description

Function to relationship matrix for founder individuals that is used for any calculation of the pedigree

Usage

```
add.founder.kinship(population, founder.kinship = "vanRaden", gen = 1)
```

Arguments

population	population list
founder.kinship	Default is to use vanRaden relationship. Alternative is to enter a pedigree-matrix (order of individuals is first male then female)
gen	Generation for which to enter the pedigree-matrix

Value

Population list

Examples

```
data(ex_pop)
population <- add.founder.kinship(ex_pop)
```

alpha_to_beta	<i>Moore-Penrose-Transformation</i>
---------------	-------------------------------------

Description

Internal transformation using Moore-Penrose

Usage

```
alpha_to_beta(alpha, G, Z)
```

Arguments

alpha	alpha
G	kinship-matrix
Z	genomic information matrix

Value

Vector with single marker effects

analyze.bv	<i>Analyze genomic values</i>
------------	-------------------------------

Description

Function to analyze correlation between bv/bve/pheno

Usage

```
analyze.bv(
  population,
  gen = NULL,
  database = NULL,
  cohorts = NULL,
  bvrow = "all",
  advanced = FALSE
)
```

Arguments

population	Population list
gen	Quick-insert for database (vector of all generations to export)
database	Groups of individuals to consider for the export
cohorts	Quick-insert for database (vector of names of cohorts to export)
bvrow	Which traits to display
advanced	Set to TRUE to also look at offspring pheno

Value

[[1]] Correlation between BV/BVE/Phenotypes [[2]] Genetic variance of the traits

Examples

```
data(ex_pop)
analyze.bv(ex_pop, gen=1)
```

analyze.population *Analyze allele frequency of a single marker*

Description

Analyze allele frequency of a single marker

Usage

```
analyze.population(
  population,
  chromosome = NULL,
  snp = NULL,
  snp.name = NULL,
  database = NULL,
  gen = NULL,
  cohorts = NULL
)
```

Arguments

population	Population list
chromosome	Number of the chromosome of the relevant SNP
snp	Number of the relevant SNP
snp.name	Name of the SNP to analyze
database	Groups of individuals to consider for the export
gen	Quick-insert for database (vector of all generations to export)
cohorts	Quick-insert for database (vector of names of cohorts to export)

Value

Frequency of AA/AB/BB in selected gen/database/cohorts

Examples

```
data(ex_pop)
analyze.population(ex_pop, snp=1, chromosome=1, gen=1:5)
```

bit.snps	<i>Decoding of bitwise-storing in R</i>
----------	---

Description

Function for decoding in bitwise-storing in R (only 30 of 32 bits are used!)

Usage

```
bit.snps(bit.seq, nbits, population = NULL, from.p.bit = 1)
```

Arguments

bit.seq	bitweise gespeicherte SNP-Sequenz
nbits	Number of usable bits (default: 30)
population	Population list
from.p.bit	Bit to start on

Value

De-coded marker sequence

bit.storing	<i>Bitwise-storing in R</i>
-------------	-----------------------------

Description

Function for bitwise-storing in R (only 30 of 32 bits are used!)

Usage

```
bit.storing(snpseq, nbits)
```

Arguments

snpseq	SNP sequence
nbits	Number of usable bits (default: 30)

Value

Bit-wise coded marker sequence

breeding.diploid *Breeding function*

Description

Function to simulate a step in a breeding scheme

Usage

```
breeding.diploid(  
  population,  
  mutation.rate = 10^-8,  
  remutation.rate = 10^-8,  
  recombination.rate = 1,  
  selection.m = NULL,  
  selection.f = NULL,  
  new.selection.calculation = TRUE,  
  selection.function.matrix = NULL,  
  selection.size = 0,  
  ignore.best = 0,  
  breeding.size = 0,  
  breeding.sex = NULL,  
  breeding.sex.random = FALSE,  
  relative.selection = FALSE,  
  class.m = 0,  
  class.f = 0,  
  add.gen = 0,  
  recom.f.indicator = NULL,  
  duplication.rate = 0,  
  duplication.length = 0.01,  
  duplication.recombination = 1,  
  new.class = 0L,  
  bve = FALSE,  
  sigma.e = NULL,  
  sigma.g = 100,  
  new.bv.child = NULL,  
  phenotyping.child = NULL,  
  relationship.matrix = "vanRaden",  
  relationship.matrix.ogc = "kinship",  
  computation.A = NULL,  
  computation.A.ogc = NULL,  
  delete.haplotypes = NULL,  
  delete.individuals = NULL,  
  fixed.breeding = NULL,  
  fixed.breeding.best = NULL,  
  max.offspring = Inf,  
  max.litter = Inf,
```

```
store.breeding.totals = FALSE,
forecast.sigma.g = TRUE,
multiple.bve = "add",
store.bve.data = FALSE,
fixed.assignment = FALSE,
reduce.group = NULL,
reduce.group.selection = "random",
selection.highest = c(TRUE, TRUE),
selection.criteria = NULL,
same.sex.activ = FALSE,
same.sex.sex = 0.5,
same.sex.selfing = FALSE,
selfing.mating = FALSE,
selfing.sex = 0.5,
praeimplantation = NULL,
heritability = NULL,
repeatability = NULL,
save.recombination.history = FALSE,
martini.selection = FALSE,
BGLR.bve = FALSE,
BGLR.model = "RKHS",
BGLR.burnin = 500,
BGLR.iteration = 5000,
BGLR.print = FALSE,
copy.individual = FALSE,
copy.individual.m = FALSE,
copy.individual.f = FALSE,
dh.mating = FALSE,
dh.sex = 0.5,
n.observation = NULL,
bve.0isNA = FALSE,
phenotype.bv = FALSE,
delete.same.origin = FALSE,
remove.effect.position = FALSE,
estimate.u = FALSE,
new.phenotype.correlation = NULL,
new.residual.correlation = NULL,
new.breeding.correlation = NULL,
estimate.add.gen.var = FALSE,
estimate.pheno.var = FALSE,
best1.from.group = NULL,
best2.from.group = NULL,
best1.from.cohort = NULL,
best2.from.cohort = NULL,
add.class.cohorts = TRUE,
store.comp.times = TRUE,
store.comp.times.bve = TRUE,
store.comp.times.generation = TRUE,
```

```
import.position.calculation = NULL,
BGLR.save = "RKHS",
BGLR.save.random = FALSE,
ogc = FALSE,
ogc.target = "min.sKin",
ogc.uniform = NULL,
ogc.ub = NULL,
ogc.lb = NULL,
ogc.ub.sKin = NULL,
ogc.lb.BV = NULL,
ogc.ub.BV = NULL,
ogc.eq.BV = NULL,
ogc.ub.sKin.increase = NULL,
ogc.lb.BV.increase = NULL,
emmreml.bve = FALSE,
rrblup.bve = FALSE,
sommer.bve = FALSE,
sommer.multi.bve = FALSE,
nr.edits = 0,
gene.editing.offspring = FALSE,
gene.editing.best = FALSE,
gene.editing.offspring.sex = c(TRUE, TRUE),
gene.editing.best.sex = c(TRUE, TRUE),
gwas.u = FALSE,
approx.residuals = TRUE,
sequenceZ = FALSE,
maxZ = 5000,
maxZtotal = 0,
delete.sex = 1:2,
gwas.group.standard = FALSE,
y.gwas.used = "pheno",
gen.architecture.m = 0,
gen.architecture.f = NULL,
add.architecture = NULL,
ncore = 1,
ncore.generation = 1,
Z.integer = FALSE,
store.effect.freq = FALSE,
backend = "doParallel",
randomSeed = NULL,
randomSeed.generation = NULL,
Rprof = FALSE,
miraculix = NULL,
miraculix.cores = 1,
miraculix.mult = NULL,
miraculix.chol = TRUE,
best.selection.ratio.m = 1,
best.selection.ratio.f = NULL,
```

```
best.selection.criteria.m = "bv",
best.selection.criteria.f = NULL,
best.selection.manual.ratio.m = NULL,
best.selection.manual.ratio.f = NULL,
best.selection.manual.reorder = TRUE,
bve.class = NULL,
parallel.generation = FALSE,
name.cohort = NULL,
display.progress = TRUE,
combine = FALSE,
repeat.mating = NULL,
repeat.mating.copy = NULL,
repeat.mating.fixed = NULL,
repeat.mating.overwrite = TRUE,
time.point = 0,
creating.type = 0,
multiple.observation = FALSE,
new.bv.observation = NULL,
new.bv.observation.gen = NULL,
new.bv.observation.cohorts = NULL,
new.bv.observation.database = NULL,
phenotyping = NULL,
phenotyping.gen = NULL,
phenotyping.cohorts = NULL,
phenotyping.database = NULL,
bve.gen = NULL,
bve.cohorts = NULL,
bve.database = NULL,
sigma.e.gen = NULL,
sigma.e.cohorts = NULL,
sigma.e.database = NULL,
sigma.g.gen = NULL,
sigma.g.cohorts = NULL,
sigma.g.database = NULL,
gwas.gen = NULL,
gwas.cohorts = NULL,
gwas.database = NULL,
bve.insert.gen = NULL,
bve.insert.cohorts = NULL,
bve.insert.database = NULL,
reduced.selection.panel.m = NULL,
reduced.selection.panel.f = NULL,
breeding.all.combination = FALSE,
depth.pedigree = 7,
depth.pedigree.ogc = 7,
copy.individual.keep.bve = TRUE,
copy.individual.keep.pheno = TRUE,
bve.avoid.duplicates = TRUE,
```

```
report.accuracy = TRUE,
share.genotyped = 1,
singlestep.active = FALSE,
remove.non.genotyped = TRUE,
added.genotyped = 0,
fast.uhat = TRUE,
offspring.bve.parents.gen = NULL,
offspring.bve.parents.database = NULL,
offspring.bve.parents.cohorts = NULL,
offspring.bve.offspring.gen = NULL,
offspring.bve.offspring.database = NULL,
offspring.bve.offspring.cohorts = NULL,
culling.gen = NULL,
culling.database = NULL,
culling.cohort = NULL,
culling.time = Inf,
culling.name = "Not_named",
culling.bv1 = 0,
culling.share1 = 0,
culling.bv2 = NULL,
culling.share2 = NULL,
culling.index = 0,
culling.single = TRUE,
culling.all.copy = TRUE,
calculate.reliability = FALSE,
selection.m.gen = NULL,
selection.f.gen = NULL,
selection.m.database = NULL,
selection.f.database = NULL,
selection.m.cohorts = NULL,
selection.f.cohorts = NULL,
selection.m.miesenberger = FALSE,
selection.f.miesenberger = NULL,
selection.miesenberger.reliability.est = "estimated",
miesenberger.trafo = 0,
multiple.bve.weights.m = 1,
multiple.bve.weights.f = NULL,
multiple.bve.scale.m = "bv_sd",
multiple.bve.scale.f = NULL,
verbose = TRUE,
bve.parent.mean = FALSE,
bve.grandparent.mean = FALSE,
bve.mean.between = "bvpheno",
bve.direct.est = TRUE,
bve.pseudo = FALSE,
bve.pseudo.accuracy = 1,
miraculix.destroyA = TRUE,
mas.bve = FALSE,
```

```

mas.markers = NULL,
mas.number = 5,
mas.effects = NULL,
threshold.selection = NULL,
threshold.sign = ">",
input.phenotype = "own",
bve.ignore.traits = NULL,
bv.ignore.traits = NULL,
genotyped.database = NULL,
genotyped.gen = NULL,
genotyped.cohorts = NULL,
genotyped.share = 1,
genotyped.array = 1,
sex.s = NULL,
bve.imputation = TRUE,
bve.imputation.errorrate = 0,
share.phenotyped = 1,
avoid.mating.fullsib = FALSE,
avoid.mating.halfsib = FALSE,
max.mating.pair = Inf,
bve.per.sample.sigma.e = TRUE,
bve.solve = "exact"
)

```

Arguments

population	Population list
mutation.rate	Mutation rate in each marker (default: 10 ⁻⁸)
remutation.rate	Remutation rate in each marker (default: 10 ⁻⁸)
recombination.rate	Average number of recombination per 1 length unit (default: 1M)
selection.m	Selection criteria for male individuals (Set to "random" to randomly select individuals - this happens automatically when no the input in selection.criteria has no input ((usually breeding values)))
selection.f	Selection criteria for female individuals (default: selection.m , alt: "random", function")
new.selection.calculation	If TRUE recalculate breeding values obtained by selection.function.matrix
selection.function.matrix	Manuel generation of a temporary selection function (Use BVs instead!)
selection.size	Number of selected individuals for breeding (default: c(0,0) - alt: positive numbers)
ignore.best	Not consider the top individuals of the selected individuals (e.g. to use 2-10 best individuals)
breeding.size	Number of individuals to generate

breeding.sex Share of female animals (if single value is used for breeding size; default: 0.5)

breeding.sex.random If TRUE randomly chose sex of new individuals (default: FALSE - use expected values)

relative.selection Use best.selection.ratio instead!

class.m Migrationlevels of male individuals to consider for mating process (default: 0)

class.f Migrationlevels of female individuals to consider for mating process (default: 0)

add.gen Generation you want to add the new individuals to (default: New generation)

recom.f.indicator Use step function for recombination map (transform snp.positions if possible instead)

duplication.rate Share of recombination points with a duplication (default: 0 - DEACTIVATED)

duplication.length Average length of a duplication (Exponentially distributed)

duplication.recombination Average number of recombinations per 1 length unit of duplication (default: 1)

new.class Migration level of newly generated individuals (default: 0)

bve If TRUE perform a breeding value estimation (default: FALSE)

sigma.e Environmental variance (default: 100)

sigma.g Genetic variance (default: 100 - only used if not computed via estimate.sigma.g² in der Zuchtwertschaetzung (Default: 100)

new.bv.child (OLD! - use phenotyping.child) Starting phenotypes of newly generated individuals (default: "mean" of both parents, "obs" - regular observation, "zero" - 0)

phenotyping.child Starting phenotypes of newly generated individuals (default: "mean" of both parents, "obs" - regular observation, "zero" - 0)

relationship.matrix Method to calculate relationship matrix for the breeding value estimation (Default: "vanRaden", alt: "kinship", "CE", "non_stand", "CE2", "CM")

relationship.matrix.ogc Method to calculate relationship matrix for OGC (Default: "kinship", alt: "vanRaden", "CE", "non_stand", "CE2", "CM")

computation.A (OLD! - use relationship.matrix) Method to calculate relationship matrix for the breeding value estimation (Default: "vanRaden", alt: "kinship", "CE", "non_stand", "CE2", "CM")

computation.A.ogc (OLD! use relationship.matrix.ogc) Method to calculate pedigree matrix in OGC (Default: "kinship", alt: "vanRaden", "CE", "non_stand", "CE2", "CM")

delete.haplotypes Generations for with haplotypes of founders can be deleted (only use if storage problem!)

delete.individuals	Generations for with individuals are completely deleted (only use if storage problem!)
fixed.breeding	Set of targeted matings to perform
fixed.breeding.best	Perform targeted matings in the group of selected individuals
max.offspring	Maximum number of offspring per individual (default: c(Inf,Inf) - (m,w))
max.litter	Maximum number of offspring per individual (default: c(Inf,Inf) - (m,w))
store.breeding.totals	If TRUE store information on selected animals in \$info\$breeding.totals
forecast.sigma.g	Set FALSE to not estimate sigma.g (Default: TRUE)
multiple.bve	Way to handle multiple traits in bv/selection (default: "add", alt: "ranking")
store.bve.data	If TRUE store information of bve in \$info\$bve.data
fixed.assignment	Set TRUE for targeted mating of best-best individual till worst-worst (of selected). set to "bestworst" for best-worst mating
reduce.group	(OLD! - use culling modules) Groups of animals for reduce to a new size (by changing class to -1)
reduce.group.selection	(OLD! - use culling modules) Selection criteria for reduction of groups (cf. selection.m / selection.f - default: "random")
selection.highest	If 0 individuals with lowest bve are selected as best individuals (default c(1,1) - (m,w))
selection.criteria	What to use in the selection proces (default: "bve", alt: "bv", "pheno")
same.sex.activ	If TRUE allow matings of individuals of same sex
same.sex.sex	Probability to use female individuals as parents (default: 0.5)
same.sex.selfing	Set to TRUE to allow for selfing when using same.sex matings
selfing.mating	If TRUE generate new individuals via selfing
selfing.sex	Share of female individuals used for selfing (default: 0.5)
praeimplantation	Only use matings the lead to a specific genotype in a specific marker
heritability	Use sigma.e to obtain a certain heritability (default: NULL)
repeatability	Set this to control the share of the residual variance (sigma.e) that is permanent (there for each observation)
save.recombination.history	If TRUE store the time point of each recombination event
martini.selection	If TRUE use the group of non-selected individuals as second parent
BGLR.bve	If TRUE use BGLR to perform breeding value estimation

BGLR.model	Select which BGLR model to use (default: "RKHS", alt: "BRR", "BL", "BayesA", "BayesB", "BayesC")
BGLR.burnin	Number of burn-in steps in BGLR (default: 1000)
BGLR.iteration	Number of iterations in BGLR (default: 5000)
BGLR.print	If TRUE set verbose to TRUE in BGLR
copy.individual	If TRUE copy the selected father for a mating
copy.individual.m	If TRUE generate exactly one copy of all selected male in a new cohort (or more by setting breeding.size)
copy.individual.f	If TRUE generate exactly one copy of all selected female in a new cohort (or more by setting breeding.size)
dh.mating	If TRUE generate a DH-line in mating process
dh.sex	Share of DH-lines generated from selected female individuals
n.observation	Number of phenotypes generated per individuals (influences enviromental variance)
bve.0isNA	Individuals with phenotype 0 are used as NA in breeding value estimation
phenotype.bv	If TRUE use phenotype as estimated breeding value
delete.same.origin	If TRUE delete recombination points when genetic origin of adjacent segments is the same
remove.effect.position	If TRUE remove real QTLs in breeding value estimation
estimate.u	If TRUE estimate u in breeding value estimation ($Y = Xb + Zu + e$)
new.phenotype.correlation	(OLD! - use new.residual.correlation!) Correlation of the simulated enviromental variance
new.residual.correlation	Correlation of the simulated enviromental variance
new.breeding.correlation	Correlation of the simulated genetic variance (child share! heritage is not influenced!)
estimate.add.gen.var	If TRUE estimate additive genetic variance and heritability based on parent model
estimate.pheno.var	If TRUE estimate total variance in breeding value estimation
best1.from.group	(OLD!- use selection.m.database) Groups of individuals to consider as First Parent / Father (also female individuals are possible)
best2.from.group	(OLD!- use selection.f.database) Groups of individuals to consider as Second Parent / Mother (also male individuals are possible)

best1.from.cohort
(OLD!- use selection.m.cohorts) Groups of individuals to consider as First Parent / Father (also female individuals are possible)

best2.from.cohort
(OLD! - use selection.f.cohorts) Groups of individuals to consider as Second Parent / Mother (also male individuals are possible)

add.class.cohorts
Migration levels of all cohorts selected for reproduction are automatically added to class.m/class.f (default: TRUE)

store.comp.times
If TRUE store computation times in \$info\$comp.times (default: TRUE)

store.comp.times.bve
If TRUE store computation times of breeding value estimation in \$info\$comp.times.bve (default: TRUE)

store.comp.times.generation
If TRUE store computation times of mating simulations in \$info\$comp.times.generation (default: TRUE)

import.position.calculation
Function to calculate recombination point into adjacent/following SNP

BGLR.save Method to use in BGLR (default: "RKHS" - alt: NON currently)

BGLR.save.random
Add random number to store location of internal BGLR computations (only needed when simulating a lot in parallel!)

ogc
If TRUE use optimal genetic contribution theory to perform selection (This requires the use of the R-package optiSel)

ogc.target
Target of OGC (default: "min.sKin" - minimize inbreeding; alt: "max.BV" / "min.BV" - maximize genetic gain; both under constrains selected below)

ogc.uniform
This corresponds to the uniform constrain in optiSel

ogc.ub
This corresponds to the ub constrain in optiSel

ogc.lb
This corresponds to the lb constrain in optiSel

ogc.ub.sKin
This corresponds to the ub.sKin constrain in optiSel

ogc.lb.BV
This corresponds to the lb.BV constrain in optiSel

ogc.ub.BV
This corresponds to the ub.BV constrain in optiSel

ogc.eq.BV
This corresponds to the eq.BV constrain in optiSel

ogc.ub.sKin.increase
This corresponds to the upper bound (current sKin + ogc.ub.sKin.increase) as ub.sKin in optiSel

ogc.lb.BV.increase
This corresponds to the lower bound (current BV + ogc.lb.BV.increase) as lb.BV in optiSel

emmreml.bve
If TRUE use REML estimator from R-package EMMREML in breeding value estimation

rrblup.bve
If TRUE use REML estimator from R-package rrBLUP in breeding value estimation

sommer.bve	If TRUE use REML estimator from R-package sommer in breeding value estimation
sommer.multi.bve	Set TRUE to use a multi-trait model in the R-package sommer for BVE
nr.edits	Number of edits to perform per individual
gene.editing.offspring	If TRUE perform gene editing on newly generated individuals
gene.editing.best	If TRUE perform gene editing on selected individuals
gene.editing.offspring.sex	Which sex to perform editing on (Default c(TRUE,TRUE), mw)
gene.editing.best.sex	Which sex to perform editing on (Default c(TRUE,TRUE), mw)
gwas.u	If TRUE estimate u via GWAS (relevant for gene editing)
approx.residuals	If FALSE calculate the variance for each marker separately instead of using a set variance (doesn't change order - only p-values)
sequenceZ	Split genomic matrix into parts (relevant if high memory usage)
maxZ	Number of SNPs to consider in each part of sequenceZ
maxZtotal	Number of matrix entries to consider jointly (maxZ = maxZtotal/number of animals)
delete.sex	Remove all individuals from these sex from generation delete.individuals (default: 1:2 ; note:delete individuals=NULL)
gwas.group.standard	If TRUE standardize phenotypes by group mean
y.gwas.used	What y value to use in GWAS study (Default: "pheno", alt: "bv", "bve")
gen.architecture.m	Genetic architecture for male animal (default: 0 - no transformation)
gen.architecture.f	Genetic architecture for female animal (default: gen.architecture.m - no transformation)
add.architecture	List with two vectors containing (A: length of chromosomes, B: position in cM of SNPs)
ncore	Cores used for parallel computing in compute.snps
ncore.generation	Number of cores to use in parallel generation
Z.integer	If TRUE save Z as an integer in parallel computing
store.effect.freq	If TRUE store the allele frequency of effect markers per generation
backend	Choose the used backend (default: "doParallel", alt: "doMPI")
randomSeed	Set random seed of the process

randomSeed.generation	Set random seed for parallel generation process
Rprof	Store computation times of each function
miraculix	If TRUE use miraculix to perform computations (ideally already generate population in creating.diploid with this; default: automatic detection from population list)
miraculix.cores	Number of cores used in miraculix applications (default: 1)
miraculix.mult	If TRUE use miraculix for matrix multiplications even if miraculix is not used for storage
miraculix.chol	Set to FALSE to deactivate miraculix based Cholesky-decomposition (default: TRUE)
best.selection.ratio.m	Ratio of the frequency of the selection of the best best animal and the worst best animal (default=1)
best.selection.ratio.f	Ratio of the frequency of the selection of the best best animal and the worst best animal (default=1)
best.selection.criteria.m	Criteria to calculate this ratio (default: "bv", alt: "bve", "pheno")
best.selection.criteria.f	Criteria to calculate this ratio (default: "bv", alt: "bve", "pheno")
best.selection.manual.ratio.m	vector containing probability to draw from for every individual (e.g. c(0.1,0.2,0.7))
best.selection.manual.ratio.f	vector containing probability to draw from for every individual (e.g. c(0.1,0.2,0.7))
best.selection.manual.reorder	Set to FALSE to not use the order from best to worst selected individual but plain order based on database-order
bve.class	Consider only animals of those class classes in breeding value estimation (default: NULL - use all)
parallel.generation	Set TRUE to active parallel computing in animal generation
name.cohort	Name of the newly added cohort
display.progress	Set FALSE to not display progress bars. Setting verbose to FALSE will automatically deactivate progress bars
combine	Copy existing individuals (e.g. to merge individuals from different groups in a joined cohort). Individuals to use are used as the first parent
repeat.mating	Generate multiple mating from the same dam/sire combination (first column: number of offspring; second column: probability)
repeat.mating.copy	Generate multiple copies from a copy action (combine / copy.individuals.m/f) (first column: number of offspring; second column: probability)

<code>repeat.mating.fixed</code>	Vector containing number of times each mating is repeated. This will overwrite sampling from <code>repeat.mating / repeat.mating.copy</code> (default: NULL)
<code>repeat.mating.override</code>	Set to FALSE to not use the current <code>repeat.mating / repeat.mating.copy</code> input as the new standard values (default: TRUE)
<code>time.point</code>	Time point at which the new individuals are generated
<code>creating.type</code>	Technique to generate new individuals (usage in web-based application)
<code>multiple.observation</code>	Set TRUE to allow for more than one phenotype observation per individual (this will decrease environmental variance!)
<code>new.bv.observation</code>	(OLD! - use <code>phenotyping</code>) Quick access to phenotyping for (all: "all", non-phenotyped: "non_obs", non-phenotyped male: "non_obs_m", non-phenotyped female: "non_obs_f")
<code>new.bv.observation.gen</code>	(OLD! use <code>phenotyping.gen</code>) Vector of generation from which to generate additional phenotypes
<code>new.bv.observation.cohorts</code>	(OLD! use <code>phenotyping.cohorts</code>) Vector of cohorts from which to generate additional phenotype
<code>new.bv.observation.database</code>	(OLD! use <code>phenotyping.database</code>) Matrix of groups from which to generate additional phenotypes
<code>phenotyping</code>	Quick access to phenotyping for (all: "all", non-phenotyped: "non_obs", non-phenotyped male: "non_obs_m", non-phenotyped female: "non_obs_f")
<code>phenotyping.gen</code>	Vector of generation from which to generate additional phenotypes
<code>phenotyping.cohorts</code>	Vector of cohorts from which to generate additional phenotype
<code>phenotyping.database</code>	Matrix of groups from which to generate additional phenotypes
<code>bve.gen</code>	Generations of individuals to consider in breeding value estimation (default: NULL)
<code>bve.cohorts</code>	Cohorts of individuals to consider in breeding value estimation (default: NULL)
<code>bve.database</code>	Groups of individuals to consider in breeding value estimation (default: NULL)
<code>sigma.e.gen</code>	Generations to consider when estimating <code>sigma.e</code> when using heritability
<code>sigma.e.cohorts</code>	Cohorts to consider when estimating <code>sigma.e</code> when using heritability
<code>sigma.e.database</code>	Groups to consider when estimating <code>sigma.e</code> when using heritability
<code>sigma.g.gen</code>	Generations to consider when estimating <code>sigma.g</code>
<code>sigma.g.cohorts</code>	Cohorts to consider when estimating <code>sigma.g</code>

sigma.g.database	Groups to consider when estimating sigma.g
gwas.gen	Generations to consider in GWAS analysis
gwas.cohorts	Cohorts to consider in GWAS analysis
gwas.database	Groups to consider in GWAS analysis
bve.insert.gen	Generations of individuals to compute breeding values for (default: all groups in bve.database)
bve.insert.cohorts	Cohorts of individuals to compute breeding values for (default: all groups in bve.database)
bve.insert.database	Groups of individuals to compute breeding values for (default: all groups in bve.database)
reduced.selection.panel.m	Use only a subset of individuals of the potential selected ones ("Split in user-interface")
reduced.selection.panel.f	Use only a subset of individuals of the potential selected ones ("Split in user-interface")
breeding.all.combination	Set to TRUE to automatically perform each mating combination possible exactly ones.
depth.pedigree	Depth of the pedigree in generations (default: 7)
depth.pedigree.ogc	Depth of the pedigree in generations (default: 7)
copy.individual.keep.bve	Set to FALSE to not keep estimated breeding value in case of use of copy.individuals
copy.individual.keep.pheno	Set to FALSE to not keep estimated breeding values in case of use of copy.individuals
bve.avoid.duplicates	If set to FALSE multiple generations of the same individual can be used in the bve (only possible by using copy.individual to generate individuals)
report.accuracy	Report the accuracy of the breeding value estimation
share.genotyped	Share of individuals newly generated individuals that are genotyped
singlestep.active	Set TRUE to use single step in breeding value estimation (only implemented for vanRaden- G matrix and without use sequenceZ) (Legarra 2014)
remove.non.genotyped	Set to FALSE to manually include non-genotyped individuals in genetic BVE, single-step will deactivate this as well
added.genotyped	Share of individuals that is additionally genotyped (only for copy.individuals)

fast.uhat Set to FALSE to derive inverse of A in rrBLUP
offspring.bve.parents.gen Generations to consider to derive phenotype from offspring phenotypes
offspring.bve.parents.database Groups to consider to derive phenotype from offspring phenotypes
offspring.bve.parents.cohorts Cohorts to consider to derive phenotype from offspring phenotypes
offspring.bve.offspring.gen Active generations for import of offspring phenotypes
offspring.bve.offspring.database Active groups for import of offspring phenotypes
offspring.bve.offspring.cohorts Active cohorts for import of offspring phenotypes
culling.gen Generations to consider to culling
culling.database Groups to consider to culling
culling.cohort Cohort to consider to culling
culling.time Age of the individuals at culling
culling.name Name of the culling action (user-interface stuff)
culling.bv1 Reference Breeding value
culling.share1 Probability of death for individuals with bv1
culling.bv2 Alternative breeding value (linear extended for other bvs)
culling.share2 Probability of death for individuals with bv2
culling.index Genomic index (default:0 - no genomic impact, use: "lastindex" to use the last selection index applied in selection)
culling.single Set to FALSE to not apply the culling module on all individuals of the cohort
culling.all.copy Set to FALSE to not kill copies of the same individual in the culling module
calculate.reliability Set TRUE to calculate a reliability when performing Direct-Mixed-Model BVE
selection.m.gen Generations available for selection of paternal parent
selection.f.gen Generations available for selection of maternal parent
selection.m.database Groups available for selection of paternal parent
selection.f.database Groups available for selection of maternal parent
selection.m.cohorts Cohorts available for selection of paternal parent
selection.f.cohorts Cohorts available for selection of maternal parent

selection.m.miesenberger	Use Weighted selection index according to Miesenberger 1997 for paternal selection
selection.f.miesenberger	Use Weighted selection index according to Miesenberger 1997 for maternal selection
selection.miesenberger.reliability.est	If available reliability estimated are used. If not use default:"estimated" (SD BVE / SD Pheno), alt: "heritability", "derived" ($\text{cor}(\text{BVE}, \text{BV})^2$) as replacement
miesenberger.trafo	Ignore all eigenvalues below this threshold and apply dimension reduction (default: 0 - use all)
multiple.bve.weights.m	Weighting between traits when using "add" (default: 1)
multiple.bve.weights.f	Weighting between traits when using "add" (default: same as multiple.bve.weights.m)
multiple.bve.scale.m	Default: "bv_sd"; Set to "pheno_sd" when using gains per phenotypic SD, "unit" when using gains per unit, "bve" when using estimated breeding values
multiple.bve.scale.f	Default: "bv_sd"; Set to "pheno_sd" when using gains per phenotypic SD, "unit" when using gains per unit, "bve" when using estimated breeding values
verbose	Set to FALSE to not display any prints
bve.parent.mean	Set to TRUE to use the average parental performance as the breeding value estimate
bve.grandparent.mean	Set to TRUE to use the average grandparental performance as the breeding value estimate
bve.mean.between	Select if you want to use the "bve", "bv", "pheno" or "bvpheno" to form the mean (default: "bvpheno" - if available bve, else pheno)
bve.direct.est	If TRUE predict BVEs in direct estimation according to vanRaden 2008 method 2 (default: TRUE)
bve.pseudo	If set to TRUE the breeding value estimation will be simulated with resulting accuracy bve.pseudo.accuracy (default: 1)
bve.pseudo.accuracy	The accuracy to be obtained in the "pseudo" - breeding value estimation
miraculix.destroyA	If FALSE A will not be destroyed in the process of inversion (less computing / more memory)
mas.bve	If TRUE use marker assisted selection in the breeding value estimation
mas.markers	Vector containing markers to be used in marker assisted selection
mas.number	If no markers are provided this nr of markers is selected (if single marker QTL are present highest effect markers are prioritized)

mas.effects	Effects assigned to the MAS markers (Default: estimated via lm())
threshold.selection	Minimum value in the selection index selected individuals have to have
threshold.sign	Pick all individuals above (" $>$ ") the threshold. Alt: (" $<$ ", "=", " $<=$ ", " $>=$ ")
input.phenotype	Select what to use in BVE (default: own phenotype ("own"), offspring phenotype ("off"), their average ("mean") or a weighted average ("weighted"))
bve.ignore.traits	Vector of traits to ignore in the breeding value estimation (default: NULL, use: "zero" to not consider traits with 0 index weight in multiple.bve.weights.m/w)
bv.ignore.traits	Vector of traits to ignore in the calculation of the genomic value (default: NULL; Only recommended for high number of traits and experienced users!)
genotyped.database	Groups to generate genotype data (that can be used in a BVE)
genotyped.gen	Generations to generate genotype data (that can be used in a BVE)
genotyped.cohorts	Cohorts to generate genotype data (that can be used in a BVE)
genotyped.share	Share of individuals in genotyped.gen/database/cohort to generate genotype data from (default: 1)
genotyped.array	Genotyping array used
sex.s	Specify which newly added individuals are male (1) or female (2)
bve.imputation	Set to FALSE to not perform imputation up to the highest marker density of genotyping data that is available
bve.imputation.errorrate	Share of errors in the imputation procedure (default: 0)
share.phenotyped	Share of the individuals to phenotype
avoid.mating.fullsib	Set to TRUE to not generate offspring of full siblings
avoid.mating.halfsib	Set to TRUE to not generate offspring from half or full siblings
max.mating.pair	Set to the maximum number of matings between two individuals (default: Inf)
bve.per.sample.sigma.e	Set to FALSE to deactivate the use of a heritability based on the number of observations generated per sample
bve.solve	Provide solver to be used in BVE (default: "exact" solution via inversion, alt: "pcg", function with inputs A, b and output $y_{\hat{}}$)

Value

Population-list

Examples

```
population <- creating.diploid(nsnp=1000, nindi=100)
population <- breeding.diploid(population, breeding.size=100, selection.size=c(25,25))
```

breeding.intern	<i>Internal function to simulate one meiosis</i>
-----------------	--

Description

Internal function to simulate one meiosis

Usage

```
breeding.intern(
  info.parent,
  parent,
  population,
  mutation.rate = 10^-5,
  remutation.rate = 10^-5,
  recombination.rate = 1,
  recom.f.indicator = NULL,
  duplication.rate = 0,
  duplication.length = 0.01,
  duplication.recombination = 1,
  delete.same.origin = FALSE,
  gene.editing = FALSE,
  nr.edits = 0,
  gen.architecture = 0,
  decodeOriginsU = MoBPS::decodeOriginsR
)
```

Arguments

info.parent	position of the parent in the dataset
parent	list of information regarding the parent
population	Population list
mutation.rate	Mutation rate in each marker (default: 10 ⁻⁵)
remutation.rate	Remutation rate in each marker (default: 10 ⁻⁵)
recombination.rate	Average number of recombination per 1 length unit (default: 1M)
recom.f.indicator	Use step function for recombination map (transform snp.positions if possible instead)

`duplication.rate` Share of recombination points with a duplication (default: 0 - DEACTIVATED)
`duplication.length` Average length of a duplication (Exponentially distributed)
`duplication.recombination` Average number of recombinations per 1 length unit of duplication (default: 1)
`delete.same.origin` If TRUE delete recombination points when genetic origin of adjacent segments is the same
`gene.editing` If TRUE perform gene editing on newly generated individual
`nr.edits` Number of edits to perform per individual
`gen.architecture` Used underlying genetic architecture (genome length in M)
`decodeOriginsU` Used function for the decoding of genetic origins `[[5]]/[[6]]`

Value

Inherited parent gamete

Examples

```

data(ex_pop)
child_gamete <- breeding.intern(info.parent = c(1,1,1), parent = ex_pop$breeding[[1]][[1]][[1]],
                             population = ex_pop)

```

 bv.development

Development of genetic/breeding value

Description

Function to plot genetic/breeding values for multiple generation/cohorts

Usage

```

bv.development(
  population,
  database = NULL,
  gen = NULL,
  cohorts = NULL,
  confidence = c(1, 2, 3),
  development = c(1, 2, 3),
  quantile = 0.95,
  bvrow = "all",
  ignore.zero = TRUE,
  json = FALSE,
  display.time.point = FALSE,

```

```

display.creating.type = FALSE,
display.cohort.name = FALSE,
display.sex = FALSE,
equal.spacing = FALSE,
time_reorder = FALSE,
display.line = TRUE,
ylim = NULL,
fix_mfrow = FALSE
)

```

Arguments

population	population list
database	Groups of individuals to consider for the export
gen	Quick-insert for database (vector of all generations to export)
cohorts	Quick-insert for database (vector of names of cohorts to export)
confidence	Draw confidence intervals for (1- bv, 2- bve, 3- pheno; default: c(1,2,3))
development	Include development of (1- bv, 2- bve, 3- pheno; default: c(1,2,3))
quantile	Quantile of the confidence interval to draw (default: 0.05)
bvrow	Which traits to display (for multiple traits separate plots (par(mfrow)))
ignore.zero	Cohorts with only 0 individuals are not displayed (default: TRUE)
json	If TRUE extract which cohorts to plot according to the json-file used in json.simulation
display.time.point	Set TRUE to use time point of generated to sort groups
display.creating.type	Set TRUE to show Breedingtype used in generation (web-interface)
display.cohort.name	Set TRUE to display the name of the cohort in the x-axis
display.sex	Set TRUE to display the creating.type (Shape of Points - web-based-application)
equal.spacing	Equal distance between groups (independent of time.point)
time_reorder	Set TRUE to order cohorts according to the time point of generation
display.line	Set FALSE to not display the line connecting cohorts
ylim	Set this to fix the y-axis of the plot
fix_mfrow	Set TRUE to not use mfrow - use for custom plots

Value

Genomic values of selected gen/database/cohort

Examples

```

data(ex_pop)
bv.development(ex_pop, gen=1:5)

```

 bv.development.box *Development of genetic/breeding value using a boxplot*

Description

Function to plot genetic/breeding values for multiple generation/cohorts using box plots

Usage

```

bv.development.box(
  population,
  database = NULL,
  gen = NULL,
  cohorts = NULL,
  bvrow = "all",
  json = FALSE,
  display = "bv",
  display.selection = FALSE,
  display.reproduction = FALSE,
  ylim = NULL,
  fix_mfrow = FALSE
)

```

Arguments

population	population list
database	Groups of individuals to consider for the export
gen	Quick-insert for database (vector of all generations to export)
cohorts	Quick-insert for database (vector of names of cohorts to export)
bvrow	Which traits to display (for multiple traits separate plots (par(mfrow)))
json	If TRUE extract which cohorts to plot according to the json-file used in json.simulation
display	Choose between "bv", "pheno", "bve" (default: "bv")
display.selection	Display lines between generated cohorts via selection (webinterface)
display.reproduction	Display lines between generated cohorts via reproduction (webinterface)
ylim	Set this to fix the y-axis of the plot
fix_mfrow	Set TRUE to not use mfrow - use for custom plots

Value

Genomic values of selected gen/database/cohort

Examples

```
data(ex_pop)
bv.development.box(ex_pop, gen=1:5)
```

bv.standardization	<i>BV standardization</i>
--------------------	---------------------------

Description

Function to get mean and genetic variance of a trait to a fixed value

Usage

```
bv.standardization(
  population,
  mean.target = 100,
  var.target = 10,
  gen = NULL,
  database = NULL,
  cohorts = NULL,
  adapt.bve = FALSE,
  adapt.pheno = FALSE,
  verbose = FALSE
)
```

Arguments

population	Population list
mean.target	Target mean
var.target	Target variance
gen	Quick-insert for database (vector of all generations to export)
database	Groups of individuals to consider for the export
cohorts	Quick-insert for database (vector of names of cohorts to export)
adapt.bve	Modify previous breeding value estimations by scaling (default: FALSE)
adapt.pheno	Modify previous phenotypes by scaling (default: FALSE)
verbose	Set to TRUE to display prints

Value

Population-list with scaled QTL-effects

Examples

```
population <- creating.diploid(nsnp=1000, nindi=100, n.additive=100)
population <- bv.standardization(population, mean.target=200, var.target=5)
```

 calculate.bv

Calculate breeding values

Description

Internal function to calculate the breeding value of a given individual

Usage

```
calculate.bv(
  population,
  gen,
  sex,
  nr,
  activ_bv,
  import.position.calculation = NULL,
  decodeOriginsU = decodeOriginsR,
  store.effect.freq = FALSE,
  bit.storing = FALSE,
  nbits = 30,
  output_compressed = FALSE,
  bv.ignore.traits = NULL
)
```

Arguments

population	Population list
gen	Generation of the individual of interest
sex	Sex of the individual of interest
nr	Number of the individual of interest
activ_bv	traits to consider
import.position.calculation	Function to calculate recombination point into adjacent/following SNP
decodeOriginsU	Used function for the decoding of genetic origins [[5]]/[[6]]
store.effect.freq	If TRUE store the allele frequency of effect markers per generation
bit.storing	Set to TRUE if the MoBPS (not-miraculix! bit-storing is used)
nbits	Bits available in MoBPS-bit-storing
output_compressed	Set to TRUE to get a miraculix-compressed genotype/haplotype
bv.ignore.traits	Vector of traits to ignore in the calculation of the genomic value (default: NULL; Only recommended for high number of traits and experienced users!)

Value

[[1]] true genomic value [[2]] allele frequency at QTL markers

Examples

```
data(ex_pop)
calculate.bv(ex_pop, gen=1, sex=1, nr=1, activ_bv = 1)
```

cattle_chip

Cattle chip

Description

Genome for cattle according to Ma et al.

Usage

```
cattle_chip
```

Author(s)

Torsten Pook <torsten.pook@uni-goettingen.de>

Source

Ma et al 2015

check.parents

Relatedness check between two individuals

Description

Internal function to check the relatedness between two individuals

Usage

```
check.parents(population, info.father, info.mother, max.rel = 2)
```

Arguments

population	Population list
info.father	position of the first parent in the dataset
info.mother	position of the second parent in the dataset
max.rel	maximal allowed relationship (default: 2, alt: 1 no full-sibs, 0 no half-sibs)

Value

logical with TRUE if relatedness does not exceed max.rel / FALSE otherwise.

Examples

```
data(ex_pop)
check.parents(ex_pop, info.father=c(4,1,1,1), info.mother=c(4,2,1,1))
```

chicken_chip	<i>chicken chip</i>
--------------	---------------------

Description

Genome for chicken according to Groenen et al.

Usage

```
chicken_chip
```

Author(s)

Torsten Pook <torsten.pook@uni-goettingen.de>

Source

Groenen et al 2009

clean.up	<i>Clean-up recombination points</i>
----------	--------------------------------------

Description

Function to remove recombination points + origins with no influence on markers

Usage

```
clean.up(population, gen = "all", database = NULL, cohorts = NULL)
```

Arguments

population	Population list
gen	Generations to clean up (default: "current")
database	Groups of individuals to consider
cohorts	Quick-insert for database (vector of names of cohorts to export)

Value

Population-list with deleted irrelevant recombination points

Examples

```
data(ex_pop)
ex_pop <- clean.up(ex_pop)
```

codeOriginsR	<i>Origins-coding(R)</i>
--------------	--------------------------

Description

R-Version of the internal bitwise-coding of origins

Usage

```
codeOriginsR(M)
```

Arguments

M Origins matrix

Value

Bit-wise coded origins

Examples

```
codeOriginsR(cbind(1,1,1,1))
```

combine.traits	<i>Combine traits</i>
----------------	-----------------------

Description

Function to combine traits in the BVE

Usage

```
combine.traits(  
  population,  
  combine.traits = NULL,  
  combine.name = NULL,  
  remove.combine = NULL,  
  remove.all = FALSE  
)
```

Arguments

population Population list
 combine.traits Vector containing the traits (numbers) to combine into a joined trait
 combine.name Name of the combined trait
 remove.combine Remove a selected previously generated combined trait
 remove.all Set TRUE to remove all previously generated combined traits

Value

Population-list

Examples

```

population <- creating.diploid(nsnp=100, nindi=100, n.additive = c(50,50))
population <- combine.traits(population, combine.traits=1:2)
population <- breeding.diploid(population, bve=TRUE, phenotyping.gen=1, heritability=0.3)

```

compute.costs	<i>Compute costs of a breeding program</i>
---------------	--

Description

Function to derive the costs of a breeding program / population-list

Usage

```

compute.costs(
  population,
  phenotyping.costs = 10,
  genotyping.costs = 100,
  fix.costs = 0,
  fix.costs.annual = 0,
  profit.per.bv = 1,
  database = NULL,
  gen = NULL,
  cohorts = NULL,
  interest.rate = 1,
  base.gen = 1
)

```

Arguments

population population-list
 phenotyping.costs
 Costs for the generation of a phenotype

genotyping.costs	Costs for the generation of a genotype
fix.costs	one time occurring fixed costs
fix.costs.annual	annually occurring fixed costs
profit.per.bv	profit generated by bv per animal
database	Groups of individuals to consider
gen	Quick-insert for database (vector of all generations to consider)
cohorts	Quick-insert for database (vector of names of cohorts to consider)
interest.rate	Applied yearly interest rate
base.gen	Base generation (application of interest rate)

Value

Cost-table for selected gen/database/cohorts of a population-list

Examples

```
data(ex_pop)
compute.costs(ex_pop, gen=1:5)
```

compute.costs.cohorts *Compute costs of a breeding program by cohorts*

Description

Function to derive the costs of a breeding program / population-list by cohorts

Usage

```
compute.costs.cohorts(
  population,
  gen = NULL,
  database = NULL,
  cohorts = NULL,
  json = TRUE,
  phenotyping.costs = NULL,
  genotyping.costs = 0,
  housing.costs = NULL,
  fix.costs = 0,
  fix.costs.annual = 0,
  profit.per.bv = 1,
  interest.rate = 1,
  verbose = TRUE
)
```

Arguments

population	population-list
gen	Quick-insert for database (vector of all generations to consider)
database	Groups of individuals to consider
cohorts	Quick-insert for database (vector of names of cohorts to consider)
json	If TRUE extract which cohorts to plot according to the json-file used in json.simulation
phenotyping.costs	Costs for the generation of a phenotype
genotyping.costs	Costs for the generation of a genotype
housing.costs	Costs for housing
fix.costs	one time occuring fixed costs
fix.costs.annual	annually occuring fixed costs
profit.per.bv	profit generated by bv per animal
interest.rate	Applied yearly interest rate
verbose	Set to FALSE to not display any prints

Value

Cost-table for selected gen/database/cohorts of a population-list

Examples

```
data(ex_pop)
compute.costs.cohorts(ex_pop, gen=1:5, genotyping.costs=25, json=FALSE)
```

compute.snps	<i>Compute genotype/haplotype</i>
--------------	-----------------------------------

Description

Internal function for the computation of genotypes & haplotypes

Usage

```
compute.snps(
  population,
  gen,
  sex,
  nr,
  faster = TRUE,
  import.position.calculation = NULL,
  from_p = 1,
```

```

    to_p = Inf,
    decodeOriginsU = decodeOriginsR,
    bit.storing = FALSE,
    nbits = 30,
    output_compressed = FALSE
  )

```

Arguments

population	Population list
gen	Generation of the individual to compute
sex	Gender of the individual to compute
nr	Number of the individual to compute
faster	If FALSE use slower version to compute markers between recombination points
import.position.calculation	Function to calculate recombination point into adjacent/following SNP
from_p	First SNP to consider
to_p	Last SNP to consider
decodeOriginsU	Used function for the decoding of genetic origins [[5]]/[[6]]
bit.storing	Set to TRUE if the MoBPS (not-miraculix! bit-storing is used)
nbits	Bits available in MoBPS-bit-storing
output_compressed	Set to TRUE to get a miraculix-compressed genotype/haplotype

Value

haplotypes for the selected individual

Examples

```

data(ex_pop)
compute.snps(ex_pop, gen=1, sex=1, nr=1)

```

compute.snps_single *Compute genotype/haplotype in gene editing application*

Description

Internal function for the computation of genotypes & haplotypes in gene editing application

Usage

```
compute.snps_single(
  population,
  current.recombi,
  current.mut,
  current.ursprung,
  faster = TRUE,
  import.position.calculation = NULL,
  decodeOriginsU = decodeOriginsR
)
```

Arguments

population Population list
current.recombi vector of currently activ recombination points
current.mut vector of currently activ mutations
current.ursprung vector of currently activ origins
faster If FALSE use slower version to compute markers between recombination points
import.position.calculation Function to calculate recombination point into adjacent/following SNP
decodeOriginsU Used function for the decoding of genetic origins [[5]]/[[6]]

Value

haplotypes for the selected individual

creating.diploid *Generation of the starting population*

Description

Generation of the starting population

Usage

```
creating.diploid(
  dataset = NULL,
  vcf = NULL,
  chr.nr = NULL,
  bp = NULL,
  snp.name = NULL,
  hom0 = NULL,
  hom1 = NULL,
```



```
bpcm.conversion = 0,
nsnp = 0,
nindi = 0,
freq = "beta",
population = NULL,
sex.s = "fixed",
add.chromosome = FALSE,
generation = 1,
class = 0L,
sex.quota = 0.5,
chromosome.length = NULL,
length.before = 5,
length.behind = 5,
real.bv.add = NULL,
real.bv.mult = NULL,
real.bv.dice = NULL,
snps.equidistant = NULL,
change.order = FALSE,
bv.total = 0,
polygenic.variance = 100,
bve.mult.factor = NULL,
bve.poly.factor = NULL,
base.bv = NULL,
add.chromosome.ends = TRUE,
new.phenotype.correlation = NULL,
new.residual.correlation = NULL,
new.breeding.correlation = NULL,
add.architecture = NULL,
snp.position = NULL,
position.scaling = FALSE,
bit.storing = FALSE,
nbits = 30,
randomSeed = NULL,
miraculix = TRUE,
miraculix.dataset = TRUE,
n.additive = 0,
n.equal.additive = 0,
n.dominant = 0,
n.equal.dominant = 0,
n.qualitative = 0,
n.quantitative = 0,
dominant.only.positive = FALSE,
var.additive.l = NULL,
var.dominant.l = NULL,
var.qualitative.l = NULL,
var.quantitative.l = NULL,
effect.size.equal.add = 1,
effect.size.equal.dom = 1,
```

```

exclude.snps = NULL,
replace.real.bv = FALSE,
shuffle.traits = NULL,
shuffle.cor = NULL,
skip.rest = FALSE,
enter.bv = TRUE,
name.cohort = NULL,
template.chip = NULL,
beta.shape1 = 1,
beta.shape2 = 1,
time.point = 0,
creating.type = 0,
trait.name = NULL,
share.genotyped = 1,
genotyped.s = NULL,
map = NULL,
remove.invalid.qtl = TRUE,
verbose = TRUE,
bv.standard = FALSE,
mean.target = NULL,
var.target = NULL,
is.maternal = NULL,
is.paternal = NULL,
vcf.maxsnp = Inf,
internal = FALSE
)

```

Arguments

dataset	SNP dataset, use "random", "allhetero" "all0" when generating a dataset via nsnp,nindi
vcf	Path to a vcf-file used as input genotypes (correct haplotype phase is assumed!)
chr.nr	Vector containing the associated chromosome for each marker (default: all on the same)
bp	Vector containing the physical position (bp) for each marker (default: 1,2,3...)
snp.name	Vector containing the name of each marker (default ChrXSNPY - XY chosen accordingly)
hom0	Vector containing the first allelic variant in each marker (default: 0)
hom1	Vector containing the second allelic variant in each marker (default: 1)
bpcm.conversion	Convert physical position (bp) into a cM position (default: 0 - not done)
nsnp	number of markers to generate in a random dataset
nindi	number of individuals to generate in a random dataset
freq	frequency of allele 1 when randomly generating a dataset
population	Population list

sex.s	Specify which newly added individuals are male (1) or female (2)
add.chromosome	If TRUE add an additional chromosome to the dataset
generation	Generation of the newly added individuals (default: 1)
class	Migration level of the newly added individuals
sex.quota	Share of newly added female individuals (deterministic if sex.s="fixed", alt: sex.s="random")
chromosome.length	Length of the newly added chromosome (default: 5)
length.before	Length before the first SNP of the dataset (default: 5)
length.behind	Length after the last SNP of the dataset (default: 5)
real.bv.add	Single Marker effects
real.bv.mult	Two Marker effects
real.bv.dice	Multi-marker effects
snps.equidistant	Use equidistant markers (computationally faster! ; default: TRUE)
change.order	If TRUE sort markers according to given marker positions
bv.total	Number of traits (If more than traits via real.bv.X use traits with no directly underlying QTL)
polygenic.variance	Genetic variance of traits with no underlying QTL
bve.mult.factor	Multiplicate trait value times this
bve.poly.factor	Potency trait value over this
base.bv	Average genetic value of a trait
add.chromosome.ends	Add chromosome ends as recombination points
new.phenotype.correlation	(OLD! - use new.residual.correlation) Correlation of the simulated enviromental variance
new.residual.correlation	Correlation of the simulated enviromental variance
new.breeding.correlation	Correlation of the simulated genetic variance (child share! heritage is not influenced!)
add.architecture	Add genetic architecture (marker positions)
snp.position	Location of each marker on the genetic map
position.scaling	Manual scaling of snp.position
bit.storing	Set to TRUE if the MoBPS (not-miraculix! bit-storing is used)
nbits	Bits available in MoBPS-bit-storing

randomSeed	Set random seed of the process
miraculix	If TRUE use miraculix package for data storage, computations and dataset generation
miraculix.dataset	Set FALSE to deactivate miraculix package for dataset generation
n.additive	Number of additive QTL with effect size drawn from a gaussian distribution
n.equal.additive	Number of additive QTL with equal effect size (effect.size)
n.dominant	Number of dominant QTL with effect size drawn from a gaussian distribution
n.equal.dominant	Number of n.equal.dominant QTL with equal effect size
n.qualitative	Number of qualitative epistatic QTL
n.quantitative	Number of quantitative epistatic QTL
dominant.only.positive	Set to TRUE to always assign the heterozygous variant with the higher of the two homozygous effects (e.g. hybrid breeding); default: FALSE
var.additive.l	Variance of additive QTL
var.dominant.l	Variance of dominante QTL
var.qualitative.l	Variance of qualitative epistatic QTL
var.quantitative.l	Variance of quantitative epistatic QTL
effect.size.equal.add	Effect size of the QTLs in n.equal.additive
effect.size.equal.dom	Effect size of the QTLs in n.equal.dominant
exclude.snps	Marker were no QTL are simulated on
replace.real.bv	If TRUE delete the simulated traits added before
shuffle.traits	Combine different traits into a joined trait
shuffle.cor	Target Correlation between shuffeled traits
skip.rest	Internal variable needed when adding multipe chromosomes jointly
enter.bv	Internal parameter
name.cohort	Name of the newly added cohort
template.chip	Import genetic map and chip from a species ("cattle", "chicken", "pig")
beta.shape1	First parameter of the beta distribution for simulating allele frequencies
beta.shape2	Second parameter of the beta distribution for simulating allele frequencies
time.point	Time point at which the new individuals are generated
creating.type	Technique to generate new individuals (usage in web-based application)
trait.name	Name of the trait generated

share.genotyped	Share of individuals genotyped in the founders
genotyped.s	Specify with newly added individuals are genotyped (1) or not (0)
map	map-file that contains up to 5 colums (Chromosome, SNP-id, M-position, Bp-position, allele freq - Everything not provides it set to NA). A map can be imported via MoBPSmaps::ensembl.map()
remove.invalid.qtl	Set to FALSE to deactivate the automatic removal of QTLs on markers that do not exist
verbose	Set to FALSE to not display any prints
bv.standard	Set TRUE to standardize trait mean and variance via bv.standardization() - automatically set to TRUE when mean/var.target are used
mean.target	Target mean
var.target	Target variance
is.maternal	Vector coding if a trait is caused by a maternal effect (Default: all FALSE)
is.paternal	Vector coding if a trait is caused by a paternal effect (Default: all FALSE)
vcf.maxsnp	Maximum number of SNPs to include in the genotype file (default: Inf)
internal	Dont touch!

Value

Population-list

Examples

```
population <- creating.diploid(nsnp=1000, nindi=100)
```

```
creating.phenotypic.transform
```

Create a phenotypic transformation

Description

Function to perform create a transformation of phenotypes

Usage

```
creating.phenotypic.transform(  
  population,  
  phenotypic.transform.function = NULL,  
  trait = 1  
)
```

Arguments

population Population list
 phenotypic.transform.function
 Phenotypic transformation to apply
 trait Trait for which a transformation is to be applied data(ex_pop) trafo <- function(x) return(x^2) ex_pop <- creating.phenotypic.transform(ex_pop, phenotypic.transform.function=trafo)

Value

Population-list with a new phenotypic transformation function

creating.trait *Generation of genomic traits*

Description

Generation of the trait in a starting population

Usage

```

creating.trait(
  population,
  real.bv.add = NULL,
  real.bv.mult = NULL,
  real.bv.dice = NULL,
  bv.total = 0,
  polygenic.variance = 100,
  bve.mult.factor = NULL,
  bve.poly.factor = NULL,
  base.bv = NULL,
  new.phenotype.correlation = NULL,
  new.residual.correlation = NULL,
  new.breeding.correlation = NULL,
  n.additive = 0,
  n.equal.additive = 0,
  n.dominant = 0,
  n.equal.dominant = 0,
  n.qualitative = 0,
  n.quantitative = 0,
  dominant.only.positive = FALSE,
  var.additive.l = NULL,
  var.dominant.l = NULL,
  var.qualitative.l = NULL,
  var.quantitative.l = NULL,
  effect.size.equal.add = 1,
  effect.size.equal.dom = 1,

```

```

exclude.snps = NULL,
randomSeed = NULL,
shuffle.traits = NULL,
shuffle.cor = NULL,
replace.traits = FALSE,
trait.name = NULL,
remove.invalid.qtl = TRUE,
bv.standard = FALSE,
mean.target = NULL,
var.target = NULL,
verbose = TRUE,
is.maternal = NULL,
is.paternal = NULL
)

```

Arguments

population	Population list
real.bv.add	Single Marker effects
real.bv.mult	Two Marker effects
real.bv.dice	Multi-marker effects
bv.total	Number of traits (If more than traits via real.bv.X use traits with no directly underlying QTL)
polygenic.variance	Genetic variance of traits with no underlying QTL
bve.mult.factor	Multiply trait value times this
bve.poly.factor	Potency trait value over this
base.bv	Average genetic value of a trait
new.phenotype.correlation	(OLD! - use new.residual.correlation) Correlation of the simulated environmental variance
new.residual.correlation	Correlation of the simulated environmental variance
new.breeding.correlation	Correlation of the simulated genetic variance (child share! heritage is not influenced!)
n.additive	Number of additive QTL with effect size drawn from a gaussian distribution
n.equal.additive	Number of additive QTL with equal effect size (effect.size)
n.dominant	Number of dominant QTL with effect size drawn from a gaussian distribution
n.equal.dominant	Number of n.equal.dominant QTL with equal effect size
n.qualitative	Number of qualitative epistatic QTL

`n.quantitative` Number of quantitative epistatic QTL
`dominant.only.positive`
 Set to TRUE to always assign the heterozygous variant with the higher of the two
 homozygous effects (e.g. hybrid breeding); default: FALSE
`var.additive.l` Variance of additive QTL
`var.dominant.l` Variance of dominant QTL
`var.qualitative.l`
 Variance of qualitative epistatic QTL
`var.quantitative.l`
 Variance of quantitative epistatic QTL
`effect.size.equal.add`
 Effect size of the QTLs in `n.equal.additive`
`effect.size.equal.dom`
 Effect size of the QTLs in `n.equal.dominant`
`exclude.snps` Marker were no QTL are simulated on
`randomSeed` Set random seed of the process
`shuffle.traits` Combine different traits into a joined trait
`shuffle.cor` Target Correlation between shuffled traits
`replace.traits` If TRUE delete the simulated traits added before
`trait.name` Name of the trait generated
`remove.invalid.qtl`
 Set to FALSE to deactivate the automatic removal of QTLs on markers that do not
 exist
`bv.standard` Set TRUE to standardize trait mean and variance via `bv.standardization()`
`mean.target` Target mean
`var.target` Target variance
`verbose` Set to FALSE to not display any prints
`is.maternal` Vector coding if a trait is caused by a maternal effect (Default: all FALSE)
`is.paternal` Vector coding if a trait is caused by a paternal effect (Default: all FALSE)

Value

Population-list with one or more additional new traits

Examples

```

population <- creating.diploid(nsnp=1000, nindi=100)
population <- creating.trait(population, n.additive=100)

```

decodeOriginsR	<i>Origins-Decoding(R)</i>
----------------	----------------------------

Description

R-Version of the internal bitwise-decoding of origins

Usage

```
decodeOriginsR(P, row)
```

Arguments

P	coded origins vector
row	row to decode

Value

de-coded origins

Examples

```
decodeOriginsR(0L)
```

demiraculix	<i>Remove miraculix-coding for genotypes</i>
-------------	--

Description

Internal function to decode all genotypes to non-miraculix objects

Usage

```
demiraculix(population)
```

Arguments

population	Population list
------------	-----------------

Value

Population list

Examples

```
# This is only relevant with the package miraculix is installed and used
population <- creating.diploid(nsnp=100, nindi=50)
population <- demiraculix(population)
```

derive.loop.elements *Derive loop elements*

Description

Internal function to derive the position of all individuals to consider for BVE/GWAS

Usage

```
derive.loop.elements(
  population,
  bve.database,
  bve.class,
  bve.avoid.duplicates,
  store.adding = FALSE,
  store.which.adding = FALSE,
  list.of.copys = FALSE
)
```

Arguments

population	Population list
bve.database	Groups of individuals to consider in breeding value estimation
bve.class	Consider only animals of those class classes in breeding value estimation (default: NULL - use all)
bve.avoid.duplicates	If set to FALSE multiple generatations of the same individual can be used in the bve (only possible by using copy.individual to generate individuals)
store.adding	Internal parameter to derive number of added individuals per database entry (only relevant internally for GWAS)
store.which.adding	Internal parameter to derive which individuals are copy entries
list.of.copys	Internal parameter to derive further information on the copies individuals

Value

Matrix of individuals in the entered database

Examples

```
data(ex_pop)
derive.loop.elements(ex_pop, bve.database=get.database(ex_pop, gen=2),
  bve.class=NULL, bve.avoid.duplicates=TRUE)
```

diag.mobps	<i>Add a genotyping array</i>
------------	-------------------------------

Description

Function to add a genotyping array for the population

Usage

```
diag.mobps(elements)
```

Arguments

elements vector with entries to put on the diagonal of a matrix

Value

Diagonal matrix

Examples

```
diag.mobps(5)
```

edges.fromto	<i>Detection of parental/child nodes</i>
--------------	--

Description

Internal function to extract parental/child node of an edge

Usage

```
edges.fromto(edges)
```

Arguments

edges Edges of the json-file generated via the web-interface

Value

Matrix of Parent/Child-nodes for the considered edges

edit_animal	<i>Internal gene editing function</i>
-------------	---------------------------------------

Description

Internal function to perform gene editing

Usage

```
edit_animal(  
  population,  
  gen,  
  sex,  
  nr,  
  nr.edits,  
  decodeOriginsU = decodeOriginsR,  
  bit.storing = FALSE,  
  nbits = 30  
)
```

Arguments

population	Population list
gen	Generation of the individual to edit
sex	Gender of the individual to edit
nr	Number of the individual to edit
nr.edits	Number of edits to perform
decodeOriginsU	Used function for the decoding of genetic origins [[5]]/[[6]]
bit.storing	Set to TRUE if the MoBPS (not-miraculix! bit-storing is used)
nbits	Bits available in MoBPS-bit-storing

Value

animal after genome editing

effect.estimate.add *Estimation of marker effects*

Description

Function to estimate marker effects

Usage

```
effect.estimate.add(geno, pheno, map = NULL, scaling = TRUE)
```

Arguments

geno	genotype dataset (marker x individuals)
pheno	phenotype dataset (each phenotype in a row)
map	genomic map
scaling	Set FALSE to not perform variance scaling

Value

Empirical kinship matrix (IBD-based since Founders)

Examples

```
data(ex_pop)
pheno <- get.pheno(ex_pop, gen=1:5)
geno <- get.geno(ex_pop, gen=1:5)
map <- get.map(ex_pop, use.snp.nr=TRUE)
real.bv.add <- effect.estimate.add(geno, pheno, map)
```

effective.size *Estimate effective population size*

Description

Internal function to estimate the effective population size

Usage

```
effective.size(ld, dist, n)
```

Arguments

ld	ld between markers
dist	distance between markers in Morgan
n	Population size

Value

Estimated effective population size

epi	<i>Martini-Test function</i>
-----	------------------------------

Description

Internal function to perform martini test

Usage

```
epi(y, Z, G = NULL)
```

Arguments

y	y
Z	genomic information matrix
G	kinship matrix

Value

Estimated breeding values

ex_json	<i>ex_json</i>
---------	----------------

Description

Exemplary json-data

Usage

```
ex_json
```

Author(s)

Torsten Pook <torsten.pook@uni-goettingen.de>

Source

Web-interface

ex_pop	<i>ex_pop</i>
--------	---------------

Description

Exemplary population-list

Usage

ex_pop

Author(s)

Torsten Pook <torsten.pook@uni-goettingen.de>

Source

MoBPS

find.chromo	<i>Position detection (chromosome)</i>
-------------	--

Description

Internal function for the detection on which chromosome each marker is

Usage

```
find.chromo(position, length.total)
```

Arguments

position	position in the genome
length.total	Length of each chromosome

Value

Chromosome the marker is part of

find.snpbefore *Position detection (SNPs)*

Description

Internal function for the detection on which position each marker is

Usage

```
find.snpbefore(position, snp.position)
```

Arguments

position Position on the genome
snp.position Position of the SNPs on the genome

Value

SNP-position of the target position

founder.simulation *Founder simulation*

Description

Function to generate founder genotypes

Usage

```
founder.simulation(  
  nindi = 100,  
  sex.quota = 0.5,  
  nsnp = 0,  
  n.gen = 100,  
  nfinal = NULL,  
  sex.quota.final = NULL,  
  big.output = FALSE,  
  plot = TRUE,  
  display.progress = TRUE,  
  depth.pedigree = 7,  
  dataset = NULL,  
  vcf = NULL,  
  chr.nr = NULL,  
  bp = NULL,  
  snp.name = NULL,
```



```

hom0 = NULL,
hom1 = NULL,
bpcm.conversion = 0,
freq = "beta",
sex.s = "fixed",
chromosome.length = NULL,
length.before = 5,
length.behind = 5,
snps.equidistant = NULL,
change.order = FALSE,
snp.position = NULL,
position.scaling = FALSE,
bit.storing = FALSE,
nbits = 30,
randomSeed = NULL,
miraculix = TRUE,
miraculix.dataset = TRUE,
template.chip = NULL,
beta.shape1 = 1,
beta.shape2 = 1,
map = NULL,
verbose = TRUE,
vcf.maxsnp = Inf
)

```

Arguments

nindi	number of individuals to generate in a random dataset
sex.quota	Share of newly added female individuals (deterministic if sex.s="fixed", alt: sex.s="random")
nsnp	number of markers to generate in a random dataset
n.gen	Number of generations to simulate (default: 100)
nfinal	Number of final individuals to include (default: nindi)
sex.quota.final	Share of female individuals in the final generation
big.output	Set to TRUE to export map, population list and pedigree relationship
plot	Set to FALSE to not generate LD-decay plot and allele frequency spectrum
display.progress	Set FALSE to not display progress bars. Setting verbose to FALSE will automatically deactivate progress bars
depth.pedigree	Depth of the pedigree in generations (default: 7)
dataset	SNP dataset, use "random", "allhetero" "all0" when generating a dataset via nsnp,nindi
vcf	Path to a vcf-file used as input genotypes (correct haplotype phase is assumed!)
chr.nr	Vector containing the associated chromosome for each marker (default: all on the same)

bp	Vector containing the physical position (bp) for each marker (default: 1,2,3...)
snp.name	Vector containing the name of each marker (default ChrXSNPY - XY chosen accordingly)
hom0	Vector containing the first allelic variant in each marker (default: 0)
hom1	Vector containing the second allelic variant in each marker (default: 1)
bpcm.conversion	Convert physical position (bp) into a cM position (default: 0 - not done)
freq	frequency of allele 1 when randomly generating a dataset
sex.s	Specify which newly added individuals are male (1) or female (2)
chromosome.length	Length of the newly added chromosome (default: 5)
length.before	Length before the first SNP of the dataset (default: 5)
length.behind	Length after the last SNP of the dataset (default: 5)
snps.equidistant	Use equidistant markers (computationally faster! ; default: TRUE)
change.order	If TRUE sort markers according to given marker positions
snp.position	Location of each marker on the genetic map
position.scaling	Manual scaling of snp.position
bit.storing	Set to TRUE if the MoBPS (not-miraculix! bit-storing is used)
nbits	Bits available in MoBPS-bit-storing
randomSeed	Set random seed of the process
miraculix	If TRUE use miraculix package for data storage, computations and dataset generation
miraculix.dataset	Set FALSE to deactivate miraculix package for dataset generation
template.chip	Import genetic map and chip from a species ("cattle", "chicken", "pig")
beta.shape1	First parameter of the beta distribution for simulating allele frequencies
beta.shape2	Second parameter of the beta distribution for simulating allele frequencies
map	map-file that contains up to 5 columns (Chromosome, SNP-id, M-position, Bp-position, allele freq - Everything not provides it set to NA). A map can be imported via MoBPSmaps::ensembl.map()
verbose	Set to FALSE to not display any prints
vcf.maxsnp	Maximum number of SNPs to include in the genotype file (default: Inf)

Examples

```
population <- founder.simulation(nindi=100, nsnp=1000, n.gen=5)
```

generation.individual *Function to generate a new individual*

Description

Function to generate a new individual

Usage

```
generation.individual(  
  indexb,  
  population,  
  info_father_list,  
  info_mother_list,  
  copy.individual,  
  mutation.rate,  
  remutation.rate,  
  recombination.rate,  
  recom.f.indicator,  
  duplication.rate,  
  duplication.length,  
  duplication.recombination,  
  delete.same.origin,  
  gene.editing,  
  nr.edits,  
  gen.architecture.m,  
  gen.architecture.f,  
  decodeOriginsU,  
  current.gen,  
  save.recombination.history,  
  new.bv.child,  
  dh.mating,  
  share.genotyped,  
  added.genotyped,  
  genotyped.array,  
  dh.sex,  
  n.observation  
)
```

Arguments

indexb	windows parallel internal test
population	windows parallel internal test
info_father_list	windows parallel internal test
info_mother_list	windows parallel internal test

```

copy.individual      windows parallel internal test
mutation.rate       windows parallel internal test
remutation.rate     windows parallel internal test
recombination.rate  windows parallel internal test
recom.f.indicator   windows parallel internal test
duplication.rate    windows parallel internal test
duplication.length  windows parallel internal test
duplication.recombination windows parallel internal test
delete.same.origin  windows parallel internal test
gene.editing        windows parallel internal test
nr.edits            windows parallel internal test
gen.architecture.m  windows parallel internal test
gen.architecture.f  windows parallel internal test
decodeOriginsU     windows parallel internal test
current.gen         windows parallel internal test
save.recombination.history windows parallel internal test
new.bv.child        windows parallel internal test
dh.mating           windows parallel internal test
share.genotyped     windows parallel internal test
added.genotyped     windows parallel internal test
genotyped.array     windows parallel internal test
dh.sex              windows parallel internal test
n.observation       windows parallel internal test

```

Value

Offspring individual

get.admixture	<i>Admixture Plot</i>
---------------	-----------------------

Description

Function to generate admixture plots

Usage

```
get.admixture(  
  population,  
  geno = NULL,  
  gen = NULL,  
  database = NULL,  
  cohorts = NULL,  
  d = NULL,  
  verbose = TRUE,  
  plot = TRUE,  
  sort = FALSE,  
  sort.cutoff = 0.01  
)
```

Arguments

population	Population list
geno	Manually provided genotype dataset to use instead of gen/database/cohorts
gen	Quick-insert for database (vector of all generations to consider)
database	Groups of individuals to consider
cohorts	Quick-insert for database (vector of names of cohorts to consider)
d	dimensions to consider in admixture plot (default: automatically estimate a reasonable number)
verbose	Set to FALSE to not display any prints
plot	Set to FALSE to not generate an admixture plot
sort	Set to TRUE to sort individuals according to contributes from the first dimension
sort.cutoff	Skip individuals with contributions under this threshold (and use next dimension instead) data(ex_pop) get.admixture(ex_pop, gen=4:6, d=2, sort=TRUE)

Value

Matrix with admixture proportion

get.age.point	<i>Derive age point</i>
---------------	-------------------------

Description

Function to derive age point for each individual (Same as time.point unless copy.individual is used for aging)

Usage

```
get.age.point(  
  population,  
  database = NULL,  
  gen = NULL,  
  cohorts = NULL,  
  use.id = FALSE  
)
```

Arguments

population	Population list
database	Groups of individuals to consider for the export
gen	Quick-insert for database (vector of all generations to export)
cohorts	Quick-insert for database (vector of names of cohorts to export)
use.id	Set to TRUE to use MoBPS ids instead of Sex_Nr_Gen based names (default: FALSE)

Value

Time point selected gen/database/cohorts-individuals are born

Examples

```
data(ex_pop)  
get.age.point(ex_pop, gen=2)
```

get.bv *Export underlying true breeding values*

Description

Function to export underlying true breeding values

Usage

```
get.bv(population, database = NULL, gen = NULL, cohorts = NULL, use.id = FALSE)
```

Arguments

population	Population list
database	Groups of individuals to consider for the export
gen	Quick-insert for database (vector of all generations to export)
cohorts	Quick-insert for database (vector of names of cohorts to export)
use.id	Set to TRUE to use MoBPS ids instead of Sex_Nr_Gen based names (default: FALSE)

Value

Genomic value of in gen/database/cohorts selected individuals

Examples

```
data(ex_pop)
get.bv(ex_pop, gen=2)
```

get.bve *Export estimated breeding values*

Description

Function to export estimated breeding values

Usage

```
get.bve(  
  population,  
  database = NULL,  
  gen = NULL,  
  cohorts = NULL,  
  use.id = FALSE  
)
```

Arguments

population	Population list
database	Groups of individuals to consider for the export
gen	Quick-insert for database (vector of all generations to export)
cohorts	Quick-insert for database (vector of names of cohorts to export)
use.id	Set to TRUE to use MoBPS ids instead of Sex_Nr_Gen based names (default: FALSE)

Value

Estimated breeding value of in gen/database/cohorts selected individuals

Examples

```
data(ex_pop)
get.bve(ex_pop, gen=2)
```

get.class	<i>Derive class</i>
-----------	---------------------

Description

Function to devide the class for each individual

Usage

```
get.class(
  population,
  database = NULL,
  gen = NULL,
  cohorts = NULL,
  use.id = FALSE
)
```

Arguments

population	Population list
database	Groups of individuals to consider for the export
gen	Quick-insert for database (vector of all generations to export)
cohorts	Quick-insert for database (vector of names of cohorts to export)
use.id	Set to TRUE to use MoBPS ids instead of Sex_Nr_Gen based names (default: FALSE)

Value

Class of in gen/database/cohorts selected individuals

Examples

```
data(ex_pop)
get.class(ex_pop, gen=2)
```

get.cohorts	<i>Export Cohort-names</i>
-------------	----------------------------

Description

Function to export cohort names for the population list

Usage

```
get.cohorts(population, extended = FALSE)
```

Arguments

population	Population list
extended	extended cohorts

Value

List of all cohorts in the population-list

Examples

```
data(ex_pop)
get.cohorts(ex_pop)
```

get.creating.type	<i>Derive creating type</i>
-------------------	-----------------------------

Description

Function to derive creating type for each individual

Usage

```
get.creating.type(
  population,
  database = NULL,
  gen = NULL,
  cohorts = NULL,
  use.id = FALSE
)
```

Arguments

population	Population list
database	Groups of individuals to consider for the export
gen	Quick-insert for database (vector of all generations to export)
cohorts	Quick-insert for database (vector of names of cohorts to export)
use.id	Set to TRUE to use MoBPS ids instead of Sex_Nr_Gen based names (default: FALSE)

Value

Creating type of in gen/database/cohorts selected individuals

Examples

```
data(ex_pop)
get.creating.type(ex_pop, gen=2)
```

get.cullingtime	<i>Derive time of culling</i>
-----------------	-------------------------------

Description

Function to devide the time of culling for all individuals

Usage

```
get.cullingtime(
  population,
  database = NULL,
  gen = NULL,
  cohorts = NULL,
  use.id = FALSE
)
```

Arguments

population	Population list
database	Groups of individuals to consider for the export
gen	Quick-insert for database (vector of all generations to export)
cohorts	Quick-insert for database (vector of names of cohorts to export)
use.id	Set to TRUE to use MoBPS ids instead of Sex_Nr_Gen based names (default: FALSE)

Value

Time of death of in gen/database/cohorts selected individuals

Examples

```
data(ex_pop)
get.cullingtime(ex_pop, gen=2)
```

get.database	<i>gen/database/cohorts conversion</i>
--------------	--

Description

Function to derive a database based on gen/database/cohorts

Usage

```
get.database(  
  population,  
  gen = NULL,  
  database = NULL,  
  cohorts = NULL,  
  avoid.merging = FALSE  
)
```

Arguments

population	Population list
gen	Quick-insert for database (vector of all generations to export)
database	Groups of individuals to consider for the export
cohorts	Quick-insert for database (vector of names of cohorts to export)
avoid.merging	Set to TRUE to avoid different cohorts to be merged in a joint group when possible

Value

Combine gen/database/cohorts to a joined database

Examples

```
data(ex_pop)
get.database(ex_pop, gen=2)
```

get.death.point *Derive death point*

Description

Function to derive the time of death for each individual (NA for individuals that are still alive)

Usage

```
get.death.point(  
  population,  
  database = NULL,  
  gen = NULL,  
  cohorts = NULL,  
  use.id = FALSE  
)
```

Arguments

population	Population list
database	Groups of individuals to consider for the export
gen	Quick-insert for database (vector of all generations to export)
cohorts	Quick-insert for database (vector of names of cohorts to export)
use.id	Set to TRUE to use MoBPS ids instead of Sex_Nr_Gen based names (default: FALSE)

Value

Time of death of in gen/database/cohorts selected individuals

Examples

```
data(ex_pop)  
get.death.point(ex_pop, gen=2)
```

get.dendrogram *Dendrogram*

Description

Function calculate a dendrogram

Usage

```
get.dendrogram(  
  population,  
  path = NULL,  
  database = NULL,  
  gen = NULL,  
  cohorts = NULL,  
  method = NULL,  
  individual.names = NULL  
)
```

Arguments

population	Population list
path	provide a path if the dendrogram would be saved as a png-file
database	Groups of individuals to consider
gen	Quick-insert for database (vector of all generations to consider)
cohorts	Quick-insert for database (vector of names of cohorts to consider)
method	Method used to calculate genetic distances (default: "Nei", alt: "Rogers", "Pre- vosti", "Modified Rogers")
individual.names	Names of the individuals in the database ((default are MoBPS internal names based on position))

Value

Dendrogram plot for genotypes

Examples

```
data(ex_pop)  
get.dendrogram(ex_pop, gen=2)
```

get.dendrogram.heatmap

Dendrogram Heatmap

Description

Function calculate a dendrogram

Usage

```
get.dendrogram.heatmap(
  population,
  path = NULL,
  database = NULL,
  gen = NULL,
  cohorts = NULL,
  method = NULL,
  individual.names = NULL,
  traits = NULL,
  type = "pheno"
)
```

Arguments

population	Population list
path	provide a path if the dendrogram would be saved as a png-file
database	Groups of individuals to consider
gen	Quick-insert for database (vector of all generations to consider)
cohorts	Quick-insert for database (vector of names of cohorts to consider)
method	Method used to calculate genetic distances (default: "Nei", alt: "Rogers", "Pre- vosti", "Modified Rogers")
individual.names	Names of the individuals in the database ((default are MoBPS internal names based on position))
traits	Traits to include in the dendrogram (default: all traits)
type	Which traits values to consider (default: "pheno", alt: "bv", "bve")

Value

Dendrogram plot of genotypes vs phenotypes

Examples

```
population <- creating.diploid(nsnp=1000, nindi=40, n.additive = c(100,100,100),
  shuffle.cor = matrix(c(1,0.8,0.2,0.8,1,0.2,0.2,0.2,1), ncol=3), shuffle.traits = 1:3)
population <- breeding.diploid(population, phenotyping = "all", heritability = 0.5)
get.dendrogram.heatmap(population, gen=1, type="pheno")
```

`get.dendrogram.trait` *Dendrogram*

Description

Function calculate a dendrogram for the traits

Usage

```
get.dendrogram.trait(  
  population,  
  path = NULL,  
  database = NULL,  
  gen = NULL,  
  cohorts = NULL,  
  traits = NULL,  
  type = "pheno"  
)
```

Arguments

population	Population list
path	provide a path if the dendrogram would be saved as a png-file
database	Groups of individuals to consider
gen	Quick-insert for database (vector of all generations to consider)
cohorts	Quick-insert for database (vector of names of cohorts to consider)
traits	Traits to include in the dendrogram (default: all traits)
type	Which traits values to consider (default: "pheno", alt: "bv", "bve")

Value

Dendrogram plot for traits

Examples

```
population <- creating.diploid(nsnp=1000, nindi=100, n.additive = c(100,100,100),  
  shuffle.cor = matrix(c(1,0.8,0.2,0.8,1,0.2,0.2,0.2,1), ncol=3), shuffle.traits = 1:3)  
population <- breeding.diploid(population, phenotyping = "all", heritability = 0.5)  
get.dendrogram.trait(population, gen=1, type="pheno")
```

get.distance

Calculate Nei distance between two or more population

Description

Function to calculate Nei's distance between two or more population

Usage

```
get.distance(
  population,
  type = "nei",
  marker = "all",
  per.marker = FALSE,
  gen1 = NULL,
  database1 = NULL,
  cohorts1 = NULL,
  gen2 = NULL,
  database2 = NULL,
  cohorts2 = NULL,
  database.list = NULL,
  gen.list = NULL,
  cohorts.list = NULL
)
```

Arguments

population	population list
type	Chose type of distance to compute (default: Neis standard genetic distance "nei"). Alt: Reynolds distance ("reynold"), Cavalli-Sforza ("cavalli"), Neis distance ("nei_distance"), Neis minimum distance ("nei_minimum")
marker	Vector with SNPs to consider (Default: "all" - use of all markers)
per.marker	Set to TRUE to return per marker statistics on genetic distances
gen1	Quick-insert for database (vector of all generations to consider)
database1	First Groups of individuals to consider
cohorts1	Quick-insert for database (vector of names of cohorts to consider)
gen2	Quick-insert for database (vector of all generations to consider)
database2	Second Groups of individuals to consider
cohorts2	Quick-insert for database (vector of names of cohorts to consider)
database.list	List of databases to consider (use when working with more than 2 populations)
gen.list	Quick-insert for database (vector of all generations to consider)
cohorts.list	Quick-insert for database (vector of names of cohorts to consider)

Value

Population list

Examples

```
data(ex_pop)
get.distance(ex_pop, database1 = cbind(1,1), database2 = cbind(1,2))
```

get.effect.freq *Compute marker frequency in QTL-markers*

Description

Function to compute marker frequency in QTL-markers

Usage

```
get.effect.freq(
  population,
  database = NULL,
  gen = NULL,
  cohorts = NULL,
  sort = FALSE
)
```

Arguments

- population Population list
- database Groups of individuals to consider for the export
- gen Quick-insert for database (vector of all generations to export)
- cohorts Quick-insert for database (vector of names of cohorts to export)
- sort Set to FALSE to not sort markers according to position on the genome

Value

Matrix with allele frequencies in the QTLs

Examples

```
data(ex_pop)
get.effect.freq(ex_pop, gen=1)
```

get.effective.size *Estimate effective population size*

Description

Function to estimate the effective population size

Usage

```
get.effective.size(population, gen = NULL, database = NULL, cohorts = NULL)
```

Arguments

population	Population list
gen	Quick-insert for database (vector of all generations to export)
database	Groups of individuals to consider for the export
cohorts	Quick-insert for database (vector of names of cohorts to export)

Value

Estimated effective population size

Examples

```
data(ex_pop)
get.effective.size(population=ex_pop, gen=5)
```

get.geno *Derive genotypes of selected individuals*

Description

Function to derive genotypes of selected individuals

Usage

```
get.geno(
  population,
  database = NULL,
  gen = NULL,
  cohorts = NULL,
  chromosomen = "all",
  export.alleles = FALSE,
  non.genotyped.as.missing = FALSE,
  use.id = FALSE
)
```

Arguments

population	Population list
database	Groups of individuals to consider for the export
gen	Quick-insert for database (vector of all generations to export)
cohorts	Quick-insert for database (vector of names of cohorts to export)
chromosomen	Beschränkung des Genotypen auf bestimmte Chromosomen (default: 1)
export.alleles	If TRUE export underlying alleles instead of just 012
non.genotyped.as.missing	Set to TRUE to replace non-genotyped markers with NA
use.id	Set to TRUE to use MoBPS ids instead of Sex_Nr_Gen based names (default: FALSE)

Value

Genotype data for in gen/database/cohorts selected individuals

Examples

```
data(ex_pop)
geno <- get.geno(ex_pop, gen=2)
```

get.genotyped *Derive genotyping status*

Description

Function to if selected individuals are genotyped

Usage

```
get.genotyped(
  population,
  database = NULL,
  gen = NULL,
  cohorts = NULL,
  use.id = FALSE
)
```

Arguments

population	Population list
database	Groups of individuals to consider for the export
gen	Quick-insert for database (vector of all generations to export)
cohorts	Quick-insert for database (vector of names of cohorts to export)
use.id	Set to TRUE to use MoBPS ids instead of Sex_Nr_Gen based names (default: FALSE)

Value

Check if in gen/database/cohorts selected individuals are genotyped

Examples

```
data(ex_pop)
get.genotyped(ex_pop, gen=2)
```

get.genotyped.snp *Derive which markers are genotyped of selected individuals*

Description

Function to decide which markers are genotyped for the selected individuals

Usage

```
get.genotyped.snp(
  population,
  database = NULL,
  gen = NULL,
  cohorts = NULL,
  export.alleles = FALSE,
  use.id = FALSE
)
```

Arguments

population	Population list
database	Groups of individuals to consider for the export
gen	Quick-insert for database (vector of all generations to export)
cohorts	Quick-insert for database (vector of names of cohorts to export)
export.alleles	If TRUE export underlying alleles instead of just 012
use.id	Set to TRUE to use MoBPS ids instead of Sex_Nr_Gen based names (default: FALSE)

Value

Binary Coded is/isnot genotyped level for in gen/database/cohorts selected individuals

Examples

```
data(ex_pop)
genotyped.snps <- get.genotyped.snp(ex_pop, gen=2)
```

`get.haplo`*Derive haplotypes of selected individuals*

Description

Function to derive haplotypes of selected individuals

Usage

```
get.haplo(  
  population,  
  database = NULL,  
  gen = NULL,  
  cohorts = NULL,  
  chromosomen = "all",  
  export.alleles = FALSE,  
  non.genotyped.as.missing = FALSE,  
  use.id = FALSE  
)
```

Arguments

<code>population</code>	Population list
<code>database</code>	Groups of individuals to consider for the export
<code>gen</code>	Quick-insert for database (vector of all generations to export)
<code>cohorts</code>	Quick-insert for database (vector of names of cohorts to export)
<code>chromosomen</code>	Beschränkung der Haplotypen auf bestimmte Chromosomen (default: 1)
<code>export.alleles</code>	If TRUE export underlying alleles instead of just 012
<code>non.genotyped.as.missing</code>	Set to TRUE to replace non-genotyped markers with NA
<code>use.id</code>	Set to TRUE to use MoBPS ids instead of Sex_Nr_Gen based names (default: FALSE)

Value

Haplotype data for in `gen/database/cohorts` selected individuals

Examples

```
data(ex_pop)  
haplo <- get.haplo(ex_pop, gen=2)
```

get.id *Derive ID on an individual*

Description

Function to derive the internal ID given to each individual

Usage

```
get.id(population, database = NULL, gen = NULL, cohorts = NULL, use.id = FALSE)
```

Arguments

population	Population list
database	Groups of individuals to consider for the export
gen	Quick-insert for database (vector of all generations to export)
cohorts	Quick-insert for database (vector of names of cohorts to export)
use.id	Set to TRUE to use MoBPS ids instead of Sex_Nr_Gen based names

Value

Individual ID for in gen/database/cohorts selected individuals

Examples

```
data(ex_pop)
get.id(ex_pop, gen=2)
```

get.individual.loc *Export location of individuals from the population list*

Description

Export location of individuals from the population list

Usage

```
get.individual.loc(population, database = NULL, gen = NULL, cohorts = NULL)
```

Arguments

population	Population list
database	Groups of individuals to consider for the export
gen	Quick-insert for database (vector of all generations to export)
cohorts	Quick-insert for database (vector of names of cohorts to export)

Value

Storage Position for in gen/database/cohorts selected individuals (Generation/Sex/IndividualNr)

Examples

```
data(ex_pop)
get.individual.loc(ex_pop, gen=2)
```

<code>get.infos</code>	<i>Extract bv/pheno/geno of selected individuals</i>
------------------------	--

Description

Function to extract bv/pheno/geno of selected individuals

Usage

```
get.infos(
  population,
  database = NULL,
  gen = NULL,
  cohorts = NULL,
  use.id = FALSE
)
```

Arguments

population	Population list
database	Groups of individuals to consider for the export
gen	Quick-insert for database (vector of all generations to export)
cohorts	Quick-insert for database (vector of names of cohorts to export)
use.id	Set to TRUE to use MoBPS ids instead of Sex_Nr_Gen based names (default: FALSE)

Value

Info list [[1]] phenotypes [[2]] genomic values [[3]] Z [[4/5/6]] additive/epistatic/dice marker effects

Examples

```
data(ex_pop)
get.infos(ex_pop, gen=2)
```

get.map *Map generation*

Description

Function to derive the genomic map for a given population list

Usage

```
get.map(population, use.snp.nr = FALSE)
```

Arguments

population	Population list
use.snp.nr	Set to TRUE to display SNP number and not SNP name

Value

Genomic map of the population list

Examples

```
data(ex_pop)
map <- get.map(ex_pop)
```

get.npheno *Export underlying number of observations per phenotype*

Description

Function to export the number of observation of each underlying phenotype

Usage

```
get.npheno(
  population,
  database = NULL,
  gen = NULL,
  cohorts = NULL,
  use.all.copy = FALSE,
  use.id = FALSE
)
```


Arguments

population	Population list
database	Groups of individuals to consider for the export
gen	Quick-insert for database (vector of all generations to export)
cohorts	Quick-insert for database (vector of names of cohorts to export)
use.all.copy	Set to TRUE to extract phenotyping
use.id	Set to TRUE to use MoBPS ids instead of Sex_Nr_Gen based names (default: FALSE)

Value

Phenotypes for in gen/database/cohorts selected individuals

Examples

```
data(ex_pop)
get.pheno(ex_pop, gen=2)
```

get.pca

Principle components analysis

Description

Function to perform a principle component analysis

Usage

```
get.pca(
  population,
  path = NULL,
  database = NULL,
  gen = NULL,
  cohorts = NULL,
  coloring = "group",
  components = c(1, 2),
  plot = TRUE,
  pch = 1,
  export.color = FALSE
)
```

Arguments

population	Population list
path	Location were to save the PCA-plot
database	Groups of individuals to consider for the export
gen	Quick-insert for database (vector of all generations to export)
cohorts	Quick-insert for database (vector of names of cohorts to export)
coloring	Coloring by "group", "sex", "plain"
components	Default: c(1,2) for the first two principle components
plot	Set to FALSE to not generate a plot
pch	Point type in the PCA plot
export.color	Set to TRUE to export the per point coloring

Value

Genotype data for in gen/database/cohorts selected individuals

Examples

```
data(ex_pop)
get.pca(ex_pop, gen=2)
```

get.pedigree	<i>Derive pedigree</i>
--------------	------------------------

Description

Derive pedigree for selected individuals

Usage

```
get.pedigree(
  population,
  database = NULL,
  gen = NULL,
  cohorts = NULL,
  founder.zero = TRUE,
  raw = FALSE,
  id = FALSE
)
```

Arguments

population	Population list
database	Groups of individuals to consider for the export
gen	Quick-insert for database (vector of all generations to export)
cohorts	Quick-insert for database (vector of names of cohorts to export)
founder.zero	Parents of founders are displayed as "0" (default: TRUE)
raw	Set to TRUE to not convert numbers into Sex etc.
id	Set to TRUE to extract individual IDs

Value

Pedigree-file for in gen/database/cohorts selected individuals

Examples

```
data(ex_pop)
get.pedigree(ex_pop, gen=2)
```

get.pedigree2	<i>Derive pedigree including grandparents</i>
---------------	---

Description

Derive pedigree for selected individuals including grandparents

Usage

```
get.pedigree2(
  population,
  database = NULL,
  gen = NULL,
  cohorts = NULL,
  shares = FALSE,
  founder.zero = TRUE
)
```

Arguments

population	Population list
database	Groups of individuals to consider for the export
gen	Quick-insert for database (vector of all generations to export)
cohorts	Quick-insert for database (vector of names of cohorts to export)
shares	Determine actual inherited shares of grandparents
founder.zero	Parents of founders are displayed as "0" (default: TRUE)

Value

Pedigree-file (grandparents) for in gen/database/cohorts selected individuals

Examples

```
data(ex_pop)
get.pedigree2(ex_pop, gen=2)
```

get.pedigree3	<i>Derive pedigree parents and grandparents</i>
---------------	---

Description

Derive pedigree for selected individuals including parents/grandparents

Usage

```
get.pedigree3(
  population,
  database = NULL,
  gen = NULL,
  cohorts = NULL,
  founder.zero = TRUE
)
```

Arguments

population	Population list
database	Groups of individuals to consider for the export
gen	Quick-insert for database (vector of all generations to export)
cohorts	Quick-insert for database (vector of names of cohorts to export)
founder.zero	Parents of founders are displayed as "0" (default: TRUE)

Value

Pedigree-file (parents + grandparents) for in gen/database/cohorts selected individuals

Examples

```
data(ex_pop)
get.pedigree3(ex_pop, gen=3)
```

get.pedmap *Generate plink-file (pedmap)*

Description

Generate a ped and map file (PLINK format) for selected groups and chromosome

Usage

```
get.pedmap(  
  population,  
  path = NULL,  
  database = NULL,  
  gen = NULL,  
  cohorts = NULL,  
  non.genotyped.as.missing = FALSE,  
  use.id = FALSE  
)
```

Arguments

population	Population list
path	Location to save pedmap-file
database	Groups of individuals to consider for the export
gen	Quick-insert for database (vector of all generations to export)
cohorts	Quick-insert for database (vector of names of cohorts to export)
non.genotyped.as.missing	Set to TRUE to replaced non-genotyped entries with "./."
use.id	Set to TRUE to use MoBPS ids instead of Sex_Nr_Gen based names

Value

Ped and map-file for in gen/database/cohorts selected individuals

Examples

```
data(ex_pop)  
  
file_path <- tempdir()  
get.pedmap(path=file_path, ex_pop, gen=2)  
file.remove(paste0(file_path, ".ped"))  
file.remove(paste0(file_path, ".map"))
```

get.pheno	<i>Export underlying phenotypes</i>
-----------	-------------------------------------

Description

Function to export underlying phenotypes

Usage

```
get.pheno(  
  population,  
  database = NULL,  
  gen = NULL,  
  cohorts = NULL,  
  use.all.copy = FALSE,  
  use.id = FALSE  
)
```

Arguments

population	Population list
database	Groups of individuals to consider for the export
gen	Quick-insert for database (vector of all generations to export)
cohorts	Quick-insert for database (vector of names of cohorts to export)
use.all.copy	Set to TRUE to extract phenotyping
use.id	Set to TRUE to use MoBPS ids instead of Sex_Nr_Gen based names (default: FALSE)

Value

Phenotypes for in gen/database/cohorts selected individuals

Examples

```
data(ex_pop)  
get.pheno(ex_pop, gen=2)
```

get.pheno.off *Export underlying offspring phenotypes*

Description

Function to export offspring phenotypes

Usage

```
get.pheno.off(
  population,
  database = NULL,
  gen = NULL,
  cohorts = NULL,
  use.id = FALSE
)
```

Arguments

population	Population list
database	Groups of individuals to consider for the export
gen	Quick-insert for database (vector of all generations to export)
cohorts	Quick-insert for database (vector of names of cohorts to export)
use.id	Set to TRUE to use MoBPS ids instead of Sex_Nr_Gen based names (default: FALSE)

Value

Avg. phenotype of the offspring of in gen/database/cohorts selected individuals

Examples

```
data(ex_pop)
get.pheno.off(ex_pop, gen=2)
```

get.pheno.off.count *Export underlying number of used offspring for offspring phenotypes*

Description

Function to export number of observations used for offspring phenotypes

Usage

```
get.pheno.off.count(population, database = NULL, gen = NULL, cohorts = NULL)
```

Arguments

population	Population list
database	Groups of individuals to consider for the export
gen	Quick-insert for database (vector of all generations to export)
cohorts	Quick-insert for database (vector of names of cohorts to export)

Value

Number of offspring with phenotypes for in gen/database/cohorts selected individuals

Examples

```
data(ex_pop)
get.pheno.off.count(ex_pop, gen=2)
```

`get.phylogenetic.tree` *Phylogenetic Tree*

Description

Function calculate a phylogenetic tree

Usage

```
get.phylogenetic.tree(
  population,
  path = NULL,
  database = NULL,
  gen = NULL,
  cohorts = NULL,
  method = NULL,
  individual.names = NULL,
  circular = FALSE
)
```

Arguments

population	Population list
path	provide a path if the dendrogram would be saved as a png-file
database	Groups of individuals to consider
gen	Quick-insert for database (vector of all generations to consider)
cohorts	Quick-insert for database (vector of names of cohorts to consider)
method	Method used to calculate genetic distances (default: "Nei", alt: "Rogers", "Pre- vosti", "Modified Rogers")

`individual.names` Names of the individuals in the database ((default are MoBPS internal names based on position))

`circular` Set to TRUE to generate a fan/circular layout tree

Value

Dendrogram plot for traits

Examples

```
data(ex_pop)
get.phylogenetic.tree(ex_pop, gen=1, circular=TRUE)
```

`get.qlt` *QTL extraction*

Description

Function to the position of QTLs (for snp/chr use `get.qlt.effects()`)

Usage

```
get.qlt(population)
```

Arguments

`population` Population list

Value

Vector of SNP positions

Examples

```
data(ex_pop)
positions <- get.qlt(ex_pop)
```

get.qtl.effects *QTL effect extraction*

Description

Function to extract QTL effect sizes

Usage

```
get.qtl.effects(population)
```

Arguments

population Population list

Value

List with [[1]] single SNP QTLs [[2]] epistatic SNP QTLs [[3]] dice QTL

Examples

```
data(ex_pop)
effects <- get.qtl.effects(ex_pop)
```

get.qtl.variance *QTL effect variance extraction*

Description

Function to extract QTL effect variance for single SNP QTLs in a given gen/database/cohort

Usage

```
get.qtl.variance(population, gen = NULL, database = NULL, cohorts = NULL)
```

Arguments

population Population list
gen Quick-insert for database (vector of all generations to consider)
database Groups of individuals to consider
cohorts Quick-insert for database (vector of names of cohorts to consider)

Value

matrix with SNP / Chr / estimated effect variance

Examples

```
data(ex_pop)
effects <- get.qtl.variance(ex_pop)
```

get.recombi *Derive genetic origins*

Description

Function to derive genetic origin

Usage

```
get.recombi(
  population,
  database = NULL,
  gen = NULL,
  cohorts = NULL,
  use.id = FALSE
)
```

Arguments

population	Population list
database	Groups of individuals to consider for the export
gen	Quick-insert for database (vector of all generations to export)
cohorts	Quick-insert for database (vector of names of cohorts to export)
use.id	Set to TRUE to use MoBPS ids instead of Sex_Nr_Gen based names (default: FALSE)

Value

Recombination points for in gen/database/cohorts selected individuals

Examples

```
data(ex_pop)
get.recombi(ex_pop, gen=2)
```

get.reliabilities *Export underlying reliabilities*

Description

Function to export underlying reliabilities

Usage

```
get.reliabilities(
  population,
  database = NULL,
  gen = NULL,
  cohorts = NULL,
  use.id = FALSE
)
```

Arguments

population	Population list
database	Groups of individuals to consider for the export
gen	Quick-insert for database (vector of all generations to export)
cohorts	Quick-insert for database (vector of names of cohorts to export)
use.id	Set to TRUE to use MoBPS ids instead of Sex_Nr_Gen based names (default: FALSE)

Value

Estimated reliability for BVE for in gen/database/cohorts selected individuals

Examples

```
data(ex_pop)
get.reliabilities(ex_pop, gen=2)
```

get.selectionbve *Export derived breeding values based on the selection index*

Description

Function to export last breeding values based on the selection index

Usage

```
get.selectionbve(
  population,
  database = NULL,
  gen = NULL,
  cohorts = NULL,
  use.id = FALSE
)
```

Arguments

population	Population list
database	Groups of individuals to consider for the export
gen	Quick-insert for database (vector of all generations to export)
cohorts	Quick-insert for database (vector of names of cohorts to export)
use.id	Set to TRUE to use MoBPS ids instead of Sex_Nr_Gen based names (default: FALSE)

Value

Last applied selection index for in gen/database/cohorts selected individuals

Examples

```
data(ex_pop)
get.selectionindex(ex_pop, gen=2)
```

get.selectionindex *Export underlying last used selection index*

Description

Function to export last used selection index (mostly relevant for Miesenberger 1997 stuff)

Usage

```
get.selectionindex(
  population,
  database = NULL,
  gen = NULL,
  cohorts = NULL,
  use.id = FALSE
)
```

Arguments

population	Population list
database	Groups of individuals to consider for the export
gen	Quick-insert for database (vector of all generations to export)
cohorts	Quick-insert for database (vector of names of cohorts to export)
use.id	Set to TRUE to use MoBPS ids instead of Sex_Nr_Gen based names (default: FALSE)

Value

Last applied selection index for in gen/database/cohorts selected individuals

Examples

```
data(ex_pop)
get.selectionindex(ex_pop, gen=2)
```

<i>get.time.point</i>	<i>Derive time point</i>
-----------------------	--------------------------

Description

Function to devide time point for each individual

Usage

```
get.time.point(
  population,
  database = NULL,
  gen = NULL,
  cohorts = NULL,
  use.id = FALSE
)
```

Arguments

population	Population list
database	Groups of individuals to consider for the export
gen	Quick-insert for database (vector of all generations to export)
cohorts	Quick-insert for database (vector of names of cohorts to export)
use.id	Set to TRUE to use MoBPS ids instead of Sex_Nr_Gen based names (default: FALSE)

Value

Time point of generation for in gen/database/cohorts selected individuals

Examples

```
data(ex_pop)
get.time.point(ex_pop, gen=2)
```

get.vcf	<i>Generate vcf-file</i>
---------	--------------------------

Description

Generate a vcf-file for selected groups and chromosome

Usage

```
get.vcf(
  population,
  path = NULL,
  database = NULL,
  gen = NULL,
  cohorts = NULL,
  chromosomen = "all",
  non.genotyped.as.missing = FALSE,
  use.id = FALSE
)
```

Arguments

population	Population list
path	Location to save vcf-file
database	Groups of individuals to consider for the export
gen	Quick-insert for database (vector of all generations to export)
cohorts	Quick-insert for database (vector of names of cohorts to export)
chromosomen	Beschaenkung des Genotypen auf bestimmte Chromosomen (default: 1)
non.genotyped.as.missing	Set to TRUE to replaced non-genotyped entries with "./."
use.id	Set to TRUE to use MoBPS ids instead of Sex_Nr_Gen based names

Value

VCF-file for in gen/database/cohorts selected individuals

Examples

```

data(ex_pop)
data(ex_pop)

file_path <- tempdir()
get.vcf(path=file_path, ex_pop, gen=2)
file.remove(paste0(file_path, ".vcf"))

```

group.diff

Function to exclude individuals from a database

Description

Function to exclude individuals from a database

Usage

```

group.diff(
  population,
  database = NULL,
  gen = NULL,
  cohorts = NULL,
  remove.gen = NULL,
  remove.database = NULL,
  remove.cohorts = NULL
)

```

Arguments

population	Population list
database	Groups of individuals to consider for the export
gen	Quick-insert for database (vector of all generations to export)
cohorts	Quick-insert for database (vector of names of cohorts to export)
remove.gen	Generations of individuals to remove from the database (same IDs!)
remove.database	Groups of individuals to remove from the database (same IDs!)
remove.cohorts	Cohorts of individuals to remove from the database (same IDs!)

Value

Database excluding removals

Examples

```

data(ex_pop)
database <- group.diff(ex_pop, gen=1, remove.database=cbind(1,1))

```

 insert.bve

Manually enter estimated breeding values

Description

Function to manually enter estimated breeding values

Usage

```
insert.bve(
  population,
  bves,
  type = "bve",
  na.override = FALSE,
  count = 1,
  count.only.increase = TRUE
)
```

Arguments

population	Population list
bves	Matrix of breeding values to enter (one row per individual with 1 element coding individual name)
type	which time of values to input (default: "bve", alt: "bv", "pheno")
na.override	Set to TRUE to also enter NA values (Default: FALSE - those entries will be skipped)
count	Counting for economic cost calculation (default: 1 - (one observation (for "pheno"), one genotyping (for "bve")))
count.only.increase	Set to FALSE to reduce the number of observation for a phenotype to "count" (default: TRUE)

Value

Population-List with newly entered estimated breeding values

Examples

```
data(ex_pop)
bv <- get.bv(ex_pop, gen=2)
new.bve <- cbind( colnames(bv), bv[,1]) ## Unrealistic but you do not get better than this!
ex_pop <- insert.bve(ex_pop, bves=new.bve)
```

json.simulation	<i>Simulation of a breeding program based on a JSON-file from MoBP-Sweb</i>
-----------------	---

Description

Function to simulate a breeding program based on a JSON-file from MoBPSweb

Usage

```

json.simulation(
  file = NULL,
  log = NULL,
  total = NULL,
  fast.mode = FALSE,
  progress.bars = FALSE,
  size.scaling = NULL,
  rep.max = 1,
  verbose = TRUE,
  miraculix.cores = NULL,
  miraculix.chol = NULL,
  skip.population = FALSE,
  time.check = FALSE,
  time.max = 7200,
  export.population = FALSE,
  export.gen = NULL,
  export.timepoint = NULL,
  fixed.generation.order = NULL
)

```

Arguments

file	Path to a json-file generated by the user-interface
log	Provide Path where to write a log-file of your simulation (or false to not write a log-file)
total	Json-file imported via jsonlite::read_json
fast.mode	Set to TRUE work on a small genome with few markers
progress.bars	Set to TRUE to display progress bars
size.scaling	Scale the size of nodes by this factor (especially for testing smaller examples)
rep.max	Maximum number of repeats to use in fast.mode
verbose	Set to FALSE to not display any prints
miraculix.cores	Number of cores used in miraculix applications (default: 1)
miraculix.chol	Set to FALSE to manually deactivate the use of miraculix for any cholesky decomposition even though miraculix is activated

skip.population	Set to TRUE to not execute breeding actions (only cost/time estimation will be performed)
time.check	Set to TRUE to automatically check simulation run-time before executing breeding actions
time.max	Maximum length of the simulation in seconds when time.check is active
export.population	Path were to export the population to (at state selected in export.gen/timepoint)
export.gen	Last generation to simulate before exporting population to file
export.timepoint	Last timepoint to simulate before exporting population to file
fixed.generation.order	Vector containing the order of cohorts to generate (Advanced // Testing Parameter!)

Value

Population-list

Examples

```
data(ex_json)
population <- json.simulation(total=ex_json)
```

kinship.development *Development of genetic/breeding value*

Description

Function to plot genetic/breeding values for multiple generation/cohorts

Usage

```
kinship.development(
  population,
  database = NULL,
  gen = NULL,
  cohorts = NULL,
  json = FALSE,
  ibd.obs = 50,
  hbd.obs = 10,
  display.cohort.name = FALSE,
  display.time.point = FALSE,
  equal.spacing = FALSE,
  time_reorder = FALSE,
  display.hbd = FALSE
)
```

Arguments

population	population list
database	Groups of individuals to consider for the export
gen	Quick-insert for database (vector of all generations to export)
cohorts	Quick-insert for database (vector of names of cohorts to export)
json	If TRUE extract which cohorts to plot according to the json-file used in json.simulation
ibd.obs	Number of Individual pairs to sample for IBD estimation
hbd.obs	Number of Individuals to sample for HBD estimation
display.cohort.name	Set TRUE to display the name of the cohort in the x-axis
display.time.point	Set TRUE to use time point of generated to sort groups
equal.spacing	Equal distance between groups (independent of time.point)
time_reorder	Set TRUE to order cohorts according to the time point of generation
display.hbd	Set to TRUE to also display HBD in plot

Value

Estimated of avg. kinship/inbreeding based on IBD/HBD

Examples

```
data(ex_pop)
kinship.development(ex_pop,gen=1:5)
```

kinship.emp

Empirical kinship

Description

Function to compute empirical kinship for a set of individuals)

Usage

```
kinship.emp(
  animals = NULL,
  population = NULL,
  gen = NULL,
  database = NULL,
  cohorts = NULL,
  sym = FALSE
)
```

Arguments

animals	List of animals to compute kinship for
population	Population list
gen	Quick-insert for database (vector of all generations to export)
database	Groups of individuals to consider for the export
cohorts	Quick-insert for database (vector of names of cohorts to export)
sym	If True derive matrix entries below principle-diagonal

Value

Empirical kinship matrix (IBD-based since Founders)

Examples

```
data(ex_pop)
kinship <- kinship.emp(population=ex_pop, database=cbind(2,1,1,25))
```

kinship.emp.fast	<i>Approximate empirical kinship</i>
------------------	--------------------------------------

Description

Function to compute empirical kinship for a set of individuals (not all pairs of individuals are evaluated)

Usage

```
kinship.emp.fast(
  animals = NULL,
  population = NULL,
  gen = NULL,
  database = NULL,
  cohorts = NULL,
  sym = FALSE,
  ibd.obs = 50,
  hbd.obs = 10
)
```

Arguments

animals	List of animals to compute kinship for
population	Population list
gen	Quick-insert for database (vector of all generations to export)
database	Groups of individuals to consider for the export

cohorts	Quick-insert for database (vector of names of cohorts to export)
sym	If True derive matrix entries below principle-diagonal
ibd.obs	Number of Individual pairs to sample for IBD estimation
hbd.obs	Number of Individuals to sample for HBD estimation

Value

Empirical kinship matrix (IBD-based since Founders) per gen/database/cohort

Examples

```
data(ex_pop)
kinship.emp.fast(population=ex_pop, gen=2)
```

kinship.exp	<i>Derive expected kinship</i>
-------------	--------------------------------

Description

Function to derive expected kinship

Usage

```
kinship.exp(
  population,
  gen = NULL,
  database = NULL,
  cohorts = NULL,
  depth.pedigree = 7,
  start.kinship = NULL,
  elements = NULL,
  mult = 2,
  storage.save = 1.5,
  verbose = TRUE
)
```

Arguments

population	Population list
gen	Quick-insert for database (vector of all generations to export)
database	Groups of individuals to consider for the export
cohorts	Quick-insert for database (vector of names of cohorts to export)
depth.pedigree	Depth of the pedigree in generations
start.kinship	Relationship matrix of the individuals in the first considered generation
elements	Vector of individuals from the database to include in pedigree matrix

mult	Multiplicator of kinship matrix (default: 2)
storage.save	Lower numbers will lead to less memory but slightly higher computing time (default: 1.5, min: 1)
verbose	Set to FALSE to not display any prints

Value

Pedigree-based kinship matrix for in gen/database/cohort selected individuals

Examples

```
data(ex_pop)
kinship <- kinship.exp(population=ex_pop, gen=2)
```

ld.decay	<i>Generate LD plot</i>
----------	-------------------------

Description

Generate LD pot

Usage

```
ld.decay(
  population,
  genotype.dataset = NULL,
  chromosomen = 1,
  dist = NULL,
  step = 5,
  max = 500,
  max.cases = 100,
  database = NULL,
  gen = NULL,
  cohorts = NULL,
  type = "snp",
  plot = FALSE
)
```

Arguments

population	Population list
genotype.dataset	Genotype dataset (default: NULL - just to save computation time when get.geno was already run)
chromosomen	Only consider a specific chromosome in calculations (default: 1)
dist	Manuel input of marker distances to analyse

step	Stepsize to calculate LD
max	Maximum distance between markers to consider for LD-plot
max.cases	Maximum number of marker pairs to consider of each distance (default: 100; randomly sampled!)
database	Groups of individuals to consider for the export
gen	Quick-insert for database (vector of all generations to export)
cohorts	Quick-insert for database (vector of names of cohorts to export)
type	Compute LD decay according to following distance measure between markers (default: "snp", alt: "bp", "cM")
plot	Set to FALSE to not generate an LD plot

Value

LD-decay plot for in gen/database/cohorts selected individuals

Examples

```
data(ex_pop)
ld.decay(population=ex_pop, gen=5)
```

maize_chip

maize chip

Description

Genome for maize according to Lee et al.

Usage

```
maize_chip
```

Author(s)

Torsten Pook <torsten.pook@uni-goettingen.de>

Source

Lee et al 2002

miesenberger.index *Miesenberger Index*

Description

Function to selection index weights according to Miesenberger 1997

Usage

```
miesenberger.index(V, G, V1 = NULL, RG = NULL, r, w, zw = NULL)
```

Arguments

V	Phenotypic covarianz matrix
G	Genomic covarianz matrix
V1	Inverted phenotypic covarianz matrix
RG	Genomic correlation matrix
r	reliability for the breeding value estimation
w	relative weighting of each trait (per genetic SD)
zw	Estimated breeding value

Value

weights of the selection index

miraculix *Add miraculix-coding for genotypes*

Description

Internal function to store genotypes bit-wise

Usage

```
miraculix(population)
```

Arguments

population	Population list
------------	-----------------

Value

Population list

Examples

```
# This is only relevant with the package miraculix is installed and used
population <- creating.diploid(nsnp=100, nindi=50, miraculix=FALSE)
population <- miraculix(population)
```

mutation.intro	<i>Mutation intro</i>
----------------	-----------------------

Description

Function to change the base-pair in a specific loci

Usage

```
mutation.intro(population, gen, sex, individual.nr, qtl.posi, haplo.set = 1)
```

Arguments

population	Population list
gen	Generation of the individual to introduce a mutation in
sex	Sex of the individual to introduce a mutation in
individual.nr	Individual Nr. of the individual to introduce a mutation in
qtl.posi	Marker number to mutate
haplo.set	Select chromosome set (default: 1 , alt: 2)

Value

Population-List with mutated marker for the selected individual

Examples

```
data(ex_pop)
ex_pop <- mutation.intro(ex_pop, 1,1,1, qtl.posi=100)
```

new.base.generation *Set new base generation*

Description

Function to set a new base generation for the population

Usage

```
new.base.generation(  
  population,  
  base.gen = NULL,  
  delete.previous.gen = FALSE,  
  delete.breeding.totals = FALSE,  
  delete.bve.data = FALSE,  
  add.chromosome.ends = TRUE  
)
```

Arguments

population	Population list
base.gen	Vector containing all new base generations
delete.previous.gen	Delete all data before base.gen (default: FALSE)
delete.breeding.totals	Delete all breeding totals before base.gen (default: FALSE)
delete.bve.data	Delete all previous bve data (default: FALSE)
add.chromosome.ends	Add chromosome ends as recombination points

Value

Population-List with mutated marker for the selected individual

Examples

```
data(ex_pop)  
ex_pop <- new.base.generation(ex_pop, base.gen=2)
```

OGC

Optimal genetic contribution

Description

In this function the OGC selection according to Meuwissen 1997 is performed

Usage

```
OGC(  
  A,  
  u,  
  Q,  
  cAc = NA,  
  single = TRUE,  
  verbose = FALSE,  
  max_male = Inf,  
  max_female = Inf  
)
```

Arguments

A	relationship matrix
u	breeding values
Q	sex indicator
cAc	target gain in inbreeding
single	If FALSE multiple individuals can be removed at the same type (this is faster but potentially inaccurate!)
verbose	Set to FALSE to not display any prints
max_male	maximum number of male with positive contributions
max_female	maximum number of females with positive contributions

Value

[[1]] Contributions [[2]] expected inbreeding gain

pedigree.simulation *Simulation of a given pedigree*

Description

Function to simulate a given pedigree

Usage

```
pedigree.simulation(  
  pedigree,  
  keep.ids = FALSE,  
  plot = TRUE,  
  dataset = NULL,  
  vcf = NULL,  
  chr.nr = NULL,  
  bp = NULL,  
  snp.name = NULL,  
  hom0 = NULL,  
  hom1 = NULL,  
  bpcm.conversion = 0,  
  nsnp = 0,  
  freq = "beta",  
  sex.s = "fixed",  
  chromosome.length = NULL,  
  length.before = 5,  
  length.behind = 5,  
  real.bv.add = NULL,  
  real.bv.mult = NULL,  
  real.bv.dice = NULL,  
  snps.equidistant = NULL,  
  change.order = FALSE,  
  bv.total = 0,  
  polygenic.variance = 100,  
  bve.mult.factor = NULL,  
  bve.poly.factor = NULL,  
  base.bv = NULL,  
  add.chromosome.ends = TRUE,  
  new.phenotype.correlation = NULL,  
  new.residual.correlation = NULL,  
  new.breeding.correlation = NULL,  
  add.architecture = NULL,  
  snp.position = NULL,  
  position.scaling = FALSE,  
  bit.storing = FALSE,  
  nbits = 30,  
  randomSeed = NULL,
```

```

miraculix = TRUE,
miraculix.dataset = TRUE,
n.additive = 0,
n.dominant = 0,
n.qualitative = 0,
n.quantitative = 0,
var.additive.l = NULL,
var.dominant.l = NULL,
var.qualitative.l = NULL,
var.quantitative.l = NULL,
exclude.snps = NULL,
replace.real.bv = FALSE,
shuffle.traits = NULL,
shuffle.cor = NULL,
skip.rest = FALSE,
enter.bv = TRUE,
name.cohort = NULL,
template.chip = NULL,
beta.shape1 = 1,
beta.shape2 = 1,
time.point = 0,
creating.type = 0,
trait.name = NULL,
share.genotyped = 1,
genotyped.s = NULL,
map = NULL,
remove.invalid.qtl = TRUE,
verbose = TRUE,
bv.standard = FALSE,
mean.target = NULL,
var.target = NULL,
is.maternal = NULL,
is.paternal = NULL,
vcf.maxsnp = Inf
)

```

Arguments

pedigree	Pedigree-file (matrix with 3 columns (Individual ID, Father ID, Mother ID), optional forth columns with earliest generations to generate an individual)
keep.ids	Set to TRUE to keep the IDs from the pedigree-file instead of the default MoBPS ids
plot	Set to FALSE to not generate an overview of inbreeding and number of individuals over time
dataset	SNP dataset, use "random", "allhetero" "all0" when generating a dataset via nsnp.nindi
vcf	Path to a vcf-file used as input genotypes (correct haplotype phase is assumed!)

chr.nr	Vector containing the associated chromosome for each marker (default: all on the same)
bp	Vector containing the physical position (bp) for each marker (default: 1,2,3...)
snp.name	Vector containing the name of each marker (default ChrXSNPY - XY chosen accordingly)
hom0	Vector containing the first allelic variant in each marker (default: 0)
hom1	Vector containing the second allelic variant in each marker (default: 1)
bpcm.conversion	Convert physical position (bp) into a cM position (default: 0 - not done)
nsnp	number of markers to generate in a random dataset
freq	frequency of allele 1 when randomly generating a dataset
sex.s	Specify which newly added individuals are male (1) or female (2)
chromosome.length	Length of the newly added chromosome (default: 5)
length.before	Length before the first SNP of the dataset (default: 5)
length.behind	Length after the last SNP of the dataset (default: 5)
real.bv.add	Single Marker effects
real.bv.mult	Two Marker effects
real.bv.dice	Multi-marker effects
snps.equidistant	Use equidistant markers (computationally faster! ; default: TRUE)
change.order	If TRUE sort markers according to given marker positions
bv.total	Number of traits (If more than traits via real.bv.X use traits with no directly underlying QTL)
polygenic.variance	Genetic variance of traits with no underlying QTL
bve.mult.factor	Multiply trait value times this
bve.poly.factor	Potency trait value over this
base.bv	Average genetic value of a trait
add.chromosome.ends	Add chromosome ends as recombination points
new.phenotype.correlation	(OLD! - use new.residual.correlation) Correlation of the simulated environmental variance
new.residual.correlation	Correlation of the simulated environmental variance
new.breeding.correlation	Correlation of the simulated genetic variance (child share! heritage is not influenced!)
add.architecture	Add genetic architecture (marker positions)

snp.position	Location of each marker on the genetic map
position.scaling	Manual scaling of snp.position
bit.storing	Set to TRUE if the MoBPS (not-miraculix! bit-storing is used)
nbits	Bits available in MoBPS-bit-storing
randomSeed	Set random seed of the process
miraculix	If TRUE use miraculix package for data storage, computations and dataset generation
miraculix.dataset	Set FALSE to deactivate miraculix package for dataset generation
n.additive	Number of additive QTL
n.dominant	Number of dominante QTL
n.qualitative	Number of qualitative epistatic QTL
n.quantitative	Number of quantitative epistatic QTL
var.additive.l	Variance of additive QTL
var.dominant.l	Variance of dominante QTL
var.qualitative.l	Variance of qualitative epistatic QTL
var.quantitative.l	Variance of quantitative epistatic QTL
exclude.snps	Marker were no QTL are simulated on
replace.real.bv	If TRUE delete the simulated traits added before
shuffle.traits	Combine different traits into a joined trait
shuffle.cor	Target Correlation between shuffeled traits
skip.rest	Internal variable needed when adding multiple chromosomes jointly
enter.bv	Internal parameter
name.cohort	Name of the newly added cohort
template.chip	Import genetic map and chip from a species ("cattle", "chicken", "pig")
beta.shape1	First parameter of the beta distribution for simulating allele frequencies
beta.shape2	Second parameter of the beta distribution for simulating allele frequencies
time.point	Time point at which the new individuals are generated
creating.type	Technique to generate new individuals (usage in web-based application)
trait.name	Name of the trait generated
share.genotyped	Share of individuals genotyped in the founders
genotyped.s	Specify with newly added individuals are genotyped (1) or not (0)
map	map-file that contains up to 5 colums (Chromosome, SNP-id, M-position, Bp-position, allele freq - Everything not provides it set to NA). A map can be imported via MoBPSmaps::ensembl.map()

remove.invalid.qtl	Set to FALSE to deactivate the automatic removal of QTLs on markers that do not exist
verbose	Set to FALSE to not display any prints
bv.standard	Set TRUE to standardize trait mean and variance via bv.standardization() - automatically set to TRUE when mean/var.target are used
mean.target	Target mean
var.target	Target variance
is.maternal	Vector coding if a trait is caused by a maternal effect (Default: all FALSE)
is.paternal	Vector coding if a trait is caused by a paternal effect (Default: all FALSE)
vcf.maxsnp	Maximum number of SNPs to include in the genotype file (default: Inf)
add.chromosome	If TRUE add an additional chromosome to the dataset

Value

Population-list

Examples

```
pedigree <- matrix(c(1,0,0,
2,0,0,
3,0,0,
4,1,2,
5,1,3,
6,1,3,
7,1,3,
8,4,6,
9,4,7), ncol=3, byrow=TRUE)
population <- pedigree.simulation(pedigree, nsnp=1000)
```

pedmap.to.phasedbeaglevcf

Internal function to perform imputing/phasing

Description

Internal function to perform imputing/phasing (path chosen for the web-based application)

Usage

```
pedmap.to.phasedbeaglevcf(
  ped_path = NULL,
  map_path = NULL,
  vcf_path = NULL,
  beagle_jar = "/home/nha/beagle.03Jul18.40b.jar",
  plink_dir = "/home/nha/Plink/plink",
```

```
db_dir = "/home/nha/Plink/DB/",  
verbose = TRUE  
)
```

Arguments

ped_path	Directory of the ped-file
map_path	Directory of the map-file
vcf_path	Directory of the vcf-file (this will override any ped/map-file input)
beagle_jar	Directory of BEAGLE
plink_dir	Directory of Plink
db_dir	Directory to save newly generated files (ped/map will be stored in the original folder)
verbose	Set to FALSE to not display any prints

Value

Phased vcf file in vcf_path

pig_chip

pig chip

Description

Genome for pig according to Rohrer et al.

Usage

pig_chip

Author(s)

Torsten Pook <torsten.pook@uni-goettingen.de>

Source

Rohrer et al 1994

plot.population	<i>Plot Population</i>
-----------------	------------------------

Description

Basic plot of the population list

Usage

```
## S3 method for class 'population'
plot(x, type = "bve", gen = NULL, database = NULL, cohorts = NULL, ...)
```

Arguments

x	Population-list
type	Default "bve" - bv.development, alt: "kinship" - kinship.development(), "pca" - get.pca()
gen	generations to consider
database	groups to consider
cohorts	cohorts to consider
...	remaining stuff

Value

Summary of the population list including number of individuals, genome length and trait overview

Examples

```
data(ex_pop)
plot(ex_pop)
```

set.class	<i>Export estimated breeding values</i>
-----------	---

Description

Function to export estimated breeding values

Usage

```
set.class(
  population,
  database = NULL,
  gen = NULL,
  cohorts = NULL,
  new.class = 0
)
```

Arguments

population	Population list
database	Groups of individuals to consider for the export
gen	Quick-insert for database (vector of all generations to export)
cohorts	Quick-insert for database (vector of names of cohorts to export)
new.class	Class to change to (either single character or vector for each individual when just a single group is selected)

Value

Population-List with newly entered class values

Examples

```
data(ex_pop)
population <- set.class(ex_pop, database=cbind(1,1), new.class = 2)
```

set.default

Set defaults

Description

Set default parameter values in breeding.diploid

Usage

```
set.default(
  population,
  parameter.name = NULL,
  parameter.value = NULL,
  parameter.remove = NULL,
  reset.all = FALSE
)
```

Arguments

population	Population list
parameter.name	Number of traits (If more than traits via real.bv.X use traits with no directly underlying QTL)
parameter.value	Genetic variance of traits with no underlying QTL
parameter.remove	Remove a specific previously generated parameter default
reset.all	Set to TRUE to remove all prior parameter values

Value

Population-list with one or more additional new traits

Examples

```
data(ex_pop)
population <- set.default(ex_pop, parameter.name="heritability", parameter.value=0.3)
```

sheep_chip	<i>sheep chip</i>
------------	-------------------

Description

Genome for sheep according to Prieur et al.

Usage

```
sheep_chip
```

Author(s)

Torsten Pook <torsten.pook@uni-goettingen.de>

Source

Prieur et al 2017

sortd	<i>Apply sort and unique</i>
-------	------------------------------

Description

Efficient function to perform `sort(unique(v))`

Usage

```
sortd(v)
```

Arguments

v Vector

Value

numerical sorted vector without duplicates

Examples

```
v <- c(1,1,4,5)
sortd(v)
```

ssGBLUP

Single Step GBLUP

Description

Function to perform single step GBLUP according to Legarra 2014

Usage

```
ssGBLUP(A11, A12, A22, G)
```

Arguments

A11	pedigree relationship matrix of non-genotyped individuals
A12	pedigree relationship matrix between non-genotyped and genotyped individuals
A22	pedigree relationship matrix of genotyped individuals
G	genomic relationship matrix of genotyped individuals

Value

Single step relationship matrix

summary.population

Summary Population

Description

Summary of the population list

Usage

```
## S3 method for class 'population'
summary(object, ...)
```

Arguments

object	Population-list
...	additional arguments affecting the summary produced

Value

Summary of the population list including number of individuals, genome length and trait overview

Examples

```
data(ex_pop)
summary(ex_pop)
```

vlist

Generation of a sublist

Description

Internal function to write a couple of list entries in a new list

Usage

```
vlist(list, skip = NULL, first = NULL, select = NULL)
```

Arguments

list	list you want to print details of
skip	Skip first that many list-elements
first	Only display first that many list-elements
select	Display only selected list-elements

Value

Selected elements of a list

Examples

```
data(ex_pop)
vlist(ex_pop$breeding[[1]], select=3:10)
```

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