

# Package ‘dasper’

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**Title** Detecting aberrant splicing events from RNA-sequencing data

**Version** 1.9.0

**Date** 2021-08-05

**Description** The aim of dasper is to detect aberrant splicing events from RNA-seq data. dasper will use as input both junction and coverage data from RNA-seq to calculate the deviation of each splicing event in a patient from a set of user-defined controls. dasper uses an unsupervised outlier detection algorithm to score each splicing event in the patient with an outlier score representing the degree to which that splicing event looks abnormal.

**License** Artistic-2.0

**URL** <https://github.com/dzhang32/dasper>

**BugReports** <https://support.bioconductor.org/t/dasper>

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annotate_junc_ref	<i>Deprecated function: Annotate junctions using reference annotation</i>
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### Description

annotate\_junc\_ref is deprecated, please use junction\_annot instead. annotates junctions by whether their start and end position precisely overlaps with a known exon boundary. Using this information along with the strand, junctions are categorised into "annotated", "novel\_acceptor", "novel\_donor", "novel\_combo", "novel\_exon\_skip", "ambig\_gene" and "none".

### Usage

```
annotate_junc_ref(junc_metadata, gtf)

.get_ref_exons_annot(
  junc_metadata,
  ref_exons,
  ref_cols = c("strand", "gene_name", "gene_id", "transcript_id", "exon_id")
)
```

```

.tidy_junc_annot(
  junc_metadata,
  cols_to_merge = c("strand", "gene_name", "gene_id")
)

.classify_junc(junc_metadata, ref_junc)

.get_gr_for_start_end(gr)

.merge_lists(x, y)

```

### Arguments

`junc_metadata` junction metadata in a [GRanges-class](#) format, the essential component being the junction co-ordinates.

`gtf` either path to gtf or object of class `ensemblGenome` loaded using `refGenome`.

### Value

junction metadata as a [GRanges-class](#) object with additional columns that detail overlapping genes/transcripts/exons and junction categories.

---

coverage_norm	<i>Processing coverage</i>
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### Description

The set of functions prefixed with "coverage\_" are used to process coverage data. They are designed to be run after you have processed your junctions in the order `coverage_norm`, `coverage_score`. Or, alternatively the wrapper function `coverage_process` can be used to run the 2 functions stated above in one go. For more details of the individual functions, see "Details".

### Usage

```

coverage_norm(
  junctions,
  ref,
  unannot_width = 20,
  coverage_paths_case,
  coverage_paths_control,
  coverage_chr_control = NULL,
  load_func = .coverage_load,
  bp_param = BiocParallel::SerialParam(),
  norm_const = 1
)

coverage_process(

```

```

junctions,
ref,
unannot_width = 20,
coverage_paths_case,
coverage_paths_control,
coverage_chr_control = NULL,
load_func = .coverage_load,
bp_param = BiocParallel::SerialParam(),
norm_const = 1,
score_func = .zscore,
...
)

coverage_score(junctions, coverage, score_func = .zscore, ...)

```

### Arguments

junctions	junction data as a <a href="#">RangedSummarizedExperiment-class</a> object.
ref	either path to gtf/gff3 or object of class <a href="#">TxDb-class</a> or <a href="#">EnsDb-class</a> . <a href="#">EnsDb-class</a> is required if you intend to annotate junctions with gene symbols/names.
unannot_width	integer scalar determining the width of the region to obtain coverage from when the end of a junction does not overlap an existing exon.
coverage_paths_case	paths to the BigWig files containing the coverage of your case samples.
coverage_paths_control	paths to the BigWig files for control samples.
coverage_chr_control	either "chr" or "no_chr", indicating the chromosome format of control coverage data. Only required if the chromosome format of the control BigWig files is different to that of your cases.
load_func	a function to use to load coverage. Currently only for internal use to increase testing speed.
bp_param	a <a href="#">BiocParallelParam-class</a> instance denoting whether to parallelise the loading of coverage across BigWig files.
norm_const	numeric scaler to add to the normalisation coverage to avoid dividing by 0s and resulting NaN or Inf values.
score_func	function to score junctions by their abnormality. By default, will use a z-score but can be switched to a user-defined function. This function must take as input an x and y argument, containing case and control counts respectively. This must return a numeric vector equal to the length of x with elements corresponding to a abnormality of each junction.
...	additional arguments passed to score_func.
coverage	list containing normalised coverage data that is outputted from <a href="#">coverage_norm</a> .

## Details

coverage\_process wraps all "coverage\_" prefixed functions in [dasper](#). This is designed to simplify processing of the coverage data for those familiar or uninterested with the intermediates.

coverage\_norm obtains regions of interest for each junction where coverage disruptions would be expected. These consist of the intron itself the overlapping exon definitions (if ends of junctions are annotated), picking the shortest exon when multiple overlap one end. If ends are unannotated, coverage\_norm will use a user-defined width set by unannot\_width. Then, coverage will be loaded using [megadepth](#) and normalised to a set region per junction. By default, the boundaries of each gene associated to a junction are used as the region to normalise to.

coverage\_score will score disruptions in the coverage across the intronic/exonic regions associated with each junction. This abnormality score generated by score\_func operates by calculating the deviation of the coverage in patients to a coverage across the same regions in controls. Then, for each junction it obtains the score of the region with the greatest disruption.

## Value

junctions as [SummarizedExperiment](#) object with additional assays named "coverage\_region" and "coverage\_score". "coverage\_region" labels the region of greatest disruption (1 = exon\_start, 2 = exon\_end, 3 = intron) and "coverage\_score" contains the abnormality scores of the region with the greatest disruption.

## Functions

- coverage\_norm: Load and normalise coverage from RNA-sequencing data
- coverage\_score: Score coverage by their abnormality

## Examples

```
##### Set up txdb #####

# use GenomicState to load txdb (GENCODE v31)
ref <- GenomicState::GenomicStateHub(
  version = "31",
  genome = "hg38",
  filetype = "TxDb"
)[[1]]

##### Set up BigWig #####

# obtain path to example bw on recount2
bw_path <- recount::download_study(
  project = "SRP012682",
  type = "samples",
  download = FALSE
)[[1]]

##### junction_process #####
```

```

junctions_processed <- junction_process(
  junctions_example,
  ref,
  types = c("ambig_gene", "unannotated"),
)

##### install megadept #####

# required to load coverage in coverage_norm()
megadept::install_megadept(force = FALSE)

##### coverage_norm #####

coverage_normed <- coverage_norm(
  junctions_processed,
  ref,
  unannot_width = 20,
  coverage_paths_case = rep(bw_path, 2),
  coverage_paths_control = rep(bw_path, 2)
)

##### coverage_score #####

junctions <- coverage_score(junctions_processed, coverage_normed)

##### coverage_process #####

# this wrapper will obtain coverage scores identical to those
# obtained through running the individual wrapped functions shown below
junctions_w_coverage <- coverage_process(
  junctions_processed,
  ref,
  coverage_paths_case = rep(bw_path, 2),
  coverage_paths_control = rep(bw_path, 3)
)

# the two objects are equivalent
all.equal(junctions_w_coverage, junctions, check.attributes = FALSE)

```

---

dasper

*dasper: detecting aberrant splicing events from RNA-seq data*

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## Description

Placeholder for package description - to be updated

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junctions_example	<i>Set of example junctions</i>
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### Description

A dataset containing the example junction data for 2 case and 3 control samples outputted from [junction\\_load](#). The junctions have been filtered for only those lying on chromosome 21 or 22.

### Usage

```
junctions_example
```

### Format

[RangedSummarizedExperiment-class](#) object from [SummarizedExperiment](#) detailing the counts, co-ordinates of junctions lying on chromosome 21/22 for 2 example samples and 3 controls:

**assays** matrix with counts for junctions (rows) and 5 samples (cols)

**colData** example sample metadata

**rowRanges** [GRanges-class](#) object describing the co-ordinates and strand of each junction

### Source

generated using data-raw/junctions\_example.R

---

junction_annot	<i>Processing junctions</i>
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---

### Description

The set of functions prefixed with "junction\_" are used to process junction data. They are designed to be run in a sequential manner in the order `junction_annot`, `junction_filter`, `junction_norm`, `junction_score`. Or, alternatively the wrapper function `junction_process` can be used to run all 4 of the functions stated above in one go. For more details of the individual functions, see "Details".

### Usage

```
junction_annot(  
  junctions,  
  ref,  
  ref_cols = c("gene_id", "tx_name", "exon_id"),  
  ref_cols_to_merge = c("gene_id")  
)  
  
junction_filter(  
  junctions,  
  ref,  
  ref_cols = c("gene_id", "tx_name", "exon_id"),  
  ref_cols_to_merge = c("gene_id")  
)
```

```

junctions,
count_thresh = c(raw = 5),
n_samp = c(raw = 1),
width_range = NULL,
types = NULL,
regions = NULL
)

junction_norm(junctions)

junction_process(
  junctions,
  ref,
  ref_cols = c("gene_id", "tx_name", "exon_name"),
  ref_cols_to_merge = c("gene_id"),
  count_thresh = c(raw = 5),
  n_samp = c(raw = 1),
  width_range = NULL,
  types = NULL,
  regions = NULL,
  score_func = .zscore,
  ...
)

junction_score(junctions, score_func = .zscore, ...)

```

### Arguments

junctions	junction data as a <a href="#">RangedSummarizedExperiment-class</a> object.
ref	either path to gtf/gff3 or object of class <a href="#">TxDb-class</a> or <a href="#">EnsDb-class</a> . <a href="#">EnsDb-class</a> is required if you intend to annotate junctions with gene symbols/names.
ref_cols	character vector listing the names of the columns in ref for which to annotate junctions with. Must contain "gene_id", used for categorising junctions.
ref_cols_to_merge	character vector listing which of the annotation columns ref_cols should be merged into in columns to merge into a single column per junction. Must contain "gene_id", used for categorising junctions.
count_thresh	named vector with names matching the names of the <a href="#">assays</a> in junctions. Values denote the number of counts below which a junction will be filtered out.
n_samp	named vector with names matching the names of the <a href="#">assays</a> in junctions. Values denotes number of samples that have to express the junction above the count_thresh in order for that junction to not be filtered.
width_range	numeric vector of length 2. The first element denoting the lower limit of junction width and the second the upper limit. Junctions with widths outside this range will be filtered out.
types	any junctions matching these types, derived from <a href="#">junction_annot</a> will be filtered out.

regions	any junctions overlapping this set of regions (in a <a href="#">GRanges-class</a> format) will be filtered out.
score_func	function to score junctions by their abnormality. By default, will use a z-score but can be switched to a user-defined function. This function must take as input an x and y argument, containing case and control counts respectively. This must return a numeric vector equal to the length of x with elements corresponding to a abnormality of each junction.
...	additional arguments passed to score_func.

## Details

junction\_process wraps all "junction\_" prefixed functions in [dasper](#) except [junction\\_load](#). This is designed to simplify processing of the junction data for those familiar or uninterested with the intermediates.

junction\_annot annotates junctions by 1. whether their start and/or end position precisely overlaps with an annotated exon boundary and 2. whether that junction matches an intron definition from existing annotation. Using this information along with the strand, junctions are categorised into "annotated", "novel\_acceptor", "novel\_donor", "novel\_combo", "novel\_exon\_skip", "ambig\_gene" and "unannotated".

junction\_filter filters out "noisy" junctions based on counts, the width of junctions, annotation category of the junction returned from [junction\\_annot](#) and whether the junction overlaps with a set of (blacklist) regions.

junction\_norm normalises the raw junction counts by 1. building junction clusters by finding junctions that share an acceptor or donor position and 2. calculating a proportion-spliced-in (PSI) for each junction by dividing the raw junction count by the total number of counts in it's associated cluster.

junction\_score will use the counts contained within the "norm" [assay](#) to calculate a deviation of each patient junction from the expected distribution of control junction counts. The function used to calculate this abnormality score can be user-inputted or left as the default z-score. Junctions will also be labelled based on whether they are up-regulated (+1) or down-regulated (-1) with respect to controls junction and this information is stored in the [assay](#) "direction" for use in [outlier\\_aggregate](#).

## Value

[RangedSummarizedExperiment-class](#) object containing filtered, annotated, normalised junction data with abnormality scores.

## Functions

- junction\_annot: Annotate junctions using reference annotation
- junction\_filter: Filter junctions by count, width, annotation or region
- junction\_norm: Normalise junction counts by cluster
- junction\_score: Score patient junctions by their abnormality

**See Also**

ENCODE blacklist regions are recommended to be included as regions for `junction_filter` and can be downloaded from <https://github.com/Boyle-Lab/Blacklist/blob/master/lists/hg38-blacklist.v2.bed.gz>. Further information can be found via the publication <https://www.nature.com/articles/s41598-019-45839-z>.

**Examples**

```
##### Set up txdb #####

# use GenomicState to load txdb (GENCODE v31)
ref <- GenomicState::GenomicStateHub(
  version = "31",
  genome = "hg38",
  filetype = "TxDb"
)[[1]]

##### junction_annot #####

junctions <- junction_annot(junctions_example, ref)

##### junction_filter #####

junctions <- junction_filter(
  junctions,
  types = c("ambig_gene", "unannotated")
)

##### junction_norm #####

junctions <- junction_norm(junctions)

##### junction_score #####

junctions <- junction_score(junctions)

##### junction_process #####

junctions_processed <- junction_process(
  junctions_example,
  ref,
  types = c("ambig_gene", "unannotated")
)

# the two objects are equivalent
all.equal(junctions_processed, junctions, check.attributes = FALSE)
```

## Description

junction\_load loads in raw patient and control junction data and formats it into a [RangedSummarizedExperiment-class](#) object. Control samples can be the user's in-house samples or selected from GTEx v6 data publicly released through the [recount2](#) and downloaded through [snaptron](#). By default, junction\_load expects the junction data to be in STAR aligned format (SJ.out) but this can be modified via the argument load\_func.

## Usage

```
junction_load(
  junction_paths,
  metadata = dplyr::tibble(samp_id = stringr::str_c("samp_",
    seq_along(junction_paths))),
  controls = rep(FALSE, length(junction_paths)),
  load_func = .STAR_load,
  chrs = NULL,
  coord_system = 1
)
```

## Arguments

junction_paths	path(s) to junction data.
metadata	data.frame containing sample metadata with rows in the same order as junction_paths.
controls	either a logical vector of the same length as junction_paths with TRUE representing controls. Or, one of "fibroblasts", "lymphocytes", "skeletal_muscle", "whole_blood" representing the samples of which GTEx tissue to use as controls. By default, will assume all samples are patients.
load_func	function to load in junctions. By default, requires STAR formatted junctions (SJ.out). But this can be switched dependent on the format of the user's junction data. Function must take as input a junction path then return a data.frame with the columns "chr", "start", "end", "strand" and "count".
chrs	chromosomes to keep. By default, no filter is applied.
coord_system	1 (1-based) or 0 (0-based) denoting the co-ordinate system corresponding to the user junctions from junction_paths. Only used when controls is set to "fibroblasts" to ensure GTEx data is harmonised to match the co-ordinate system of the user's junctions.

## Value

[RangedSummarizedExperiment-class](#) object containing junction data.

## Examples

```
junctions_example_1_path <-
  system.file("extdata",
    "junctions_example_1.txt",
    package = "dasper",
    mustWork = TRUE
```

```
)
junctions_example_2_path <-
  system.file("extdata",
             "junctions_example_2.txt",
             package = "dasper",
             mustWork = TRUE
  )

junctions <-
  junction_load(
    junction_paths = c(junctions_example_1_path, junctions_example_2_path)
  )

junctions
```

---

outlier\_aggregate      *Processing outliers*

---

### Description

The set of functions prefixed with "outlier\_" are used to detect outliers. They are designed to be run after you have extracted your junctions and coverage based features, in the order `outlier_detect`, `outlier_aggregate`. Or, alternatively the wrapper function `outlier_process` can be used to run the 2 functions stated above in one go. For more details of the individual functions, see "Details".

### Usage

```
outlier_aggregate(
  junctions,
  samp_id_col = "samp_id",
  bp_param = BiocParallel::SerialParam()
)

outlier_detect(
  junctions,
  feature_names = c("score", "coverage_score"),
  bp_param = BiocParallel::SerialParam(),
  ...
)

outlier_process(
  junctions,
  feature_names = c("score", "coverage_score"),
  samp_id_col = "samp_id",
  bp_param = BiocParallel::SerialParam(),
  ...
)
```

**Arguments**

junctions	junction data as a <a href="#">RangedSummarizedExperiment-class</a> object.
samp_id_col	name of the column in the <a href="#">SummarizedExperiment</a> that details the sample ids.
bp_param	a <a href="#">BiocParallelParam-class</a> instance denoting whether to parallelise the calculating of outlier scores across samples.
feature_names	names of assays in junctions that are to be used as input into the outlier detection model.
...	additional arguments passed to the outlier detection model (isolation forest) for setting parameters.

**Details**

outlier\_process wraps all "outlier\_" prefixed functions in [dasper](#). This is designed to simplify processing of the detecting outlier junctions for those familiar or uninterested with the intermediates.

outlier\_detect will use the features in [assays](#) named feature\_names as input into an unsupervised outlier detection algorithm to score each junction based on how outlier-y it looks in relation to other junctions in the patient. The default expected\_score and coverage\_score features can be calculated using the [junction\\_process](#) and [coverage\\_process](#) respectively.

outlier\_aggregate will aggregate the outlier scores into a cluster-level. It will then rank each cluster based on this aggregated score and annotate each cluster with it's associated gene and transcript.

**Value**

DataFrame with one row per cluster detailing each cluster's associated junctions, outlier scores, ranks and genes.

**Functions**

- outlier\_aggregate: Aggregate outlier scores from per junction to cluster-level
- outlier\_detect: Detecting outlier junctions

**See Also**

for more details on the isolation forest model used: <https://scikit-learn.org/stable/modules/generated/sklearn.ensemble.IsolationForest.html>

**Examples**

```
##### Set up txdb #####

# use GenomicState to load txdb (GENCODE v31)
ref <- GenomicState::GenomicStateHub(
  version = "31",
  genome = "hg38",
  filetype = "TxDb"
)[[1]]
```

```
##### Set up BigWig #####

# obtain path to example bw on recount2
bw_path <- recount::download_study(
  project = "SRP012682",
  type = "samples",
  download = FALSE
)[[1]]

# cache the bw for speed in later
# examples/testing during R CMD Check
bw_path <- dasper:::file_cache(bw_path)

##### junction_process #####

junctions_processed <- junction_process(
  junctions_example,
  ref,
  types = c("ambig_gene", "unannotated"),
)

##### coverage_process #####

junctions_w_coverage <- coverage_process(
  junctions_processed,
  ref,
  coverage_paths_case = rep(bw_path, 2),
  coverage_paths_control = rep(bw_path, 3)
)

##### outlier_detect #####

junctions_w_outliers <- outlier_detect(junctions_w_coverage)

##### outlier_aggregate #####

outlier_scores <- outlier_aggregate(junctions_w_outliers)

##### outlier_process #####

# this wrapper will obtain outlier scores identical to those
# obtained through running the individual wrapped functions shown below
outlier_processed <- outlier_process(junctions_w_coverage)

# the two objects are equivalent
all.equal(outlier_processed, outlier_scores, check.attributes = FALSE)
```

plot\_sashimi

*Visualise RNA-seq data in the form of a sashimi plot***Description**

plot\_sashimi plots the splicing events and coverage across specific genes/transcripts/regions of interest. Unlike traditional sashimi plots, coverage and junction tracks are separated, which enables user's to choose whether they would like to plot only the junctions.

**Usage**

```
plot_sashimi(
  junctions,
  ref,
  gene_tx_id,
  gene_tx_col,
  case_id = NULL,
  sum_func = mean,
  region = NULL,
  assay_name = "norm",
  annot_colour = c(ggpubr::get_palette("jco", 1), ggpubr::get_palette("npg", 7)[c(1, 3,
    2, 5, 6)], ggpubr::get_palette("jco", 6)[c(3)]),
  digits = 2,
  count_label = TRUE,
  coverage_paths_case = NULL,
  coverage_paths_control = NULL,
  coverage_chr_control = NULL,
  load_func = .coverage_load,
  binwidth = 100
)
```

**Arguments**

junctions	junction data as a <a href="#">RangedSummarizedExperiment-class</a> object.
ref	either path to <code>gtf/gff3</code> or object of class <code>TxDb-class</code> or <code>EnsDb-class</code> . <code>EnsDb-class</code> is required if you intend to annotate junctions with gene symbols/names.
gene_tx_id	character scalar with the id of the gene. This must be a an identifier for a gene or transcript, which has a matching entry in <code>ref</code> .
gene_tx_col	character scalar with the name of the column to search for the <code>gene_tx_id</code> in <code>ref</code> .
case_id	list containing 1 element. The contents of this element must be a character vector specifying sample ids that are to be plotted. The name of this element must correspond to the column containing sample ids in the junction <code>SummarizedExperiment::mcols()</code> . By default, all cases will be plotted.
sum_func	function that will be used to aggregate the junction counts and coverage for controls. By default, <code>mean</code> will be used.

region	a <a href="#">GenomicRanges</a> of length 1 that is used to filter the exons/junctions plotted. Only those that overlap this region are plotted.
assay_name	a character scalar with the name of the <code>SummarizedExperiment::assay()</code> from which to obtain junction counts.
annot_colour	character vector length 7, representing the colours of junction types.
digits	used in <code>round()</code> , specifying the number of digits to round the junction counts to for visualisation purposes.
count_label	logical value specifying whether to add label the count of each junction.
coverage_paths_case	paths to the BigWig files containing the coverage of your case samples.
coverage_paths_control	paths to the BigWig files for control samples.
coverage_chr_control	either "chr" or "no_chr", indicating the chromosome format of control coverage data. Only required if the chromosome format of the control BigWig files is different to that of your cases.
load_func	function used to load coverage.
binwidth	the number of bases to aggregate coverage across using <code>sum_func</code> when plotting.

## Value

ggplot displaying the splicing (and coverage) surrounding the transcript/region of interest.

## Examples

```
# use GenomicState to load txdb (GENCODE v31)
ref <- GenomicState::GenomicStateHub(
  version = "31",
  genome = "hg38",
  filetype = "TxDb"
)[[1]]

junctions_processed <- junction_process(
  junctions_example,
  ref,
  types = c("ambig_gene", "unannotated")
)

sashimi_plot <- plot_sashimi(
  junctions = junction_filter(junctions_processed),
  ref = ref,
  gene_tx_id = "ENSG00000142156.14",
  gene_tx_col = "gene_id",
  sum_func = NULL
)
```

---

ref_load	<i>Load reference annotation into a TxDb format</i>
----------	---

---

**Description**

ref\_load`` loads reference annotation using [makeTxDbFromGFF][GenomicFeatures::makeTxDbFromGFF] if a c unchanged if already a [TxDb-class](#). If you would

**Usage**

```
ref_load(ref)
```

**Arguments**

ref either path to gtf/gff3 or object of class [TxDb-class](#) or [EnsDb-class](#). [EnsDb-class](#) is required if you intend to annotate junctions with gene symbols/names.

**Value**

a [TxDb-class](#) object.

**Examples**

```
# create a TxDb,  
ref <- GenomicState::GenomicStateHub(  
  version = "31",  
  genome = "hg38",  
  filetype = "TxDb"  
)[[1]]
```

```
# alternatively ref can be a character specifying a path to a GTF file  
ref <- ref_load(ref)
```

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