

# Package ‘qtlhot’

February 20, 2015

**Version** 0.9.0

**Date** 2013-09-18

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**Title** Inference for QTL Hotspots

**Description** Functions to infer co-mapping trait hotspots and causal models

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**Depends** R (>= 2.10), stats, qtl, lattice, corpcor, mnormt

**LazyLoad** yes

**LazyData** yes

**License** GPL (>= 2)

**URL** <http://www.stat.wisc.edu/~yandell/statgen>

**NeedsCompilation** yes

**Repository** CRAN

**Date/Publication** 2013-09-19 08:54:03

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add.phenos	<i>Add phenotypes to cross object.</i>
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**Description**

Add phenotypes to cross object by checking index.

**Usage**

```
add.phenos(cross, newdata = NULL, index = NULL)
```

**Arguments**

cross	object of class cross; see <a href="#">read.cross</a>
newdata	data frame with row names matching values of phenotype identified by index for object cross
index	character string name of phenotype in object cross; if NULL, then newdata must be of same size as cross with phenotypes in order

**Details**

The name index must be a phenotype in the cross object. The row names of newdata are matched with values of index.

**Value**

object of class cross with added phenotypes

**Author(s)**

Brian S. Yandell, <[byandell@wisc.edu](mailto:byandell@wisc.edu)>

**See Also**

[read.cross](#)

**Examples**

```
## Not run:
data(hyper)
x <- data.frame(x = rnorm(nind(hyper)))
hyperx <- add.phenos(hyper, x)

## End(Not run)
```

---

CMSTtests *Perform CMST Tests on cross object*

---

### Description

Performs 6 separate CMST tests (3 versions, 2 penalties).

### Usage

```
CMSTtests(cross, pheno1, pheno2, Q.chr, Q.pos,
  addcov1 = NULL, addcov2 = NULL, intcov1 = NULL, intcov2 = NULL,
  method = c("par", "non.par", "joint", "all"),
  penalty = c("bic", "aic", "both"), verbose = FALSE)
CMSTtestsList(cross, pheno1, pheno2, Q.chr, Q.pos,
  addcov1 = NULL, addcov2 = NULL, intcov1 = NULL, intcov2 = NULL,
  method = c("par", "non.par", "joint", "all"),
  penalty = c("bic", "aic", "both"), verbose = TRUE)
```

### Arguments

cross	object of class cross
pheno1	first phenotype column number or character string name
pheno2	second phenotype column number or character string name; if more than one, then all phenotypes will be tested against pheno1
Q.chr	QTL chromosome (number or label)
Q.pos	QTL position in cM
addcov1, addcov2	additive covariates for first and second phenotype, respectively
intcov1, intcov2	interactive covariates for first and second phenotype, respectively
method	test method; see details
penalty	type of penalty; see details
verbose	verbose printout if TRUE

### Details

Explain method and penalty here.

### References

Chaibub Neto E, Broman AT, Keller MP, Attie AD, Zhang B, Zhu J, Yandell BS, Causal model selection hypothesis tests in systems genetics. *Genetics* (in review).

### See Also

[CMSTCross](#), [PrecTpFpMatrix](#), [FitAllTests](#)

**Examples**

```

data(CMSTCross)
nms <- names(CMSTCross$pheno)
out1 <- CMSTtests(CMSTCross,
                  pheno1 = nms[1],
                  pheno2 = nms[2],
                  Q.chr = 1,
                  Q.pos = 55,
                  addcov1 = NULL,
                  addcov2 = NULL,
                  intcov1 = NULL,
                  intcov2 = NULL,
                  method = "all",
                  penalty = "both")

out1[1:6]
out1[7]
out1[8:12]
out1[13:17]
## list of phenotypes
out2 <- CMSTtests(CMSTCross,
                  pheno1 = nms[1],
                  pheno2 = nms[-1],
                  Q.chr = 1,
                  Q.pos = 55,
                  addcov1 = NULL,
                  addcov2 = NULL,
                  intcov1 = NULL,
                  intcov2 = NULL,
                  method = "par",
                  penalty = "bic")

out2

```

---

filter.threshold	<i>Summary of threshold results</i>
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---

**Description**

Summary of threshold results.

**Usage**

```

filter.threshold(cross, pheno.col, latent.eff, res.var, lod.thrs, drop.lod = 1.5,
                s.quant, n.perm, alpha.levels, qh.thrs, ww.thrs, addcovar = NULL,
                intcovar = NULL, verbose = FALSE, ...)

```

**Arguments**

cross	object of class cross; see <a href="#">read.cross</a>
pheno.col	phenotype columns used for filtering thresholds

<code>latent.eff</code>	ratio of latent effect SD to residual SD
<code>res.var</code>	residual variance (=SD <sup>2</sup> )
<code>lod.thrs</code>	LOD threshold values for range of significance (alpha) levels
<code>drop.lod</code>	LOD drop from max LOD to keep in analysis
<code>s.quant</code>	vector of 1:Nmax with Nmax the maximum hotspot size to be considered
<code>n.perm</code>	number of permutations
<code>alpha.levels</code>	range of significance levels; same length as <code>lod.thrs</code>
<code>qh.thrs</code>	Results of call to <a href="#">hotperm</a>
<code>ww.thrs</code>	Results of call to <a href="#">ww.perm</a>
<code>addcovar</code>	additive covariates as vector or matrix; see <a href="#">scanone</a>
<code>intcovar</code>	interactive covariates as vector or matrix; see <a href="#">scanone</a>
<code>verbose</code>	verbose output if TRUE
<code>...</code>	arguments passed along to <a href="#">scanone</a>

**Value**

List with items

`NL.thrs`

`N.thrs`

`WW.thrs`

`NL`

`N.counts`

`WW.counts`

**References**

Manichaikul A, Dupuis J, Sen S, Broman KW (2006) Poor performance of bootstrap confidence intervals for the location of a quantitative trait locus. *Genetics* 174: 481-489.

**See Also**

[hotperm](#), [ww.perm](#), [scanone](#)

---

GetCandReg	<i>Get genetic information on candidate regulators and co-mapping traits.</i>
------------	---

---

### Description

Get chromosome (phys.chr) and physical position in cM (phys.pos), along with the LOD score (peak.lod) at the peak position (peak.pos), and the chromosome where the peak is located (peak.chr). Some candidates may map to the same chromosome where they are physically located.

### Usage

```
GetCandReg(highobj, annot, traits)
GetCisCandReg(highobj, cand.reg, lod.thr = NULL)
GetCoMappingTraits(highobj, cand.reg)
```

### Arguments

highobj	data frame from <a href="#">highlod</a> , which is sparse summary of high LODs in large <a href="#">scanone</a> object
annot	data frame with annotation information; must have first column as unique identifier, third column as chromosome, and fifth column as position in cM; typically column 2 has gene name, and column 4 has position in Mb
traits	names of traits to examine as candidate regulators; names must correspond to phenotypes in cross object
cand.reg	data frame with candidate regulator; see value section below
lod.thr	LOD threshold; restrict to intervals above this value if not NULL

### Details

Traits that map to positions close to their physical locations are said to map in cis (local linkages). Traits that map to positions away from their physical locations are said to map in trans (distal linkages). There is no unambiguous way to determine how close a trait needs to map to its physical location in order to be classified as cis. Our choice is to classify a trait as cis if the 1.5-LOD support interval (Manichaikul et al. 2006) around the LOD peak contains the trait's physical location, and if the LOD score at its physical location is higher than the LOD threshold. The function `GetCisCandReg` determines which of the candidate regulators map in cis. The function `GetCoMappingTraits` returns a list with the putative targets of each trait. A trait is included in the putative target list of a trait when its LOD peak is greater than `lod.thr` and the drop LOD support interval around the peak contains the location of the trait's QTL. The function `JoinTestOutputs` currently relies on external files that contain results of [FitAllTests](#). It needs to be rewritten to save space.

**Value**

GetCoMappingTraits returns a list with each element being the names of co-mapping traits for a particular name in traits. GetCandReg returns a data frame while GetCisCandReg returns a list with a similar candidate regulator data frame as the element cis.reg, and the index of trait names as the element cis.index. The elements of the candidate regulator data frame are as follows (peak.pos.lower and peak.pos.upper only for GetCisCandReg):

gene	name of trait, which might be a gene name
phys.chr	chromosome on which gene physically resides
phys.pos	physical position (in cM)
peak.chr	chromosome where peak LOD is located
peak.pos	position of peak (in cM)
peak.lod	LOD value at peak
peak.pos.lower, peak.pos.upper	lower and upper bounds of the 1.5-LOD support interval around peak.pos

**Author(s)**

Elias Chaibub Neto

**References**

Manichaikul et al. (2006) Genetics

**See Also**

[highlod](#), [FitAllTests](#), [scanone](#)

**Examples**

```
## data(CMSTCross) is loaded lazily.
CMSTscan <- scanone(CMSTCross, pheno.col = 1:3, method = "hk")
CMSThigh <- highlod(CMSTscan)
traits <- names(CMSTCross$pheno)
annot <- data.frame(name = traits, traits = traits, chr = rep(1, 3),
  Mb.pos = c(55,10,100))
annot$cM.pos <- annot$Mb.pos
cand.reg <- GetCandReg(CMSThigh, annot, traits)
cis.cand.reg <- GetCisCandReg(CMSThigh, cand.reg)
comap.targets <- GetCoMappingTraits(CMSThigh, cand.reg)
```

---

 GetCommonQtls

*Get common QTLs for phenotypes*


---

### Description

Perform joint QTL mapping for phenotypes with marginal LOD peak positions higher than LOD threshold and within set distance of each other

### Usage

```
GetCommonQtls(cross, pheno1, pheno2, thr = 3, peak.dist = 5,
  addcov1 = NULL, addcov2 = NULL, intcov1 = NULL, intcov2 = NULL)
```

### Arguments

cross	object of class cross
pheno1	first phenotype column number or character string name
pheno2	second phenotype column number or character string name; if more than one, then all phenotypes will be tested against pheno1
thr	LOD threshold
peak.dist	maximal peak distance to be considered the same peak (in cM)
addcov1, addcov2	additive covariates for first and second phenotype, respectively
intcov1, intcov2	interactive covariates for first and second phenotype, respectively

### References

Chaibub Neto E, Broman AT, Keller MP, Attie AD, Zhang B, Zhu J, Yandell BS, Causal model selection hypothesis tests in systems genetics. *Genetics* (in review).

### See Also

[CMSTCross](#)

### Examples

```
data(CMSTCross)
commqtls <- GetCommonQtls(CMSTCross,
  pheno1 = "y1",
  pheno2 = "y3",
  thr = 3,
  peak.dist = 5,
  addcov1 = NULL,
  addcov2 = NULL,
  intcov1 = NULL,
  intcov2 = NULL)

commqtls
```



---

highlod	<i>Pull high LOD values with chr and pos.</i>
---------	---

---

### Description

Pull high LOD values with chr and pos.

### Usage

```
highlod(scans, lod.thr = 0, drop.lod = 1.5,
        extend = TRUE, restrict.lod = FALSE, ...)
pull.highlod(object, chr, pos, ...)
## S3 method for class 'highlod'
print(x, ...)
## S3 method for class 'highlod'
summary(object, ...)
## S3 method for class 'highlod'
plot(x, ..., quant.level = NULL, sliding = FALSE)
## S3 method for class 'highlod'
max(x, lod.thr = NULL, window = NULL, quant.level = NULL, ...)
## S3 method for class 'highlod'
quantile(x, probs = NULL, lod.thr = NULL, n.quant,
         n.pheno, max.quantile = TRUE, ...)
```

### Arguments

scans	object of class <a href="#">scanone</a>
lod.thr	LOD threshold
drop.lod	LOD drop from max to keep for support intervals
extend	extend support interval just past drop.lod; matches behavior of <a href="#">lodint</a> when TRUE
restrict.lod	restrict to loci above LOD threshold if TRUE; matches behavior of <a href="#">lodint</a> when FALSE (default)
chr	chromosome identifier
pos	position, or range of positions, in cM
x, object	object of class highlod
probs	probability levels for quantiles (should be > 0.5)
n.quant	maximum of s.quant
n.pheno	number of phenotypes considered
max.quantile	return only quantiles of max LOD across genome if TRUE
window	size of window for smoothing hotspot size
quant.level	vector of LOD levels for 1 up to length(quant.level) size hotspots
sliding	plot as sliding hotspot if TRUE
...	arguments passed along

**Details**

The `highlod` condenses a `scanone` object to the peaks above a `lod.thr` and/or within `drop.lod` of such peaks. The `pull.highlod` pulls out the entries at a particular genomic location or interval of locations. `Summary`, `print`, `plot`, `max` and `quantile` methods provide ways to examine a `highlod` object.

**Value**

Data frame with

<code>row</code>	row number in <code>scanone</code> object
<code>phenos</code>	phenotype column number
<code>lod</code>	LOD score for phenotype at locus indicated by row

**Author(s)**

Brian S Yandell and Elias Chaibub Neto

**See Also**

[highlod](#), [hotperm](#)

**Examples**

```
example(include.hotspots)
scan1 <- scanone(cross1, pheno.col = 1:1000, method = "hk")
high1 <- highlod(scan1, lod.thr = 2.11, drop.lod = 1.5)
pull.highlod(high1, chr = 2, pos = 24)
summary(high1, lod.thr = 2.44)
max(high1, lod.thr = seq(2.11, 3.11, by = .1))
```

---

hotperm

*Conduct NL and N permutation tests*

---

**Description**

Conduct NL and N permutation tests.

**Usage**

```
hotperm(cross, n.quant, n.perm, lod.thrs, alpha.levels, drop.lod = 1.5,
        window = NULL, verbose = FALSE, init.seed = 0,
        addcovar = NULL, intcovar = NULL, ...)
data(hotperm1)
## S3 method for class 'hotperm'
print(x, ...)
## S3 method for class 'hotperm'
```

```
summary(object, quant.levels, ...)
## S3 method for class 'hotperm'
quantile(x, probs, ..., lod.thr = NULL)
## S3 method for class 'summary.hotperm'
print(x, ...)
```

### Arguments

cross	object of class cross
n.quant	maximum of s.quant
n.perm	number of permutations
lod.thrs	vector of LOD thresholds
alpha.levels	vector of significance levels
quant.levels	quantile levels, as number of traits, to show in summary; default is 1, 2, 5, 10, ... up to maximum recorded
drop.lod	LOD drop amount for support intervals
window	window size for smoothed hotspot size
verbose	verbose output if TRUE
init.seed	initial seed for pseudo-random number generation
x,object	object of class hotperm or summary.hotperm
probs	probability levels for quantiles (1-probs if all > 0.5); default is alpha.levels
lod.thr	restrict to values above this if not NULL
addcovar	additive covariates as vector or matrix; see <a href="#">scanone</a>
intcovar	interactive covariates as vector or matrix; see <a href="#">scanone</a>
...	arguments passed along to scanone

### Author(s)

Elias Chaibub Neto and Brian S Yandell

### Examples

```
example(include.hotspots)
set.seed(123)
pt <- scanone(ncross1, method = "hk", n.perm = 1000)
alphas <- seq(0.01, 0.10, by=0.01)
lod.thrs <- summary(pt, alphas)
## Not run:
## This takes awhile, so we save the object.
set.seed(12345)
hotperm1 <- hotperm(cross = cross1,
                   n.quant = 300,
                   n.perm = 100,
                   lod.thrs = lod.thrs,
                   alpha.levels = alphas,
                   drop.lod = 1.5,
```

```

        verbose = FALSE)
save(hotperm1, file = "hotperm1.RData", compress = TRUE)

## End(Not run)
summary(hotperm1)

```

---

hotsize

*Hotspot size routines.*


---

### Description

Determine hotspot sizes and display. Use individual threshold and quantile thresholds as provided.

### Usage

```

hotsize(hotobject, ...)
## S3 method for class 'scanone'
hotsize(hotobject, lod.thr = NULL, drop.lod = 1.5, ...)
## S3 method for class 'highlod'
hotsize(hotobject, lod.thr = NULL, window = NULL,
        quant.level = NULL, ...)
## S3 method for class 'hotsize'
print(x, ...)
## S3 method for class 'hotsize'
summary(object, ...)
## S3 method for class 'hotsize'
plot(x, ylab = "counts", quant.axis = pretty(x$max.N),
     col = c("black", "red", "blue"), by.chr = FALSE, maps = NULL,
     title = "", ...)

```

### Arguments

hotobject	object of class <a href="#">scanone</a> or <a href="#">highlod</a>
lod.thr	LOD threshold
drop.lod	LOD drop from max to keep for support intervals
window	window width in cM for smoothing hotspot size; not used if 0 or NULL
quant.level	vector of LOD levels for 1 up to length(quant.level) size hotspots
x, object	object of class hotsize
ylab	label for vertical plot axis
quant.axis	hotspot sizes for quantile axis (vertical on right side of plot)
col	col of hotspot size, smoothed hotspot size, and sliding hotspot size
by.chr	separate plot by chromosome if TRUE
maps	if not NULL, list of objects of class map to use for rugs on top and bottom of plot
title	title for plot
...	arguments passed along to scanone methods

**Value**

hotsize methods return an object of class hotsize, which is essentially an object of class `summary.scanone` with additional attributes for `lod.thr`, `window`, and `quant.level`.

**Author(s)**

Brian S Yandell and Elias Chaibub Neto

**See Also**

[highlod](#), [hotperm](#)

**Examples**

```
example(highlod)
hots1 <- hotsize(high1)
summary(hots1)
plot(hots1)
```

---

parallel.qtlhot

*Code for parallelizing R/qtlhot.*

---

**Description**

Code for parallelizing R/qtlhot. See installed parallel directory for proper use. There is apparently an S3 parallel method, so doc has to be as shown below, even though it is called as parallel.qtlhot.

**Usage**

```
## S3 method for class 'qtlhot'
parallel(x, data = 1, ..., dirpath = ".")
qtlhot.phase0(dirpath, init.seed = 92387475, len = rep(400,16), n.mar = 185, n.ind = 112,
  n.phe = 100, latent.eff = 0, res.var = 1, lod.thrs, ...)
big.phase0(dirpath, cross, trait.file, trait.matrix, droptrait.names = NULL,
  keeptrait.names = NULL, lod.thrs, sex = "Sex", trait.index,
  batch.effect = NULL, size.set = 250, offset = 0, subset.sex = NULL, verbose = TRUE)
```

**Arguments**

x	phase of parallel processing (1,2,3)
data	index for parallel processing (1,2,...)
...	additional arguments passed along
dirpath	directory path as character string
init.seed	initial seed for pseudorandom number generation
len	vector of chromosome lengths for simulated map
n.mar	number of markers for simulated map

n.ind	number of individuals for simulated cross
n.phe	number of phenotypes for simulated phenotypes
latent.eff	size of latent effect
res.var	magnitude of residual variance
lod.thrs	vector of LOD thresholds to examine
cross	object of class cross
trait.file	character string name of trait file
trait.matrix	character string name of trait matrix
droptrait.names	vector of character strings for traits to drop (none if NULL)
keeptrait.names	vector of character strings for traits to keep (keep all if NULL)
sex	character string name of phenotype for sex
trait.index	vector of character strings for trait names
batch.effect	character string for batch effect (none if NULL)
size.set	maximum size of set of traits to scan at one time
offset	offset for name of trait RData files
subset.sex	string of sex to subset on (both sexes if NULL)
verbose	verbose output if TRUE

**Author(s)**

Brian S Yandell and Elias Chaibub Neto

**See Also**

[read.cross](#)

---

PrecTpFpMatrix

*Determine false positive and true positive rates for known targets.*

---

**Description**

Determine how well different tests do to predict candidates of regulation.

**Usage**

```
FitAllTests(cross, pheno1, pheno2, Q.chr, Q.pos, verbose = TRUE)
JoinTestOutputs(comap, tests, file)
PrecTpFpMatrix(alpha, val.targets, all.orfs, tests, cand.reg, cis.cand.reg)
p.adjust.np(tests, method = "BH")
```

**Arguments**

cross	object of class cross
pheno1	first phenotype column number or character string name
pheno2	second phenotype column number or character string name; if more than one, then all phenotypes will be tested against pheno1
Q.chr	QTL chromosome (number or label)
Q.pos	QTL position in cM
verbose	verbose printout if TRUE
comap	list result of GetComappingTraits
alpha	significance levels at which summaries are computed
val.targets	validated targets of candidate regulators
all.orfs	all trait names
tests	list object as list of FitAllTests results, or of joined output created by JoinTestsOutputs
file	prefix for file names when running FitAllTests in parallel and saving test results in separate files
cand.reg	object from <a href="#">GetCandReg</a>
cis.cand.reg	object from <a href="#">GetCisCandReg</a>
method	method for p-value adjustment; see <a href="#">p.adjust</a>

**Details**

FitAllTests invokes 7 tests. The hidden routine CitTests is invoked by call to FitAllTests; this is hidden because we do not recommend its use.

JoinTestOutputs joins results of [FitAllTests](#), either from a list tests or from a collection of files prefixed by file. The joined tests from JoinTestOutputs are summarized with PrecTpFpMatrix using the biologically validated true positives, false positives and precision, for the inferred causal relations. We define a true positive as a statistically significant causal relation between a gene and a putative target gene when the putative target gene belongs to the known signature of the gene. Similarly, we define a false positive as a statistically significant causal relation between a gene and a putative target gene when the target gene does not belong to the signature. (For the AIC and BIC methods that do not provide a p-value measuring the significance of the causal call, we simply use the detected causal relations in the computation of true and false positives). The validated precision is computed as the ratio of true positives by the sum of true and false positives. The PrecTpFpMatrix computes these measures to both all genes, and to cis genes only. Simulations suggest only non-parametric tests need to be adjusted using Benjamini-Hochberg via [p.adjust.np](#).

**Value**

List containing	
Prec1, Prec2	matrix of precision with rows for significance level and columns for test; first is for all, second is for cis candidates only
Tp1, Tp2	matrix of true positive rate with rows for significance level and columns for test; first is for all, second is for cis candidates only
Fp1, Fp2	matrix of false positive rate with rows for significance level and columns for test; first is for all, second is for cis candidates only

**Author(s)**

Elias Chaibub Neto

**See Also**[GetCandReg](#), [CMSTtests](#), [p.adjust](#)**Examples**

```

example(GetCandReg)
## Suppose y1 is causal with targets y2 and y3.
targets <- list(y1 = c("y2","y3"))

tests <- list()
for(k in seq(names(comap.targets))) {
  tests[[k]] <- FitAllTests(CMSTCross, pheno1 = names(comap.targets)[k],
                           pheno2 = comap.targets[[k]],
                           Q.chr = cand.reg[k, 4],
                           Q.pos = cand.reg[k, 5])
}
names(tests) <- names(comap.targets)
tests <- JoinTestOutputs(comap.targets, tests)

PrecTpFpMatrix(alpha = seq(0.01, 0.10, by = 0.01),
               val.targets = targets, all.orfs = CMSThigh$names, tests = tests,
               cand.reg = cand.reg, cis.cand.reg = cis.cand.reg)

```

---

sim.hotspot

*Wrapper routine for simulations.*


---

**Description**

Wrapper routine for simulations

**Usage**

```

sim.hotspot(nSim, cross, n.pheno, latent.eff, res.var = 1, n.quant, n.perm,
           alpha.levels, lod.thrs, drop.lod = 1.5, verbose = FALSE)
mySimulations(...)
sim.null.cross(chr.len = rep(400, 16), n.mar = 185, n.ind = 112,
              type = "bc", n.pheno = 6000, latent.eff = 1.5, res.var = 1,
              init.seed = 92387475)
sim.null.pheno.data(cross, n.pheno, latent.eff, res.var)
include.hotspots(cross, hchr, hpos, hsize, Q.eff, latent.eff,
                 lod.range.1, lod.range.2, lod.range.3, res.var=1, n.pheno, init.seed)

```



**Arguments**

nSim	Number of simulated sets of phenotypes to create. See details.
cross	Object of class cross. See <a href="#">read.cross</a> .
n.pheno	Number of traits, or phenotypes, to simulate for cross object.
latent.eff	Strength of latent effect, which is included in all traits. See <a href="#">sim.null.cross</a> .
res.var	Residual variance for traits. Should not affect results.
n.quant	maximum size of hotspots examined; ideally large enough to exceed the largest Breitling alpha critical value.
n.perm	Number of permutations to perform per realization. Good idea to do 1000, but this takes time.
alpha.levels	Vector of significance levels.
lod.thrs	Vector of LOD thresholds, typically single-trait permutation thresholds for various significance levels.
drop.lod	Drop in LOD score examined. LODs below this drop from the maximum for a chromosome will not be scored.
init.seed	initial seed for pseudo-random number generation
chr.len	vector of chromosome lengths
n.mar	number of markers
n.ind	number of individuals
type	type of cross
hchr,hpos,hsize	vectors for hotspot chromosomes, positions, and sizes
Q.eff	QTL effect
lod.range.1,lod.range.2,lod.range.3	2-vectors of LOD ranges for multiple purposes
verbose	Verbose output if TRUE. More detailed output if 2.
...	Arguments passed directly to sim.hotspot.

**Details**

Simulate nSim realizations of cross object with n.pheno phenotypes with correlation latent.eff. All simulations use the same genotypes in the cross object.

**Value**

sim.null.cross simulates an object of class cross. sim.null.pheno.data simulates a data frame of phenotypes. sim.hotspot uses these other routines to simulate a hotspot, returning an list object.

**Author(s)**

Elias Chaibub Neto and Brian S. Yandell

**See Also**

[sim.null.cross](#), [read.cross](#).

**Examples**

```
ncross1 <- sim.null.cross(chr.len = rep(100, 4),
  n.mar = 51,
  n.ind = 100,
  type = "bc",
  n.phe = 1000,
  latent.eff = 3,
  res.var = 1,
  init.seed = 123457)
cross1 <- include.hotspots(cross = ncross1,
  hchr = c(2, 3, 4),
  hpos = c(25, 75, 50),
  hsize = c(100, 50, 20),
  Q.eff = 2,
  latent.eff = 3,
  lod.range.1 = c(2.5, 2.5),
  lod.range.2 = c(5, 8),
  lod.range.3 = c(10, 15),
  res.var = 1,
  n.phe = 1000,
  init.seed = 12345)
```

---

SimCrossCausal

*Simulate Cross for Causal Tests*

---

**Description**

Creates cross with certain pattern of dependence across phenotypes.

**Usage**

```
SimCrossCausal(n.ind, len, n.mar, beta, add.eff, dom.eff,
  sig2.1 = 1, sig2.2 = 1, eq.spacing = FALSE,
  cross.type = c("bc", "f2"), normalize = FALSE)
SimCrossIndep(n.ind, len, n.mar, beta, add.eff.1, dom.eff.1,
  add.eff.h, dom.eff.h, sig2.1 = 1, sig2.2 = 1, sig2.h = 1,
  eq.spacing = FALSE, cross.type = "f2", normalize = FALSE)
data(CMSTCross)
```

**Arguments**

n.ind	number of individuals to simulate
len	vector specifying the chromosome lengths (in cM)
n.mar	vector specifying the number of markers per chromosome

beta                   causal effect (slope) of first phenotype on others  
 add.eff, add.eff.1, add.eff.h  
                           additive genetic effect  
 dom.eff, dom.eff.1, dom.eff.h  
                           dominance genetic effect  
 sig2.1                 residual variance for first phenotype  
 sig2.2, sig2.h         residual variance for all other phenotypes  
 eq.spacing            if TRUE, markers will be equally spaced  
 cross.type            type of cross (bc and f2 for now)  
 normalize             normalize values if TRUE

## References

Chaibub Neto E, Broman AT, Keller MP, Attie AD, Zhang B, Zhu J, Yandell BS, Causal model selection hypothesis tests in systems genetics. *Genetics* (in review).

## Examples

```

set.seed(987654321)
CMSTCross <- SimCrossCausal(n.ind = 100,
  len = rep(100, 3), n.mar = 101,
  beta = rep(0.5, 2), add.eff = 1, dom.eff = 0,
  sig2.1 = 0.4, sig2.2 = 0.1, eq.spacing = FALSE,
  cross.type = "bc", normalize = TRUE)
CMSTCross <- calc.genoprob(CMSTCross, step = 1)
## Not run:
save(CMSTCross, file = "CMSTCross.RData", compress = TRUE)

## End(Not run)

```

---

 ww.perm

*Conduct West-Wu (Q) permutation tests*


---

## Description

Conduct West-Wu (Q) permutation tests.

## Usage

```

ww.perm(highobj, n.perm, lod.thrs, alpha.levels, verbose = FALSE)
## S3 method for class 'ww.perm'
print(x, ...)
## S3 method for class 'ww.perm'
summary(object, alpha.levels, ...)

```

**Arguments**

highobj	object of class <code>highlod</code>
n.perm	number of permutations
lod.thrs	vector of LOD thresholds
alpha.levels	vector of significance levels
x,object	object of class <code>ww.perm</code>
...	ignored
verbose	verbose output if TRUE

**Details**

Perform permutation tests to assess the statistical significance of the hotspots detected using the West-Wu Q-method permutations. The `ww.perm` function implements the Q-method's permutation scheme (see the Method's section of Chaibub Neto et al. 2012, for details). The `n.perm` parameter specifies the number of simulations. Here we set it to 100 in order to save time. In practice, we recommend at least 1,000 permutations. The function's output is a matrix with 100 rows representing the permutations, and 10 columns representing the QTL mapping thresholds. Each entry  $ij$ , represents the maximum number of significant linkages across the entire genome detected at permutation  $i$ , using the LOD threshold  $j$ . The `ww.summary` function computes the Q-method's hotspot size permutation thresholds, that is, the  $1-\alpha$  quantiles for each one of the QTL mapping LOD thresholds in `lod.thrs`. For instance, the entry at row 10 and column 1 of the `Q.1.thr` matrix tells us that the 99% percentile of the permutation distribution of genome wide maximum hotspot size based on a QTL mapping threshold of 2.11 is 27.00. In other words, any hotspot greater than 27 is considered statistically significant at a 0.01 significance level when QTL mapping is done using a 2.11 LOD threshold. In general, we are often interested in using the same error rates for the QTL mapping and hotspot analysis. That is, if we adopt a QTL mapping threshold that controls GWER at a 1% level (in our case, 3.11) we will also want to consider  $\alpha = 0.01$  for the hotspot analysis, leading to a hotspot threshold of 12.00. Therefore, we are usually more interested in the diagonal of `Q.1.thr`. We adopted a GWER of 5%, and the corresponding Q-method's permutation threshold is 18. According to this threshold, all hotspots are significant.

**Author(s)**

Elias Chaibub Neto and Brian S Yandell

**Examples**

```
## Not run:
## All unspecified objects come from vignette qtlhot.
set.seed(12345)
Q.1 <- ww.perm(high1, n.perm = 100, lod.thrs, alphas)
Q.1.thr <- summary(Q.1, alphas)
Q.1.thr
diag(Q.1.thr)

set.seed(12345)
Q.2 <- ww.perm(high2, 100, lod.thrs, alphas)
Q.2.thr <- summary(Q.2, alphas)
```

## End(Not run)

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