

# Package: maser (via r-universe)

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**Type** Package

**Title** Mapping Alternative Splicing Events to pRoteins

**Version** 1.23.0

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

**Depends** R (>= 3.5.0), ggplot2, GenomicRanges

**License** MIT + file LICENSE

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**VignetteBuilder** knitr

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**BugReports** <https://github.com/DiogoVeiga/maser/issues>

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**Repository** <https://bioc.r-universe.dev>

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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     *Retrieve raw read counts values from a maser object.*

---

**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            *Visualization of splicing events annotation using an interactive data table.*

---

**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      *Filter splicing events based on event identifier and type.*

---

**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

*Retrieve splicing events for a given gene.*

---

**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 *Retrieve genomic ranges of splicing events from a maser object.*

---

### 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). 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

*Create a maser object by importing rMATS splicing events.*

---

**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. `f` type 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>Principial component analysis of PSI distributions.</i>
-----	------------------------------------------------------------

---

**Description**

Principial component analysis of PSI distributions.

**Usage**

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

**Arguments**

<code>events</code>	a maser object.
<code>type</code>	character indicating splice type. Possible values are <code>c("A3SS", "A5SS", "SE", "RI", "MXE")</code>

**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                      *Boxplots of Percent spliced-in levels for gene events.*

---

**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                *Mapping and visualization of Ensembl transcripts affected by splicing.*

---

**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 *Volcano plot of splicing events.*

---

**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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