# Package 'snapcount'

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```
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     querying of expression counts
Version 1.14.0
Description snapcount is a client interface to the Snaptron webservices
     which support querying by gene name or genomic region.
     Results include raw expression counts derived from alignment of RNA-seq samples
     and/or various summarized measures of expression across one or more regions/genes
     per-sample (e.g. percent spliced in).
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## Description

snapcount is a client interface to the Snaptron webservice which supports querying by gene name or genomic region.

## **Details**

Results include raw expression counts derived from alignment of RNA-seq samples and/or various summarized measures of expression across one or more regions/genes per-sample (e.g. percent spliced in).

To learn more about snapcount, check out the vignette: browseVignettes(package = "snapcount")

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#### Package options

snapcount.host Change the host that snapcount uses when connecting to Snaptron. Default: snaptron.cs.jhu.edu

snapcount.port Change the port that snapcount uses when connecting to Snaptron. Default: 80

#### Author(s)

Maintainer: Rone Charles < rcharle8@jh.edu>

#### See Also

Useful links:

- https://github.com/langmead-lab/snapcount
- Report bugs at https://github.com/langmead-lab/snapcount/issues

Compilation

Enum for Snaptron compilations

## **Description**

The variants for this enum will be populated dynamically after the package has been loaded. If the package cannot connect to the internet the variants will default to:

## Usage

Compilation

#### **Format**

An object of class environment of length 21.

#### **Details**

- gtex
- tcga
- srav2
- sra

#### See Also

http://snaptron.cs.jhu.edu/data.html for more information about Snaptron compilations.

```
qb <- QueryBuilder(compilation = Compilation$gtex, regions = "KCNIP4")
query_jx(qb)</pre>
```

from\_url

Coordinates

Enum for Snaptron Coordinate modifiers

#### **Description**

Enum for Snaptron Coordinate modifiers

## Usage

Coordinates

#### **Format**

An object of class environment of length 4.

#### **Fields**

Exact Return junctions whose start and end coordinates match the boundaries of the region requested.

Within Return junctions whose start and end coordinates are within the boundaries of the region requested.

StartIsExactOrWithin Return junctions whose start coordinate matches, or is within, the boundaries of the region requested.

EndIsExactOrWithin Return junctions whose end coordinate matches, or is within, the boundaries of the region requested.

## **Examples**

```
qb <- QueryBuilder(compilation = "gtex", regions = "CD99")
qb <- set_coordinate_modifier(qb, Coordinates$Exact)
qb</pre>
```

from\_url

Constructs a QueryBuilder object from the given url

## **Description**

Constructs a QueryBuilder object from the given url

## Usage

```
from_url(url)
```

get\_column\_filters 5

#### **Arguments**

url

a well-formed url preferably obtained from a call to the  $\verb"uri_of_last_successful_request" function$ 

#### Value

Returns a QueryBuilder object with attributes set from the parsed url.

## **Examples**

```
sb <- from_url("http://snaptron.cs.jhu.edu/gtex/snaptron?regions=CD99")
get_regions(sb)
get_compilation(sb)</pre>
```

get\_column\_filters

Get or set sample-related contraints for query

## **Description**

Get or set sample-related contraints for query

#### Usage

```
get_column_filters(qb)
set_column_filters(qb, ...)
```

#### **Arguments**

qb a QueryBuilder object constructed using the QueryBuilder function... one or more boolean predicates as either strings or unevaluated expressions

#### Value

get\_column\_filters returns the current filters as a list of strings. set\_column\_filters returns a new QueryBuilder object with the column filters set to the value of column\_filters.

```
qb <- QueryBuilder(compilation = "gtex", regions = "CD99")
# column filters set using a string
qb <- set_column_filters(qb, "SMTS == Brain")
get_column_filters(qb)
# column filters set using unevaluated expression
qb <- set_column_filters(qb, SMTS == "Spleen")
get_column_filters(qb)</pre>
```

get\_compilation

Get or set query compilation

## **Description**

Get or set query compilation

#### Usage

```
get_compilation(qb)
set_compilation(qb, compilation)
```

## **Arguments**

qb A QueryBuilder object constructed using the QueryBuilder function.

compilation A single string containing the name of the Snaptron data source. Any variant of

the Compilation enum can also be passed as an argument.

#### Value

get\_compilation returns the current compilation as string. set\_compilation returns a new QueryBuilder object with the compilation set to the value of compilation.

## **Examples**

```
qb <- QueryBuilder(compilation = "gtex", regions = "CD99")
get_compilation(qb)
qb <- set_compilation(qb, Compilation$tcga)
get_compilation(qb)</pre>
```

```
get_coordinate_modifier
```

Get or set coordinate modifiers for the query.

## **Description**

Get or set coordinate modifiers for the query.

## Usage

```
get_coordinate_modifier(qb)
set_coordinate_modifier(qb, coordinate_modifier)
```

get\_regions 7

#### **Arguments**

```
qb a QueryBuilder object constructed using the QueryBuilder function. coordinate_modifier any of the variants of the Coordinates enum.
```

#### Value

get\_coordinate\_modifier returns the current coodinate modifier as a string. set\_coordinate\_modifier returns a new QueryBuilder object with the coordinate modifier set to the value of coordinate\_modifier.

#### **Examples**

```
qb <- QueryBuilder(compilation = "gtex", regions = "CD99")
qb <- set_coordinate_modifier(qb, Coordinates$Within)
get_coordinate_modifier(qb)</pre>
```

get\_regions

Get or set query regions

#### **Description**

Get or set query regions

#### Usage

```
get_regions(qb)
set_regions(qb, regions)
```

#### **Arguments**

qb A QueryBuilder object constructed using the QueryBuilder function.

regions Either a list of 1 more HUGO gene names as strings e.g. "BRCA1" or a Granges class object containing one or more geneomic intervals (e.g. "chr1:1-1000").

#### Value

get\_regions returns the current regions as a list of strings. set\_regions returns a new QueryBuilder object with the regions set to the value of regions.

```
qb <- QueryBuilder(compilation = "gtex", regions = "CD99")
get_regions(qb)
qb <- set_regions(qb, "chr1:1-1000")
get_regions(qb)
qb <- set_regions(qb, GenomicRanges::GRanges("chr1", "1-1000"))
get_regions(qb)</pre>
```

get\_sids

get\_row\_filters

Get or set range-related contraints for query

#### **Description**

Get or set range-related contraints for query

## Usage

```
get_row_filters(qb)
set_row_filters(qb, ...)
```

## Arguments

qb a QueryBuilder object constructed using the QueryBuilder function.

... one or more boolean predicates as either strings or unevaluated expressions.

#### Value

get\_row\_filters returns the current row filters as list of strings. set\_row\_filters returns a new QueryBuilder object with the row filters set to the value of row\_filters.

## **Examples**

```
qb <- QueryBuilder(compilation = "gtex", regions = "CD99")
# row filters set as a string
qb <- set_row_filters(qb, "strand == +")
get_row_filters(qb)
# row filters set using unevaluated expression
qb <- set_row_filters(qb, strand == "+")
get_row_filters(qb)</pre>
```

get\_sids

Get or set query sample ids

## **Description**

Get or set query sample ids

## Usage

```
get_sids(qb)
set_sids(qb, sids)
```

#### **Arguments**

qb a QueryBuilder object constructed using the QueryBuilder function. sids a vector or 1 or more whole numbers to filter results on.

#### Value

get\_sids returns the current sample ids as a vector of integers. set\_sids returns a new QueryBuilder object with the sample ids set to the value of sids.

#### **Examples**

```
qb <- QueryBuilder(compilation = "gtex", regions = "CD99")
qb <- set_sids(qb, c(1, 2, 3))
get_sids(qb)</pre>
```

junction\_inclusion\_ratio

Relative measure of splice variant usage similar to PSI that allows for 2 arbitrarily defined groups of junctions (not limited to cassette exons).

## Description

Calculates a coverage summary statistic per sample of the normalized coverage difference between two sets of separate junctions defined by at least two basic queries and organized into two groups.

#### Usage

```
junction_inclusion_ratio(group1, group2, group_names = NULL)
```

## **Arguments**

```
group1, group2 Each group is a list of 1 or more QueryBuilder objects
group_names Optional vector of strings representing the group names
```

#### **Details**

```
The summary statistic is as follows: If the coverage of the first group is "A" and the second is "B": JIR(A,B)=(A-B)/(A+B+1)
```

This is calculated for every sample that occurs in one or the other (or both) groups results.

#### Value

A DataFrame of samples, with their JIR score and metadata, which had > 0 coverage in at least one resulting row in at least one of the groups

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#### **Examples**

```
sb1 <- QueryBuilder(compilation = "srav2", regions = "chr2:29446395-30142858")
sb1 <- set_coordinate_modifier(sb1, Coordinates$Within)
sb1 <- set_row_filters(sb1, strand == "-")
sb2 <- QueryBuilder(compilation = "srav2", regions = "chr2:29416789-29446394")
sb2 <- set_coordinate_modifier(sb2, Coordinates$Within)
sb2 <- set_row_filters(sb2, strand == "-")
junction_inclusion_ratio(list(sb1), list(sb2))</pre>
```

junction\_intersection Get the intersection of junctions from 2 or more compilations which are on the same reference

#### **Description**

This function operates similar to the junction\_union() function, i.e it is cross compilation and merging of the same junction from multiple compilations will be handled exactly the same way. But instead of every junction which appears in at least one compilation, only the junctions which appear in *every* compilation will be returned.

#### Usage

```
junction_intersection(...)
```

## Arguments

.. One or more QueryBuilder objects

#### Value

A RangedSummarizedExperiment of junctions common across compilations

#### See Also

```
junction_union()
```

```
# Using query builder wrappers
sb1 <- QueryBuilder(compilation = "gtex", regions = "chr1:1879786-1879786")
sb1 <- set_coordinate_modifier(sb1, Coordinates$EndIsExactOrWithin)
sb1 <- set_row_filters(sb1, strand == "-")
sb2 <- QueryBuilder(compilation = "tcga", regions = "chr1:1879786-1879786")
sb2 <- set_coordinate_modifier(sb2, Coordinates$EndIsExactOrWithin)
sb2 <- set_row_filters(sb2, strand == "-")
junction_intersection(sb1, sb2)</pre>
```

junction\_union 11

junction_union	Get the union of junctions from 2 or more compilations which are on the same reference
	·

#### **Description**

This function queries 2 or more compilations which are on the same reference version (e.g. hg38) and merges the resulting junctions into a single output table, unioning the sample coverage columns and the snaptron\_id (jx ID) columns (the latter delimiter will be ":"). All sample IDs will be disjoint between compilations.

## Usage

```
junction_union(...)
```

#### **Arguments**

... One or more QueryBuilder objects

#### **Details**

Union is based on the following fields (combined into a comparison key):

- group
- · chromosome
- start
- end
- strand

The goal is to have a single list of junctions where every junction occurs in at least one compilation and if a junction occurs in > 1 compilations it still only has a single row representing all the samples across compilations that it appears in. Sample aggregate statistics will be recalculated for junctions which are merged across *all* samples from all compilations:

- · sample\_count
- · coverage\_sum
- coverage\_avg
- · coverage\_median

#### Value

A RangedSummarizedExperiment of junctions appearing in at least one compilation

#### See Also

```
junction_intersection()
```

12 percent\_spliced\_in

#### **Examples**

```
# Using query builder wrappers
sb1 <- QueryBuilder(compilation = "gtex", regions = "chr1:1879786-1879786")
sb1 <- set_coordinate_modifier(sb1, Coordinates$EndIsExactOrWithin)
sb1 <- set_row_filters(sb1, strand == "-")
sb2 <- QueryBuilder(compilation = "tcga", regions = "chr1:1879786-1879786")
sb2 <- set_coordinate_modifier(sb2, Coordinates$EndIsExactOrWithin)
sb2 <- set_row_filters(sb2, strand == "-")
junction_union(sb1, sb2)</pre>
```

percent\_spliced\_in

Relative measure of splice variant usage, limited currently to cassette exon splice variants

#### Description

Similar to the JIR, this calculates Percent Spliced In (PSI) statistics for the definition of 2 different groups: inclusion and exclusion. Currently this function only supports the cassette exon use case.

## Usage

```
percent_spliced_in(
   inclusion_group1,
   inclusion_group2,
   exclusion_group,
   min_count = 20,
   group_names = NULL
)
```

#### **Arguments**

```
inclusion_group1, inclusion_group2, exclusion_group

Where each is a list of 1 or more QueryBuilder objects

min_count

minimum total count (denominator) required to not be assigned -1
group_names

Optional vector of strings representing the group names
```

#### **Details**

Inclusion typically defines 2 basic queries, one for the junction preceding the cassette exon, and the second for the junction following the cassette exon. The exclusion group contains one basic query which defines the junction which skips the cassette exon.

The PSI itself is implemented as:

```
PSI(inclusion1, inclusion2, exclusion) = mean(inclusion1, inclusion2) / (mean(inclusion1, inclusion2) + exclusion)
```

where each term denotes the coverage of junctions that resulted from the basic queries in that group in the current sample.

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## Value

A DataFrame of samples, with their PSI score and metadata, which had > 0 coverage in at least one resulting row in at least one of the groups

#### **Examples**

```
in1 <- QueryBuilder(compilation = "srav2", regions = "chr1:94468008-94472172")
in1 <- set_coordinate_modifier(in1, Coordinates$Exact)
in1 <- set_row_filters(in1, strand == "+")

in2 <- QueryBuilder(compilation = "srav2", regions = "chr1:94468008-94472172")
in2 <- set_coordinate_modifier(in2, Coordinates$Exact)
in2 <- set_row_filters(in2, strand == "+")

ex <- QueryBuilder(compilation = "srav2", regions = "chr1:94468008-94475142")
ex <- set_coordinate_modifier(ex, Coordinates$Exact)
ex <- set_row_filters(ex, strand == "+")

percent_spliced_in(list(in1), list(in2), list(ex))</pre>
```

QueryBuilder Co

Construct a QueryBuilder object given a compilation and one or regions.

## **Description**

Construct a QueryBuilder object given a compilation and one or regions.

## Usage

```
QueryBuilder(compilation, regions)
```

## **Arguments**

compilation A single string containing the name of the Snaptron data source. Any variant of

the Compilation enum can also be passed an argument.

regions Either a list of 1 more HUGO gene names as strings e.g. "BRCA1" or a Granges

class object containing one or more geneomic intervals (e.g. "chr1:1-1000").

#### Value

A QueryBuilder object.

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#### **Examples**

```
# contruct a query builder for GTEX data source and BRAC1 gene
qb <- QueryBuilder(compilation = Compilation$gtex, regions = "BRCA1")

# contruct a query builder for TCGA data source and chromosome region
qb <- QueryBuilder(compilation = "tcga", regions = "chr1:1-1000")

# construct a query builder for TCGA data source using GRanges object
library(GenomicRanges)
qb <- QueryBuilder(compilation = "tcga", regions = GRanges("chr1", "1-1000"))</pre>
```

query\_jx

Query Junctions/Genes/Exons

#### **Description**

Given one or more gene names or genomic range intervals it will return a list of 0 or more genes, junctions, or exons (depending on which query form is used) which overlap the ranges.

#### Usage

```
query_jx(sb, return_rse = TRUE, split_by_region = FALSE)
query_gene(sb, return_rse = TRUE, split_by_region = FALSE)
query_exon(sb, return_rse = TRUE, split_by_region = FALSE)
```

## **Arguments**

sb A SnaptronQueryBuilder object

return\_rse Should the query data be returned as a simple data frame or converted to a

Ranged Summarized Experiment.

split\_by\_region

By default the results from multiple queries will be returned in a RangedSummarizedExperiment object with a rowData entry for each, labeling each result row according to the query it resulted from. However, if this is set to TRUE, the result will be a list of RangedSummarizedExperiment objects, one per original interval/gene. This latter option may be useful, but it requires a separate copy of the sample metadata for each original interval/gene.

#### Value

Functions will return either a RangedSummarizedExperiment or data.table depending on whether the return\_rse parameter is set to TRUE or FALSE.

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#### **Examples**

```
# Contruct a QueryBuilder object
qb <- QueryBuilder(compilation = "gtex", regions = "chr1:1-100000")
qb <- set_row_filters(qb, samples_count >= 20)
query_jx(qb)

qb <- set_row_filters(qb, NULL)
qb <- set_column_filters(qb, SMTS == "Brain")
query_gene(qb)</pre>
```

shared\_sample\_counts

Shared Sample Count (SSC): counts total number of samples in which 2 different junctions both occur in.

#### **Description**

This produces a list of user-specified groups and the read coverage of the junctions in all the samples which were shared across all the basic queries occurring in each group.

## Usage

```
shared_sample_counts(..., group_names = NULL)
```

## **Arguments**

... One or more lists of QueryBuilder objects

group\_names Optional vector of character strings representing group names

#### **Details**

Example: User defines a single group of junctions "GroupA" made up of 2 separate regions (two basic queries).

An SSC query will return a single line for GroupA which will have the total number of samples which had at least one junction which was returned from both basic queries. It will also report a summary statistic of the total number of groups which had one or more samples that were shared across the basic queries, in this case it would be 1. Also, it will report the number of groups which had at least one shared sample and which had matching junctions (from the query) which were fully annotated.

This function can be used to determine how much cross-sample support there is for a particular junction configuration (typically a cassette exon).

#### Value

A DataFrame of results based on the list of groups passed in via "group\_names". Each group is reported with the # of unique samples which occurred in all of its defined set of related basic queries (e.g. two inclusion basic queries in a cassette exon scenario).

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#### **Examples**

```
g1 <- QueryBuilder(compilation = "gtex", regions = "chr1:1879786-1879786")
g1 <- set_coordinate_modifier(g1, Coordinates$EndIsExactOrWithin)
g1 <- set_row_filters(g1, strand == "-")
g2 <- QueryBuilder(compilation = "gtex", regions = "chr1:1879903-1879903")
g2 <- set_coordinate_modifier(g2, Coordinates$StartIsExactOrWithin)
g2 <- set_row_filters(g2, strand == "-")
ssc<-shared_sample_counts(list(g1, g2))</pre>
```

tissue\_specificity

Tissue Specificity (TS): produces a list of samples with their tissues marked which either contain queried junctions (1) or not (0); can be used as input to significance testing methods such as Kruskal-Wallis to look for tissue enrichment (currently only works for the GTEx compilation).

#### **Description**

Lists the number of samples labeled with a specific tissue type. Samples are filtered for ones which have junctions across all the user-specified groups. That is, if a sample only appears in the results of some of the groups (from their basic queries) it will be assigned a 0, otherwise if it is in all of the groups' results it will be assigned a 1. This is similar to the SSC high level query type, but doesn't sum the coverage.

#### Usage

```
tissue_specificity(..., group_names = NULL)
```

#### **Arguments**

... One or more QueryBuilder objects
group\_names Optional vector of strings representing the group names

#### **Details**

The samples are then grouped by their tissue type (e.g. Brain). This is useful for determining if there's an enrichment for a specific tissue in the set of junctions queried. Results from this can be fed to a statistical test, such as the Kruskal-wallis non-parametric rank test. This query is limited to GTEx only, due to the fact that GTEx is one of the few compilations that has consistent and complete tissue metadata.

#### Value

A DataFrame of all samples in the compilation with either a 0 or 1 indicating their occurrence and shared status (if > 1 group passed in). Occurrence here is if the sample has at least one result with > 0 coverage, and further, if > 1 group is passed in, then if it occurs in the results of all groups. Also includes the sample tissue type and sample\_id.

#### **Examples**

```
in1 <- QueryBuilder(compilation = "gtex", regions = "chr4:20763023-20763023")
in1 <- set_coordinate_modifier(in1, Coordinates$EndIsExactOrWithin)
in1 <- set_row_filters(in1, strand == "-")

in2 <- QueryBuilder(compilation = "gtex", regions = "chr4:20763098-20763098")
in2 <- set_coordinate_modifier(in2, Coordinates$StartIsExactOrWithin)
in2 <- set_row_filters(in2, strand == "-")

tissue_specificity(list(in1, in2))</pre>
```

```
uri_of_last_successful_request
```

Return the URI of the last successful request to Snaptron

## **Description**

This function can be paired with the from\_url method from the QueryBuilder class, allowing users to share sources of data from Snaptron.

#### Usage

```
uri_of_last_successful_request()
```

#### Value

URI of last successful request to Snaptron or NULL if there have not been any successful requests.

```
qb <- QueryBuilder(compilation = "gtex", regions = "CD99")
query_jx(qb)
uri_of_last_successful_request()</pre>
```

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