

# Package ‘maser’

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**Type** Package

**Title** Mapping Alternative Splicing Events to pRoteins

**Version** 1.27.0

**Description** This package provides functionalities for downstream analysis, annotation and visualizaton of alternative splicing events generated by rMATS.

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---

annotation,Maser-method

*Retrieve annotation of splicing events from a maser object.*

---

## Description

Retrieve annotation of splicing events from a maser object.

## Usage

```
## S4 method for signature 'Maser'
annotation(object, type = c("A3SS", "A5SS", "SE", "RI",
                             "MXE"))
```

## Arguments

|        |                                                                                                   |
|--------|---------------------------------------------------------------------------------------------------|
| object | a maser object.                                                                                   |
| type   | a character indicating the splice type. Possible values are c("A3SS", "A5SS", "SE", "RI", "MXE"). |

## Value

a data.frame.

**Examples**

```
path <- system.file("extdata", file.path("MATS_output"), package = "maser")
hypoxia <- maser(path, c("Hypoxia 0h", "Hypoxia 24h"))
head(annotation(hypoxia, "SE"))
```

---

availableFeaturesUniprotKB

*Query available human protein features in UniprotKB.*

---

**Description**

Query available human protein features in UniprotKB.

**Usage**

```
availableFeaturesUniprotKB()
```

**Value**

a data.frame.

**Examples**

```
head(availableFeaturesUniprotKB(), 10)
```

---

boxplot\_PSI\_levels

*Boxplots of PSI distributions by splicing type.*

---

**Description**

Boxplots of PSI distributions by splicing type.

**Usage**

```
boxplot_PSI_levels(events, type = c("A3SS", "A5SS", "SE", "RI", "MXE"))
```

**Arguments**

|        |                                                                                            |
|--------|--------------------------------------------------------------------------------------------|
| events | a maser object.                                                                            |
| type   | character indicating splice type. Possible values are c("A3SS", "A5SS", "SE", "RI", "MXE") |

**Value**

a ggplot object.

**Examples**

```
path <- system.file("extdata", file.path("MATS_output"), package = "maser")
hypoxia <- maser(path, c("Hypoxia 0h", "Hypoxia 24h"))
hypoxia_filt <- filterByCoverage(hypoxia, avg_reads = 5)
boxplot_PSI_levels(hypoxia_filt, type = "RI")
```

---

|                     |                                                             |
|---------------------|-------------------------------------------------------------|
| counts,Maser-method | <i>Retrieve raw read counts values from a maser object.</i> |
|---------------------|-------------------------------------------------------------|

---

### Description

Retrieve raw read counts values from a maser object.

### Usage

```
## S4 method for signature 'Maser'
counts(object, type = c("A3SS", "A5SS", "SE", "RI", "MXE"))
```

### Arguments

|        |                                                                                                   |
|--------|---------------------------------------------------------------------------------------------------|
| object | a maser object.                                                                                   |
| type   | a character indicating the splice type. Possible values are c("A3SS", "A5SS", "SE", "RI", "MXE"). |

### Value

a matrix.

### Examples

```
path <- system.file("extdata", file.path("MATS_output"), package = "maser")
hypoxia <- maser(path, c("Hypoxia 0h", "Hypoxia 24h"))
head(counts(hypoxia, "SE"))
```

---

|         |                                                                                     |
|---------|-------------------------------------------------------------------------------------|
| display | <i>Visualization of splicing events annotation using an interactive data table.</i> |
|---------|-------------------------------------------------------------------------------------|

---

### Description

Visualization of splicing events annotation using an interactive data table.

### Usage

```
display(events, type = c("A3SS", "A5SS", "SE", "RI", "MXE"))
```

### Arguments

|        |                                                                                            |
|--------|--------------------------------------------------------------------------------------------|
| events | a maser object.                                                                            |
| type   | character indicating splice type. Possible values are c("A3SS", "A5SS", "SE", "RI", "MXE") |

### Value

a datatables object.

**Examples**

```
path <- system.file("extdata", file.path("MATS_output"), package = "maser")
hypoxia <- maser(path, c("Hypoxia 0h", "Hypoxia 24h"))
hypoxia_filt <- filterByCoverage(hypoxia, avg_reads = 5)
hypoxia_top <- topEvents(hypoxia_filt)
display(hypoxia_top, type = "SE")
```

dotplot

*Dotplot representation of splicing events.***Description**

Dotplot representation of splicing events.

**Usage**

```
dotplot(events, type = c("A3SS", "A5SS", "SE", "RI", "MXE"), fdr = 0.05,
  deltaPSI = 0.1)
```

**Arguments**

|          |                                                                                            |
|----------|--------------------------------------------------------------------------------------------|
| events   | a maser object.                                                                            |
| type     | character indicating splice type. Possible values are c("A3SS", "A5SS", "SE", "RI", "MXE") |
| fdr      | numeric, FDR (False Discovery Rate) cutoff.                                                |
| deltaPSI | numeric, absolute minimum PSI (Percent spliced-in) change                                  |

**Value**

a ggplot object.

**Examples**

```
path <- system.file("extdata", file.path("MATS_output"), package = "maser")
hypoxia <- maser(path, c("Hypoxia 0h", "Hypoxia 24h"))
hypoxia_filt <- filterByCoverage(hypoxia, avg_reads = 5)
dotplot(hypoxia_filt, type = "SE")
```

filterByCoverage

*Filter splicing events based on coverage.***Description**

Filter splicing events based on coverage.

**Usage**

```
filterByCoverage(events, avg_reads = 5)
```

**Arguments**

events            a maser object.

avg\_reads        numeric, average number of reads covering the splice event.

**Value**

a maser object.

**Examples**

```
path <- system.file("extdata", file.path("MATS_output"), package = "maser")
hypoxia <- maser(path, c("Hypoxia 0h", "Hypoxia 24h"))
hypoxia_filt <- filterByCoverage(hypoxia, avg_reads = 5)
```

---

|                 |                                                                   |
|-----------------|-------------------------------------------------------------------|
| filterByEventId | <i>Filter splicing events based on event identifier and type.</i> |
|-----------------|-------------------------------------------------------------------|

---

**Description**

Filter splicing events based on event identifier and type.

**Usage**

```
filterByEventId(events, event_id, type = c("A3SS", "A5SS", "SE", "RI", "MXE"))
```

**Arguments**

events            a maser object.

event\_id          numeric vector of event identifiers.

type              character indicating splice type. Possible values are c("A3SS", "A5SS", "SE", "RI", "MXE").

**Value**

a maser object.

**Examples**

```
path <- system.file("extdata", file.path("MATS_output"), package = "maser")
hypoxia <- maser(path, c("Hypoxia 0h", "Hypoxia 24h"))
filterByEventId(hypoxia, 33208, "SE")
```

---

|            |                                                   |
|------------|---------------------------------------------------|
| geneEvents | <i>Retrieve splicing events for a given gene.</i> |
|------------|---------------------------------------------------|

---

**Description**

Retrieve splicing events for a given gene.

**Usage**

```
geneEvents(events, geneS, fdr = 0.05, deltaPSI = 0.1)
```

**Arguments**

|          |                                         |
|----------|-----------------------------------------|
| events   | a maser object.                         |
| geneS    | a character indicating the gene symbol. |
| fdr      | numeric, FDR cutoff.                    |
| deltaPSI | numeric, minimum PSI change.            |

**Value**

a maser object.

**Examples**

```
path <- system.file("extdata", file.path("MATS_output"), package = "maser")
hypoxia <- maser(path, c("Hypoxia 0h", "Hypoxia 24h"))
hypoxia_mib2 <- geneEvents(hypoxia, "MIB2")
```

---

|                       |                                                                        |
|-----------------------|------------------------------------------------------------------------|
| granges, Maser-method | <i>Retrieve genomic ranges of splicing events from a maser object.</i> |
|-----------------------|------------------------------------------------------------------------|

---

**Description**

Retrieve genomic ranges of splicing events from a maser object.

**Usage**

```
## S4 method for signature 'Maser'
granges(x, type = c("A3SS", "A5SS", "SE", "RI", "MXE"), ...)
```

**Arguments**

|      |                                                                                                   |
|------|---------------------------------------------------------------------------------------------------|
| x    | a maser object.                                                                                   |
| type | a character indicating the splice type. Possible values are c("A3SS", "A5SS", "SE", "RI", "MXE"). |
| ...  | additional arguments.                                                                             |

**Value**

a GRangesList.

**Examples**

```
path <- system.file("extdata", file.path("MATS_output"), package = "maser")
hypoxia <- maser(path, c("Hypoxia 0h", "Hypoxia 24h"))
head(granges(hypoxia, type = "SE"))
```

---

mapProteinFeaturesToEvents

*Mapping of splice events to UniprotKB protein features.*


---

**Description**

Mapping of splice events to UniprotKB protein features.

**Usage**

```
mapProteinFeaturesToEvents(events, tracks, by = c("feature", "category"),
  ncores = 1)
```

**Arguments**

|        |                                                                                                                                      |
|--------|--------------------------------------------------------------------------------------------------------------------------------------|
| events | a maser object with transcript and protein identifiers.                                                                              |
| tracks | a character vector indicating valid UniprotKB features or categories.                                                                |
| by     | a character vector, possible values are c("feature", "category").                                                                    |
| ncores | number of cores for multithreading (available only in OSX and Linux machines).<br>If Windows, ncores will be set to 1 automatically. |

**Details**

This function performs mapping of splicing events to protein features available in the UniprotKB database. Annotation tracks of protein features mapped to the hg38 build of the human genome are retrieved from the public UniprotKB FTP. The function will overlap exons involved in the splice event with the feature genomic coordinates retrieved from UniprotKB.

Annotation can be executed either by feature or category. If categories are provided, all features within the category group will be included for annotation.

Thus, batch annotation is enabled either by using by = category or by providing multiple features in the tracks argument.

Visualization of protein features can be done using [plotUniprotKBFeatures](#).

**Value**

a maser object with protein feature annotation.

**See Also**

[plotUniprotKBFeatures](#)

**Examples**

```
## Create the maser object
path <- system.file("extdata", file.path("MATS_output"), package = "maser")
hypoxia <- maser(path, c("Hypoxia 0h", "Hypoxia 24h"))
hypoxia_filt <- filterByCoverage(hypoxia, avg_reads = 5)

## Ensembl GTF annotation for SRSF6
gtf_path <- system.file("extdata", file.path("GTF", "SRSF6_Ensembl85.gtf"),
  package = "maser")
ens_gtf <- rtracklayer::import.gff(gtf_path)

## Retrieve gene specific splice events
srsf6_events <- geneEvents(hypoxia_filt, geneS = "SRSF6")

## Map splicing events to transcripts
srsf6_mapped <- mapTranscriptsToEvents(srsf6_events, ens_gtf)

## Annotate splice events with protein domains
srsf6_annot <- mapProteinFeaturesToEvents(srsf6_mapped, tracks = "domain")
head(annotation(srsf6_annot, "SE"))
```

---

mapTranscriptsToEvents

*Mapping of splice events to Ensembl transcripts.*


---

**Description**

Mapping of splice events to Ensembl transcripts.

**Usage**

```
mapTranscriptsToEvents(events, gtf, ncores = 1)
```

**Arguments**

|        |                                                                                                                                   |
|--------|-----------------------------------------------------------------------------------------------------------------------------------|
| events | a maser object.                                                                                                                   |
| gtf    | a GRanges object obtained from an Ensembl or Gencode GTF file using the hg38 build of the human genome.                           |
| ncores | number of cores for multithreading (available only in OSX and Linux machines). If Windows, ncores will be set to 1 automatically. |

**Details**

This function performs mapping of splice events in the maser object to Ensembl transcripts by overlapping exons involved in the splice event to the transcript models provided in the GTF.

Each type of splice event requires a specific mapping procedure (described below).

The mapping will also add Uniprot identifiers when the ENST transcript encodes for a protein.

Visualization of affected transcripts can be done using [plotTranscripts](#).

**Exon skipping**

**Inclusion transcript(s)** Transcript(s) overlapping the cassette exon, as well both flanking exons (i.e upstream and downstream exons).

**Skipping transcript(s)** Transcript(s) overlapping both flanking exons but not the cassette exon.

### **Intron retention**

**Retention transcript(s)** Transcript(s) overlapping exactly the retained intron.

**Skipping transcript(s)** Transcript(s) where intron is spliced out and overlapping both flanking exons.

### **Mutually exclusive exons**

**Exon1 transcript(s)** Transcript(s) overlapping the first exon and both flanking exons.

**Exon2 transcript(s)** Transcript(s) overlapping the second exon and both flanking exons.

### **Alternative 3' and 5' splice sites**

**Short exon transcript(s)** Transcript(s) overlapping both short and downstream exons.

**Long exon transcript(s)** Transcript(s) overlapping both long and downstream exons.

### **Value**

a maser object with transcript and protein identifiers.

### **See Also**

[plotTranscripts](#)

### **Examples**

```
## Create the maser object
path <- system.file("extdata", file.path("MATS_output"), package = "maser")
hypoxia <- maser(path, c("Hypoxia 0h", "Hypoxia 24h"))
hypoxia_filt <- filterByCoverage(hypoxia, avg_reads = 5)

## Ensembl GTF annotation for SRSF6
gtf_path <- system.file("extdata", file.path("GTF",
  "Ensembl85_examples.gtf.gz"), package = "maser")
ens_gtf <- rtracklayer::import.gff(gtf_path)

## Retrieve gene specific splice events
srsf6_events <- geneEvents(hypoxia_filt, geneS = "SRSF6")

## Map splicing events to transcripts
srsf6_mapped <- mapTranscriptsToEvents(srsf6_events, ens_gtf)
head(annotation(srsf6_mapped, "SE"))
```

---

|       |                                                                  |
|-------|------------------------------------------------------------------|
| maser | <i>Create a maser object by importing rMATS splicing events.</i> |
|-------|------------------------------------------------------------------|

---

## Description

Create a maser object by importing rMATS splicing events.

## Usage

```
maser(path, cond_labels, ftype = c("ReadsOnTargetAndJunctionCounts",
  "JunctionCountOnly", "JCEC", "JC"))
```

## Arguments

|             |                                                                                                                    |
|-------------|--------------------------------------------------------------------------------------------------------------------|
| path        | a character specifying the folder containing rMATS output files.                                                   |
| cond_labels | a character vector of length 2 describing labels for experimental conditions.                                      |
| ftype       | a character indicating the rMATS file type. Possible values are c("ReadsOnTargetAndJunctionCounts", "JCEC", "JC"). |

## Details

This function creates a maser object by importing rMATS output. ftype indicates which rMATS files to import. ReadsOnTargetandJunction or JunctionCountOnly are used in rMATS 3.2.5 or lower. Newer versions (>4.0.1) use "JCEC" or "JC" nomenclature.

## Value

A maser object.

## Examples

```
path <- system.file("extdata", file.path("MATS_output"), package = "maser")
hypoxia <- maser(path, c("Hypoxia 0h", "Hypoxia 24h"))
```

---

|             |                                                                   |
|-------------|-------------------------------------------------------------------|
| Maser-class | <i>S4 class to represent splicing events imported from rMATS.</i> |
|-------------|-------------------------------------------------------------------|

---

## Description

S4 class to represent splicing events imported from rMATS.

---

|     |                                                            |
|-----|------------------------------------------------------------|
| pca | <i>Prinicipal component analysis of PSI distributions.</i> |
|-----|------------------------------------------------------------|

---

**Description**

Prinicipal component analysis of PSI distributions.

**Usage**

```
pca(events, type = c("A3SS", "A5SS", "SE", "RI", "MXE"))
```

**Arguments**

|        |                                                                                            |
|--------|--------------------------------------------------------------------------------------------|
| events | a maser object.                                                                            |
| type   | character indicating splice type. Possible values are c("A3SS", "A5SS", "SE", "RI", "MXE") |

**Value**

a ggplot object.

**Examples**

```
path <- system.file("extdata", file.path("MATS_output"), package = "maser")
hypoxia <- maser(path, c("Hypoxia 0h", "Hypoxia 24h"))
hypoxia_filt <- filterByCoverage(hypoxia, avg_reads = 5)
pca(hypoxia_filt, type = "RI")
```

---

|             |                                                               |
|-------------|---------------------------------------------------------------|
| plotGenePSI | <i>Boxplots of Percent spliced-in levels for gene events.</i> |
|-------------|---------------------------------------------------------------|

---

**Description**

Boxplots of Percent spliced-in levels for gene events.

**Usage**

```
plotGenePSI(events, type = c("A3SS", "A5SS", "SE", "RI", "MXE"),
  show_replicates = TRUE)
```

**Arguments**

|                 |                                                                                            |
|-----------------|--------------------------------------------------------------------------------------------|
| events          | a maser object.                                                                            |
| type            | character indicating splice type. Possible values are c("A3SS", "A5SS", "SE", "RI", "MXE") |
| show_replicates | logical, add data points for individual replicates                                         |

**Value**

a ggplot object.

**Examples**

```
path <- system.file("extdata", file.path("MATS_output"), package = "maser")
hypoxia <- maser(path, c("Hypoxia 0h", "Hypoxia 24h"))
hypoxia_filt <- filterByCoverage(hypoxia, avg_reads = 5)
hypoxia_mib2 <- geneEvents(hypoxia_filt, geneS = "MIB2")
plotGenePSI(hypoxia_mib2, type = "SE", show_replicates = TRUE)
```

---

|                 |                                                                               |
|-----------------|-------------------------------------------------------------------------------|
| plotTranscripts | <i>Mapping and visualization of Ensembl transcripts affected by splicing.</i> |
|-----------------|-------------------------------------------------------------------------------|

---

**Description**

Mapping and visualization of Ensembl transcripts affected by splicing.

**Usage**

```
plotTranscripts(events, type = c("A3SS", "A5SS", "SE", "RI", "MXE"), event_id,
  gtf, zoom = FALSE, show_PSI = TRUE)
```

**Arguments**

|          |                                                                                             |
|----------|---------------------------------------------------------------------------------------------|
| events   | a maser object.                                                                             |
| type     | character indicating splice type. Possible values are c("A3SS", "A5SS", "SE", "RI", "MXE"). |
| event_id | numeric, event identifier.                                                                  |
| gtf      | a GRanges, Ensembl or Gencode GTF using the hg38 build of the human genome.                 |
| zoom     | logical, zoom to the genomic coordinates of the splice event.                               |
| show_PSI | logical, display the PSI track.                                                             |

**Details**

This is a wrapper function for performing both mapping and visualization of Ensembl transcripts that are compatible with the splice event. This function calls [mapTranscriptsToEvents](#) for transcript mapping, which in turn uses `findOverlaps` for transcript overlapping. The [GViz](#) package is used for creating annotation tracks for genomic visualization of splicing events.

Each type of splice event requires a specific overlapping rule (described below), #’ and a customized Gviz plot is created for each splicing type.

**Exon skipping**

**Inclusion track** Transcript(s) overlapping the cassette exon, as well both flanking exons (i.e upstream and downstream exons).

**Skipping track** Transcript(s) overlapping both flanking exons but not the cassette exon.

**Intron retention**

**Retention track** Transcript(s) overlapping exactly the retained intron.

**Skipping track** Transcript(s) where intron is spliced out and overlapping both flanking exons.

**Mutually exclusive exons**

**Exon1 track** Transcript(s) overlapping the first exon and both flanking exons.

**Exon2 track** Transcript(s) overlapping the second exon and both flanking exons.

**Alternative 3' and 5' splice sites**

**Short exon track** Transcript(s) overlapping both short and downstream exons.

**Long exon track** Transcript(s) overlapping both long and downstream exons.

### Value

a Gviz object.

### See Also

[mapTranscriptsToEvents](#)

### Examples

```
## Create the maser object
path <- system.file("extdata", file.path("MATS_output"), package = "maser")
hypoxia <- maser(path, c("Hypoxia 0h", "Hypoxia 24h"))
hypoxia_filt <- filterByCoverage(hypoxia, avg_reads = 5)

## Ensembl GTF annotation for SRSF6
gtf_path <- system.file("extdata", file.path("GTF",
  "SRSF6_Ensembl85.gtf"), package = "maser")
ens_gtf <- rtracklayer::import.gff(gtf_path)

## Retrieve gene specific splicing events
srsf6_events <- geneEvents(hypoxia_filt, geneS = "SRSF6")

## Plot exon skipping event
plotTranscripts(srsf6_events, type = "SE", event_id = 33209, gtf = ens_gtf)
```

---

plotUniprotKBFeatures *Mapping and visualization of UniprotKB protein features affected by splicing.*

---

### Description

Mapping and visualization of UniprotKB protein features affected by splicing.

### Usage

```
plotUniprotKBFeatures(events, type = c("A3SS", "A5SS", "SE", "RI", "MXE"),
  event_id, gtf, features, zoom = FALSE, show_transcripts = FALSE,
  show_PSI = TRUE, ncores = 1)
```

**Arguments**

|                  |                                                                                                                                   |
|------------------|-----------------------------------------------------------------------------------------------------------------------------------|
| events           | a maser object.                                                                                                                   |
| type             | character indicating splice type. Possible values are c("A3SS", "A5SS", "SE", "RI", "MXE").                                       |
| event_id         | numeric, event identifier.                                                                                                        |
| gtf              | a GRanges, Ensembl or Gencode GTF using the hg38 build of the human genome.                                                       |
| features         | a character vector indicating valid UniprotKB features.                                                                           |
| zoom             | logical, zoom to the genomic coordinates of the splice event.                                                                     |
| show_transcripts | logical, display transcripts track.                                                                                               |
| show_PSI         | logical, display the PSI track.                                                                                                   |
| ncores           | number of cores for multithreading (available only in OSX and Linux machines). If Windows, ncores will be set to 1 automatically. |

**Details**

This is a wrapper function for performing both mapping and visualization of protein features affected by the splice event. This function calls [mapProteinFeaturesToEvents](#) for mapping of protein features to splicing events.

The [GViz](#) package is used for creating annotation tracks for genomic visualization.

Multiple protein annotation tracks can be created using the features argument.

**Value**

a Gviz object.

**See Also**

[mapProteinFeaturesToEvents](#)

**Examples**

```
## Create the maser object
path <- system.file("extdata", file.path("MATS_output"), package = "maser")
hypoxia <- maser(path, c("Hypoxia 0h", "Hypoxia 24h"))
hypoxia_filt <- filterByCoverage(hypoxia, avg_reads = 5)

## Ensembl GTF annotation for SRSF6
gtf_path <- system.file("extdata", file.path("GTF",
  "SRSF6_Ensembl85.gtf"), package = "maser")
ens_gtf <- rtracklayer::import.gff(gtf_path)

## Retrieve gene specific splicing events
srsf6_events <- geneEvents(hypoxia_filt, geneS = "SRSF6")

## Map splicing events to transcripts
srsf6_mapped <- mapTranscriptsToEvents(srsf6_events, ens_gtf)

## Plot splice event, transcripts and protein features
plotUniprotKBFeatures(srsf6_mapped, "SE", event_id = 33209, gtf = ens_gtf,
  features = c("domain"), show_transcripts = TRUE)
```

PSI

*Retrieve PSI (percent spliced in) values from a maser object.***Description**

Retrieve PSI (percent spliced in) values from a maser object.

**Usage**

```
PSI(events, type)
```

**Arguments**

**events** a maser object.

**type** a character indicating the splice type. Possible values are c("A3SS", "A5SS", "SE", "RI", "MXE").

**Value**

a matrix.

**Examples**

```
path <- system.file("extdata", file.path("MATS_output"), package = "maser")
hypoxia <- maser(path, c("Hypoxia 0h", "Hypoxia 24h"))
head(PSI(hypoxia, "SE"))
```

PSI,Maser,character-method

*Retrieve PSI (percent spliced in) values from a maser object.***Description**

Retrieve PSI (percent spliced in) values from a maser object.

**Usage**

```
## S4 method for signature 'Maser,character'
PSI(events, type = c("A3SS", "A5SS", "SE", "RI",
  "MXE"))
```

**Arguments**

**events** a maser object.

**type** a character indicating the splice type. Possible values are c("A3SS", "A5SS", "SE", "RI", "MXE").

**Value**

a matrix.

**Examples**

```
path <- system.file("extdata", file.path("MATS_output"), package = "maser")
hypoxia <- maser(path, c("Hypoxia 0h", "Hypoxia 24h"))
head(PSI(hypoxia, "SE"))
```

---

splicingDistribution    *Proportion of events by splicing type.*

---

**Description**

Proportion of events by splicing type.

**Usage**

```
splicingDistribution(events, fdr = 0.05, deltaPSI = 0.1)
```

**Arguments**

|          |                                                           |
|----------|-----------------------------------------------------------|
| events   | a maser object.                                           |
| fdr      | numeric, FDR (False Discovery Rate) cutoff.               |
| deltaPSI | numeric, absolute minimum PSI (Percent spliced-in) change |

**Value**

a ggplot object.

**Examples**

```
path <- system.file("extdata", file.path("MATS_output"), package = "maser")
hypoxia <- maser(path, c("Hypoxia 0h", "Hypoxia 24h"))
hypoxia_filt <- filterByCoverage(hypoxia, avg_reads = 5)
splicingDistribution(hypoxia_filt)
```

---

summary,Maser-method    *Retrieve rMATS stats of differential splicing from a maser object.*

---

**Description**

Retrieve rMATS stats of differential splicing from a maser object.

**Usage**

```
## S4 method for signature 'Maser'
summary(object, type = c("A3SS", "A5SS", "SE", "RI", "MXE"))
```

**Arguments**

|        |                                                                                                   |
|--------|---------------------------------------------------------------------------------------------------|
| object | a maser object.                                                                                   |
| type   | a character indicating the splice type. Possible values are c("A3SS", "A5SS", "SE", "RI", "MXE"). |

**Value**

a data.frame.

**Examples**

```
path <- system.file("extdata", file.path("MATS_output"), package = "maser")
hypoxia <- maser(path, c("Hypoxia 0h", "Hypoxia 24h"))
head(summary(hypoxia, "SE"))
```

---

|           |                                                                             |
|-----------|-----------------------------------------------------------------------------|
| topEvents | <i>Filter splicing events based on false discovery rate and PSI change.</i> |
|-----------|-----------------------------------------------------------------------------|

---

**Description**

Filter splicing events based on false discovery rate and PSI change.

**Usage**

```
topEvents(events, fdr = 0.05, deltaPSI = 0.1)
```

**Arguments**

|          |                                                           |
|----------|-----------------------------------------------------------|
| events   | a maser object.                                           |
| fdr      | numeric, FDR (False Discovery Rate) cutoff.               |
| deltaPSI | numeric, absolute minimum PSI (Percent spliced-in) change |

**Value**

a maser object.

**Examples**

```
path <- system.file("extdata", file.path("MATS_output"), package = "maser")
hypoxia <- maser(path, c("Hypoxia 0h", "Hypoxia 24h"))

## To select all events with minimum 10% change in PSI, and FDR < 0.01
hypoxia_top <- topEvents(hypoxia, fdr = 0.01, deltaPSI = 0.1)
```

---

|         |                                         |
|---------|-----------------------------------------|
| volcano | <i>Volcano plot of splicing events.</i> |
|---------|-----------------------------------------|

---

**Description**

Volcano plot of splicing events.

**Usage**

```
volcano(events, type = c("A3SS", "A5SS", "SE", "RI", "MXE"), fdr = 0.05,  
        deltaPSI = 0.1)
```

**Arguments**

|          |                                                                                            |
|----------|--------------------------------------------------------------------------------------------|
| events   | a maser object.                                                                            |
| type     | character indicating splice type. Possible values are c("A3SS", "A5SS", "SE", "RI", "MXE") |
| fdr      | numeric, FDR (False Discovery Rate) cutoff.                                                |
| deltaPSI | numeric, absolute minimum PSI (Percent spliced-in) change                                  |

**Value**

a ggplot object.

**Examples**

```
path <- system.file("extdata", file.path("MATS_output"), package = "maser")  
hypoxia <- maser(path, c("Hypoxia 0h", "Hypoxia 24h"))  
hypoxia_filt <- filterByCoverage(hypoxia, avg_reads = 5)  
volcano(hypoxia_filt, type = "SE")
```

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