

# Package ‘forensIT’

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**Title** Information Theory Tools for Forensic Analysis

**Version** 1.1.1

**Description** The ‘forensIT’ package is a comprehensive statistical toolkit tailored for handling missing person cases. By leveraging information theory metrics, it enables accurate assessment of kinship, particularly when limited genetic evidence is available. With a focus on optimizing statistical power, ‘forensIT’ empowers investigators to effectively prioritize family members, enhancing the reliability and efficiency of missing person investigations.

**License** GPL (>= 3)

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**Depends** R (>= 2.10)

**NeedsCompilation** no

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buildEnsembleCPTs	<i>buildEnsembleCPTs</i>
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## Description

Build ensemble of CPTs from a list of simulations

## Usage

```
buildEnsembleCPTs(lsimu, lminimalProbGenoMOI)
```

## Arguments

lsimu	list of simulations
lminimalProbGenoMOI	list of minimal probabilities of genotypes given MOI # nolint

## Value

list of CPTs

**Examples**

```

library(forrel)
library(mispitools)
freqs <- lapply(getfreqs(Argentina)[1:15], function(x) {x[x!=0]})
fam <- linearPed(2)
fam <- addChildren(fam, father = 1, mother = 2)
fam <- pedtools::setMarkers(fam, locusAttributes = freqs)
ped <- profileSim(fam, N = 1, ids = c(6) , numCores = 1,seed=123)
lsimEnsemble <- simTestIDMarkers(ped,2,numSim=5,seed=123)
lensembleIT <- buildEnsembleITValues(lsimu=lsimEnsemble,ITtab=simME$ITtable,bFullIT = TRUE)
lensembleCPTs <- buildEnsembleCPTs(lsimu=lsimEnsemble,lminimalProbGenoMOI=simME$lprobGenoMOI)

```

---

buildEnsembleITValues *buildEnsembleITValues*

---

**Description**

Build ensemble of IT values from a list of simulations

**Usage**

```

buildEnsembleITValues(
  lsimu = lsimulation,
  ITtab = sim$ITtable,
  bFullIT = FALSE
)

```

**Arguments**

lsimu	list of simulations
ITtab	IT table
bFullIT	boolean to return full IT table

**Value**

list of IT values

**Examples**

```

library(forrel)
library(mispitools)
freqs <- lapply(getfreqs(Argentina)[1:15], function(x) {x[x!=0]})
fam <- linearPed(2)
fam <- addChildren(fam, father = 1, mother = 2)
fam <- pedtools::setMarkers(fam, locusAttributes = freqs)
ped <- profileSim(fam, N = 1, ids = c(6) , numCores = 1,seed=123)
lsimEnsemble <- simTestIDMarkers(ped,2,numSim=5,seed=123)
lensembleIT <- buildEnsembleITValues(lsimu=lsimEnsemble,ITtab=simME$ITtable,bFullIT = TRUE)

```

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compareBnetPopGenoPDFs

*Compare population and Bayesian network genotype probability density functions # nolint*

---

### Description

Compare population and Bayesian network genotype probability density functions # nolint

### Usage

compareBnetPopGenoPDFs(lprobTable)

### Arguments

lprobTable      list of probability tables

### Value

list of KL divergences

---

crossH

*Cross entropy*

---

### Description

Cross entropy

### Usage

crossH(px, py, epsilon = 1e-20)

### Arguments

px                  probability distribution  
 py                  probability distribution  
 epsilon            small number to avoid log(0)

### Value

cross entropy

---

distKL	<i>distKL: KL distribution obtained for specific relative contributor</i>
--------	---

---

**Description**

distKL: KL distribution obtained for specific relative contributor

**Usage**

```
distKL(ped, missing, relative, frequency, numsims = 100, cores = 1)
```

**Arguments**

ped	Reference pedigree. It could be an input from read_fam() function or a pedigree built with pedtools. # nolint
missing	Missing person
relative	Selected relative.
frequency	Allele frequency database.
numsims	Number of simulated genotypes.
cores	Enables parallelization.

**Value**

An object of class data.frame with KLS.

**Examples**

```
library(forrel)
x = linearPed(2)
x = setMarkers(x, locusAttributes = NorwegianFrequencies[1:2])
x = profileSim(x, N = 1, ids = 2)
distKL(ped = x, missing = 5, relative = 1, cores = 1,
frequency = NorwegianFrequencies[1:2], numsims = 3)
```

---

elimLangeGoradia	<i>Eliminate Mendelian errors using Lange-Goradia algorithm</i>
------------------	---

---

**Description**

Eliminate Mendelian errors using Lange-Goradia algorithm

**Usage**

```
elimLangeGoradia(ped, iMarker = 1, bitera = TRUE, bverbose = TRUE)
```

**Arguments**

ped	pedigree
iMarker	index of marker to be used
bitera	iterate until no more errors are found
bverbose	print progress

**Value**

pedigree with Mendelian errors eliminated

---

exportPed	<i>Export a pedigree to a file</i>
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---

**Description**

Export a pedigree to a file

**Usage**

```
exportPed(ped, fname, iMarker = 1)
```

**Arguments**

ped	pedigree
fname	file name
iMarker	index of marker to be used

**Value**

pedigree with Mendelian errors eliminated

---

forensIT	<i>forensIT: Information Theory Tools for Forensic Analysis</i>
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---

**Description**

The 'forensIT' package, available on CRAN, is a comprehensive statistical toolkit tailored for handling missing person cases. By leveraging information theory metrics, it enables accurate assessment of kinship, particularly when limited genetic evidence is available. With a focus on optimizing statistical power, 'forensIT' empowers investigators to effectively prioritize family members, enhancing the reliability and efficiency of missing person investigations. Experience the power of information theory in kinship testing with the user-friendly 'forensIT' package, freely accessible on CRAN. # nolint

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Authors:

- Ariel Chernomoretz <ariel@df.uba.ar>

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genotypeProbs            *Genotype probabilities*

---

**Description**

Calculate genotype probabilities from parental probabilities

**Usage**

```
genotypeProbs(probP, probM)
```

**Arguments**

probP            vector of parental probabilities  
probM            vector of parental probabilities

**Value**

matrix of genotype probabilities

---

genotypeProbTable        *Genotype Probability Table*

---

**Description**

Genotype Probability Table

**Usage**

```
genotypeProbTable(bbn1, resQQ, bplot = FALSE, numMarkers = 4, lLoci)
```

**Arguments**

bbn1            Bayesian network  
resQQ           results from bn  
bplot           boolean to plot  
numMarkers     number of markers  
lLoci           list of loci

**Value**

Genotype Probability Table

---

genotypeProbTable\_bis *genotypeProbTable\_bis*

---

**Description**

function to calculate the probability of genotypes given the MOI

**Usage**

```
genotypeProbTable_bis(bbn1, resQQ, bplot = FALSE, numMarkers = 4, freq)
```

**Arguments**

bbn1	bayesian network
resQQ	list of results from the inference
bplot	plot results
numMarkers	number of markers
freq	allele frequencies

**Value**

matrix of genotype probabilities

---

getAllelesFromGenotypes  
*getAllelesFromGenotypes*

---

**Description**

Get alleles from genotypes

**Usage**

```
getAllelesFromGenotypes(g)
```

**Arguments**

g	genotypes
---	-----------

**Value**

alleles



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H	<i>Entropy of a discrete probability distribution</i>
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---

**Description**

Entropy of a discrete probability distribution

**Usage**

H(px, epsilon = 1e-20, normalized = FALSE)

**Arguments**

px	probability distribution
epsilon	small number to avoid log(0)
normalized	boolean to normalize entropy

**Value**

entropy

---

index2Genotypes2	<i>index2Genotypes2</i>
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---

**Description**

index2Genotypes2

**Usage**

index2Genotypes2(ped, id, iMarker, alleleSet)

**Arguments**

ped	pedigree
id	individual id
iMarker	marker index
alleleSet	allele set

**Value**

genotypes

---

index2Genotypes2.pedtools  
*index2Genotypes*

---

**Description**

index2Genotypes

**Usage**

index2Genotypes2.pedtools(ped, id, iMarker, alleleSet)

**Arguments**

ped	pedigree
id	individual id
iMarker	marker index
alleleSet	allele set

**Value**

genotypes

---

KLd                      *KL divergence*

---

**Description**

KL divergence

**Usage**

KLd(ppx, ppy, epsilon = 1e-20, bsigma = FALSE)

**Arguments**

ppx	probability distribution
ppy	probability distribution
epsilon	small number to avoid log(0)
bsigma	boolean to compute sigma

**Value**

KL divergence

---

KLde	<i>KL divergence</i>
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---

**Description**

KL divergence

**Usage**

KLde(px, py, epsilon = 1e-20)

**Arguments**

px	probability distribution
py	probability distribution
epsilon	small number to avoid log(0)

**Value**

KL divergence

---

perMarkerKLs	<i>perMarkerKLs</i>
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---

**Description**

perMarkerKLs

**Usage**

perMarkerKLs(ped, MP, frequency)

**Arguments**

ped	Reference pedigree.
MP	missing person
frequency	Allele frequency database.

**Value**

An object of class data.frame with KLs.

**Examples**

```
library(forrel)
x = linearPed(2)
plot(x)
x = setMarkers(x, locusAttributes = NorwegianFrequencies[1:5])
x = profileSim(x, N = 1, ids = 2)
perMarkerKLs(x, MP = 5 , NorwegianFrequencies[1:5])
```

---

plotKL

*Plot KL distances.*

---

**Description**

Plot KL distances.

**Usage**

```
plotKL(res)
```

**Arguments**

res                    output from distKL function.

**Value**

A scatterplot.

**Examples**

```
library(forrel)
x = linearPed(2)
plot(x)
x = setMarkers(x, locusAttributes = NorwegianFrequencies[1:5])
x = profileSim(x, N = 1, ids = 2)
res <- distKL(ped = x, missing = 5, relative = 1,
cores = 1, frequency = NorwegianFrequencies[1:5], numsims = 5)
plotKL(res)
```

---

Px

*Px*


---

**Description**

Px

**Usage**

Px(p1, p0, dbg = FALSE)

**Arguments**

p1	probability distribution
p0	probability distribution
dbg	boolean to compute sigma

**Value**

Px

---

runIT

*runIT*


---

**Description**

run information theory (IT) metrics

**Usage**

```
runIT(
  lped = NULL,
  freqs,
  QP,
  dbg,
  numCores,
  bOnlyIT = FALSE,
  lprobq_ped = NULL,
  bsigma = FALSE,
  blog = FALSE,
  dep = TRUE
)
```

**Arguments**

lped	list of pedigree objects
freqs	list of allele frequencies
QP	QP
dbg	debug
numCores	number of cores
bOnlyIT	boolean to only run IT
lprobG_ped	list of probG
bsigma	boolean to compute sigma
blog	boolean to write log
dep	check fbnet dependency

**Value**

runIT

simLR

*Simulate LR***Description**

Simulate LR

**Usage**

```
simLR(
  lprobG_ped,
  numSim = 10000,
  epsilon = 1e-20,
  bplot = FALSE,
  bLRs = FALSE,
  seed = 123457
)
```

**Arguments**

lprobG_ped	list of probability distributions
numSim	number of simulations
epsilon	small number to avoid log(0)
bplot	boolean to plot
bLRs	boolean to return LRs
seed	seed

**Value**

LRs

---

simME	<i>simME: output from simMinimalEnsemble considering an uncle</i>
-------	---

---

**Description**

simME: output from simMinimalEnsemble considering an uncle

**Usage**

```
simME
```

**Format**

A list with minimalEnsemble of genotypes

---

simMinimalEnsemble	<i>simMinimalEnsemble</i>
--------------------	---------------------------

---

**Description**

It performs simulations of minimal ensembles of genotypes

**Usage**

```
simMinimalEnsemble(  
  ped,  
  QP,  
  testID,  
  freqs,  
  numCores = 1,  
  seed = 123457,  
  bVerbose = TRUE,  
  bJustGetNumber = FALSE,  
  bdbg = FALSE,  
  dep = TRUE  
)
```

**Arguments**

ped	pedigree
QP	QP
testID	test ID
freqs	frequencies
numCores	number of cores

seed	seed
bVerbose	boolean to print information
bJustGetNumber	boolean to just get the number of runs
bdbg	boolean to debug
dep	check dependency fbnet

**Value**

list of results

---

simTestIDMarkers	<i>Simulate testID markers</i>
------------------	--------------------------------

---

**Description**

Simulate testID markers

**Usage**

```
simTestIDMarkers(ped, testID, numSim = 10, seed = 123457)
```

**Arguments**

ped	pedigree
testID	test ID
numSim	number of simulations
seed	seed

**Value**

list of simulations

**Examples**

```
library(forrel)
library(mispitools)
freqs <- lapply(getfreqs(Argentina)[1:15], function(x) {x[x!=0]})
fam <- linearPed(2)
fam <- addChildren(fam, father = 1, mother = 2)
fam <- pedtools::setMarkers(fam, locusAttributes = freqs)
ped <- profileSim(fam, N = 1, ids = c(6) , numCores = 1,seed=123)
lsimEnsemble <- simTestIDMarkers(ped,2,numSim=5,seed=123)
```



---

`strsplit2`*strsplit2*

---

**Description**

strsplit2

**Usage**`strsplit2(x, split)`**Arguments**`x` character vector`split` character**Value**matrix

---

`trioCheckFast`*trioCheckFast*

---

**Description**

Check for Mendelian errors in trios

**Usage**`trioCheckFast(ffa, mmo, oof)`**Arguments**`ffa` father's alleles`mmo` mother's alleles`oof` offspring's alleles**Value**

TRUE if there is a Mendelian error

---

unidimKLplot	<i>unidimKLplot: KL distributions presented in the same units (Log10(LR))</i>
--------------	---

---

**Description**

unidimKLplot: KL distributions presented in the same units (Log10(LR))

**Usage**

```
unidimKLplot(res)
```

**Arguments**

res                    output from distKL function.

**Value**

A scatterplot.

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