

# Package ‘IntEREst’

January 9, 2025

**Title** Intron-Exon Retention Estimator

**Version** 1.30.0

**Date** 2024-03-19

**Author** Ali Oghabian <Ali.Oghabian@Helsinki.Fi>, Dario Greco  
<dario.greco@helsinki.fi>, Mikko Frilander  
<Mikko.Frilander@helsinki.fi>

**Maintainer** Ali Oghabian <Ali.Oghabian@Helsinki.Fi>

**Description** This package performs Intron-Exon Retention analysis on RNA-seq data (.bam files).

**Depends** R (>= 3.5.0), GenomicRanges, Rsamtools, SummarizedExperiment, edgeR, S4Vectors, GenomicFiles

**Imports** seqLogo, Biostrings, GenomicFeatures (>= 1.39.4), txdbmaker, IRanges, seqinr, graphics, grDevices, stats, utils, grid, methods, DBI, RMariaDB, GenomicAlignments, BiocParallel, BiocGenerics, DEXSeq, DESeq2

**Suggests** clinfun, knitr, rmarkdown, BSgenome.Hsapiens.UCSC.hg19

**VignetteBuilder** knitr

**LazyData** true

**biocViews** Software, AlternativeSplicing, Coverage, DifferentialSplicing, Sequencing, RNASeq, Alignment, Normalization, DifferentialExpression, ImmunoOncology

**License** GPL-2

**NeedsCompilation** no

**git\_url** <https://git.bioconductor.org/packages/IntEREst>

**git\_branch** RELEASE\_3\_20

**git\_last\_commit** 806b1f1

**git\_last\_commit\_date** 2024-10-29

**Repository** Bioconductor 3.20

**Date/Publication** 2025-01-09

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 IntREEst-package

*IntREEst*


---

## Description

Intron/Exon retention estimator quantifies and normalizes Intron retention and Exon junction read levels by analyzing mapped reads (.bam) files.

**Details**

Package: IntREst  
Type: Package  
Version: 1.0  
Date: 2015-11-18  
License: GPL-2

To run the pipeline use functions `interest()` or `interest.sequential()`, i.e. wrapper functions that run all the necessary functions.

**Author(s)**

Ali Oghabian <Ali.Oghabian@Helsinki.Fi>, Dario Greco <dario.greco@helsinki.fi>, Mikko Frilander <Mikko.Frilander@helsinki.fi>

Maintainer: Ali Oghabian <Ali.Oghabian@Helsinki.Fi>, Mikko Frilander <Mikko.Frilander@helsinki.fi>

---

addAnnotation      *Adding sample annotations to a SummarizedExperiment object*

---

**Description**

Adds a new sample annotation to the `SummarizedExperiment` object. In other words it adds and column with sample annotations to the `colData` of the `SummarizedExperiment` object.

**Usage**

```
addAnnotation(x, sampleAnnotationType, sampleAnnotation)
```

**Arguments**

`x`                    Object of type `SummarizedExperiment`.  
`sampleAnnotationType`  
                      The name of the new column to be added to the `colData` table of `SummarizedExperiment` object.  
`sampleAnnotation`  
                      Vector with the same length as the row-size of the `colData` attribute of the `SummarizedExperiment` object, which includes the sample annotations.

**Value**

An `InterestResult` object.

**Author(s)**

Ali Oghabian

**See Also**

[getAnnotation](#)

**Examples**

```
# Check the annotation table of mdsChr220bj data
getAnnotation(mdsChr220bj)

# Add a new sample annotation
newMdsChr220bj <- addAnnotation(x=mdsChr220bj,
  sampleAnnotationType="sample_number",
  sampleAnnotation=1:16
)

# Retrieve annotations of the new object
getAnnotation(newMdsChr220bj)
```

---

 annotateU12

*Annotate the U12 (and U2) type introns*


---

**Description**

Receives coordinates, a reference genome and PWMs of splice site of U12 and U2 type introns, and returns a data.frame with 2 columns. The first column shows whether the corresponding sequences matches U12, U2 or both (U12/U2) consensus sequences (based on their score when fitting the PWMs). The second column shows whether the match is on positive strand or negative when fitting the PWMs to the sequences.

**Usage**

```
annotateU12(pwmU12U2=c(), pwmSsIndex=c(), referenceChr, referenceBegin,
  referenceEnd, referenceIntronExon, intronExon='intron',
  matchWindowRelativeUpstreamPos=c() , matchWindowRelativeDownstreamPos=c(),
  minMatchScore='80%', refGenome='', setNaAs='U2', annotateU12Subtype=TRUE,
  includeMatchScores=FALSE, ignoreHybrid=TRUE, filterReference)
```

**Arguments**

pwmU12U2	A list containing position weight matrices of (in order): Donor site, branch point, and acceptor site of U12-type introns, and donor site and acceptor site of U2-type introns. If not provided, the information related to pwmU12db data is used.
pwmSsIndex	A list (or vector) that contains the column number in each element of pwmU12U2 that represents the 5' or 3' Splice Site; The order should be equivalent to the pwmU12U2. If not provided the information from pwmU12db data is used, i.e. <code>pwmSsIndex=list(indexDonU12=1, indexBpU12=1, indexAccU12=3, indexDonU2=1, indexAccU2=3)</code>
referenceChr	Chromosome names of the references (e.g. introns).
referenceBegin	A vector that corresponds to the begin coordinates of the reference (e.g. introns).
referenceEnd	A vector that corresponds to the end coordinates of the reference (e.g. introns). <code>referenceEnd</code> should be greater than or equal to <code>referenceBegin</code> .
referenceIntronExon	A vector with the same size as the <code>referenceChr</code> , <code>referenceBegin</code> and <code>referenceEnd</code> which contains 'intron' and 'exon' describing what (either intron or exon) each element of the 3 vectors represents.

intronExon	Should be assigned either 'intron' or 'exon' or c('intron','exon') based on whether match the PWM to the intronic, exonic, or intronic and exonic regions of the reference. By default it seeks matches in intronic regions (intronExon='intron').
matchWindowRelativeUpstreamPos	A vector the same size as the pwmU12U2 (and the same order of donor/acceptor sites' information in pwmU12U2) which consists of the upstream distance from the donor/acceptor site from which each PWM should be tested (to see if they match). If not provided, the information from pwmU12db data is used i.e. matchWindowRelativeUpstreamPos=c(NA, -29, NA, NA, NA).
matchWindowRelativeDownstreamPos	A vector the same size as the pwmU12U2 (and the same order of donor/acceptor sites' information in pwmU12U2) which consists of the downstream distance from the donor/acceptor site to which each PWM should be tested (to see if they match). If not provided, the information from pwmU12db data is used i.e. matchWindowRelativeDownstreamPos=c(NA, -9, NA, NA, NA).
minMatchScore	Min percentage match score, when scoring matching of a sequence to pwm. Different score thresholds could also be defined for the various sites (U12/U2 donors, the U12 branch point and U12/U2 acceptors); A vector with 5 elements can be assigned which each shows the match score to use for each PWM in pwmU12U2.
refGenome	The reference genome; Object of class BSgenome. Use available.genome() from the BSgenome package to see the available genomes. DNASTringSet objects (from Biostrings package) and fasta files are also accepted as input.
setNaAs	Defines that if reference (e.g. intron) did not match any of U12 or U2 type introns based on the scores obtained from PWM what should the function return. If an intron was not proven to be U12 or U2 based on PWM scores it can be considered as U2-type since the U12 type introns constitute for about 1% of introns in human genome and they are much more conserved than the U2 type introns, hence the default is 'U2'; otherwise it is also possible to set it as NA or nan or 'U12/U2'.
annotateU12Subtype	Whether annotate the subtypes of the U12 type Introns. The value is TRUE by default.
includeMatchScores	If set as TRUE the final data frame result includes the PWM match scores (FALSE by default).
ignoreHybrid	Whether ignore the U12 hybrid subtypes, i.e. GT-AC and AT-AG (TRUE by default).
filterReference	Optional parameter that can be defined either as a GRanges or SummarizedExperiment object. If defined as the latter, the first 3 columns of the rowData must be: chr name, start and end of the coordinates. If the parameter is defined the introns/exon coordinates will be mapped against it and the intron type of all those that do not match will be set as NA.

**Value**

Data frame containing 3 columns representing (in order): intron type (U12, U2 or none), strand match indicating whether the PWM matches to the sequence (+ strand) or the reverse complement of the sequence (- strand) or none (NA), and the U12 subtype (GT-AG or AT-AC). If includeMatchScores is set as TRUE further columns that include the PWM match scores will also be included.

**Author(s)**

Ali Oghabian

**See Also**[buildSsTypePwms.](#)**Examples**

```
# Improting genome
BSgenome.Hsapiens.UCSC.hg19 <-
  BSgenome.Hsapiens.UCSC.hg19::BSgenome.Hsapiens.UCSC.hg19
#Choosing subset of rows
ind<- 69:94
# Annotate U12 introns with strong U12 donor site, branch point
# and acceptor site from the u12 data in the package
annoU12<-
  annotateU12(pwmU12U2=list(pwmU12db[[1]][,11:17],pwmU12db[[2]]
    ,pwmU12db[[3]][,38:40],pwmU12db[[4]][,11:17],
    pwmU12db[[5]][,38:40]),
  pwmSsIndex=list(indexDonU12=1, indexBpU12=1, indexAccU12=3,
    indexDonU2=1, indexAccU2=3),
  referenceChr=u12[ind,'chr'],
  referenceBegin=u12[ind,'begin'],
  referenceEnd=u12[ind,'end'],
  referenceIntronExon=u12[ind,"int_ex"],
  intronExon="intron",
  matchWindowRelativeUpstreamPos=c(NA,-29,NA,NA,NA),
  matchWindowRelativeDownstreamPos=c(NA,-9,NA,NA,NA),
  minMatchScore=c(rep(paste(80,"%",sep=""),2), "60%",
    paste(80,"%",sep=""), "60%"),
  refGenome=BSgenome.Hsapiens.UCSC.hg19,
  setNaAs="U2",
  annotateU12Subtype=TRUE)

# How many U12 and U2 type introns with strong U12 donor sites,
# acceptor sites (and branch points for U12-type) are there?
table(annoU12[,1])
```

---

 applyOverlap

*Apply function over counts*


---

**Description**

Runs a function on columns of the counts (assay) of a 'SummarizedExperiment' object (resulted by `interest()`, `interest.sequential()` or `readInterestResults()`) based on the overlap of its exon/intron coordinates with those of another 'SummarizedExperiment' object. The number of the rows and the dimensions of the counts of the result are equal to those of the subject. The function is applied on the query based on it's overlap to the subject.

**Usage**

```

applyOverlap(
  query,
  subject,
  type="any",
  replaceValues=FALSE,
  intExCol="int_ex",
  intronExon="intron",
  subjectGeneNamesCol,
  repeatsTableToFilter=c(),
  scaleFragment=TRUE,
  scaleLength=TRUE,
  unmapValue=0,
  FUN=mean,
  ...
)

```

**Arguments**

query, subject	SummarizedExperiment objects resulted by <code>interest()</code> , <code>interest.sequential()</code> or <code>readInterestResults()</code> functions.
type	The type of overlap. By default it considers any overlap. See <a href="#">findOverlaps-methods</a> for more info.
replaceValues	Whether return a 'SummarizedExperiment' object with new counts (resulted by running function) replaced.
intExCol	Column name (or number) in the rowData of the objects that represents whether each row of the assay is "intron" or "exon".
intronExon	Should be assigned either 'intron' or 'exon' or <code>c('intron', 'exon')</code> based on whether match the PWM to the intronic, exonic, or intronic and exonic regions of the reference. By default it seeks matches in intronic regions ( <code>intronExon='intron'</code> ).
subjectGeneNamesCol	The column in the row data of the subject that includes the gene names.
repeatsTableToFilter	A data.frame table that includes chr,begin and end columns. If defined, all reads mapped to the described regions will be ignored.
scaleFragment	Logical value, indicating whether the retention levels must be scaled by (genome-wide) fragment levels.
scaleLength	Logical value, indicating whether the retention levels must be scaled by length of the introns/exons.
unmapValue	The value to assign to unmapped rows (i.e. introns/exons).
FUN	The function to apply.
...	Other parameter settings from <code>aggregate()</code> function.

**Value**

The returned value is a data frame if `replaceValues` is `FALSE` and it is `SummarizedExperiment` if `replaceValues` is `TRUE`.

**Author(s)**

Ali Oghabian

**See Also**[readInterestResults](#) [interest](#) [interest.sequential](#)**Examples**

```

mdsChr220bj

tmp<- applyOverlap(
  query=mdsChr220bj,
  subject=mdsChr220bj,
  type="equal",
  replaceValues=FALSE,
  intExCol="int_ex",
  intronExon="intron",
  subjectGeneNamesCol="collapsed_transcripts",
  scaleFragment=TRUE,
  scaleLength=TRUE,
  unmapValue=0,
  FUN=head,
  n=1
)

```

attributes

*Extracting values of useful attributes of SummarizedExperiment objects***Description**

Several functions are provided that can extract various attributes from an object of class `SummarizedExperiment` generated by `IntREEst` functions, e.g. `interest()`, `interest`, and `readInterestResults`. It is possible to extract sample annotations using `getAnnotation` function. One can also extract the scaled retention levels of the introns/exons using `scaledRetention()` function. Notes that `colData` and `rowData` methods of `SummarizedExperiment` class can also be used to extract row and column data.

**Usage**

```

getAnnotation(x)
scaledRetention(x)

```

**Arguments**

x                    Object of type `SummarizedExperiment`.

**Value**

Various data types (`data.frame`/`vector`) dependent on the function used. See the "Description" for more information.



**Author(s)**

Ali Oghabian

**See Also**[SummarizedExperiment-class addAnnotation counts-method plot-method](#)**Examples**

```
# Retrieve the sample annotations from mdsChr22Obj
getAnnotation(mdsChr22Obj)
# Retrieving the scaled retention levels from mdsChr22Obj
head(scaledRetention(mdsChr22Obj))

#for row and column data SummarizedExperiment methods can be used
head(rowData(mdsChr22Obj))
colData(mdsChr22Obj)
```

boxplot-method

*boxplot - method***Description**

boxplot method for SummarizedExperiment objects.

**Usage**

```
## S4 method for signature 'SummarizedExperiment'
boxplot(x, sampleAnnoCol=NA,
        intexTypeCol="int_type", intexType=c(), col="white", boxplotNames=c(),
        lasNames=3, outline=FALSE, addGrid=FALSE, ...)
```

**Arguments**

<code>x</code>	Object of type SummarizedExperiment generated by either <code>interest()</code> , <code>interest.sequential()</code> or <code>readInterestResults()</code> .
<code>sampleAnnoCol</code>	Which column of <code>colData</code> in <code>x</code> to consider for plotting.
<code>intexTypeCol</code>	Column name (or number) that represents what type of intron/exon each row of <code>x</code> assays represents.
<code>intexType</code>	A vector of characters describing types of introns/exons to be plotted. They must be elements in the <code>intexTypeCol</code> column of the <code>rowData</code> of <code>x</code> . <code>rowData</code> of <code>x</code> is a dataframe that includes various annotations of the introns/exons.
<code>col</code>	Vector showing box colours. It is either of size 1 or the same size as the number of groups to be plotted.
<code>boxplotNames</code>	Names to write under boxes. If not defined, as names, it pastes the row (intron/exon) annotation names to the sample group annotations separated by a space " ".
<code>lasNames</code>	Orientation of the box names.

outline	If outline is TRUE the outlier points are drawn otherwise if FALSE (default) they are not.
addGrid	Whether add a grid under the boxplots (FALSE by default).
...	Other arguments to pass to the boxplot() and axis function.

**Value**

Returns NULL.

**Author(s)**

Ali Oghabian

**See Also**

Class: [SummarizedExperiment-class](#) Method: [counts-method](#) [plot-method](#)

**Examples**

```
#Plotting U12- vs U2-type introns
par(mar=c(8,4,2,1))
boxplot(x=mdsChr220bj, sampleAnnoCol="type", intexTypeCol="intron_type",
  intexType=c("U2", "U12"),
  col=rep(c("yellow", "orange"),3),
  boxplotNames=c(), lasNames=3, outline=FALSE,
  addGrid=TRUE)
```

---

buildSsTypePwms	<i>Building Position Weight Matrices for Splice Sites of U12 and U2 type introns.</i>
-----------------	---

---

**Description**

Builds position Weigh Matrices for the donor and acceptor sites of the U12 and U2 type introns, and the branchpoint of the U12 type introns. if pdfFileSeqLogos is defined a pdf is also produced that contains the sequence logos of the results. The result is a list that contains PWMs of the splice sites of U12 and U2 dependent introns.

**Usage**

```
buildSsTypePwms( cexSeqLogo=1, pdfWidth=35, pdfHeight=10, tmpDir="./",
  u12dbSpecies="Homo_sapiens",
  pwmSource="U12DB",
  u12DonorBegin, u12BranchpointBegin, u12AcceptorBegin,
  u2DonorBegin, u2AcceptorBegin, u12DonorEnd,
  u12BranchpointEnd, u12AcceptorEnd, u2DonorEnd,
  u2AcceptorEnd, pasteSites=FALSE,
  splicerackSsLinks=list(
  U12_AT_AC_donor=
    "http://katahdin.mssm.edu/splice/out/9606_logo_file.25",
  U12_AT_AC_branchpoint=
```

```

    "http://katahdin.mssm.edu/splice/out/9606_logo_file.26",
    U12_AT_AC_acceptor=
    "http://katahdin.mssm.edu/splice/out/9606_logo_file.29",
    U12_GT_AG_donor=
    "http://katahdin.mssm.edu/splice/out/9606_logo_file.22",
    U12_GT_AG_branchpoint=
    "http://katahdin.mssm.edu/splice/out/9606_logo_file.27",
    U12_GT_AG_acceptor=
    "http://katahdin.mssm.edu/splice/out/9606_logo_file.21",
    U2_GC_AG_donor="http://katahdin.mssm.edu/splice/out/9606_logo_file.24",
    U2_GC_AG_acceptor=
    "http://katahdin.mssm.edu/splice/out/9606_logo_file.30",
    U2_GT_AG_donor="http://katahdin.mssm.edu/splice/out/9606_logo_file.23",
    U2_GT_AG_acceptor=
    "http://katahdin.mssm.edu/splice/out/9606_logo_file.28"),
    u12dbLink="https://genome.crg.cat/pub/software/u12/u12db_v1_0.sql.gz",
    u12dbDbName="u12db", u12dbDropDb=TRUE, pdfFileSeqLogos="",
    removeTempFiles=TRUE, ...)

```

### Arguments

cexSeqLogo	Font size of sequence logo plots; used only if pdfFileSeqLogos is defined.
pdfWidth, pdfHeight	The width and height of the graphics region of the pdf in inches. The default values are 35 and 10.
tmpDir	Path to directory used for storing temporary files.
u12dbSpecies	What species data to use when getting the data from the U12DB database (pwmSource="U12DB").
pwmSource	The source used to buildSplice Sites of U12 and U2 type introns the PWM for U12 and U2 dependent introns. Default is U12DB; but also accepts SpliceRack.
u12DonorBegin, u12DonorEnd	Integer values. They correspond to the begin and end point of the donor sequences of U12-type introns to consider (optional).
u12BranchpointBegin, u12BranchpointEnd	Integer values. Begin and end points of the branch point sequences of U12-type introns (optional).
u12AcceptorBegin, u12AcceptorEnd	Integer values. Begin and end points of the acceptor sequences of U12-type introns (optional).
u2DonorBegin, u2DonorEnd	Integer values. Begin and end points of the donor sequences of U2-type introns (optional).
u2AcceptorBegin, u2AcceptorEnd	Integer values. Begin and end points of the acceptor sequences of U2-type introns (optional).
pasteSites	Logical. If TRUE the donor, branch point and acceptor seqs are pasted before a PWM is built; then the PWMs of each (donor, acceptor and bp) are assigned. If FALSE (default) the PWMs for each is built separately.
splicerackSsLinks	A list (or vector) that contains the SpliceRack URL links to the text files that contain Position Weigh Matrices of the splice sites of U12 and U2 introns. This

parameter is used only when `pwmSource="SpliceRack"`. You can get the links to PWM files from this URL (choose logo files with "File" links): [http://katahdin.mssm.edu/splice/splice\\_matrix.cgi?database=spliceNew](http://katahdin.mssm.edu/splice/splice_matrix.cgi?database=spliceNew). The links should be defined in the following order: U12\_AT\_AC\_donor, U12\_AT\_AC\_branchpoint, U12\_AT\_AC\_acceptor, U12\_GT\_AG\_donor, U12\_GT\_AG\_branchpoint, U12\_GT\_AG\_acceptor, U2\_GC\_AG\_donor, U2\_GC\_AG\_acceptor, U2\_GT\_AG\_donor, and U2\_GT\_AG\_acceptor.

<code>u12dbLink</code>	A character string containing the URL for downloading the zipped MySQL dump file of the U12DB. Used when <code>pwmSource="U12DB"</code> .
<code>u12dbDbName</code>	Name of the database copy of the U12DB that is build locally. Used when <code>pwmSource="U12DB"</code> .
<code>u12dbDropDb</code>	Drop (or remove) the local copy of the U12DB database at the end of the run. Used when <code>pwmSource="U12DB"</code> .
<code>pdfFileSeqLogos</code>	Path to PDF file containing the sequence logos of the results. By default it does not produce a file.
<code>removeTempFiles</code>	Whether remove temporary files at the end of the run; accepts TRUE or FALSE values (default is TRUE).
<code>...</code>	Authorization arguments needed by the DBMS instance. See the manual for <code>dbConnect</code> of the DBI package for more info.

### Value

<code>pwmDonorU12</code>	Matrix (with 4 rows representing A, C, G, T and n columns representing the genomic coordinates) representing the Position Weight Matrix of donor site of U12-type introns.
<code>pwmBpU12</code>	Position Weight Matrix of branchpoint of U12-type introns.
<code>pwmAccU12</code>	Position Weight Matrix of acceptor site of U12-type introns.
<code>pwmDonU2</code>	Position Weight Matrix of donor site of U2-type introns.
<code>pwmAccU2</code>	Position Weight Matrix of acceptor site of U2-type introns.

### Author(s)

Ali Oghabian

### See Also

[annotateU12](#).

### Examples

```
# Time demanding function
## Not run:
#Build temp directory
tmpDir<- tempdir()

# Creating subdirectory for storing u12db temp files
dir.create(paste(tmpDir, "u12dbTmp", sep="/"))

# Extracting PWMs of Splice Sites of U12 and U2 type introns -
```

```
# based on u12db
u12dbPwm<-buildSsTypePwms(
  tmpDir=paste(tmpDir, "u12dbTmp", sep="/"),
  u12dbSpecies="Homo_sapiens",
  resource="U12DB",
  u12dbDbName="u12db",
  u12dbDropDb=TRUE,
  removeTempFiles=TRUE)

# Creating subdirectory for storing SpliceRack temp files
dir.create(paste(tmpDir, "splicerackTmp", sep="/"))

# Extracting PWMs of Splice Sites of U12 and U2 type introns -
# based on SpliceRack
spliceRackPwm<- buildSsTypePwms(
  tmpDir= paste(tmpDir, "splicerackTmp", sep="/"),
  resource="SpliceRack",
  removeTempFiles=TRUE)

## End(Not run)
```

---

counts-method

*Counts - method*

---

## Description

Returns the (row) number of reads that are mapped to introns/exons in various samples.

## Usage

```
## S4 method for signature 'SummarizedExperiment'
counts(object)
```

## Arguments

object            Object of type SummarizedExperiment.

## Value

Returns a numeric matrix.

## Author(s)

Ali Oghabian

## See Also

Class: [SummarizedExperiment-class](#)

Method: [plot-method](#).

**Examples**

```

#Show contents of a InterestResults object included in IntEREst
head(counts(mdsChr22Obj))

#Make a test InterestResults object
geneId<- paste("gene", c(rep(1,5), rep(2,5), rep(3,5), rep(4,5)),
  sep="_")
readCnt1<- sample(1:100, 20)
readCnt2<- sample(1:100, 20)
readCnt3<- sample(1:100, 20)
readCnt4<- sample(1:100, 20)
fpkm1<- readCnt1/(tapply(readCnt1, geneId, sum))[geneId]
fpkm2<- readCnt2/(tapply(readCnt2, geneId, sum))[geneId]
fpkm3<- readCnt3/(tapply(readCnt3, geneId, sum))[geneId]
fpkm4<- readCnt4/(tapply(readCnt4, geneId, sum))[geneId]

# Creating object using test data
interestDat<- data.frame(
  int_ex=rep(c(rep(c("exon", "intron"),2), "exon"),4),
  int_ex_num= rep(c(1,1,2,2,3),4),
  gene_id= geneId,
  sam1_readCnt=readCnt1,
  sam2_readCnt=readCnt2,
  sam3_readCnt=readCnt3,
  sam4_readCnt=readCnt4,
  sam1_fpkm=fpkm1,
  sam2_fpkm=fpkm2,
  sam3_fpkm=fpkm3,
  sam4_fpkm=fpkm4
)
readFreqColIndex<- grep("_readCnt$", colnames(interestDat))
scaledRetentionColIndex<- grep("_fpkm$", colnames(interestDat))

scalRetTmp<- as.matrix(interestDat[, scaledRetentionColIndex])
colnames(scalRetTmp)<-gsub("_fpkm$", "", colnames(scalRetTmp))

frqTmp<- as.matrix(interestDat[, readFreqColIndex])
colnames(frqTmp)<-gsub("_readCnt$", "", colnames(frqTmp))

InterestResultObj<- InterestResult(
  resultFiles=paste("file", 1:4, sep="_"),
  rowData= interestDat[, -c(readFreqColIndex,
  scaledRetentionColIndex)],
  counts= frqTmp,
  scaledRetention= scalRetTmp,
  scaleLength=TRUE,
  scaleFragment=FALSE,
  sampleAnnotation=data.frame(
    sampleName=paste("sam", 1:4, sep=""),
    gender=c("M", "M", "F", "F"),
    health=c("healthy", "unhealthy", "healthy", "unhealthy")
  , row.names=paste("sam", 1:4, sep="")
  )
)

```

```
#Show
head(counts(InterestResultObj))
```

---

```
deseqInterest      DESeq2 analysis for IntERESt object
```

---

## Description

Differential intron retention test adapted from the DESeq2 package.

## Usage

```
deseqInterest(x, design, pAdjustMethod = "BH",
              sizeFactor=c(), contrast, bparam, ...)
```

## Arguments

x	Object of type SummarizedExperiment.
design	Formula specifying the design of the experiment. It must specify an interaction term between variables from column names of <code>sampleData(x)</code> .
pAdjustMethod	What adjustment method to be used on the p-values. See <a href="#">p.adjust</a> for more information.
sizeFactor	Numeric vector with the same size as the column size of the count matrix in x, if defined it will be used for scaling of the count matrix.
contrast	Argument specifying the comparison to extract from x. See <a href="#">results</a> function in the DESeq2 package for more information.
bparam	An optional BiocParallelParam instance defining the parallel back-end to be used. If not defined the function will run sequentially (on a single computing core).
...	Other parameter settings for the <a href="#">results</a> function in the DESeq2 package.

## Value

a DESeqResults object.

## Author(s)

Ali Oghabian

## See Also

[exactTestInterest](#) [qlfInterest](#), [treatInterest](#) [DEXSeqIntERESt](#)

## Examples

```
mCh22IntObj<- mCh22Obj[rowData(mCh22Obj)$int_ex=="intron",]
deseqRes<- deseqInterest(x=mCh22IntObj,
                        design=~test_ctrl, contrast=list("test_ctrl_test_vs_ctrl"))

# Number of U12/U2 type significantly differential retained introns in chr22
table(rowData(mCh22Obj)[which(deseqRes$padj<.01), "intron_type"])
```

---

DEXSeqIntEREst                      *DEXSeq test for IntEREst object*

---

### Description

Genewise differential exon usage or intron retention test adapted from the DEXSeq package.

### Usage

```
DEXSeqIntEREst (x, design, reducedModel = ~ sample + intex, fitExpToVar,
intExCol, geneIdCol, bpparam, silent=TRUE,...)
```

### Arguments

x	Object of type SummarizedExperiment.
design	Formula specifying the design of the experiment. It must specify an interaction term between a variable from columns of <code>sampleData(x)</code> with one of the 'exon', 'intron' or 'intex' (i.e. intron and exon) variables; based on which of these variables are used (exon, intron, or 'intex') the x will be filtered relatively to include exons, introns, or introns and exons. See <a href="#">DEXSeqDataSet</a> for more information.
reducedModel	The null model formula. By default it is '~ sample + intex'.
fitExpToVar	A variable name contained in the column data (i.e. column names of <code>colData(x)</code> ). See <a href="#">DEXSeq</a> for more information.
intExCol	Column name (or number) that represents whether each row is "intron" or "exon" in <code>rowData</code> of x.
geneIdCol	Column name (or number of column) in <code>rowData</code> of x, i.e. SummarizedExperiment object, that represents the gene ID of the introns and exons in x.
bpparam	An optional <code>BiocParallelParam</code> instance defining the parallel back-end to be used.
silent	Whether run the DEXSeq function silently (if TRUE) or allow it to print messages at each step (if FALSE).
...	Other parameter settings for the <a href="#">DEXSeqDataSet</a> function in the DEXSeq package.

### Details

The `design` and `reducedModel` accept formula that specify the design of the experiment. The formula must describe an interaction between variables from columns of `sampleData(x)` with one of the 'exon', 'intron' or 'intex' (i.e. intron and exon) variables; Based on which of these variables are used (exon, intron, or 'intex') the input object (x) will be filtered relatively to include exons, introns, or introns and exons. Hence the number of the rows of the returned value is equal to the number of the rows of the filtered object, i.e. the number of the exons, introns or both based on the design formula.

### Value

A `DEXSeqResults` object.



**Author(s)**

Ali Oghabian

**See Also**[exactTestInterest](#)**Examples**

```
dexseqExRes<-DEXSeqIntERESt (x=mdsChr22ExObj,
  design= ~ sample + exon + test_ctrl:exon,
  reducedModel = ~ sample + exon, fitExpToVar="test_ctrl",
  intExCol="int_ex", geneIdCol="transcripts_id", silent=TRUE)
head(dexseqExRes)
```

---

exactTestInterest	<i>Exact test</i>
-------------------	-------------------

---

**Description**

Compute genewise exact test between two groups of read counts, using the edgeR package.

**Usage**

```
exactTestInterest(x, sampleAnnoCol=c(), sampleAnnotation=c(),
  geneIdCol, silent=TRUE, group=c(), rejection.region="doubletail",
  big.count=900, prior.count=0.125, disp="common", ...)
```

**Arguments**

x	Object of type SummarizedExperiment.
sampleAnnoCol	Which column of colData of x to consider for the analysis.
sampleAnnotation	A vector of size 2 which contains values from colData of SummarizedExperiment object; e.g. if <code>getAnnotation(x)[,sampleAnnoCol]= c("test", "test", "ctrl", "ctrl", ...)</code> , and the goal is to compare "test" and "ctrl" samples, sampleAnnotation should either be <code>c("test", "ctrl")</code> or <code>c("ctrl", "test")</code> .
geneIdCol	Column name (or number of column) in rowData of x, i.e. SummarizedExperiment object, that represents the gene ID of the introns and exons in x.
silent	Whether run the function silently, i.e. without printing the top differential expression tags.
group	Vector to manually define the sample groups (or annotations). It is ignored if sampleAnnoCol is defined.
rejection.region	The rejection.region parameter in <a href="#">exactTest</a> from edgeR package.
big.count	The big.count parameter in <a href="#">exactTest</a> from edgeR package.
prior.count	The prior.count parameter in <a href="#">exactTest</a> from edgeR package.

disp	The type of estimating the dispersion in the data. Available options are: "tag-wise", "trended", "common" and "genewise". It is also possible to assign a number for manually setting the disp.
...	Other parameter settings for the <code>estimateDisp</code> function (e.g. the design parameter) in the edgeR package.

**Value**

table	Data frame containing columns for the log2 fold-change (logFC), the average of log2 counts-per-million (logCPM), and the two-sided p-value (PValue).
comparison	The name of the two compared groups.
dispersionType	The name of the type of dispersion used.
dispersion	The estimated dispersion values.

**Author(s)**

Ali Oghabian

**See Also**

[lfc](#), [glmInterest](#), [qlfInterest](#), [treatInterest](#), [DEXSeqIntERESt](#)

**Examples**

```
geneId<- paste("gene", c(rep(1,5), rep(2,5), rep(3,5), rep(4,5)),
  sep="_")
readCnt1<- sample(1:100, 20)
readCnt2<- sample(1:100, 20)
readCnt3<- sample(1:100, 20)
readCnt4<- sample(1:100, 20)
fpkm1<- readCnt1/(tapply(readCnt1, geneId, sum))[geneId]
fpkm2<- readCnt2/(tapply(readCnt2, geneId, sum))[geneId]
fpkm3<- readCnt3/(tapply(readCnt3, geneId, sum))[geneId]
fpkm4<- readCnt4/(tapply(readCnt4, geneId, sum))[geneId]

# Creating object using test data
interestDat<- data.frame(
  int_ex=rep(c(rep(c("exon","intron"),2),"exon"),4),
  int_ex_num= rep(c(1,1,2,2,3),4),
  gene_id= geneId,
  sam1_readCnt=readCnt1,
  sam2_readCnt=readCnt2,
  sam3_readCnt=readCnt3,
  sam4_readCnt=readCnt4,
  sam1_fpkm=fpkm1,
  sam2_fpkm=fpkm2,
  sam3_fpkm=fpkm3,
  sam4_fpkm=fpkm4
)
readFreqColIndex<- grep("_readCnt$",colnames(interestDat))
scaledRetentionColIndex<- grep("_fpkm$",colnames(interestDat))

scalRetTmp<- as.matrix(interestDat[,scaledRetentionColIndex])
colnames(scalRetTmp)<-gsub("_fpkm$","", colnames(scalRetTmp))
```

```

frqTmp<- as.matrix(interestDat[,readFreqColIndex])
colnames(frqTmp)<-gsub("_readCnt$", "", colnames(frqTmp))

InterestResultObj<- InterestResult(
  resultFiles=paste("file",1:4, sep="_"),
  rowData= interestDat[, -c(readFreqColIndex,
  scaledRetentionColIndex)],
  counts= frqTmp,
  scaledRetention= scalRetTmp,
  scaleLength=TRUE,
  scaleFragment=FALSE,
  sampleAnnotation=data.frame(
    sampleName=paste("sam",1:4, sep=""),
    gender=c("M","M","F","F"), row.names=paste("sam", 1:4, sep="")
  )
)

res<- exactTestInterest(InterestResultObj, sampleAnnoCol="gender",
  sampleAnnotation=c("F","M"), geneIdCol= "gene_id",
  silent=TRUE, disp="common")

```

---

getRepeatTable

*Get table of regions with repetitive DNA sequences*


---

### Description

This function returns a data.frame that includes regions with repetitive DNA sequences. These sequences can bias the mapping of the reads to the genome excluding them will remove the bias.

### Usage

```

getRepeatTable( dbUser="genome",
  dbHost="genome-mysql.cse.ucsc.edu", ucscGenome="hg19",
  ucscTable="rmsk", minLength=0, repFamilyFil="Alu",
  repFamilyCol="repFamily", repChrCol="genoName",
  repBegCol="genoStart", repEndCol="genoEnd",
  repStrandCol="strand", repNameCol="repName",
  repClassCol="repClass")

```

### Arguments

dbUser	Database user name; set as "genome" by default.
dbHost	Database host address; set as "genome-mysql.cse.ucsc.edu" by default.
ucscGenome	The UCSC genome.
ucscTable	The UCSC table name. The table with repetitive sequences by default it is set as "rmsk".
minLength	the minimum length criteria to consider the repetitive sequences. the default setting is 0.

repFamilyFil	A vector including the repeats family to consider. By default the "Alu" elements are considered.
repFamilyCol	The name of the column of the input table (ucscTable) that represents the repeats family.
repChrCol	The column (either name or the number of the column) of the input table that represents the Chromosome names.
repBegCol	The column of the table that represents the start coordinates.
repEndCol	The column of the table that represents the end coordinates.
repStrandCol	The column of the table that represents the strand.
repNameCol	The column of the table representing the repeats' names.
repClassCol	The column of the table representing the repeats' classes.

**Value**

Data frame with columns representing coordinates and annotations of repetitive DNA elements.

**Author(s)**

Ali Oghabian

**Examples**

```
## Not run:
# Download table for Alu elemnts in the human genome
suppressWarnings(repTable<- getRepeatTable(repFamilyFil="Alu",
  ucscGenome="hg19"))

## End(Not run)
```

---

glmInterest

*generalized linear model likelihood ratio tests*

---

**Description**

Compute generalized linear model likelihood ratio tests using edgeR package. For more information see [glmfit](#) and [glmLRT\(\)](#) functions in edgeR package.

**Usage**

```
glmInterest(x, design=c(), silent=TRUE, disp="common",
  coef=c(), contrast=NULL, ...)
```

**Arguments**

x	Object of type SummarizedExperiment.
design	Design matrix.
silent	Whether run the function silently, i.e. without printing the top differential expression tags. Default is TRUE.

disp	The method of estimating the dispersion in the data. Available options are: "common", "trended", "tagwiseInitCommon" and "tagwiseInitTrended". It is also possible to assign a number.
coef	Integer or character vector indicating which coefficients of the linear model are to be tested equal to zero. See <code>glmLRT()</code> in <code>edgeR</code> for more information.
contrast	Numeric vector or matrix specifying contrasts of the linear model coefficients to be tested equal to zero. See <code>glmLRT()</code> in <code>edgeR</code> for more information.
...	Other parameter settings for the <code>glmLRT()</code> function in the <code>edgeR</code> package.

**Value**

All values produced by `glmLRT` in `edgeR` package plus following:

dispersionType	The name of the type of dispersion used.
dispersion	The estimated dispersion values.

**Author(s)**

Ali Oghabian

**See Also**

[exactTestInterest](#), [qlfInterest](#), [treatInterest](#)

**Examples**

```
#Test retention differentiation across the 3 types of sampels
group <- getAnnotation(mdsChr220bj)[,"type"]
glmRes<- glmInterest(x=mdsChr220bj,
  design=model.matrix(~group), silent=TRUE,
  disp="tagwiseInitTrended", coef=2:3, contrast=NULL)
```

---

interest

*Wrapper function: Parallel run*

---

**Description**

A read summarization function that counts all the reads mapping to the introns/exons based on the users detailed parameter settings. The process can be run in parallel on multiple computing cores to improve its performance.

**Usage**

```
interest( bamFileYieldSize=1000000, bamFile, isPaired,
  isPairedDuplicate=FALSE, isSingleReadDuplicate= NA, reference,
  referenceGeneNames, referenceIntronExon, repeatsTableToFilter=c(),
  junctionReadsOnly=FALSE, outFile, logFile="",
  returnObj= FALSE, method=c("ExEx", "IntRet", "IntSpan", "ExSkip"),
  strandSpecific,
  bpparam, appendLogFile=FALSE, sampleName="")
```

```

scaleLength= c(TRUE,FALSE), scaleFragment= c(TRUE,TRUE),
limitRanges=GRanges(),
excludeFusionReads=FALSE,
loadLimitRangesReads=FALSE, ...)

```

## Arguments

<code>bamFileYieldSize</code>	Maximum number of pair reads in the temporary files created as the result of dividing the input .bam file.
<code>bamFile</code>	Path of the input bam file.
<code>isPaired</code>	Whether the bam file is the result of a paired end sequencing read mapping (TRUE) or not (FALSE).
<code>isPairedDuplicate</code>	Whether extract only (if set TRUE), filter (FALSE) or include (if set NA) PCR duplicates for paired mapped reads. It uses the FLAG field in the bam file to filter the duplicate read. If the mapping software does not support detection and flagging the duplicate reads dedup tool of BamUtil or MarkDuplicates of Picard tools could be used.
<code>isSingleReadDuplicate</code>	Whether extract only (if set TRUE), filter (FALSE) or include (if set NA) PCR duplicates for single mapped reads.
<code>reference</code>	Dataframe to be used as reference; It should at least contain three same-size vectors with the tag names chr, begin, and end which describe the exons and introns genome coordinates. It also accepts a GRanges object. To build a new reference check the <a href="#">referencePrepare</a> function.
<code>referenceGeneNames</code>	A vector with the same size as the row-size of the reference which includes the gene names of the reference.
<code>referenceIntronExon</code>	A vector with the same size as the row-size of the reference with values "intron" and "exon" describing which (intron or exon) each row of the reference represents.
<code>repeatsTableToFilter</code>	A dataframe table with similar structure to the reference. It includes chr, begin, and end columns. If defined, all reads mapped to the described regions would be ignored and the Intron/exon lengths would be corrected to exclude the to exclude the regions with repetitive DNA sequences. See <a href="#">getRepeatTable</a> .
<code>junctionReadsOnly</code>	The parameter is considered if the method is set as IntRet or ExEx (NOT IntSpan). It declares whether only consider the Intron-Exon or Exon-Exon junction reads and ignore the reads that fully map to exons or introns. By default this argument is set as FALSE.
<code>outFile</code>	The name or path of the result file.
<code>logFile</code>	The log file path; if defined log information are written to the log file.
<code>returnObj</code>	If set TRUE in addition to making result text files, the results would also be returned as an object of class SummarizedExperiment.
<code>method</code>	A vector describing the summarization methods to use; i.e. whether count reads mapping to the introns (IntRet), reads mapping to the exons (ExEx), reads spanning the introns (IntSpan), or reads that skip the exons (ExSkip). In IntSpan

mode the introns in the reference are taken into account only; whilst in `IntRet` the introns and their spanning exons, and in `ExEx` and `ExSkip` mode only the exons in the reference are taken into account.

<code>strandSpecific</code>	The description for strand specificity of the RNAseq data. The values are either "unstranded", "stranded", or "reverse". If the reads are not strand specific or directional use "unstranded". If the first read in paired-read sequencing or the reads single-read sequencing is in the same direction as the the transcript strand use "stranded". If the first read in paired-read sequencing or the reads in single-read sequencing is in the oposite direction to the transcript strand use "reverse".
<code>bpparam</code>	An optional <code>BiocParallelParam</code> instance defining the parallel back-end to be used.
<code>appendLogFile</code>	Whether log information should be appended to the <code>logfile</code> . It is set <code>FALSE</code> by default.
<code>sampleName</code>	The name of the sample being analyzed. It will be included in the returned object if <code>returnObj</code> is <code>TRUE</code> .
<code>scaleLength</code>	A vector constructed of <code>TRUE/FALSE</code> values, same size as the method argument. It indicates whether the retention levels of the intron/exons should be scaled to their lengths.
<code>scaleFragment</code>	A vector constructed of <code>TRUE/FALSE</code> values, same size as the method argument. It indicates whether the retention levels of the intron/exons should be scaled to the sum of retention levels (i.e. mapped fragments) over the genes.
<code>limitRanges</code>	A <code>GRanges</code> object. If defined it loads sequencing reads that fall in the defined coordinates. It is similar to which parameter in <a href="#">ScanBamParam</a> .
<code>excludeFusionReads</code>	Only valid if <code>limitRanges</code> is defined. It filters the defined by <code>limitRanges</code> . It also filters the read pairs if each read pair maps reads pairs where one of the reads either do not fall into one of the regions to a different region defined in <code>limitRanges</code> . It is useful to ignore analyzing the chimeric reads and fusion reads, i.e. reads that map to fusion genes. To filter properly, <code>limitRanges</code> must include coordinates of all genes.
<code>loadLimitRangesReads</code>	Boolean ( <code>TRUE/FALSE</code> ) variable. If set as <code>TRUE</code> only the reads in the <code>limitRanges</code> are loaded from bam file (and <code>bamFileYieldSize</code> parameter will be ignored).
...	Other parameter settings specific to <a href="#">BamFile-class</a> function in the <code>Rsamtools</code> package. Parameters <code>qnamePrefixEnd</code> and <code>qnameSuffixStart</code> are in particular useful to modify <code>qnames</code> in the BAM files.

**Value**

If `returnObj` is set `TRUE` in addition to making result text files, dependant on whether a single or two method is defined, the results would be returned as a single object of class `SummarizedExperiment` or as a list of size 2 which includes 2 objects of class `SummarizedExperiment` one for `IntRet` and the other for `ExEx`.

**Author(s)**

Ali Oghabian

**See Also**

[interest.sequential](#).

**Examples**

```
# Creating temp directory to store the results
outDir<- file.path(tempdir(),"interestFolder")
dir.create(outDir)
outDir<- normalizePath(outDir)

# Loading suitable bam file
bamF <- system.file("extdata", "small_test_SRR1691637_ZRSR2Mut_RHBDD3.bam",
  package="IntERESt", mustWork=TRUE)

# Choosing reference for the gene RHBDD3
ref= u12[u12[,"gene_name"]=="RHBDD3",]

test= interest(
  bamFileYieldSize=10000,
  bamFile=bamF,
  isPaired=TRUE,
  isPairedDuplicate=FALSE,
  isSingleReadDuplicate=NA,
  reference=ref,
  referenceGeneNames=ref[, "ens_gene_id"],
  referenceIntronExon=ref[, "int_ex"],
  repeatsTableToFilter=c(),
  outFile=paste(outDir,
    "interestRes.tsv", sep="/"),
  logFile=paste(outDir,
    "log.txt", sep="/"),
  method=c("IntRet", "IntSpan"),
  strandSpecific="unstranded",
  junctionReadsOnly=FALSE,
  returnObj=TRUE,
  scaleLength= c(TRUE,FALSE),
  scaleFragment= c(TRUE,TRUE)
)

test
```

---

interest.sequential      *Wrapup function: Sequential running*

---

**Description**

A read summarization function that counts all the reads mapping to the introns/exons based on the users detailed parameter settings. The process runs on a single computing core.

**Usage**

```
interest.sequential( bamFileYieldSize=1000000, bamFile, isPaired,
  isPairedDuplicate=FALSE, isSingleReadDuplicate=NA,
```



```
reference, referenceGeneNames,
referenceIntronExon, repeatsTableToFilter=c(),
junctionReadsOnly=FALSE, outFile, logFile="",
returnObj= FALSE, method=c("ExEx", "IntRet", "IntSpan", "ExSkip"),
strandSpecific, appendLogFile=FALSE, sampleName="",
scaleLength= c(TRUE,FALSE), scaleFragment= c(TRUE,TRUE),
limitRanges=GRanges(),
excludeFusionReads=FALSE,
loadLimitRangesReads=FALSE, ...)
```

## Arguments

bamFileYieldSize	Maximum number of paired Reads in the temporary files created as the result of dividing the input .bam file.
bamFile	Path of the input bam file.
isPaired	Whether the bam file is the result of a paired end sequencing read mapping (TRUE) or not (FALSE).
isPairedDuplicate	Whether extract only (if set TRUE), filter (FALSE) or include (if set NA) PCR duplicates for paired mapped reads. It uses the FLAG field in the bam file to filter the duplicate read. If the mapping software does not support detection and flagging the duplicate reads dedup tool of BamUtil or MarkDuplicates of Picard tools could be used.
isSingleReadDuplicate	Whether extract only (if set TRUE), filter (FALSE) or include (if set NA) PCR duplicates for single mapped reads.
reference	Dataframe to be used as reference; It should at least contain three same-size vectors with the tag names chr, begin, and end which describe the genome coordinates of the introns and exons. It also accepts a GRanges object as input. To build a new reference check the <a href="#">referencePrepare</a> function.
referenceGeneNames	A vector with the same size as the row-size of the reference which include the gene names.
referenceIntronExon	A vector with the same size as the row-size of the reference with values "intron" and "exon" describing which (intron or exon) each row of the reference represents.
repeatsTableToFilter	A data frame with similar structure as the reference, i.e. includes chr, begin, and end columns. If defined, all reads mapped to the described regions would be ignored and the Intron/exon lengths would be corrected to exclude the regions with repetitive DNA sequences. See <a href="#">getRepeatTable</a> .
junctionReadsOnly	The parameter is considered if the method is set as IntRet or ExEx (NOT IntSpan). It declares whether only consider the Intron-Exon or Exon-Exon junction reads and ignore the reads that fully map to exons or introns. By default this argument is set as FALSE.
outFile	The name or path of the result file.
logFile	The log file path; if defined log information are written to the log file.

returnObj	If set TRUE in addition to producing result text files, the results would also be returned as an object of class SummarizedExperiment.
method	A vector describing the summarization methods to use; i.e. whether count reads mapping to the introns (IntRet), reads mapping to the exons (ExEx), reads spanning the introns (IntSpan), or reads that skip the exons (ExSkip). In IntSpan mode the introns in the reference are taken into account only; whilst in IntRet the introns and their spanning exons, and in ExEx and ExSkip mode only the exons in the reference are taken into account.
strandSpecific	The description for strand specificity of the RNAseq data. The values are either "unstranded", "stranded", or "reverse". If the reads are not strand specific or directional use "unstranded". If the first read in paired-read sequencing or the reads single-read sequencing is in the same direction as the the transcript strand use "stranded". If the first read in paired-read sequencing or the reads in single-read sequencing is in the oposite direction to the transcript strand use "reverse".
appendLogFile	Whether log information should be appended to the logFile. It is FALSE by default.
sampleName	The name of the sample being analyzed. It will be included in the returned object if returnObj is TRUE.
scaleLength	A vector constructed of TRUE/FALSE values, same size as the method argument. It indicates whether the retention levels of the intron/exons should be scaled to their lengths.
scaleFragment	A vector constructed of TRUE/FALSE values, same size as the method argument. It indicates whether the retention levels of the intron/exons should be scaled to the sum of retention levels (i.e. mapped fragments) over the genes.
limitRanges	A GRanges object. If defined it only loads sequencing read if they fall in the defined coordinates. It is similar to which parameter in <a href="#">ScanBamParam</a> .
excludeFusionReads	Only valid if limitRanges is defined. It filters the defined by limitRanges. It also filters the read pairs if each read pair maps reads pairs where one of the reads either do not fall into one of the regions to a different region defined in limitRanges. It is useful to ignore analyzing the chimeric reads and fusion reads, i.e. reads that map to fusion genes. To filter properly, limitRanges must include coordinates of all genes.
loadLimitRangesReads	Boolean (TRUE/FALSE) variable. If set as TRUE only the reads in the limitRanges are loaded from bam file (and bamFileYieldSize parameter will be ignored).
...	Other parameter settings specific to <a href="#">BamFile-class</a> function in the Rsamtools package. Parameters qnamePrefixEnd and qnameSuffixStart are in particular useful to modify qnames in the BAM files.

## Value

If returnObj is set TRUE in addition to making result text files, dependant on whether a single or two method is defined, the results would be returned as a single object of class SummarizedExperiment or as a list of size 2 which includes 2 objects of class SummarizedExperiment one for IntRet and the other for ExEx.

**Author(s)**

Ali Oghabian

**See Also**[interest](#).**Examples**

```

# Creating temp directory to store the results
outDir<- file.path(tempdir(),"interestFolder")
dir.create(outDir)
outDir<- normalizePath(outDir)

# Loading suitable bam file
bamF <- system.file("extdata", "small_test_SRR1691637_ZRSR2Mut_RHBDD3.bam",
  package="IntEREst", mustWork=TRUE)

# Choosing reference for the gene RHBDD3
ref=u12[u12[, "gene_name"]=="RHBDD3",]

test= interest.sequential(
  bamFileYieldSize=10000,
  bamFile=bamF,
  isPaired=TRUE,
  isPairedDuplicate=FALSE,
  isSingleReadDuplicate=NA,
  reference=ref,
  referenceGeneNames=ref[, "ens_gene_id"],
  referenceIntronExon=ref[, "int_ex"],
  repeatsTableToFilter=c(),
  outFile=paste(outDir,
    "interestRes.tsv", sep="/"),
  logFile=paste(outDir,
    "log.txt", sep="/"),
  method=c("IntRet", "IntSpan"),
  strandSpecific="unstranded",
  returnObj=TRUE,
  scaleLength= c(TRUE,FALSE),
  scaleFragment= c(TRUE,TRUE)
)

test

```

InterestResult

*Building SummarizedExperiment object from results in IntEREst.***Description**

Calls the constructors and creates a SummarizedExperiment object. For more information on the resulted object and the class see [SummarizedExperiment-class](#).

**Usage**

```
InterestResult(resultFiles=c(), counts, scaledRetention,
               scaleLength, scaleFragment, sampleAnnotation, rowData)
```

**Arguments**

<code>resultFiles</code>	Vector of link to the result files of interest.
<code>counts</code>	Numeric Matrix that includes the read counts.
<code>scaledRetention</code>	Matrix that includes the scaled retention values.
<code>scaleLength</code>	Logical value, indicating whether the intron/exon retention levels are scaled to the length of the introns/exons.
<code>scaleFragment</code>	Logical value, indicating whether the intron/exon retention levels are scaled to the fragments mapped to the genes.
<code>sampleAnnotation</code>	Data frame with the row-size equal to the size of <code>resultFiles</code> and <code>sampleAnnotation</code> . Each column of the matrix represents annotations for the samples. Column name represents annotation name.
<code>rowData</code>	Data frame with Intron/Exon annotations and read count and scaled retention values for each sample.

**Value**

Returns an object of class `SummarizedExperiment`.

**Author(s)**

Ali Oghabian

**See Also**

[SummarizedExperiment-class attributes](#) [addAnnotation](#) [counts-method](#) [plot-method](#)

**Examples**

```
geneId<- paste("gene", c(rep(1,5), rep(2,5), rep(3,5), rep(4,5)),
               sep="_")
readCnt1<- sample(1:100, 20)
readCnt2<- sample(1:100, 20)
readCnt3<- sample(1:100, 20)
readCnt4<- sample(1:100, 20)
fpkm1<- readCnt1/(tapply(readCnt1, geneId, sum))[geneId]
fpkm2<- readCnt2/(tapply(readCnt2, geneId, sum))[geneId]
fpkm3<- readCnt3/(tapply(readCnt3, geneId, sum))[geneId]
fpkm4<- readCnt4/(tapply(readCnt4, geneId, sum))[geneId]

# Creating object using test data
interestDat<- data.frame(
  int_ex=rep(c(rep(c("exon", "intron"),2),"exon"),4),
  int_ex_num= rep(c(1,1,2,2,3),4),
  gene_id= geneId,
```

```

sam1_readCnt=readCnt1,
sam2_readCnt=readCnt2,
sam3_readCnt=readCnt3,
sam4_readCnt=readCnt4,
sam1_fpkm=fpkm1,
sam2_fpkm=fpkm2,
sam3_fpkm=fpkm3,
sam4_fpkm=fpkm4
)
readFreqColIndex<- grep("_readCnt$",colnames(interestDat))
scaledRetentionColIndex<- grep("_fpkm$",colnames(interestDat))

scalRetTmp<- as.matrix(interestDat[ ,scaledRetentionColIndex])
colnames(scalRetTmp)<-gsub("_fpkm$","", colnames(scalRetTmp))

frqTmp<- as.matrix(interestDat[ ,readFreqColIndex])
colnames(frqTmp)<-gsub("_readCnt$","", colnames(frqTmp))

InterestResultObj<- InterestResult(
  resultFiles=paste("file",1:4, sep="_"),
  rowData= interestDat[ , -c(readFreqColIndex,
  scaledRetentionColIndex)],
  counts= frqTmp,
  scaledRetention= scalRetTmp,
  scaleLength=TRUE,
  scaleFragment=FALSE,
  sampleAnnotation=data.frame(
    sampleName=paste("sam",1:4, sep=""),
    gender=c("M","M","F","F"), row.names=paste("sam", 1:4, sep="")
  )
)

# View object
InterestResultObj

```

---

interestResultIntEx     *Building results object that contains Intron-retention and exon-exon junction information*

---

## Description

Building [SummarizedExperiment-class](#) object from an intron retention and an exon-exon junction results in IntERESt. The average of the junction levels are added to the SummerizedExperiment object of the intron retentions.

## Usage

```

interestResultIntEx (intObj, exObj, intExCol=c(),
  mean.na.rm=TRUE, postExName="ex_junc" )

```

**Arguments**

intObj	A SummarizedExperiment including intron retention information.
exObj	A SummarizedExperiment including exon-exon junction information.
intExCol	Column name (or number) in the rowData of the intron object that represents whether each row of x assays is "intron" or "exon".
mean.na.rm	Whether exclude missing values when measuring the mean.
postExName	The postfix to use for the column names of the exons junction values in the

**Value**

Returns an object of class SummarizedExperiment.

**Author(s)**

Ali Oghabian

**See Also**

[SummarizedExperiment-class attributes addAnnotation counts-method plot-method](#)

**Examples**

```
testIntObj<- InterestResult(
  resultFiles= paste(paste("testFile",1:3, sep="_"),"bam", sep="."),
  counts= matrix(1:15, ncol=3, nrow=5, byrow=TRUE,
    dimnames= list(c(), paste("s", 1:3, sep="_"))),
  scaledRetention= matrix(1:15, ncol=3, nrow=5, byrow=TRUE,
    dimnames= list(c(), paste("s", 1:3, sep="_"))),
  scaleLength= FALSE,
  scaleFragment= FALSE,
  sampleAnnotation= data.frame(
    files=paste(paste("testFile",1:3, sep="_"),"bam", sep="."),
    names=paste("s", 1:3, sep="_"),
    row.names=paste("s", 1:3, sep="_")),
  rowData=data.frame(id= paste("i", 1:5, sep="_"),
    chr= rep("chr1", 5),
    begin=seq(100, by=100, length.out=5 ),
    end=seq(110, by=100, length.out=5 ),
    strand=rep("+",5))
)

testExObj<- InterestResult(
  resultFiles= paste(paste("testFile",1:3, sep="_"),"bam", sep="."),
  counts= matrix(1:30, ncol=3, nrow=10, byrow=TRUE,
    dimnames= list(c(), paste("s", 1:3, sep="_"))),
  scaledRetention= matrix(1:30, ncol=3, nrow=10, byrow=TRUE,
    dimnames= list(c(), paste("s", 1:3, sep="_"))),
  scaleLength= FALSE,
  scaleFragment= FALSE,
  sampleAnnotation= data.frame(
    files=paste(paste("testFile",1:3, sep="_"),"bam", sep="."),
    names=paste("s", 1:3, sep="_"),
    row.names=paste("s", 1:3, sep="_")),
  rowData=data.frame(id= paste("e", 1:10, sep="_"),
```

```

chr= rep("chr1", 10),
begin= c(seq(90, by=100, length.out=5),
  seq(111, by=100, length.out=5)),
end= c(seq(99, by=100, length.out=5),
  seq(120, by=100, length.out=5 )),
strand=rep("+",10))
)

(testIntExObj<- interestResultIntEx(intObj=testIntObj, exObj=testExObj,
  mean.na.rm=TRUE, postExName="ex_junc" ) )

```

---

intexIndex

*Extract index of intron or exon rows*


---

### Description

Extract row numbers where introns (or exons dependant on user's request) are located in an object of type SummarizedExperiment.

### Usage

```
intexIndex(x, intExCol="int_ex", what="intron")
```

### Arguments

x	Object of type SummarizedExperiment.
intExCol	Column name (or number) that represents whether each row is "intron" or "exon" in rowData of x.
what	A character string that defines whether the index for the introns or exons should be returned. Accepts either "exon" or "intron" (default) as values.

### Value

A numeric vector which includes the index of the introns/exons.

### Author(s)

Ali Oghabian

### See Also

[u12NbIndex](#)

### Examples

```
# Show the few first index of rows that represent the introns
head(intexIndex(mdsChr22Obj, what="intron"))
```

---

lfc *Log fold change*

---

### Description

Log fold change estimation and normalized log fold change using edgeR package.

### Usage

```
lfc(x, fcType="edgeR", sampleAnnoCol=c(), sampleAnnotation=c(),
    silent=TRUE, group=c(), rejection.region="doubletail",
    pseudoCnt=1, log2=TRUE, ...)
```

### Arguments

x	Object of type SummarizedExperiment.
fcType	Available as "scaledRetention" or "edgeR" (as default) corresponding to either log fold change of scaled retention values or edgeR normalized log fold change values.
sampleAnnoCol	Which column of colData of x to consider for the analysis.
sampleAnnotation	A vector of size 2 which contains values from colData of SummarizedExperiment object; e.g. if <code>getAnnotation(x)[,sampleAnnoCol]=c("test", "test", "ctrl", "ctrl", ...)</code> , and the goal is to compare "test" and "ctrl" samples, sampleAnnotation should either be <code>c("test", "ctrl")</code> or <code>c("ctrl", "test")</code> .
silent	Whether run <code>exactTestInterest</code> silently, without warnings.
group	Vector to manually define the sample groups (or annotations). It is ignored if sampleAnnoCol is defined.
rejection.region	The rejection.region parameter in <code>exactTest</code> , considered only if fcType is "edgeR".
pseudoCnt	Pseudo count for log transformation (default=1).
log2	Logical value either TRUE (default) or FALSE indicating whether the fold-changes should be log 2 transformed.
...	Other parameter settings from the <code>exactTestInterest</code> function.

### Value

Vector including fold change values.

### Author(s)

Ali Oghabian

### See Also

[exactTestInterest](#), [u12DensityPlotIntron](#)



**Examples**

```
lfcFpkm<- lfc(mdsChr22Obj, fcType="scaledRetention",
  sampleAnnoCol="test_ctrl",
  sampleAnnotation=c("ctrl", "test"),
  silent=TRUE, group=c(), pseudoFpkm=1, log2=TRUE)
```

```
lfcEdgeRFpkm<- lfc(mdsChr22Obj, fcType="edgeR",
  sampleAnnoCol="test_ctrl",
  sampleAnnotation=c("ctrl", "test"),
  silent=TRUE, group=c(), pseudoFpkm=1, log2=TRUE)
```

---

mdsChr22ExObj	<i>Object of SummarizedExperiment type for exon-exon junction of MDS data</i>
---------------	---

---

**Description**

The Results of `interest()` analysis in exon-exon junction mode, for the genes that feature U12-type introns and are located on Chr22 in MDS data.

**Usage**

```
data(mdsChr22ExObj)
```

**Format**

An Object of class `SummarizedExperiment` that contains intron retention results generated by `interest()` function on MDS data consisting of bone-marrow samples of 8 MDS patients with ZRSR2 mutations, 4 patients without the mutation and 4 healthy individuals.

`@colData` A "DataFrame" (from "S4Vectors" package) that its rownames can be set as the sample identification names and the other columns are various annotations for the samples. Its column names are characters that describe the annotations.

`@assays` List of size 2 that includes two numeric matrices: counts that includes raw read counts of the sequencing reads mapped to introns and exons, and (2) `scaledRetention`, i.e. the normalized read counts.

`@NAMES` A NULL value.

`@elementMetadata` A "DataFrame" (from "S4Vectors" package) that include intron and exon annotations.

`@metadata` A list of size 2 that includes parameter settings for the `interest()` and `interest.sequential()` runs.

**Value**

Object of class `SummarizedExperiment`.

**Source**

Madan, V., et al., Aberrant splicing of U12-type introns is the hallmark of ZRSR2 mutant myelodysplastic syndrome. *Nat Communication* 2015 Jan 14;6:6042. doi: 10.1038/ncomms7042.

---

mdsChr22IntSpObj	<i>Object of SummarizedExperiment type for intron spanning reads of MDS data</i>
------------------	--

---

### Description

The Results of `interest()` analysis in intron-spanning mode, for the genes that feature U12-type introns and are located on Chr22 in MDS data.

### Usage

```
data(mdsChr22ExObj)
```

### Format

An Object of class `SummarizedExperiment` that contains intron retention results generated by `interest()` function on MDS data consisting of bone-marrow samples of 8 MDS patients with ZRSR2 mutations, 4 patients without the mutation and 4 healthy individuals.

`@colData` A "DataFrame" (from "S4Vectors" package) that its rownames can be set as the sample identification names and the other columns are various annotations for the samples. Its column names are characters that describe the annotations.

`@assays` List of size 2 that includes two numeric matrices: `counts` that includes raw read counts of the sequencing reads mapped to introns and exons, and (2) `scaledRetention`, i.e. the normalized read counts.

`@NAMES` A NULL value.

`@elementMetadata` A "DataFrame" (from "S4Vectors" package) that include intron and exon annotations.

`@metadata` A list of size 2 that includes parameter settings for the `interest()` and `interest.sequential()` runs.

### Value

Object of class `SummarizedExperiment`.

### Source

Madan, V., et.al., Aberrant splicing of U12-type introns is the hallmark of ZRSR2 mutant myelodysplastic syndrome. *Nat Communication* 2015 Jan 14;6:6042. doi: 10.1038/ncomms7042.

---

mdsChr22Obj

*Object of SummarizedExperiment type for intron retention MDS data*


---

**Description**

The Results of `interest()` analysis in Intron-retention mode, for the genes that feature U12-type introns and are located on Chr22 in MDS data.

**Usage**

```
data(mdsChr22Obj)
```

**Format**

An Object of class `SummarizedExperiment` that contains intron retention results generated by `interest()` function on MDS data consisting of bone-marrow samples of 8 MDS patients with ZRSR2 mutations, 4 patients without the mutation and 4 healthy individuals.

`@colData` A "DataFrame" (from "S4Vectors" package) that its rownames can be set as the sample identification names and the other columns are various annotations for the samples. Its column names are characters that describe the annotations.

`@assays` List of size 2 that includes two numeric matrices: counts that includes raw read counts of the sequencing reads mapped to introns and exons, and (2) `scaledRetention`, i.e. the normalized read counts.

`@NAMES` A NULL value.

`@elementMetadata` A "DataFrame" (from "S4Vectors" package) that include intron and exon annotations.

`@metadata` A list of size 2 that includes parameter settings for the `interest()` and `interest.sequential()` runs.

**Value**

Object of class `SummarizedExperiment`.

**Source**

Madan, V., et.al., Aberrant splicing of U12-type introns is the hallmark of ZRSR2 mutant myelodysplastic syndrome. *Nat Communication* 2015 Jan 14;6:6042. doi: 10.1038/ncomms7042.

---

mergeInterestResult

*merge two SummarizedExperiment objects into one*


---

**Description**

Build a new object by merging data of two `SummarizedExperiment` objects.

**Usage**

```
mergeInterestResult(x, y)
```

**Arguments**

x                    Object of type SummarizedExperiment.  
 y                    Object of type SummarizedExperiment.

**Value**

An object of class SummarizedExperiment.

**Author(s)**

Ali Oghabian

**See Also**

[interest](#), [InterestResult](#).

**Examples**

```
geneId<- paste("gene", c(rep(1,5), rep(2,5), rep(3,5), rep(4,5)),
  sep="_")
readCnt1<- sample(1:100, 20)
readCnt2<- sample(1:100, 20)
readCnt3<- sample(1:100, 20)
readCnt4<- sample(1:100, 20)
fpkm1<- readCnt1/(tapply(readCnt1, geneId, sum))[geneId]
fpkm2<- readCnt2/(tapply(readCnt2, geneId, sum))[geneId]
fpkm3<- readCnt3/(tapply(readCnt3, geneId, sum))[geneId]
fpkm4<- readCnt4/(tapply(readCnt4, geneId, sum))[geneId]

# Creating object using test data
interestDat<- data.frame(
  int_ex=rep(c(rep(c("exon", "intron"),2),"exon"),4),
  int_ex_num= rep(c(1,1,2,2,3),4),
  gene_id= geneId,
  sam1_readCnt=readCnt1,
  sam2_readCnt=readCnt2,
  sam3_readCnt=readCnt3,
  sam4_readCnt=readCnt4,
  sam1_fpkm=fpkm1,
  sam2_fpkm=fpkm2,
  sam3_fpkm=fpkm3,
  sam4_fpkm=fpkm4
)
readFreqColIndex<- grep("_readCnt$", colnames(interestDat))
scaledRetentionColIndex<- grep("_fpkm$", colnames(interestDat))

scalRetTmp<- as.matrix(interestDat[, scaledRetentionColIndex])
colnames(scalRetTmp)<-gsub("_fpkm$", "", colnames(scalRetTmp))

frqTmp<- as.matrix(interestDat[, readFreqColIndex])
colnames(frqTmp)<-gsub("_readCnt$", "", colnames(frqTmp))

#Object including data for Males
interestResObjM<-InterestResult(
  resultFiles=paste("file",1:2, sep="_"),
```

```

rowData= interestDat[, -c(readFreqColIndex,
  scaledRetentionColIndex)],
counts= frqTmp[,1:2],
scaledRetention= scalRetTmp[,1:2],
scaleLength=TRUE,
scaleFragment=FALSE,
sampleAnnotation=data.frame(
  sampleName=paste("sam",1:2, sep=""),
  gender=c("M", "M"),
  health=c("healthy", "unhealthy"),
  row.names=paste("sam", 1:2, sep="")
)
)

#Object including data for Females
interestResObjF<-InterestResult(
  resultFiles=paste("file",3:4, sep="_"),
  rowData= interestDat[, -c(readFreqColIndex,
  scaledRetentionColIndex)],
  counts= frqTmp[,3:4],
  scaledRetention= scalRetTmp[,3:4],
  scaleLength=TRUE,
  scaleFragment=FALSE,
  sampleAnnotation=data.frame(
    sampleName=paste("sam",3:4, sep=""),
    gender=c("F", "F"),
    health=c("healthy", "unhealthy"),
    row.names=paste("sam", 3:4, sep="")
  )
)

#Build new object
newObj<- mergeInterestResult(interestResObjM, interestResObjF)

#View newObj
print(newObj)

```

---

plot-method

*plot - method*


---

## Description

plot method for SummarizedExperiment objects.

## Usage

```

## S4 method for signature 'SummarizedExperiment,ANY'
plot(x, summary="none",
  subsetRows=NULL, what="scaled", intronExon="intron",
  logScaleBase=NULL, logPseudoCnt=1, plotLoess=TRUE,
  loessCol="red", loessLwd=1, loessLty=1, cexText=1,
  marPlot=c(2,2,2,2), mgpPlot=c(1, 1, 0), cexAxis=1,
  writeCor=TRUE, corCex=1, corMethod="pearson", corCol="grey63",

```

```
upperCorXY=c("topleft", NULL), lowerCorXY=c("topleft", NULL),
na.rm=TRUE, cex=1, sampleAnnoCol=c(), lowerPlot=FALSE,
upperPlot=TRUE, ...)
```

### Arguments

x	Object of type SummarizedExperiment generated by either interest(), interest.sequential() or readInterestResults().
summary	Whether to plot the mean or median of the values over the sample with the same annotations, or plot the values for each individual sample separately. The available options are "mean", "median", or "none".
subsetRows	Vector either constructed of TRUE/FALSE values or constructed of numeric values that could be used to choose rows of x i.e. the SummarizedExperiment object.
what	Whether plot "scaled" (default) or read counts ("counts").
intronExon	Whether plot intron retention, i.e. "intron" (default) or exon-junction "exon".
logScaleBase	Base of the log transform of the values, if defined. By default the value is NULL meaning that the values would not be log transformed.
logPseudoCnt	Pseudocount for the log transformation (default=1).
plotLoess	Whether fit and plot LOESS curve line (default="red").
loessCol	loess line colour (default="red").
loessLwd	loess line width (default=1).
loessLty	loess line type (default=1).
cexText	Size of the text for sample names or annotations (default=1).
marPlot	Plot margins (default=c(2,2,2,2)). See ?par for more information.
mgpPlot	Plotting mgp parameter (default=c(1, 1, 0)). See ?par for more information.
cexAxis	Size of the text for the axis (default=1).
writeCor	Write correlation values (default=TRUE).
corCex	Text size of correlation values (default=1).
corMethod	Method used for correlation calculation. For more information see <a href="#">cor</a> from stats package of R.
corCol	Color of the text of correlation (default="grey").
upperCorXY	The coordinates of the correlation text in the upper panel plots ( default= c("topleft", NULL) ).
lowerCorXY	The coordinates of the correlation text in the lower panel plots ( default= c("topleft", NULL) ).
na.rm	whether remove the rows with missing values (default=TRUE).
cex	size of the plot text and symbols (default=1).
sampleAnnoCol	Which column of colData of object SummarizedExperiment to consider for plotting.
lowerPlot	Whether plot the lower panel (default=FALSE).
upperPlot	Whether plot the upper panel (default=TRUE).
...	Other arguments to pass to the plot() function.

**Value**

Returns NULL.

**Author(s)**

Ali Oghabian

**See Also**

Class: [SummarizedExperiment-class](#) Method: [counts-method](#) [boxplot-method](#)

**Examples**

```
geneId<- paste("gene", c(rep(1,5), rep(2,5), rep(3,5), rep(4,5)),
  sep="_")
readCnt1<- sample(1:100, 20)
readCnt2<- sample(1:100, 20)
readCnt3<- sample(1:100, 20)
readCnt4<- sample(1:100, 20)
fpkm1<- readCnt1/(tapply(readCnt1, geneId, sum))[geneId]
fpkm2<- readCnt2/(tapply(readCnt2, geneId, sum))[geneId]
fpkm3<- readCnt3/(tapply(readCnt3, geneId, sum))[geneId]
fpkm4<- readCnt4/(tapply(readCnt4, geneId, sum))[geneId]

# Creating object using test data
interestDat<- data.frame(
  int_ex=rep(c(rep(c("exon", "intron"),2), "exon"),4),
  int_ex_num= rep(c(1,1,2,2,3),4),
  gene_id= geneId,
  sam1_readCnt=readCnt1,
  sam2_readCnt=readCnt2,
  sam3_readCnt=readCnt3,
  sam4_readCnt=readCnt4,
  sam1_fpkm=fpkm1,
  sam2_fpkm=fpkm2,
  sam3_fpkm=fpkm3,
  sam4_fpkm=fpkm4
)
readFreqColIndex<- grep("_readCnt$", colnames(interestDat))
scaledRetentionColIndex<- grep("_fpkm$", colnames(interestDat))

scalRetTmp<- as.matrix(interestDat[, scaledRetentionColIndex])
colnames(scalRetTmp)<-gsub("_fpkm$", "", colnames(scalRetTmp))

frqTmp<- as.matrix(interestDat[, readFreqColIndex])
colnames(frqTmp)<-gsub("_readCnt$", "", colnames(frqTmp))

InterestResultObj<- InterestResult(
  resultFiles=paste("file", 1:4, sep="_"),
  rowData= interestDat[, -c(readFreqColIndex,
  scaledRetentionColIndex)],
  counts= frqTmp,
  scaledRetention= scalRetTmp,
  scaleLength=TRUE,
```

```

scaleFragment=FALSE,
sampleAnnotation=data.frame(
  sampleName=paste("sam",1:4, sep=""),
  gender=c("M","M","F","F"), row.names=paste("sam", 1:4, sep="")
)
)

InterestResultObj2<- addAnnotation(x=InterestResultObj,
  sampleAnnotationType="health",
  sampleAnnotation=c("healthy","unhealthy","healthy","unhealthy")
)

#Plotting
plot(InterestResultObj)
plot(InterestResultObj, sampleAnnoCol="gender", summary="mean")
plot(InterestResultObj2, sampleAnnoCol=3, summary="mean")
plot(InterestResultObj2, summary="none")

```

---

psi

*Psi values estimation*


---

### Description

Calculating the relative inclusion level of intron or Psi values base on two count matrices from a single or two separate objects. The values for each intron is in the range of [0,1], where 0 means complete splicing or no retention of the intron and 1 represnet complete 100

### Usage

```
psi (x, y, intCol, exCol, pseudoCnt=0)
```

### Arguments

x	Object of type SummarizedExperiment.
y	Optional; i.e. an object of type SummarizedExperiment.
intCol	Column numbers or column names in counts matrix of x which include the number of reads mapped to the introns.
exCol	Column numbers or column names in counts matrix of x (or if defined y) which include the number of reads spanning the introns (or mapping exons flanking the introns).
pseudoCnt	Pseudo counts to sum to the denominator of the devision to avoid devision to zero.

### Value

data.frame with column size equal to the size of intCol parameter, and row size equal to the number of rows in x. It contains the psi values (i.e.values between 0 and 1 showing the fraction of spliced in transcripts).

### Author(s)

Ali Oghabian



**See Also**[interestResultIntEx](#)**Examples**

```

mdsChr22IntObj<- mdsChr22Obj[which(rowData(mdsChr22Obj)$int_ex=="intron"), ]

#Build object including intron-retention and exon-junction results
mdsChr22RefIntExObj<- interestResultIntEx(intObj=mdsChr22Obj,
  exObj=mdsChr22ExObj, mean.na.rm=TRUE, postExName="ex_junc",
  intExCol="int_ex" )
# Calculate Psi
psiRes<- psi(mdsChr22RefIntExObj,
  intCol=which(colData(mdsChr22RefIntExObj)$intronExon=="intron"),
  exCol=which(colData(mdsChr22RefIntExObj)$intronExon=="exon"))
# show Psi results
head(psiRes)

```

pwmU12db

*PWM of U12 and U2-type introns splice sites***Description**

PWM of U12 and U2-type introns splice sites and it is based on the U12DB database.

**Usage**

```
data("pwmU12db")
```

**Format**

A list that contains Position Weight Matrices (PWM) of donor site, branch point and acceptor site of U12-type introns and the PWMs of donor site and acceptor site of U2-type introns. It is based on the U12DB database.

**pwmDonU12** A position weigh matrix for the donor site of the U12-type introns, with 4 rows and 46 columns. The rows of the matrix represent "A", "C", "G", and "T" nucleotides and the columns represent the postions in the genome. Each position in the matrix include a weight (i.e. number between 0 and 1) which indicates how common the corresponding base (represented by the row of the matrix) is observed in the coresponding position (represented by the colum of the matrix).

**pwmBpU12** A position weigh matrix for the branch point of the U12-type introns, with 4 rows and 9 columns.

**pwmAccU12** A position weigh matrix for the acceptor site of the U12-type introns, with 4 rows and 46 columns.

**pwmDonU2** A position weigh matrix for the donor site of the U2-type introns, with 4 rows and 25 columns.

**pwmAccU2** A position weigh matrix for the acceptor site of the U12-type introns, with 4 rows and 46 columns.

**Value**

List of 5 numeric matrices representing the PWMs of donor site of U12-type introns, branch point site of U12-type introns, acceptor site of U12-type introns, donor site of U2-type introns, and acceptor site of U2-type introns.

**Source**

Alioto, T.S. U12DB: a database of orthologous U12-type spliceosomal introns. *Nucleic Acids Research* 2006, doi: 10.1093/nar/gkl796

---

qlfInterest	<i>quasi-likelihood F-test</i>
-------------	--------------------------------

---

**Description**

Compute quasi-likelihood F-test using edgeR package. For more information see [glmQLFit](#) and [glmQLFTest](#) functions in edgeR package.

**Usage**

```
qlfInterest(x, design=c(), silent=TRUE, disp="common",
            coef=c(), contrast=NULL,
            poisson.bound=TRUE, ...)
```

**Arguments**

x	Object of type SummarizedExperiment.
design	Design matrix.
silent	Whether run silently, i.e. without printing the top differential expression tags. The default is TRUE.
disp	The method of estimating the dispersion in the data. Available options are: "common", "trended", "tagwiseInitCommon" and "tagwiseInitTrended". It is also possible to assign a number.
coef	Integer or character vector indicating which coefficients of the linear model are to be tested equal to zero. See <a href="#">glmQLFTest</a> for more information.
contrast	Numeric vector or matrix specifying contrasts of the linear model coefficients to be tested equal to zero. See <a href="#">glmQLFTest</a> for more information.
poisson.bound	Logical value, if TRUE (i.e. default) the pvalue would be higher than when obtained from likelihood ratio test while Negative Binomial dispersion is zero.
...	Other parameter settings for the <a href="#">glmQLFTest</a> function in the edgeR package.

**Value**

All values produced by [glmQLFTest](#) plus the following :

dispersionType	The name of the type of dispersion used.
dispersion	The estimated dispersion values.

**Author(s)**

Ali Oghabian

**See Also**[exactTestInterest](#), [glmInterest](#), [treatInterest](#)**Examples**

```
#Test retention differentiation across the 3 types of sampels
group <- getAnnotation(mdsChr220bj)[,"type"]
qlfRes<- qlfInterest(x=mdsChr220bj,
  design=model.matrix(~group), silent=TRUE,
  disp="tagwiseInitTrended", coef=2:3, contrast=NULL)

qlfRes
```

---

readInterestResults     *Read interest/interest.sequential results text files*

---

**Description**

Reads one or multiple text file results generated by the [interest](#) or [interest.sequential](#) functions and builds an object of [SummarizedExperiment-class](#) class.

**Usage**

```
readInterestResults(resultFiles, sampleNames,
  sampleAnnotation, commonColumns, freqCol, scaledRetentionCol,
  scaleLength, scaleFragment, reScale=FALSE, geneIdCol,
  repeatsTableToFilter=c())
```

**Arguments**

<code>resultFiles</code>	Vector of character strings which includes the path to the tab-separated files resulted by the <a href="#">interest</a> function.
<code>sampleNames</code>	Vector of character strings which includes the name of the samples. It should be the same size as the <code>resultFiles</code> parameter.
<code>sampleAnnotation</code>	Data frame with the same row number as the size of <code>resultFiles</code> and <code>sampleNames</code> parameter. The column names represent the annotation names and values in each column represent the annotations of the samples.
<code>commonColumns</code>	Columns in the result file which include intron/exon annotations and are common across all files defined in <code>resultFiles</code> .
<code>freqCol</code>	Column in the result file which include the read counts for introns/exons.
<code>scaledRetentionCol</code>	Column in the result file which include the scaled retention values for introns/exons.
<code>scaleLength</code>	Logical value, indicating whether the intron/exon retention levels are scaled to the length of the introns/exons. If <code>reScale</code> is TRUE the scaled retention levels would be recalculated when reading the data.

scaleFragment	Logical value, indicating whether the intron/exon retention levels are scaled to the fragments mapped to the genes. If reScale is TRUE the scaled retention levels would be recalculated when reading the data.
reScale	Logical value, indicating whether the scaled retention levels would be recalculated when reading the data. By default it does not calculate and trusts the user to set the scaleLength and scaleFragment parameters correctly, i.e. as it was set in the interest() or interest.sequential() analysis.
geneIdCol	The number or name of the column in resultFiles which represents the gene/transcript names. It would be used for summing up the number of mapped fragments to the genes when scaling the retention levels. It is only used if reScale and scaleFragment arguments are set TRUE.
repeatsTableToFilter	A data.frame table with similar structure to the reference. It includes chr, begin, and end columns. If defined, all reads mapped to the described regions would be ignored and the Intron/exon lengths would be corrected to exclude the to exclude the regions with repetitive DNA sequences. See <a href="#">getRepeatTable</a> . It is only used if reScale and scaleLength arguments are set TRUE.

**Value**

An object of class [SummarizedExperiment-class](#).

**Author(s)**

Ali Oghabian

**See Also**

[interest](#), [InterestResult](#).

**Examples**

```
geneId<- paste("gene", c(rep(1,7), rep(2,7), rep(3,7), rep(4,7)),
  sep="_")
readCnt1<- sample(1:100, 28)
readCnt2<- sample(1:100, 28)
readCnt3<- sample(1:100, 28)
readCnt4<- sample(1:100, 28)
fpkm1<- readCnt1/(tapply(readCnt1, geneId, sum))[geneId]
fpkm2<- readCnt2/(tapply(readCnt2, geneId, sum))[geneId]
fpkm3<- readCnt3/(tapply(readCnt3, geneId, sum))[geneId]
fpkm4<- readCnt4/(tapply(readCnt4, geneId, sum))[geneId]

#Create tmp director
tmpDir=file.path(tempdir(),"InterestResult")
dir.create(tmpDir)

# Build text files similar to files resulted by interest
dfTmp=data.frame(
  int_ex=rep(c(rep(c("exon","intron"),3),"exon"),4),
  int_ex_num= rep(c(1,1,2,2,3,3,4),4),
  int_type=rep(c(NA,"U2",NA,"U12",NA,"U2",NA),4),
  strand=rep("*",28),
```

```

    gene_id= geneId,
    sam1_readCnt=readCnt1,
    sam2_readCnt=readCnt2,
    sam3_readCnt=readCnt3,
    sam4_readCnt=readCnt4,
    sam1_fpkm=fpkm1,
    sam2_fpkm=fpkm2,
    sam3_fpkm=fpkm3,
    sam4_fpkm=fpkm4
  )

writeDf<-function(df, file){
  write.table(df, file, col.names=TRUE,
    row.names=FALSE, quote=FALSE, sep='\t')
}

writeDf(dfTmp[, c(1:5,6,10)], paste(tmpDir, "df1.tsv", sep="/"))
writeDf(dfTmp[, c(1:5,7,11)], paste(tmpDir, "df2.tsv", sep="/"))
writeDf(dfTmp[, c(1:5,8,12)], paste(tmpDir, "df3.tsv", sep="/"))
writeDf(dfTmp[, c(1:5,9,13)], paste(tmpDir, "df4.tsv", sep="/"))

# Build object from generated text file results
testObj<-readInterestResults(
  resultFiles=paste(tmpDir,
    c("df1.tsv", "df2.tsv", "df3.tsv", "df4.tsv"), sep="/"),
  sampleNames=c("sam1", "sam2", "sam3", "sam4"),
  sampleAnnotation= data.frame( gender=c("M", "M", "F", "F"),
    health=c("healthy", "unhealthy", "healthy", "unhealthy")),
  commonColumns=1:5, freqCol=6, scaledRetentionCol=7,
  scaleLength=FALSE, scaleFragment=TRUE, reScale=FALSE)

#View object
testObj

```

---

referencePrepare	<i>Creates reference file</i>
------------------	-------------------------------

---

## Description

Creates reference file for IntEREst functions, e.g. `interest()`. The function uses functions of `biomaRt` library.

## Usage

```

referencePrepare( outFileTranscriptsAnnotation="",
  annotateGeneIds=TRUE,
  u12IntronsChr=c(), u12IntronsBeg=c(), u12IntronsEnd=c(),
  u12IntronsRef, collapseExons=TRUE, sourceBuild="UCSC",
  ucscGenome="hg19", ucscTableName="knownGene",
  ucscUrl="http://genome-euro.ucsc.edu/cgi-bin/",
  biomart="ENSEMBL_MART_ENSEMBL",
  biomartDataset="hsapiens_gene_ensembl",

```

```

biomartTranscriptIds=NULL, biomartExtraFilters=NULL,
biomartIdPrefix="ensembl_", biomartHost="www.ensembl.org",
biomartPort=80, circSeqs="", miRBaseBuild=NA, taxonomyId=NA,
filePath="", fileFormat=c("auto", "gff3", "gtf"), fileDatSrc=NA,
fileOrganism=NA, fileChrInf=NULL,
fileDbXrefTag=c(), addCollapsedTranscripts=TRUE,
ignore.strand=FALSE )

```

## Arguments

- outFileTranscriptsAnnotation** If defined outputs transcripts annotations.
- annotateGeneIds** Whether annotate and add the gene ids information.
- collapseExons** Whether collapse (i.e. reduce) the exonic regions. TRUE by default.
- sourceBuild** The source to use to build the reference data, "UCSC", "biomaRt", and "file" (for GFF3 or GTF files) are supported.
- ucscGenome** The genome to use. "hg19" is the default. See genome parameter of [makeTxDbFromUCSC](#) function of txdbmaker library for more information.
- ucscTableName** The UCSC table name to use. See tablename parameter of [makeTxDbFromUCSC](#) function of txdbmaker library for more information.
- ucscUrl** The UCSC URL address. See url parameter of [makeTxDbFromUCSC](#) function of txdbmaker library for more information.
- u12IntronsChr** A vector of character strings that includes chromosomal locations of the U12 type introns. If defined together with u12IntronsBeg and u12IntronsEnd, they would be used to annotate the U12-type introns.
- u12IntronsBeg** A vector of numbers that defines the begin (or start) coordinates of the u12-type introns.
- u12IntronsEnd** A vector of numbers that defines the end coordinates of the u12-type introns.
- u12IntronsRef** A GRanges object that includes the coordinates of the U12 type introns. If defined, it would be used to annotate the U12-type introns.
- biomart** BioMart database name. See biomart parameter of [makeTxDbFromBiomart](#) function of txdbmaker library for more information.
- biomartDataset** BioMart dataset name; default is "hsapiens\_gene\_ensembl". See dataset parameter of [makeTxDbFromBiomart](#) function of txdbmaker library for more information.
- biomartTranscriptIds** optional parameter to only retrieve transcript annotation results for a defined set of transcript ids. See transcript\_ids parameter of [makeTxDbFromBiomart](#) function of txdbmaker library for more information.
- biomartExtraFilters** A list of names; i.e. additional filters to use in the BioMart query. See filters parameter of [makeTxDbFromBiomart](#) function of txdbmaker library for more information.
- biomartIdPrefix** A list of names; i.e. additional filters to use in the BioMart query. See id\_prefix parameter of [makeTxDbFromBiomart](#) function of txdbmaker library for more information.

biomartHost	Host to connect to; the default is "www.ensembl.org". For older versions of the GRCH you can provide the archive websites, e.g. for GRCH37 you can use "grch37.ensembl.org".
biomartPort	The port to use in the HTTP communication with the host. Default is 80.
circSeqs	A character vector that includes chromosomes that should be marked as circular. See circ_seqs parameter of <a href="#">makeTxDbFromBiomart</a> and <a href="#">makeTxDbFromUCSC</a> functions of txdbmaker library for more information.
miRBaseBuild	Set appropriate build Information from mirbase.db to use for microRNAs (default=NA). See miRBaseBuild parameter of <a href="#">makeTxDbFromBiomart</a> and <a href="#">makeTxDbFromUCSC</a> functions of txdbmaker library for more information.
taxonomyId	This parameter can be used to provide taxonomy Ids. It is set to NA by default. You can check the taxonomy Ids with the available.species() function in GenomeInfoDb package. For more information see taxonomyId parameter of <a href="#">makeTxDbFromBiomart</a> and <a href="#">makeTxDbFromUCSC</a> functions of txdbmaker library.
filePath	Character string i.e. the path to file. Used if sourceBuild is "file".
fileFormat	The format of the input file. "auto", "gff3" and "gtf" is supported.
fileDatSrc	Character string describing the source of the data file. Used if sourceBuild is "file".
fileOrganism	The genus and species name of the organism. Used if sourceBuild is "file".
fileChrInf	Dataframe that includes information about the chromosome. The first column represents the chromosome name and the second column is the length of the chromosome. Used if sourceBuild is "file".
fileDbXrefTag	A vector of character strings which if defined it would be used as feature names. Used if sourceBuild is "file".
addCollapsedTranscripts	Whether add a column that includes the collapsed transcripts information. Used if collapseExons is TRUE.
ignore.strand	Whether consider the strands in the reference. If set TRUE the strands would be ignored.

**Value**

Data frame that includes the coordinates and annotations of the introns and exons of the transcripts, i.e. the reference.

**Author(s)**

Ali Oghabian

**Examples**

```
# Build test gff3 data
tmpGen<- u12[u12["ens_trans_id"]=="ENST00000413811",]
tmpEx<-tmpGen[tmpGen["int_ex"]=="exon",]
exonDat<- cbind(tmpEx[,3], ".",
  tmpEx[,c(7,4,5)], ".", tmpEx[,6], ".", paste("ID=exon",
  tmpEx[,11], "; Parent=ENST00000413811", sep="")) )
trDat<- c(tmpEx[1,3], ".", "mRNA", as.numeric(min(tmpEx[,4])),
  as.numeric(max(tmpEx[,5])), ".", tmpEx[1,6], ".",
```

```

"ID=ENST00000413811")

outDir<- file.path(tempdir(),"tmpFolder")
dir.create(outDir)
outDir<- normalizePath(outDir)

gff3File=paste(outDir, "gffFile.gff", sep="/")

cat("##gff-version 3\n",file=gff3File, append=FALSE)
cat(paste(paste(trDat, collapse="\t"),"\n", sep=""),
    file=gff3File, append=TRUE)

write.table(exonDat, gff3File,
            row.names=FALSE, col.names=FALSE,
            sep='\t', quote=FALSE, append=TRUE)

# Selecting U12 introns info from 'u12' data
u12Int<-u12[u12$int_ex=="intron"&u12$int_type=="U12",]

# Test the function
refseqRef<- referencePrepare (sourceBuild="file",
                             filePath=gff3File, u12IntronsChr=u12Int[, "chr"],
                             u12IntronsBeg=u12Int[, "begin"],
                             u12IntronsEnd=u12Int[, "end"], collapseExons=TRUE,
                             fileFormat="gff3", annotateGeneIds=FALSE)

```

---

subInterestResult      *Extract subset of object*

---

## Description

Build a new object using subset of data in an SummarizedExperiment object.

## Usage

```
subInterestResult(x, selectRow, selectCol,
                 sampleAnnoCol, sampleAnnotation=c())
```

## Arguments

x	Object of type SummarizedExperiment.
selectRow	Numeric or TRUE/FALSE Vector indicating what rows to extract.
selectCol	A vector with Numeric values, character strings (sample names) or TRUE/FALSE Vector indicating what columns to extract.
sampleAnnoCol	Which column of colData of object x to consider for subset data extraction.
sampleAnnotation	Vector including the annotations to consider for subset data extraction. They should be present in the sampleAnnoCol column of the colData of x.

## Value

An object of class SummarizedExperiment.



**Author(s)**

Ali Oghabian

**See Also**[interest](#), [InterestResult](#).**Examples**

```

geneId<- paste("gene", c(rep(1,7), rep(2,7), rep(3,7), rep(4,7)),
  sep="_")
readCnt1<- sample(1:100, 28)
readCnt2<- sample(1:100, 28)
readCnt3<- sample(1:100, 28)
readCnt4<- sample(1:100, 28)
fpkm1<- readCnt1/(tapply(readCnt1, geneId, sum))[geneId]
fpkm2<- readCnt2/(tapply(readCnt2, geneId, sum))[geneId]
fpkm3<- readCnt3/(tapply(readCnt3, geneId, sum))[geneId]
fpkm4<- readCnt4/(tapply(readCnt4, geneId, sum))[geneId]

# Creating object using test data
interestDat<-data.frame(
  int_ex=rep(c(rep(c("exon","intron"),3),"exon"),4),
  int_ex_num= rep(c(1,1,2,2,3,3,4),4),
  int_type=rep(c(NA,"U2",NA,"U12",NA,"U2",NA),4),
  strand=rep("*",28),
  gene_id= geneId,
  sam1_readCnt=readCnt1,
  sam2_readCnt=readCnt2,
  sam3_readCnt=readCnt3,
  sam4_readCnt=readCnt4,
  sam1_fpkm=fpkm1,
  sam2_fpkm=fpkm2,
  sam3_fpkm=fpkm3,
  sam4_fpkm=fpkm4
)
readFreqColIndex<- grep("_readCnt$",colnames(interestDat))
scaledRetentionColIndex<- grep("_fpkm$",colnames(interestDat))
samNames<-paste("sam", 1:4, sep="")
frqTmp<-as.matrix(interestDat[, readFreqColIndex])
sclTmp<-as.matrix(interestDat[, scaledRetentionColIndex])
colnames(frqTmp)<- samNames
colnames(sclTmp)<- samNames
interestResObj<- InterestResult(
  resultFiles=paste("file",1:4, sep="_"),
  rowData= interestDat[, -c(readFreqColIndex,
  scaledRetentionColIndex)],
  counts= frqTmp,
  scaledRetention= sclTmp ,
  scaleLength=TRUE,
  scaleFragment=FALSE,
  sampleAnnotation=data.frame(
    sampleName=paste("sam",1:4, sep=""),
    gender=c("M","M","F","F"),
    health=c("healthy","unhealthy","healthy","unhealthy"),
    row.names=samNames
  )
)

```

```

)
)

#Build new object
newObj<- subInterestResult(interestResObj, selectRow=1:20)

#View newObj
print(newObj)

```

---

treatInterest	<i>Differential retention test relative to a threshold</i>
---------------	--

---

### Description

Compute a genewise statistical test relative to a fold-change threshold using edgeR package. For more information see [glmTreat](#) function in edgeR package.

### Usage

```
treatInterest(x, design=c(), silent=TRUE, disp="common",
             coef=c(), contrast=NULL, lfc=0, ...)
```

### Arguments

x	Object of class SummarizedExperiment.
design	Design matrix.
silent	Whether run silently, i.e. without printing the top differential expression tags. Default is TRUE.
disp	The method of estimating the dispersion in the data. Available options are: "common", "trended", "tagwiseInitCommon" and "tagwiseInitTrended". It is also possible to assign a number.
coef	Integer or character vector indicating which coefficients of the linear model are to be tested equal to zero. See <a href="#">glmTreat</a> for more information.
contrast	Numeric vector or matrix specifying contrasts of the linear model coefficients to be tested equal to zero. See <a href="#">glmTreat</a> for more information.
lfc	Numeric scalar i.e. the log fold change threshold.
...	Other parameter settings for the <code>glmFit</code> function in the edgeR package.

### Value

All values produced by [glmTreat](#) plus the following :

dispersionType	The name of the type of dispersion used.
dispersion	The estimated dispersion values.

### Author(s)

Ali Oghabian

**See Also**

[exactTestInterest](#), [qlfInterest](#), [glmInterest](#)

**Examples**

```
group <- getAnnotation(mdsChr220bj)[,"type"]

#Test retention differentiation across the 3 types of sampels
# The log fold change threshold is 0
treatRes<- treatInterest(x=mdsChr220bj,
  design=model.matrix(~group), silent=TRUE,
  disp="tagwiseInitTrended", coef=2:3, contrast=NULL, lfc=0)
treatRes
```

u12

*U12 data***Description**

Intron/exon annotations of genes featuring U12 introns. It is based on HG19/GRCh37 (converted from hg17/NCBI35). Moreover the u12 genes are based on the U12DB database.

**Usage**

```
data("u12")
```

**Format**

A data frame with 22713 observations on the following 17 variables.

id a numeric vector  
 int\_ex\_id a character vector  
 chr a character vector  
 begin a numeric vector  
 end a numeric vector  
 strand a numeric vector  
 int\_ex a character vector  
 trans\_type a character vector  
 ens\_gene\_id a character vector  
 ens\_trans\_id a character vector  
 int\_ex\_num a numeric vector  
 gene\_name a character vector  
 trans\_name a character vector  
 overlap\_no a numeric vector  
 int\_type a character vector  
 int\_subtype a character vector

**Value**

Data frame that includes the coordinates and annotations of the introns and exons of the transcripts, i.e. the reference.

**Source**

Alioto, T.S. U12DB: a database of orthologous U12-type spliceosomal introns. *Nucleic Acids Research* 2006, doi: 10.1093/nar/gkl796

---

u12Boxplot	<i>U12 boxplot</i>
------------	--------------------

---

**Description**

A boxplot method for U12 and U2-type introns of SummarizedExperiment objects.

**Usage**

```
u12Boxplot(x, sampleAnnoCol=NA, intExCol="int_ex",
  intTypeCol="int_type", intronExon, col="white",
  boxplotNames=c(), lasNames=3, outline=FALSE, addGrid=FALSE, ...)
```

**Arguments**

x	Object of type SummarizedExperiment.
sampleAnnoCol	Which column of colData in x to consider for plotting.
intExCol	Column name (or number) that represents whether each row of x assays is "intron" or "exon".
intTypeCol	Column name (or number) that represents what type of intron each row of x assays represents.
intronExon	Whether plot intron retention (set intronExon="intron") or exon-exon junction (set intronExon="exon") levels.
col	Vector showing box colours. It is either of size 1 or the same size as the number of groups to be plotted.
boxplotNames	Names to write under boxes. If not defined, as names, it pastes U12/U2 (intron annotation) to the sample group annotations separated by a space " ".
lasNames	Orientation of the box names.
outline	If outline is TRUE the outlier points are drawn otherwise if FALSE (default) they are not.
addGrid	Whether add a grid under the boxplots (FALSE by default).
...	Other arguments to pass to the boxplot() function.

**Value**

A SummarizedExperiment object.

**Author(s)**

Ali Oghabian

**See Also**[u12BoxplotNb](#)**Examples**

```
u12Boxplot(mdsChr220bj, sampleAnnoCol="type",
  intExCol="int_ex", intTypeCol="intron_type", intronExon="intron",
  col=rep(c("orange", "yellow"),3) , lasNames=3,
  outline=FALSE, ylab="FPKM", cex.axis=0.8)
```

---

u12BoxplotNb	<i>boxplot U12 introns retention levels (or flanking exons junction levels) and (up/down)stream U2 introns (or exons junction levels)</i>
--------------	---

---

**Description**

boxplot U12 introns and (Up/Down)stream U2 introns in SummarizedExperiment objects.

**Usage**

```
u12BoxplotNb(x, sampleAnnoCol=2, intExCol="int_ex",
  intTypeCol="int_type", intronExon, strandCol="strand", geneIdCol,
  col=c(), names=c(), lasNames=1, outline=FALSE, plotLegend=TRUE,
  cexLegend=1, xLegend="topright", yLegend=NULL, bgLegend="transparent",
  legend=c(), addGrid=FALSE, ...)
```

**Arguments**

x	Object of type SummarizedExperiment.
sampleAnnoCol	Which column of colData of x to consider for plotting.
intExCol	Column name (or number) that represents whether each row of x assays is "intron" or "exon".
intTypeCol	Column name (or number) that represents what type of intron each row of x assays represents.
intronExon	Whether plot intron retention (set intronExon="intron") or exon-exon junction (set intronExon="exon") levels.
strandCol	Column name (or number) that represents the strand of each row of assays in x. The values in the column are either "+", "-" or "*".
geneIdCol	Column name (or number) that represents the gene ID of each row of assays in x.
col	Vector containing box colours. It is either of size 1 or the same size as the number of boxes resulted based on the grouping of the samples defined by sampleAnnoCol.
names	Names to write under group of boxes.
lasNames	Orientation of the box names.
outline	If outline is TRUE the outlier points are drawn otherwise if FALSE (default) they are not.

plotLegend	Whether show legend (TRUE by default).
cexLegend	Size of the text in legend .
xLegend, yLegend	Position of legend in the plot. For more info see x and y parameters in <a href="#">legend</a> .
bgLegend	Bakcground colour of the legend box. It is "transparent" by default.
legend	The replacement texts to be used in legend.
addGrid	Whether add a grid under the boxplots (FALSE by default).
...	Other arguments to pass to the boxplot() function.

**Value**

Returns NULL

**Author(s)**

Ali Oghabian

**See Also**

[u12Boxplot](#)

**Examples**

```
u12BoxplotNb(mdsChr220bj, sampleAnnoCol="type", lasNames=1,
  intExCol="int_ex", intTypeCol="intron_type", intronExon="intron",
  boxplotNames=c(), outline=FALSE, plotLegend=TRUE,
  geneIdCol="collapsed_transcripts_id", xLegend="topleft",
  col=c("pink", "lightblue", "lightyellow"), ylim=c(0,600000),
  ylab="FPKM", cex.axis=0.8)
```

---

u12DensityPlot	<i>Density plot of fld changes of intron retention and exon-exon junction levels</i>
----------------	--

---

**Description**

Density plot of fold change of the retention levels of U12- vs U2- type intron, or exon-exon junction levels of the flanking exons. For the density plot of the foldchange of intron retention levels the `u12DensityPlotIntron()` function or `u12DensityPlot()` function with `intronExon= "intron"` can be used. For density plot of the foldchange of exon-exon junction levels use `u12DensityPlot()` function with `intronExon= "exon"`.

**Usage**

```
u12DensityPlot(x,
  type=c("U12", "U2Up", "U2Dn", "U2UpDn", "U2Rand"),
  fcType="edgeR", sampleAnnotation=c(), sampleAnnoCol=c(),
  group=c(), intExCol="int_ex", intTypeCol="int_type", intronExon,
  strandCol="strand", geneIdCol="collapsed_transcripts",
  naUnstrand=FALSE, col=1, lty=1, lwd=1, plotLegend=TRUE,
```

```
cexLegend=1, xLegend="topright", yLegend=NULL, legend=c(),
randomSeed=NULL, xlab="", ...)
```

```
u12DensityPlotIntron(x,
  type= c("U12", "U2Up", "U2Dn", "U2UpDn", "U2Rand"),
  fcType= "edgeR", sampleAnnotation=c(), sampleAnnoCol=c(),
  group=c(), intExCol="int_ex", intTypeCol="int_type",
  strandCol= "strand", geneIdCol= "collapsed_transcripts",
  naUnstrand=FALSE, col=1, lty=1, lwd=1, plotLegend=TRUE,
  cexLegend=1, xLegend="topright", yLegend=NULL, legend=c(),
  randomSeed=NULL, xlab="", ...)
```

## Arguments

x	Object of type SummarizedExperiment.
type	A vector that includes the type of introns to plot. Available options are U12 introns "U12", U2 introns at downstream of U12 introns "U2Dn", U2 introns at upstream of U12 introns "U2Up", U2 introns at upstream or downstream of U12 introns suitable for when the coordinates in object x are unstranded (their strand is "*") "U2UpDn", random U2 introns from object x "U2Rand". Settings "U2Up", "U2Dn" and "U2UpDn" are useful only if the reference is linearly ordered. References with exons only resulted by referencePrepare and unionRefTr are NOT necessarily linearly ordered.
fcType	Available as "fpkm" or "edgeR" (as default) corresponding to either log fold change of fpkm values or edgeR normalized log fold change values.
sampleAnnoCol	Which column of colData of x to consider for plotting.
sampleAnnotation	A vector of size 2 which contains values from colData of SummarizedExperiment object; e.g. if getAnnotation(x)[, sampleAnnoCol]= c("test", "test", "ctrl", "ctrl", ...), and the goal is to compare "test" and "ctrl" samples, sampleAnnotation should either be c("test", "ctrl") or c("ctrl", "test").
group	Vector to manually define the sample groups (or annotations). It is ignored if sampleAnnoCol is defined.
intExCol	Column name (or number) that represents whether each row of x assays is "intron" or "exon".
intTypeCol	Column name (or number) that represents what type of intron each row of x assays represents.
intronExon	Whether plot intron retention (set intronExon="intron") or exon-exon junction (set intronExon="exon") levels.
strandCol	Column name (or number) that represents the strand of each row of assays in x. The values in the column are either "+", "-" or "*".
geneIdCol	Column name (or number) that represents the gene ID of each row of assays in x.
naUnstrand	Replace unstranded results, i.e. introns or exon with "*" strand, with NA (to be excluded).
col	A vector with the size of 1 or the same size as the type parameter which includes the colour/colours of the plotted density lines (default=1).
lty	A vector with the size of 1 or the same size as the type parameter which includes the type of the plotted density lines (default=1).

lwd	A vector with the size of 1 or the same size as the type parameter which includes the width of the plotted density lines (default=1).
plotLegend	Whether show legend (TRUE by default).
cexLegend	Size of the text in legend .
xLegend, yLegend	Position of legend in the plot. For more info see x and y parameters in <a href="#">legend</a> .
legend	The replacement texts to be used in legend.
randomSeed	Seed value for random number generator.
xlab	The lable of the X axis of the plot; by default it is "".
...	Other parameter settings from the <a href="#">plot</a> function.

**Value**

Returns NULL.

**Author(s)**

Ali Oghabian

**See Also**

[exactTestInterest](#), [lfc](#)

**Examples**

```
u12DensityPlotIntron(mdsChr22Obj,
  type= c("U12", "U2Up", "U2Dn", "U2UpDn", "U2Rand"),
  fcType= "edgeR", sampleAnnoCol="test_ctrl",
  sampleAnnotation=c("ctrl","test"), intExCol="int_ex",
  intTypeCol="intron_type", strandCol= "strand",
  geneIdCol= "collapsed_transcripts_id", naUnstrand=FALSE, col=c(2,3,4,5,6),
  lty=c(1,2,3,4,5), lwd=1, plotLegend=TRUE, cexLegend=0.7,
  xLegend="topright", yLegend=NULL, legend=c(), randomSeed=10,
  ylim=c(0,0.6), xlab=expression("log"[2]*" fold change FPKM"))
```

---

u12Index

*Extract index of U12 introns rows*

---

**Description**

Extract row numbers of U12 introns in an object of class SummarizedExperiment.

**Usage**

```
u12Index(x, intExCol="int_ex", intTypeCol="int_type", intronExon="intron")
```



**Arguments**

x	Object of type SummarizedExperiment.
intExCol	Column name (or number) that represents whether each row of x assays is "intron" or "exon".
intTypeCol	Column name (or number) that represents what type of intron each row of x assays represents.
intronExon	Whether extract U12 type introns (set intronExon="intron") or exon-exon junction (set intronExon="exon") flanking U12 introns.

**Value**

A numeric vector which includes the index of U12 introns.

**Author(s)**

Ali Oghabian

**See Also**

[u12NbIndex](#)

**Examples**

```
head(u12Index(mdsChr220bj, intTypeCol="intron_type"))
```

---

u12NbIndex	<i>Extract index of U2 introns (up/down)stream of U12 introns rows</i>
------------	--

---

**Description**

Extract row numbers of U2-type introns (up/down)stream of U12-type introns (in the @interestDf attribute of an object of class SummarizedExperiment).

**Usage**

```
u12NbIndex(x, intExCol="int_ex", intTypeCol="int_type",
  strandCol="strand", geneIdCol="collapsed_transcripts",
  naUnstrand=FALSE)
```

**Arguments**

x	Object of type SummarizedExperiment.
intExCol	Column name (or number) that represents whether each row of x assays is "intron" or "exon".
intTypeCol	Column name (or number) that represents what type of intron each row of x assays represents.
strandCol	Column name (or number) that represents the strand of each row of assays in x. The values in the column are either "+", "-" or "*".

geneIdCol	Column name (or number) that represents the gene ID of each row of assays in x.
naUnstrand	Replace unstranded results, i.e. introns or exon with "*" strand, with NA. If set as FALSE (default) "*" strand would be same as "+" strand.

**Value**

upIntron	A numeric vector which includes the index of U2-type intron upstream the U12-type introns.
downIntron	A numeric vector which includes the index of U2-type intron downstream the U12-type introns.
upExon	A numeric vector which includes the index of exon upstream the U12-type introns.
downExon	A numeric vector which includes the index of exon downstream the U12-type introns.

**Author(s)**

Ali Oghabian

**See Also**

[u12Index](#)

**Examples**

```
head(u12NbIndex(mdsChr220bj, intExCol="int_ex",
  intTypeCol="intron_type", strandCol="strand",
  geneIdCol="collapsed_transcripts_id", naUnstrand=FALSE))
# Return NA if no strand information available
head(u12NbIndex(mdsChr220bj, intExCol="int_ex",
  intTypeCol="intron_type", strandCol="strand",
  geneIdCol="collapsed_transcripts_id", naUnstrand=TRUE))
```

---

unionRefTr

*Union introns/exons of transcripts*

---

**Description**

Performs union on the overlapping introns/exons so that the final merged transcripts would feature from each exon or intron, one copy.

**Usage**

```
unionRefTr( referenceChr, referenceBegin, referenceEnd, referenceTr,
  referenceIntronExon, intronExon="exon", silent=FALSE)
```

**Arguments**

referenceChr	Chromosome names of the references (e.g. introns).
referenceBegin	A vector that corresponds to the begin coordinates of the reference.
referenceEnd	A vector that corresponds to the end coordinates of the reference.
referenceTr	A character vector that includes transcription IDs.
referenceIntronExon	A vector with the same size as the referenceChr, referenceBegin and referenceEnd which contains 'intron' and 'exon' describing what (either intron or exon) each element of the 3 vectors represents.
intronExon	Should be assigned either 'intron' or 'exon' or c('intron','exon') based on whether match the PWM to the intronic, exonic, or intronic and exonic regions of the reference. By default it seeks matches in intronic regions (intronExon='intron').
silent	Whether run silently.

**Value**

Data frame containing merged transcripts structure. The merged transcripts feature from each intron or exon, one copy ONLY.

**Author(s)**

Ali Oghabian

**See Also**

[annotateU12](#).

**Examples**

```
unU12Ex<-unionRefTr( referenceChr=u12[1:94,"chr"],
  referenceBegin=u12[1:94,"begin"], referenceEnd=u12[1:94,"end"],
  referenceTr=u12[1:94,"trans_name"],
  referenceIntronExon=u12[1:94,"int_ex"], intronExon="exon", silent=TRUE)

unU12Int<-unionRefTr( referenceChr=u12[1:94,"chr"],
  referenceBegin=u12[1:94,"begin"], referenceEnd=u12[1:94,"end"],
  referenceTr=u12[1:94,"trans_name"],
  referenceIntronExon=u12[1:94,"int_ex"], intronExon="intron", silent=TRUE)

unU12IntEx<-unionRefTr( referenceChr=u12[1:94,"chr"],
  referenceBegin=u12[1:94,"begin"], referenceEnd=u12[1:94,"end"],
  referenceTr=u12[1:94,"trans_name"],
  referenceIntronExon=u12[1:94,"int_ex"], intronExon=c("intron","exon"),
  silent=TRUE)
```

---

updateRowDataCol	<i>Updating contents of rowData of SummarizedExperiment objects</i>
------------------	---

---

### Description

Updates the values in a single column of the rowData of SummarizedExperiment objects.

### Usage

```
updateRowDataCol(x, updateCol, value)
```

### Arguments

x	Object of type SummarizedExperiment.
updateCol	Name or the number of the column in the rowData of x to be updated with the new values. if the updateCol does not match to any column names it will be added as a new column.
value	The new Replacing values.

### Value

Returns an object of type SummarizedExperiment.

### Author(s)

Ali Oghabian

### See Also

[annotateU12](#)

### Examples

```
test<- mdsChr220bj
# See the the frequency of each intron type annotation
table(rowData(test)$intron_type)

#Change U2 to u2
newIntType<- as.character(rowData(test)$intron_type)
newIntType[newIntType=="U2" &
 !is.na(newIntType=="U2")]<- "u2"
#Updating values
test<- updateRowDataCol(test, updateCol="intron_type",
 value=newIntType)
#See the frequency of the updated intron type annotations
table(rowData(test)$intron_type)

#Adding a new column
test<- updateRowDataCol(test, updateCol="new_column",
 value=rep(NA, nrow(rowData(test))) )
head(rowData(test))
```

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