

# Package ‘consensusSeeker’

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**Version** 1.33.0

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**Title** Detection of consensus regions inside a group of experiences using genomic positions and genomic ranges

**Description** This package compares genomic positions and genomic ranges from multiple experiments to extract common regions. The size of the analyzed region is adjustable as well as the number of experiences in which a feature must be present in a potential region to tag this region as a consensus region. In genomic analysis where feature identification generates a position value surrounded by a genomic range, such as ChIP-Seq peaks and nucleosome positions, the replication of an experiment may result in slight differences between predicted values. This package enables the conciliation of the results into consensus regions.

**Depends** R (>= 3.5.0), BiocGenerics, IRanges, GenomicRanges, BiocParallel

**Imports** GenomeInfoDb, rtracklayer, stringr, S4Vectors, methods

**Suggests** BiocStyle, ggplot2, knitr, rmarkdown, RUnit

**License** Artistic-2.0

**Encoding** UTF-8

**URL** <https://github.com/adeschen/consensusSeeker>

**BugReports** <https://github.com/adeschen/consensusSeeker/issues>

**VignetteBuilder** knitr

**NeedsCompilation** no

**biocViews** BiologicalQuestion, ChIPSeq, Genetics, MultipleComparison, Transcription, PeakDetection, Sequencing, Coverage

**RoxygenNote** 7.2.3

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consensusSeeker-package

*consensusSeeker: Detection of consensus peak regions inside a group of experiments using narrowPeak files*

---

## Description

This package compares positions and ranges data from multiple experiments to extract common consensus regions. The size of the analyzed region is adjustable as well as the number of experiments in which a peak must be detected to mark a potential region as a consensus peak region.

## Author(s)

Astrid Deschênes, Fabien Claude Lamaze, Pascal Belleau and Arnaud Droit

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## See Also

- [readNarrowPeakFile](#) for extracting regions and peaks from a narrowPeak file.
- [findConsensusPeakRegions](#) for extracting regions sharing the same features in more than one experiment.

---

A549\_CTCF\_MYJ\_NarrowPeaks\_partial

*Sites with the greatest evidence of transcription factor binding for the CTCF transcription factor (for demonstration purpose)*

---

## Description

Sites representing the greatest evidence of enrichment for the CTCF transcription factor (DCC accession: ENCFF000MYJ) for regions chr1:246000000-249250621 and chr10:100000000-125000000 from the Encyclopedia of DNA Elements (ENCODE) data (Dunham I et al. 2012).

## Usage

```
data(A549_CTCF_MYJ_NarrowPeaks_partial)
```

## Format

A GRanges containing one entry per site.

## Source

The Encyclopedia of DNA Elements (ENCODE) (DCC accession: ENCFF000MYJ)

## References

- Dunham I, Kundaje A, Aldred SF, et al. An integrated encyclopedia of DNA elements in the human genome. *Nature*. 2012 Sep 6;489(7414):57-74.

## See Also

- [A549\\_CTCF\\_MYJ\\_Peaks\\_partial](#) the associate genomic peaks dataset.
- [findConsensusPeakRegions](#) for extracting regions sharing the same features in more than one experiment.

A549\_CTCF\_MYJ\_NarrowPeaks\_partial

## Examples

```
## Loading datasets
data(A549_CTCF_MYJ_NarrowPeaks_partial)
data(A549_CTCF_MYJ_Peaks_partial)
data(A549_CTCF_MYN_NarrowPeaks_partial)
data(A549_CTCF_MYN_Peaks_partial)

## Assigning experiment name to each row of the dataset.
## NarrowPeak and Peak datasets from the same experiment must
## have identical names.
names(A549_CTCF_MYJ_Peaks_partial) <- rep("CTCF_MYJ",
                                          length(A549_CTCF_MYJ_Peaks_partial))
names(A549_CTCF_MYJ_NarrowPeaks_partial) <- rep("CTCF_MYJ",
                                                  length(A549_CTCF_MYJ_NarrowPeaks_partial))
names(A549_CTCF_MYN_Peaks_partial) <- rep("CTCF_MYN",
                                           length(A549_CTCF_MYN_Peaks_partial))
names(A549_CTCF_MYN_NarrowPeaks_partial) <- rep("CTCF_MYN",
                                                  length(A549_CTCF_MYN_NarrowPeaks_partial))

## Calculating consensus regions for chromosome 10
## with a default region size of 100 bp (2 * extendingSize)
## which is extended to include all genomic regions for the closest
## peak to the median position of all peaks included in the region (for each
## experiment).
## A peak from both experiments must be present in a region to
## be retained as a consensus region.
chrList <- Seqinfo(c("chr10"), c(135534747), NA)
findConsensusPeakRegions(
  narrowPeaks = c(A549_CTCF_MYJ_NarrowPeaks_partial,
                  A549_CTCF_MYN_NarrowPeaks_partial),
  peaks = c(A549_CTCF_MYJ_Peaks_partial,
            A549_CTCF_MYN_Peaks_partial),
  chrInfo = chrList,
  extendingSize = 50,
  expandToFitPeakRegion = TRUE,
  shrinkToFitPeakRegion = TRUE,
  minNbrExp = 2,
  nbrThreads = 1)
```

---

A549\_CTCF\_MYJ\_Peaks\_partial

*Sites with the greatest evidence of transcription factor binding for the CTCF transcription factor (for demonstration purpose)*

---

## Description

Sites representing the greatest evidence of enrichment for the CTCF transcription factor (DCC accession: ENCFF000MYJ) for regions chr1:246000000-249250621 and chr10:100000000-125000000 from the Encyclopedia of DNA Elements (ENCODE) data (Dunham I et al. 2012).

## Usage

```
data(A549_CTCF_MYJ_Peaks_partial)
```

## Format

A GRanges containing one entry per site.

## Source

The Encyclopedia of DNA Elements (ENCODE) (DCC accession: ENCFF000MYJ)

## References

- Dunham I, Kundaje A, Aldred SF, et al. An integrated encyclopedia of DNA elements in the human genome. *Nature*. 2012 Sep 6;489(7414):57-74.

## See Also

- [A549\\_CTCF\\_MYJ\\_NarrowPeaks\\_partial](#) the associate genomic regions dataset.
- [findConsensusPeakRegions](#) for extracting regions sharing the same features in more than one experiment.

## Examples

```
## Loading datasets
data(A549_CTCF_MYJ_NarrowPeaks_partial)
data(A549_CTCF_MYJ_Peaks_partial)
data(A549_CTCF_MYN_NarrowPeaks_partial)
data(A549_CTCF_MYN_Peaks_partial)

## Assigning experiment name to each row of the dataset.
## NarrowPeak and Peak datasets from the same experiment must
## have identical names.
names(A549_CTCF_MYJ_Peaks_partial) <- rep("CTCF_MYJ",
                                          length(A549_CTCF_MYJ_Peaks_partial))
names(A549_CTCF_MYJ_NarrowPeaks_partial) <- rep("CTCF_MYJ",
```

```

                                length(A549_CTCF_MYJ_NarrowPeaks_partial))
names(A549_CTCF_MYN_Peaks_partial) <- rep("CTCF_MYN",
                                length(A549_CTCF_MYN_Peaks_partial))
names(A549_CTCF_MYN_NarrowPeaks_partial) <- rep("CTCF_MYN",
                                length(A549_CTCF_MYN_NarrowPeaks_partial))

## Calculating consensus regions for chromosome 10
## with a default region size of 40 bp (2 * extendingSize)
## which is extended to include all genomic regions for the closest
## peak to the median position of all peaks included in the region (for each
## experiment).
## A peak from both experiments must be present in a region to
## be retained as a consensus region.
chrList <- Seqinfo(c("chr10"), c(135534747), NA)
findConsensusPeakRegions(
  narrowPeaks = c(A549_CTCF_MYJ_NarrowPeaks_partial,
                  A549_CTCF_MYN_NarrowPeaks_partial),
  peaks = c(A549_CTCF_MYJ_Peaks_partial,
            A549_CTCF_MYN_Peaks_partial),

  chrInfo = chrList,
  extendingSize = 20,
  expandToFitPeakRegion = FALSE,
  shrinkToFitPeakRegion = FALSE,
  minNbrExp = 2,
  nbrThreads = 1)

```

---

A549\_CTCF\_MYN\_NarrowPeaks\_partial

*Sites with the greatest evidence of transcription factor binding for the  
CTCF transcription factor (for demonstration purpose)*

---

## Description

Sites representing the greatest evidence of enrichment for the CTCF transcription factor (DCC accession: ENCFF000MYN) for regions chr1:246000000-249250621 and chr10:100000000-125000000 from the Encyclopedia of DNA Elements (ENCODE) data (Dunham I et al. 2012).

## Usage

```
data(A549_CTCF_MYN_NarrowPeaks_partial)
```

## Format

A GRanges containing one entry per site.

## Source

The Encyclopedia of DNA Elements (ENCODE) (DCC accession: ENCFF000MYN)

## References

- Dunham I, Kundaje A, Aldred SF, et al. An integrated encyclopedia of DNA elements in the human genome. Nature. 2012 Sep 6;489(7414):57-74.

## See Also

- [A549\\_CTCF\\_MYN\\_Peaks\\_partial](#) the associate genomic regions dataset.
- [findConsensusPeakRegions](#) for extracting regions sharing the same features in more than one experiment.

## Examples

```
## Loading datasets
data(A549_CTCF_MYJ_NarrowPeaks_partial)
data(A549_CTCF_MYJ_Peaks_partial)
data(A549_CTCF_MYN_NarrowPeaks_partial)
data(A549_CTCF_MYN_Peaks_partial)

## Assigning experiment name to each row of the dataset.
## NarrowPeak and Peak datasets from the same experiment must
## have identical names.
names(A549_CTCF_MYJ_Peaks_partial) <- rep("CTCF_MYJ",
                                          length(A549_CTCF_MYJ_Peaks_partial))
names(A549_CTCF_MYJ_NarrowPeaks_partial) <- rep("CTCF_MYJ",
                                                  length(A549_CTCF_MYJ_NarrowPeaks_partial))
names(A549_CTCF_MYN_Peaks_partial) <- rep("CTCF_MYN",
                                           length(A549_CTCF_MYN_Peaks_partial))
names(A549_CTCF_MYN_NarrowPeaks_partial) <- rep("CTCF_MYN",
                                                  length(A549_CTCF_MYN_NarrowPeaks_partial))

## Calculating consensus regions for chromosome 10
## with a default region size of 40 bp (2 * extendingSize)
## which is extended to include all genomic regions for the closest
## peak to the median position of all peaks included in the region (for each
## experiment).
## A peak from both experiments must be present in a region to
## be retained as a consensus region.
chrList <- Seqinfo("chr10", 135534747, NA)
findConsensusPeakRegions(
  narrowPeaks = c(A549_CTCF_MYJ_NarrowPeaks_partial,
                  A549_CTCF_MYN_NarrowPeaks_partial),
  peaks = c(A549_CTCF_MYJ_Peaks_partial,
            A549_CTCF_MYN_Peaks_partial),
  chrInfo = chrList,
  extendingSize = 20,
  expandToFitPeakRegion = FALSE,
  shrinkToFitPeakRegion = FALSE,
  minNbrExp = 2,
  nbrThreads = 1)
```

---

A549\_CTCF\_MYN\_Peaks\_partial

*Sites with the greatest evidence of transcription factor binding for the CTCF transcription factor (for demonstration purpose)*

---

## Description

Sites representing the greatest evidence of enrichment for the CTCF transcription factor (DCC accession: ENCFF000MYN) for regions chr1:246000000-249250621 and chr10:100000000-125000000 from the Encyclopedia of DNA Elements (ENCODE) data (Dunham I et al. 2012).

## Usage

```
data(A549_CTCF_MYN_Peaks_partial)
```

## Format

A GRanges containing one entry per site.

## Source

The Encyclopedia of DNA Elements (ENCODE) (DCC accession: ENCFF000MYN)

## References

- Dunham I, Kundaje A, Aldred SF, et al. An integrated encyclopedia of DNA elements in the human genome. *Nature*. 2012 Sep 6;489(7414):57-74.

## See Also

- [A549\\_CTCF\\_MYN\\_NarrowPeaks\\_partial](#) the associate genomic regions dataset.
- [findConsensusPeakRegions](#) for extracting regions sharing the same features in more than one experiment.

## Examples

```
## Loading datasets
data(A549_CTCF_MYJ_NarrowPeaks_partial)
data(A549_CTCF_MYJ_Peaks_partial)
data(A549_CTCF_MYN_NarrowPeaks_partial)
data(A549_CTCF_MYN_Peaks_partial)

## Assigning experiment name to each row of the dataset.
## NarrowPeak and Peak datasets from the same experiment must
## have identical names.
names(A549_CTCF_MYJ_Peaks_partial) <- rep("CTCF_MYJ",
                                          length(A549_CTCF_MYJ_Peaks_partial))
names(A549_CTCF_MYJ_NarrowPeaks_partial) <- rep("CTCF_MYJ",
```



```

                                length(A549_CTCF_MYJ_NarrowPeaks_partial))
names(A549_CTCF_MYN_Peaks_partial) <- rep("CTCF_MYN",
                                length(A549_CTCF_MYN_Peaks_partial))
names(A549_CTCF_MYN_NarrowPeaks_partial) <- rep("CTCF_MYN",
                                length(A549_CTCF_MYN_NarrowPeaks_partial))

## Calculating consensus regions for chromosomes 1
## with a default region size of 40 bp (2 * extendingSize)
## which is extended to include all genomic regions for the closest
## peak to the median position of all peaks included in the region (for each
## experiment).
## A peak from both experiments must be present in a region to
## be retained as a consensus region.
chrList <- Seqinfo(c("chr1"), c(249250621), NA)
findConsensusPeakRegions(
  narrowPeaks = c(A549_CTCF_MYJ_NarrowPeaks_partial,
                  A549_CTCF_MYN_NarrowPeaks_partial),
  peaks = c(A549_CTCF_MYJ_Peaks_partial,
            A549_CTCF_MYN_Peaks_partial),

  chrInfo = chrList,
  extendingSize = 20,
  expandToFitPeakRegion = FALSE,
  shrinkToFitPeakRegion = FALSE,
  minNbrExp = 2,
  nbrThreads = 1)

```

---

A549\_FOSL2\_01\_NarrowPeaks\_partial

*Genomic regions with the greatest evidence of transcription factor binding for the FOSL2 transcription factor (for demonstration purpose)*

---

## Description

Genomic regions representing the greatest evidence of enrichment for the FOSL2 transcription factor (DCC accession: ENCF000MZT) for regions chr1:249120200-249250621 and chr10:1-370100 from the Encyclopedia of DNA Elements (ENCODE) data (Dunham I et al. 2012).

## Usage

```
data(A549_FOSL2_01_NarrowPeaks_partial)
```

## Format

A GRanges containing one entry per genomic regions. Each row of GRanges has a name which represent the name of the experiment.

**Source**

The Encyclopedia of DNA Elements (ENCODE) (DCC accession: ENCF000MZT)

**References**

- Dunham I, Kundaje A, Aldred SF, et al. An integrated encyclopedia of DNA elements in the human genome. *Nature*. 2012 Sep 6;489(7414):57-74.

**See Also**

- [A549\\_FOSL2\\_01\\_Peaks\\_partial](#) the associate sites dataset.
- [findConsensusPeakRegions](#) for extracting regions sharing the same features in more than one experiment.

**Examples**

```
## Loading datasets
data(A549_FOSL2_01_NarrowPeaks_partial)
data(A549_FOSL2_01_Peaks_partial)
data(A549_FOXA1_01_NarrowPeaks_partial)
data(A549_FOXA1_01_Peaks_partial)

## Assigning experiment name to each row of the dataset.
## NarrowPeak and Peak datasets from the same experiment must
## have identical names.
names(A549_FOXA1_01_Peaks_partial) <- rep("FOXA1_01",
                                         length(A549_FOXA1_01_Peaks_partial))
names(A549_FOXA1_01_NarrowPeaks_partial) <- rep("FOXA1_01",
                                                length(A549_FOXA1_01_NarrowPeaks_partial))
names(A549_FOSL2_01_Peaks_partial) <- rep("FOSL2_01",
                                           length(A549_FOSL2_01_Peaks_partial))
names(A549_FOSL2_01_NarrowPeaks_partial) <- rep("FOSL2_01",
                                                length(A549_FOSL2_01_NarrowPeaks_partial))

## Calculating consensus regions for chromosome 10 only
## with a default region size of 200 bp (2 * extendingSize)
## which is not extended to include all genomic regions.
## A peak from both experiments must be present in a region to
## be retained as a consensus region.
chrList <- Seqinfo("chr10", 135534747, NA)
findConsensusPeakRegions(
  narrowPeaks = c(A549_FOXA1_01_NarrowPeaks_partial,
                 A549_FOSL2_01_NarrowPeaks_partial),
  peaks = c(A549_FOXA1_01_Peaks_partial,
            A549_FOSL2_01_Peaks_partial),
  chrInfo = chrList,
  extendingSize = 100,
  expandToFitPeakRegion = FALSE,
  shrinkToFitPeakRegion = TRUE,
  minNbrExp = 2,
  nbrThreads = 1)
```

---

A549\_FOSL2\_01\_Peaks\_partial

*Sites with the greatest evidence of transcription factor binding for the FOSL2 transcription factor (for demonstration purpose)*

---

## Description

Sites representing the greatest evidence of enrichment for the FOSL2 transcription factor (DCC accession: ENCFF000MZT) for regions chr1:249120200-249250621 and chr10:1-370100 from the Encyclopedia of DNA Elements (ENCODE) data (Dunham I et al. 2012).

## Usage

```
data(A549_FOSL2_01_Peaks_partial)
```

## Format

A GRanges containing one entry per site. Each row of GRanges has the same row name which represent the name of the experiment.

## Source

The Encyclopedia of DNA Elements (ENCODE) (DCC accession: ENCFF000MZT)

## References

- Dunham I, Kundaje A, Aldred SF, et al. An integrated encyclopedia of DNA elements in the human genome. Nature. 2012 Sep 6;489(7414):57-74.

## See Also

- [A549\\_FOSL2\\_01\\_NarrowPeaks\\_partial](#) the associate genomic regions dataset.
- [findConsensusPeakRegions](#) for extracting regions sharing the same features in more than one experiment.

## Examples

```
## Loading datasets
data(A549_FOSL2_01_NarrowPeaks_partial)
data(A549_FOXA1_01_NarrowPeaks_partial)
data(A549_FOSL2_01_Peaks_partial)
data(A549_FOXA1_01_Peaks_partial)

## Assigning experiment name to each row of the dataset.
## NarrowPeak and Peak datasets from the same experiment must
## have identical names.
```

```

names(A549_FOXA1_01_Peaks_partial) <- rep("FOXA1_01",
                                          length(A549_FOXA1_01_Peaks_partial))
names(A549_FOXA1_01_NarrowPeaks_partial) <- rep("FOXA1_01",
                                                length(A549_FOXA1_01_NarrowPeaks_partial))
names(A549_FOSL2_01_Peaks_partial) <- rep("FOSL2_01",
                                           length(A549_FOSL2_01_Peaks_partial))
names(A549_FOSL2_01_NarrowPeaks_partial) <- rep("FOSL2_01",
                                                length(A549_FOSL2_01_NarrowPeaks_partial))

## Calculating consensus regions for chromosome 1 only
## with a default region size of 400 bp (2 * extendingSize)
## which is extended to include all genomic regions of the
## closest peak (for each experiment).
## A peak from both experiments must be present in a region to
## be retained as a consensus region.
chrList <- Seqinfo("chr1", 249250621, NA)
findConsensusPeakRegions(
  narrowPeaks = c(A549_FOXA1_01_NarrowPeaks_partial,
                 A549_FOSL2_01_NarrowPeaks_partial),
  peaks = c(A549_FOXA1_01_Peaks_partial,
            A549_FOSL2_01_Peaks_partial),

  chrInfo = chrList,
  extendingSize = 200,
  expandToFitPeakRegion = TRUE,
  shrinkToFitPeakRegion = FALSE,
  minNbrExp = 2,
  nbrThreads = 1)

```

---

A549\_FOXA1\_01\_NarrowPeaks\_partial

*Genomic regions with the greatest evidence of transcription factor binding for the FOXA1 transcription factor (for demonstration purpose)*

---

## Description

Genomic regions representing the greatest evidence of enrichment for the FOXA1 transcription factor (DCC accession: ENCF000NAH) for regions chr1:249120200-249250621 and chr10:1-370100 from the Encyclopedia of DNA Elements (ENCODE) data (Dunham I et al. 2012).

## Usage

```
data(A549_FOXA1_01_NarrowPeaks_partial)
```

## Format

A GRanges containing one entry per genomic regions. Each row of GRanges has a name which represent the name of the experiment.

**Source**

The Encyclopedia of DNA Elements (ENCODE) (DCC accession: ENCFF000NAH)

**References**

- Dunham I, Kundaje A, Aldred SF, et al. An integrated encyclopedia of DNA elements in the human genome. Nature. 2012 Sep 6;489(7414):57-74.

**See Also**

- [A549\\_FOXA1\\_01\\_Peaks\\_partial](#) the associate sites dataset.
- [findConsensusPeakRegions](#) for extracting regions sharing the same features in more than one experiment.

**Examples**

```
## Loading datasets
data(A549_FOSL2_01_NarrowPeaks_partial)
data(A549_FOSL2_01_Peaks_partial)
data(A549_FOXA1_01_NarrowPeaks_partial)
data(A549_FOXA1_01_Peaks_partial)

## Assigning experiment name to each row of the dataset.
## NarrowPeak and Peak datasets from the same experiment must
## have identical names.
names(A549_FOXA1_01_Peaks_partial) <- rep("FOXA1_01",
    length(A549_FOXA1_01_Peaks_partial))
names(A549_FOXA1_01_NarrowPeaks_partial) <- rep("FOXA1_01",
    length(A549_FOXA1_01_NarrowPeaks_partial))
names(A549_FOSL2_01_Peaks_partial) <- rep("FOSL2_01",
    length(A549_FOSL2_01_Peaks_partial))
names(A549_FOSL2_01_NarrowPeaks_partial) <- rep("FOSL2_01",
    length(A549_FOSL2_01_NarrowPeaks_partial))

## Calculating consensus regions for both chromosomes 1 and 10
## with a default region size of 300 bp (2 * extendingSize)
## which is not extended to include all genomic regions.
## A peak from both experiments must be present in a region to
## be retained as a consensus region.
chrList <- Seqinfo(c("chr1", "chr10"), c(249250621, 135534747), NA)
findConsensusPeakRegions(
  narrowPeaks = c(A549_FOXA1_01_NarrowPeaks_partial,
    A549_FOSL2_01_NarrowPeaks_partial),
  peaks = c(A549_FOXA1_01_Peaks_partial,
    A549_FOSL2_01_Peaks_partial),
  chrInfo = chrList,
  extendingSize = 150,
  expandToFitPeakRegion = FALSE,
  minNbrExp = 2,
  nbrThreads = 1)
```

---

A549\_FOXA1\_01\_Peaks\_partial

*Sites with the greatest evidence of transcription factor binding for the FOXA1 transcription factor (for demonstration purpose)*

---

## Description

Sites representing the greatest evidence of enrichment for the FOXA1 transcription factor (DCC accession: ENCFF000NAH) for regions chr1:249120200-249250621 and chr10:1-370100 from the Encyclopedia of DNA Elements (ENCODE) data (Dunham I et al. 2012).

## Usage

```
data(A549_FOXA1_01_Peaks_partial)
```

## Format

A GRanges containing one entry per site . Each row of GRanges has a name which represent the name of the experiment.

## Source

The Encyclopedia of DNA Elements (ENCODE) (DCC accession: ENCFF000NAH)

## References

- Dunham I, Kundaje A, Aldred SF, et al. An integrated encyclopedia of DNA elements in the human genome. Nature. 2012 Sep 6;489(7414):57-74.

## See Also

- [A549\\_FOXA1\\_01\\_NarrowPeaks\\_partial](#) the associate genomic regions dataset.
- [findConsensusPeakRegions](#) for extracting regions sharing the same features in more than one experiment.

## Examples

```
## Loading datasets
data(A549_FOSL2_01_NarrowPeaks_partial)
data(A549_FOSL2_01_Peaks_partial)
data(A549_FOXA1_01_NarrowPeaks_partial)
data(A549_FOXA1_01_Peaks_partial)

## Assigning experiment name to each row of the dataset.
## NarrowPeak and Peak datasets from the same experiment must
## have identical names.
names(A549_FOXA1_01_Peaks_partial) <- rep("FOXA1_01",
    length(A549_FOXA1_01_Peaks_partial))
```

```

names(A549_FOXA1_01_NarrowPeaks_partial) <- rep("FOXA1_01",
  length(A549_FOXA1_01_NarrowPeaks_partial))
names(A549_FOSL2_01_Peaks_partial) <-rep("FOSL2_01",
  length(A549_FOSL2_01_Peaks_partial))
names(A549_FOSL2_01_NarrowPeaks_partial) <- rep("FOSL2_01",
  length(A549_FOSL2_01_NarrowPeaks_partial))

## Calculating consensus regions for both chromosomes 1 and 10
## with a default region size of 100 bp (2 * extendingSize)
## which is extended to include all genomic regions for the closest
## peak to the median position of all peaks included in the region (for each
## experiment).
## A peak from both experiments must be present in a region to
## be retained as a consensus region.
chrList <- Seqinfo(c("chr1", "chr10"), c(249250621, 135534747), NA)
findConsensusPeakRegions(
  narrowPeaks = c(A549_FOXA1_01_NarrowPeaks_partial,
    A549_FOSL2_01_NarrowPeaks_partial),
  peaks = c(A549_FOXA1_01_Peaks_partial,
    A549_FOSL2_01_Peaks_partial),
  chrInfo = chrList,
  extendingSize = 50,
  expandToFitPeakRegion = TRUE,
  shrinkToFitPeakRegion = TRUE,
  minNbrExp = 2,
  nbrThreads = 1)

```

---

A549\_NR3C1\_CFQ\_NarrowPeaks\_partial

*Ranges with the greatest evidence of transcription factor binding for the NR3C1 transcription factor from ENCODE (DDC accession: ENCFF002CFQ). For demonstration purpose.*

---

## Description

Ranges representing the greatest evidence of enrichment for the NR3C1 transcription factor (DCC accession: ENCFF002CFQ) for regions chr2:40000000-50000000 and chr3:10000000-13000000 from the Encyclopedia of DNA Elements (ENCODE) data (Dunham I et al. 2012).

## Usage

```
data(A549_NR3C1_CFQ_NarrowPeaks_partial)
```

## Format

A GRanges containing one entry per site. The ranges are surrounding the peaks present in the dataset A549\_NR3C1\_CFQ\_Peaks\_partial.

**Details**

The peaks and ranges have been obtained using an optimal IDR analysis done on all replicates.

**Source**

The Encyclopedia of DNA Elements (ENCODE) (DCC accession: ENCFF002CFQ)

**References**

- Dunham I, Kundaje A, Aldred SF, et al. An integrated encyclopedia of DNA elements in the human genome. *Nature*. 2012 Sep 6;489(7414):57-74.

**See Also**

- [A549\\_NR3C1\\_CFQ\\_Peaks\\_partial](#) the associate genomic regions dataset.
- [findConsensusPeakRegions](#) for extracting regions sharing the same features in more than one experiment.

**Examples**

```
## Loading datasets
data(A549_NR3C1_CFQ_NarrowPeaks_partial)
data(A549_NR3C1_CFQ_Peaks_partial)
data(A549_NR3C1_CFS_NarrowPeaks_partial)
data(A549_NR3C1_CFS_Peaks_partial)

## Assigning experiment name to each row of the dataset.
## NarrowPeak and Peak datasets from the same experiment must
## have identical names.
names(A549_NR3C1_CFQ_NarrowPeaks_partial) <- rep("NR3C1_CFQ",
                                                length(A549_NR3C1_CFQ_NarrowPeaks_partial))
names(A549_NR3C1_CFQ_Peaks_partial) <- rep("NR3C1_CFQ",
                                           length(A549_NR3C1_CFQ_Peaks_partial))
names(A549_NR3C1_CFS_NarrowPeaks_partial) <- rep("NR3C1_CFS",
                                                  length(A549_NR3C1_CFS_NarrowPeaks_partial))
names(A549_NR3C1_CFS_Peaks_partial) <- rep("NR3C1_CFS",
                                           length(A549_NR3C1_CFS_Peaks_partial))

## Calculating consensus regions for chromosome 3
## with a default region size of 300 bp (2 * extendingSize)
## which is extended to include all genomic regions for the closest
## peak to the median position of all peaks included in the region (for
## each experiment).
## Peaks from both experiments must be present in a region to
## be retained as a consensus region.
chrList <- Seqinfo(c("chr3"), c(198022430), NA)
findConsensusPeakRegions(
  narrowPeaks = c(A549_NR3C1_CFQ_NarrowPeaks_partial,
                 A549_NR3C1_CFS_NarrowPeaks_partial),
  peaks = c(A549_NR3C1_CFQ_Peaks_partial,
            A549_NR3C1_CFS_Peaks_partial),
```



```
chrInfo = chrList,  
extendingSize = 150,  
expandToFitPeakRegion = FALSE,  
shrinkToFitPeakRegion = TRUE,  
minNbrExp = 2,  
nbrThreads = 1)
```

---

A549\_NR3C1\_CFQ\_Peaks\_partial

*Sites with the greatest evidence of transcription factor binding for the NR3C1 transcription factor from ENCODE (DCC accession: ENCFF002CFQ). For demonstration purpose.*

---

### Description

Sites representing the greatest evidence of enrichment for the NR3C1 transcription factor (DCC accession: ENCFF002CFQ) for regions chr2:40000000-50000000 and chr3:10000000-13000000 from the Encyclopedia of DNA Elements (ENCODE) data (Dunham I et al. 2012).

### Usage

```
data(A549_NR3C1_CFQ_Peaks_partial)
```

### Format

A GRanges containing one entry per site. The peaks are surrounded by ranges present in the dataset A549\_NR3C1\_CFQ\_NarrowPeaks\_partial.

### Details

The peaks and ranges have been obtained using an optimal IDR analysis done on all replicates.

### Source

The Encyclopedia of DNA Elements (ENCODE) (DCC accession: ENCFF002CFQ)

### References

- Dunham I, Kundaje A, Aldred SF, et al. An integrated encyclopedia of DNA elements in the human genome. Nature. 2012 Sep 6;489(7414):57-74.

### See Also

- [A549\\_NR3C1\\_CFQ\\_NarrowPeaks\\_partial](#) the associate genomic regions dataset.
- [findConsensusPeakRegions](#) for extracting regions sharing the same features in more than one experiment.

## Examples

```
## Loading datasets
data(A549_NR3C1_CFQ_NarrowPeaks_partial)
data(A549_NR3C1_CFQ_Peaks_partial)
data(A549_NR3C1_CFS_NarrowPeaks_partial)
data(A549_NR3C1_CFS_Peaks_partial)
data(A549_NR3C1_CFR_NarrowPeaks_partial)
data(A549_NR3C1_CFR_Peaks_partial)

## Assigning experiment name to each row of the dataset.
## NarrowPeak and Peak datasets from the same experiment must
## have identical names.
names(A549_NR3C1_CFQ_NarrowPeaks_partial) <- rep("NR3C1_CFQ",
        length(A549_NR3C1_CFQ_NarrowPeaks_partial))
names(A549_NR3C1_CFQ_Peaks_partial) <- rep("NR3C1_CFQ",
        length(A549_NR3C1_CFQ_Peaks_partial))
names(A549_NR3C1_CFS_NarrowPeaks_partial) <- rep("NR3C1_CFS",
        length(A549_NR3C1_CFS_NarrowPeaks_partial))
names(A549_NR3C1_CFS_Peaks_partial) <- rep("NR3C1_CFS",
        length(A549_NR3C1_CFS_Peaks_partial))
names(A549_NR3C1_CFR_NarrowPeaks_partial) <- rep("NR3C1_CFR",
        length(A549_NR3C1_CFR_NarrowPeaks_partial))
names(A549_NR3C1_CFR_Peaks_partial) <- rep("NR3C1_CFR",
        length(A549_NR3C1_CFR_Peaks_partial))

## Calculating consensus regions for chromosome 3
## with a default region size of 140 bp (2 * extendingSize)
## which is extended to include all genomic regions for the closest
## peak to the median position of all peaks included in the region (for
## each experiment).
## Peaks from at least 2 experiments must be present in a region to
## be retained as a consensus region.
chrList <- Seqinfo(c("chr3"), c(198022430), NA)
findConsensusPeakRegions(
  narrowPeaks = c(A549_NR3C1_CFQ_NarrowPeaks_partial,
        A549_NR3C1_CFS_NarrowPeaks_partial,
        A549_NR3C1_CFR_NarrowPeaks_partial),
  peaks = c(A549_NR3C1_CFQ_Peaks_partial,
        A549_NR3C1_CFS_Peaks_partial,
        A549_NR3C1_CFR_Peaks_partial),
  chrInfo = chrList,
  extendingSize = 70,
  expandToFitPeakRegion = FALSE,
  shrinkToFitPeakRegion = FALSE,
  minNbrExp = 2,
  nbrThreads = 1)
```

---

A549\_NR3C1\_CFR\_NarrowPeaks\_partial

*Ranges with the greatest evidence of transcription factor binding for the NR3C1 transcription factor from ENCODE (DCC accession: ENCFF002CFR). For demonstration purpose.*

---

## Description

Ranges representing the greatest evidence of enrichment for the NR3C1 transcription factor (DCC accession: ENCFF002CFR) for regions chr2:40000000-50000000 and chr3:10000000-13000000 from the Encyclopedia of DNA Elements (ENCODE) data (Dunham I et al. 2012).

## Usage

```
data(A549_NR3C1_CFR_NarrowPeaks_partial)
```

## Format

A GRanges containing one entry per site. The ranges are surrounding the peaks present in the dataset A549\_NR3C1\_CFR\_Peaks\_partial.

## Details

The peaks and ranges have been obtained using an optimal IDR analysis done on all replicates.

## Source

The Encyclopedia of DNA Elements (ENCODE) (DCC accession: ENCFF002CFR)

## References

- Dunham I, Kundaje A, Aldred SF, et al. An integrated encyclopedia of DNA elements in the human genome. Nature. 2012 Sep 6;489(7414):57-74.

## See Also

- [A549\\_NR3C1\\_CFR\\_Peaks\\_partial](#) the associate genomic regions dataset.
- [findConsensusPeakRegions](#) for extracting regions sharing the same features in more than one experiment.

## Examples

```
## Loading datasets
data(A549_NR3C1_CFR_NarrowPeaks_partial)
data(A549_NR3C1_CFR_Peaks_partial)
data(A549_NR3C1_CFR_NarrowPeaks_partial)
data(A549_NR3C1_CFR_Peaks_partial)

## Assigning experiment name to each row of the dataset.
## NarrowPeak and Peak datasets from the same experiment must
```

```

## have identical names.
names(A549_NR3C1_CFQ_NarrowPeaks_partial) <- rep("NR3C1_CFQ",
                                                length(A549_NR3C1_CFQ_NarrowPeaks_partial))
names(A549_NR3C1_CFQ_Peaks_partial) <- rep("NR3C1_CFQ",
                                           length(A549_NR3C1_CFQ_Peaks_partial))
names(A549_NR3C1_CFR_NarrowPeaks_partial) <- rep("NR3C1_CFR",
                                                  length(A549_NR3C1_CFR_NarrowPeaks_partial))
names(A549_NR3C1_CFR_Peaks_partial) <- rep("NR3C1_CFR",
                                           length(A549_NR3C1_CFR_Peaks_partial))

## Calculating consensus regions for chromosome 2
## with a default region size of 250 bp (2 * extendingSize)
## which is extended to include all genomic regions for the closest
## peak to the median position of all peaks included in the region (for
## each experiment).
## Peaks from both experiments must be present in a region to
## be retained as a consensus region.
chrList <- Seqinfo(c("chr2"), c(243199373), NA)
findConsensusPeakRegions(
  narrowPeaks = c(A549_NR3C1_CFQ_NarrowPeaks_partial,
                 A549_NR3C1_CFR_NarrowPeaks_partial),
  peaks = c(A549_NR3C1_CFQ_Peaks_partial,
            A549_NR3C1_CFR_Peaks_partial),
  chrInfo = chrList,
  extendingSize = 125,
  expandToFitPeakRegion = TRUE,
  shrinkToFitPeakRegion = TRUE,
  minNbrExp = 2,
  nbrThreads = 1)

```

---

A549\_NR3C1\_CFR\_Peaks\_partial

*Sites with the greatest evidence of transcription factor binding for the NR3C1 transcription factor from ENCODE (DCC accession: ENCFF002CFR). For demonstration purpose.*

---

## Description

Sites representing the greatest evidence of enrichment for the NR3C1 transcription factor (DCC accession: ENCFF002CFR) for regions chr2:40000000-50000000 and chr3:10000000-13000000 from the Encyclopedia of DNA Elements (ENCODE) data (Dunham I et al. 2012).

## Usage

```
data(A549_NR3C1_CFR_Peaks_partial)
```

## Format

A GRanges containing one entry per site. The peaks are surrounded by ranges present in the dataset A549\_NR3C1\_CFR\_NarrowPeaks\_partial.

**Details**

The peaks and ranges have been obtained using an optimal IDR analysis done on all replicates.

**Source**

The Encyclopedia of DNA Elements (ENCODE) (DCC accession: ENCFF002CFR)

**References**

- Dunham I, Kundaje A, Aldred SF, et al. An integrated encyclopedia of DNA elements in the human genome. *Nature*. 2012 Sep 6;489(7414):57-74.

**See Also**

- [A549\\_NR3C1\\_CFR\\_NarrowPeaks\\_partial](#) the associate genomic regions dataset.
- [findConsensusPeakRegions](#) for extracting regions sharing the same features in more than one experiment.

**Examples**

```
## Loading datasets
data(A549_NR3C1_CFQ_NarrowPeaks_partial)
data(A549_NR3C1_CFQ_Peaks_partial)
data(A549_NR3C1_CFR_NarrowPeaks_partial)
data(A549_NR3C1_CFR_Peaks_partial)

## Assigning experiment name to each row of the dataset.
## NarrowPeak and Peak datasets from the same experiment must
## have identical names.
names(A549_NR3C1_CFQ_NarrowPeaks_partial) <- rep("NR3C1_CFQ",
                                                length(A549_NR3C1_CFQ_NarrowPeaks_partial))
names(A549_NR3C1_CFQ_Peaks_partial) <- rep("NR3C1_CFQ",
                                            length(A549_NR3C1_CFQ_Peaks_partial))
names(A549_NR3C1_CFR_NarrowPeaks_partial) <- rep("NR3C1_CFR",
                                                  length(A549_NR3C1_CFR_NarrowPeaks_partial))
names(A549_NR3C1_CFR_Peaks_partial) <- rep("NR3C1_CFR",
                                            length(A549_NR3C1_CFR_Peaks_partial))

## Calculating consensus regions for chromosome 2
## with a default region size of 40 bp (2 * extendingSize)
## which is extended to include all genomic regions for the closest
## peak to the median position of all peaks included in the region (for
## each experiment).
## Peaks from both experiments must be present in a region to
## be retained as a consensus region.
chrList <- Seqinfo(c("chr2"), c(243199373), NA)
findConsensusPeakRegions(
  narrowPeaks = c(A549_NR3C1_CFQ_NarrowPeaks_partial,
                  A549_NR3C1_CFR_NarrowPeaks_partial),
  peaks = c(A549_NR3C1_CFQ_Peaks_partial,
            A549_NR3C1_CFR_Peaks_partial),
```

```
chrInfo = chrList,  
extendingSize = 20,  
expandToFitPeakRegion = TRUE,  
shrinkToFitPeakRegion = FALSE,  
minNbrExp = 2,  
nbrThreads = 1)
```

---

A549\_NR3C1\_CFS\_NarrowPeaks\_partial

*Ranges with the greatest evidence of transcription factor binding for the NR3C1 transcription factor from ENCODE (DCC accession: ENCFF002CFS). For demonstration purpose.*

---

### Description

Ranges representing the greatest evidence of enrichment for the NR3C1 transcription factor (DCC accession: ENCFF002CFS) for regions chr2:40000000-50000000 and chr3:10000000-13000000 from the Encyclopedia of DNA Elements (ENCODE) data (Dunham I et al. 2012).

### Usage

```
data(A549_NR3C1_CFS_NarrowPeaks_partial)
```

### Format

A GRanges containing one entry per site. The ranges are surrounding the peaks present in the dataset A549\_NR3C1\_CFS\_Peaks\_partial.

### Details

The peaks and ranges have been obtained using an optimal IDR analysis done on all replicates.

### References

- Dunham I, Kundaje A, Aldred SF, et al. An integrated encyclopedia of DNA elements in the human genome. Nature. 2012 Sep 6;489(7414):57-74.

### See Also

- [A549\\_NR3C1\\_CFS\\_Peaks\\_partial](#) the associate genomic regions dataset.
- [findConsensusPeakRegions](#) for extracting regions sharing the same features in more than one experiment.

**Examples**

```
## Loading datasets
data(A549_NR3C1_CFQ_NarrowPeaks_partial)
data(A549_NR3C1_CFQ_Peaks_partial)
data(A549_NR3C1_CFS_NarrowPeaks_partial)
data(A549_NR3C1_CFS_Peaks_partial)

## Assigning experiment name to each row of the dataset.
## NarrowPeak and Peak datasets from the same experiment must
## have identical names.
names(A549_NR3C1_CFQ_NarrowPeaks_partial) <- rep("NR3C1_CFQ",
                                                length(A549_NR3C1_CFQ_NarrowPeaks_partial))
names(A549_NR3C1_CFQ_Peaks_partial) <- rep("NR3C1_CFQ",
                                            length(A549_NR3C1_CFQ_Peaks_partial))
names(A549_NR3C1_CFS_NarrowPeaks_partial) <- rep("NR3C1_CFS",
                                                  length(A549_NR3C1_CFS_NarrowPeaks_partial))
names(A549_NR3C1_CFS_Peaks_partial) <- rep("NR3C1_CFS",
                                            length(A549_NR3C1_CFS_Peaks_partial))

## Calculating consensus regions for chromosome 2
## with a default region size of 400 bp (2 * extendingSize).
## The consensus regions are not resized to fit the narrowPeak regions.
## Peaks from both experiments must be present in a region to
## be retained as a consensus region.
chrList <- Seqinfo(c("chr2"), c(243199373), NA)
findConsensusPeakRegions(
  narrowPeaks = c(A549_NR3C1_CFQ_NarrowPeaks_partial,
                 A549_NR3C1_CFS_NarrowPeaks_partial),
  peaks = c(A549_NR3C1_CFQ_Peaks_partial,
            A549_NR3C1_CFS_Peaks_partial),
  chrInfo = chrList,
  extendingSize = 200,
  expandToFitPeakRegion = FALSE,
  shrinkToFitPeakRegion = FALSE,
  minNbrExp = 2,
  nbrThreads = 1)
```

---

A549\_NR3C1\_CFS\_Peaks\_partial

*Sites with the greatest evidence of transcription factor binding for the NR3C1 transcription factor from ENCODE (DCC accession: ENCFF002CFS). For demonstration purpose.*

---

**Description**

Sites representing the greatest evidence of enrichment for the NR3C1 transcription factor (DCC accession: ENCFF002CFS) for regions chr2:40000000-50000000 and chr3:10000000-13000000 from the Encyclopedia of DNA Elements (ENCODE) data (Dunham I et al. 2012).

**Usage**

```
data(A549_NR3C1_CFS_Peaks_partial)
```

**Format**

A GRanges containing one entry per site. The peaks are surrounded by ranges present in the dataset A549\_NR3C1\_CFS\_NarrowPeaks\_partial.

**Details**

The peaks and ranges have been obtained using an optimal IDR analysis done on all replicates.

**Source**

The Encyclopedia of DNA Elements (ENCODE) (DCC accession: ENCFF002CFS)

**References**

- Dunham I, Kundaje A, Aldred SF, et al. An integrated encyclopedia of DNA elements in the human genome. Nature. 2012 Sep 6;489(7414):57-74.

**See Also**

- [A549\\_NR3C1\\_CFS\\_NarrowPeaks\\_partial](#) the associate genomic regions dataset.
- [findConsensusPeakRegions](#) for extracting regions sharing the same features in more than one experiment.

**Examples**

```
## Loading datasets
data(A549_NR3C1_CFQ_NarrowPeaks_partial)
data(A549_NR3C1_CFQ_Peaks_partial)
data(A549_NR3C1_CFS_NarrowPeaks_partial)
data(A549_NR3C1_CFS_Peaks_partial)

## Assigning experiment name to each row of the dataset.
## NarrowPeak and Peak datasets from the same experiment must
## have identical names.
names(A549_NR3C1_CFQ_NarrowPeaks_partial) <- rep("NR3C1_CFQ",
                                                length(A549_NR3C1_CFQ_NarrowPeaks_partial))
names(A549_NR3C1_CFQ_Peaks_partial) <- rep("NR3C1_CFQ",
                                           length(A549_NR3C1_CFQ_Peaks_partial))
names(A549_NR3C1_CFS_NarrowPeaks_partial) <- rep("NR3C1_CFS",
                                                  length(A549_NR3C1_CFS_NarrowPeaks_partial))
names(A549_NR3C1_CFS_Peaks_partial) <- rep("NR3C1_CFS",
                                           length(A549_NR3C1_CFS_Peaks_partial))

## Calculating consensus regions for chromosome 2
## with a default region size of 80 bp (2 * extendingSize).
## The consensus regions are not resized to fit the narrowPeak regions.
## Peaks from both experiments must be present in a region to
```



```
## be retained as a consensus region.
chrList <- Seqinfo(c("chr2"), c(243199373), NA)
findConsensusPeakRegions(
  narrowPeaks = c(A549_NR3C1_CFQ_NarrowPeaks_partial,
                  A549_NR3C1_CFS_NarrowPeaks_partial),
  peaks = c(A549_NR3C1_CFQ_Peaks_partial,
            A549_NR3C1_CFS_Peaks_partial),
  chrInfo = chrList,
  extendingSize = 40,
  expandToFitPeakRegion = FALSE,
  shrinkToFitPeakRegion = FALSE,
  minNbrExp = 2,
  nbrThreads = 1)
```

---

findConsensusPeakRegions

*Extract regions sharing features in more than one experiment*

---

## Description

Find regions sharing the same features for a minimum number of experiments using called peaks of signal enrichment based on pooled, normalized data (mainly coming from narrowPeak files). The peaks and narrow peaks are used to identify the consensus regions. The minimum number of experiments that must have at least one peak in a region so that it is retained as a consensus region is specified by user, as well as the size of mining regions. Only the chromosomes specified by the user are treated. The function can be paralled by specifying a number of threads superior to 1.

When the padding is small, the detected regions are smaller than the one that could be obtained by doing an overlap of the narrow regions. Even more, the parameter specifying the minimum number of experiments needed to retain a region add versatility to the function.

Beware that the side of the padding can have a large effect on the detected consensus regions. It is recommended to test more than one size and to do some manual validation of the resulting consensus regions before selecting the final padding size.

## Usage

```
findConsensusPeakRegions(
  narrowPeaks,
  peaks,
  chrInfo,
  extendingSize = 250,
  expandToFitPeakRegion = FALSE,
  shrinkToFitPeakRegion = FALSE,
  minNbrExp = 1L,
  nbrThreads = 1L
)
```

**Arguments**

narrowPeaks	a GRanges containing called peak regions of signal enrichment based on pooled, normalized data for all analyzed experiments. All GRanges entries must have a metadata field called "name" which identifies the region to the called peak. All GRanges entries must also have a row name which identifies the experiment of origin. Each peaks entry must have an associated narrowPeaks entry. A GRanges entry is associated to a narrowPeaks entry by having a identical metadata "name" field and a identical row name.
peaks	a GRanges containing called peaks of signal enrichment based on pooled, normalized data for all analyzed experiments. All GRanges entries must have a metadata field called "name" which identifies the called peak. All GRanges entries must have a row name which identifies the experiment of origin. Each peaks entry must have an associated narrowPeaks entry. A GRanges entry is associated to a narrowPeaks entry by having a identical metadata "name" field and a identical row name.
chrInfo	a Seqinfo containing the name and the length of the chromosomes to analyze. Only the chromosomes contained in this Seqinfo will be analyzed.
extendingSize	a numeric value indicating the size of padding on both sides of the position of the peaks median to create the consensus region. The minimum size of the consensus region is equal to twice the value of the extendingSize parameter. The size of the extendingSize must be a positive integer. Default = 250.
expandToFitPeakRegion	a logical indicating if the region size, which is set by the extendingSize parameter is extended to include the entire narrow peak regions of all peaks included in the unextended consensus region. The narrow peak regions of the peaks added because of the extension are not considered for the extension. Default: FALSE.
shrinkToFitPeakRegion	a logical indicating if the region size, which is set by the extendingSize parameter is shrunk to fit the narrow peak regions of the peaks when all those regions are smaller than the consensus region. Default: FALSE.
minNbrExp	a positive numeric or a positive integer indicating the minimum number of experiments in which at least one peak must be present for a potential consensus region. The numeric must be a positive integer inferior or equal to the number of experiments present in the narrowPeaks and peaks parameters. Default = 1.
nbrThreads	a numeric or a integer indicating the number of threads to use in parallel. The nbrThreads must be a positive integer. Default = 1.

**Value**

an list of class "consensusRanges" containing :

- call the matched call.
- consensusRanges a GRanges containing the consensus regions.

**Author(s)**

Astrid Deschênes

**Examples**

```
## Loading datasets
data(A549_CTCF_MYN_NarrowPeaks_partial)
data(A549_CTCF_MYN_Peaks_partial)
data(A549_CTCF_MYJ_NarrowPeaks_partial)
data(A549_CTCF_MYJ_Peaks_partial)

## Assigning experiment name "CTCF_MYJ" to first experiment
names(A549_CTCF_MYJ_NarrowPeaks_partial) <- rep("CTCF_MYJ",
  length(A549_CTCF_MYJ_NarrowPeaks_partial))
names(A549_CTCF_MYJ_Peaks_partial) <- rep("CTCF_MYJ",
  length(A549_CTCF_MYJ_Peaks_partial))

## Assigning experiment name "CTCF_MYN" to second experiment
names(A549_CTCF_MYN_NarrowPeaks_partial) <- rep("CTCF_MYN",
  length(A549_CTCF_MYN_NarrowPeaks_partial))
names(A549_CTCF_MYN_Peaks_partial) <- rep("CTCF_MYN",
  length(A549_CTCF_MYN_Peaks_partial))

## Only chromosome 1 is going to be analysed
chrList <- Seqinfo("chr1", 249250621, NA)

## Find consensus regions with both experiments
results <- findConsensusPeakRegions(
  narrowPeaks = c(A549_CTCF_MYJ_NarrowPeaks_partial,
    A549_CTCF_MYN_NarrowPeaks_partial),
  peaks = c(A549_CTCF_MYJ_Peaks_partial,
    A549_CTCF_MYN_Peaks_partial),
  chrInfo = chrList,
  extendingSize = 300,
  expandToFitPeakRegion = TRUE,
  shrinkToFitPeakRegion = FALSE,
  minNbrExp = 2,
  nbrThreads = 1)

## Print 2 first consensus regions
head(results$consensusRanges, 2)
```

---

```
findConsensusPeakRegionsForOneChrom
```

*Extract regions sharing features in more than one experiment for one specific chromosome.*

---

**Description**

Find regions sharing the same features for a minimum number of experiments using called peaks of signal enrichment based on pooled, normalized data (mainly coming from narrowPeak files). The analysis is limited to one chromosome. The peaks and narrow peaks are used to identify the

consensus regions. The minimum number of experiments that must have at least one peak in a region so that it is retained as a consensus region is specified by user, as well as the size of mining regions.

### Usage

```
findConsensusPeakRegionsForOneChrom(
  chrName,
  allPeaks,
  allNarrowPeaks,
  extendingSize,
  expandToFitPeakRegion,
  shrinkToFitPeakRegion,
  minNbrExp,
  chrList
)
```

### Arguments

<code>chrName</code>	a character, the name of the chromosome to analyze.
<code>allPeaks</code>	a GRanges containing all peaks from all experiments sorted by position.
<code>allNarrowPeaks</code>	a GRanges containing all narrow peaks from all experiments sorted by position.
<code>extendingSize</code>	a numeric value indicating the size of padding at each side of the peaks median position to create the consensus region. The minimum size of the consensus region will be equal to twice the value of the <code>extendingSize</code> parameter. The size of the <code>extendingSize</code> must be a positive integer. Default = 250.
<code>expandToFitPeakRegion</code>	a logical indicating if the region set by the <code>extendingSize</code> parameter is extended to include all narrow peak regions. Only the narrow peaks regions of the peaks included in the unextended region are used during the extension process. It is possible that has a side effect, adding peaks are being included in the extended region.
<code>shrinkToFitPeakRegion</code>	a logical indicating if the region set by the <code>extendingSize</code> parameter is shrunk to fit the narrow peak regions.
<code>minNbrExp</code>	a numeric indicating the minimum number of BED files in which a peak must be present for a region to be retained. The numeric must be a positive value inferior or equal to the number of files present in the <code>narrowPeakFiles</code> parameter. Default = 1.
<code>chrList</code>	a Seqinfo containing the name and the length of the chromosomes to analyze.

### Value

an object of class "commonFeatures".

### Author(s)

Astrid Deschênes

---

`findConsensusPeakRegionsValidation`*Parameter validation for the `findConsensusPeakRegions` function*

---

## Description

Validation of all parameters needed by the public `findConsensusPeakRegions` function.

## Usage

```
findConsensusPeakRegionsValidation(  
  narrowPeaks,  
  peaks,  
  chrList,  
  extendingSize,  
  expandToFitPeakRegion,  
  shrinkToFitPeakRegion,  
  minNbrExp,  
  nbrThreads  
)
```

## Arguments

<code>narrowPeaks</code>	a GRanges representing called peaks of signal for all experiments.
<code>peaks</code>	a GRanges representing peaks for all experiments.
<code>chrList</code>	a Seqinfo containing the name and the length of the chromosomes to analyze which indicate that all chromosomes must be analyzed.
<code>extendingSize</code>	a numeric value indicating the size of padding at each side of the peaks median position to create the consensus region. The minimum size of the consensus region will be equal to twice the value of the <code>extendingSize</code> parameter. The size of the <code>extendingSize</code> must be a positive integer. Default = 250.
<code>expandToFitPeakRegion</code>	a logical indicating if the region set by the <code>extendingSize</code> parameter is extended to include all region of the peak closest to the peaks median position for each experiment.
<code>shrinkToFitPeakRegion</code>	a logical indicating if the region size, which is set by the <code>extendingSize</code> parameter is shrunk to fit the narrow peak regions of the peaks when all those regions are smaller than the consensus region.
<code>minNbrExp</code>	a numeric indicating the minimum number of BED files in which a peak must be present for a region to be retained. The numeric must be a positive value inferior or equal to the number of files present in the <code>narrowPeakFiles</code> parameter. Default = 1.
<code>nbrThreads</code>	a numeric indicating the number of threads to use in parallel.

**Value**

0 indicating that all parameters validations have been successful.

**Author(s)**

Astrid Deschênes

**Examples**

```
## Loading datasets
data(A549_FOSL2_01_NarrowPeaks_partial)
data(A549_FOXA1_01_NarrowPeaks_partial)
data(A549_FOSL2_01_Peaks_partial)
data(A549_FOXA1_01_Peaks_partial)

## Assigning experiment name to each row of the dataset.
## NarrowPeak and Peak datasets from the same experiment must
## have identical names.
names(A549_FOXA1_01_Peaks_partial) <- rep("FOXA1_01",
                                         length(A549_FOXA1_01_Peaks_partial))
names(A549_FOXA1_01_NarrowPeaks_partial) <- rep("FOXA1_01",
                                                length(A549_FOXA1_01_NarrowPeaks_partial))
names(A549_FOSL2_01_Peaks_partial) <- rep("FOSL2_01",
                                           length(A549_FOSL2_01_Peaks_partial))
names(A549_FOSL2_01_NarrowPeaks_partial) <- rep("FOSL2_01",
                                                length(A549_FOSL2_01_NarrowPeaks_partial))

chrList <- Seqinfo("chr10", 135534747, NA)

consensusSeeker::findConsensusPeakRegionsValidation(
  narrowPeaks = c(A549_FOXA1_01_NarrowPeaks_partial,
                 A549_FOSL2_01_NarrowPeaks_partial),
  peaks = c(A549_FOXA1_01_Peaks_partial,
            A549_FOSL2_01_Peaks_partial),
  chrList = chrList,
  extendingSize = 110,
  expandToFitPeakRegion = FALSE,
  shrinkToFitPeakRegion = TRUE,
  minNbrExp = 2,
  nbrThreads = 1)
```

---

isInteger

*Validate if a value is an integer*

---

**Description**

Validate if the value passed to the function is an integer or not. To be considered as an integer, the value must have a length of 1. The type of value can be a integer or numerical. However, a numerical must have the same value once casted to a integer. A vector of integers will returned FALSE.

**Usage**

```
isInteger(value)
```

**Arguments**

value            an object to validate.

**Value**

TRUE is the parameter is a integer; otherwise FALSE

**Author(s)**

Astrid Deschênes

---

NOrMAL\_nucleosome\_positions

*Nucleosome positions detected by the NOrMAL software using synthetic reads generated using a normal distribution. For demonstration purpose.*

---

**Description**

Nucleosome positions detected by the NOrMAL software using synthetic reads generated using a normal distribution with a variance of 20 for regions chr1:10000-15000.

**Usage**

```
data(NOrMAL_nucleosome_positions)
```

**Format**

A GRanges containing one entry per detected nucleosome. The surrounding ranges associated to those nucleosomes are in the dataset NOrMAL\_nucleosome\_ranges.

**References**

- Polishko A, Ponts N, Le Roch KG and Lonardi S. 2012. NOrMAL: Accurate nucleosome positioning using a modified Gaussian mixture model. *Bioinformatics* 28 (12): 242-49.

**See Also**

- [NOrMAL\\_nucleosome\\_ranges](#) the associate genomic ranges dataset.
- [findConsensusPeakRegions](#) for extracting regions sharing nucleosomes from more than one experiment.

## Examples

```
## Loading datasets
data(PING_nucleosome_positions)
data(PING_nucleosome_ranges)
data(NOrMAL_nucleosome_positions)
data(NOrMAL_nucleosome_ranges)
data(NucPosSimulator_nucleosome_positions)
data(NucPosSimulator_nucleosome_ranges)

## Assigning experiment name to each row of the dataset.
## Position and range datasets from the same software must
## have identical names.
names(PING_nucleosome_positions) <- rep("PING",
                                       length(PING_nucleosome_positions))
names(PING_nucleosome_ranges) <- rep("PING",
                                       length(PING_nucleosome_ranges))
names(NOrMAL_nucleosome_positions) <- rep("NOrMAL",
                                           length(NOrMAL_nucleosome_positions))
names(NOrMAL_nucleosome_ranges) <- rep("NOrMAL",
                                       length(NOrMAL_nucleosome_ranges))
names(NucPosSimulator_nucleosome_positions) <- rep("NucPosSimulator",
                                                    length(NucPosSimulator_nucleosome_positions))
names(NucPosSimulator_nucleosome_ranges) <- rep("NucPosSimulator",
                                                length(NucPosSimulator_nucleosome_ranges))

## Calculating consensus regions for chromosome 1
## with a default region size of 40 bp (2 * extendingSize).
## The consensus regions are extended to include all genomic regions for
## all nucleosomes. However, if the consensus regions are larger than the
## genomic regions of the nucleosomes, the consensus regions are not
## shrunked.
## Nucleosomes from all software must be present in a region to
## be retained as a consensus region.
chrList <- Seqinfo(c("chr1"), c(249250621), NA)
findConsensusPeakRegions(
  narrowPeaks = c(PING_nucleosome_ranges,
                 NOrMAL_nucleosome_ranges,
                 NucPosSimulator_nucleosome_ranges),
  peaks = c(PING_nucleosome_positions,
            NOrMAL_nucleosome_positions,
            NucPosSimulator_nucleosome_positions),
  chrInfo = chrList,
  extendingSize = 20,
  expandToFitPeakRegion = TRUE,
  shrinkToFitPeakRegion = FALSE,
  minNbrExp = 3,
  nbrThreads = 1)
```

---



NOrMAL\_nucleosome\_ranges

*Ranges associated to nucleosomes detected by the NOrMAL software using synthetic reads generated using a normal distribution. For demonstration purpose.*

---

## Description

Ranges associated to nucleosomes detected by the NOrMAL software using synthetic reads generated using a normal distribution with a variance of 20 for regions chr1:10000-15000.

## Usage

```
data(NOrMAL_nucleosome_ranges)
```

## Format

A GRanges containing one entry per detected nucleosome. The ranges are surrounding the nucleosomes present in the dataset NOrMAL\_nucleosome\_positions. The genomic ranges have been obtained by adding 73 bps on each side of the detected positions.

## References

- Polishko A, Ponts N, Le Roch KG and Lonardi S. 2012. NOrMAL: Accurate nucleosome positioning using a modified Gaussian mixture model. *Bioinformatics* 28 (12): 242-49.

## See Also

- [NOrMAL\\_nucleosome\\_positions](#) the associate genomic positions dataset.
- [findConsensusPeakRegions](#) for extracting regions sharing nucleosomes from more than one experiment.

## Examples

```
## Loading datasets
data(PING_nucleosome_positions)
data(PING_nucleosome_ranges)
data(NOrMAL_nucleosome_positions)
data(NOrMAL_nucleosome_ranges)
data(NucPosSimulator_nucleosome_positions)
data(NucPosSimulator_nucleosome_ranges)

## Assigning experiment name to each row of the dataset.
## Position and range datasets from the same software must
## have identical names.
names(PING_nucleosome_positions) <- rep("PING",
                                         length(PING_nucleosome_positions))
names(PING_nucleosome_ranges) <- rep("PING",
                                       length(PING_nucleosome_ranges))
names(NOrMAL_nucleosome_positions) <- rep("NOrMAL",
```

```

                                length(NORMAL_nucleosome_positions))
names(NORMAL_nucleosome_ranges) <- rep("NORMAL",
                                length(NORMAL_nucleosome_ranges))
names(NucPosSimulator_nucleosome_positions) <-rep("NucPosSimulator",
                                length(NucPosSimulator_nucleosome_positions))
names(NucPosSimulator_nucleosome_ranges) <- rep("NucPosSimulator",
                                length(NucPosSimulator_nucleosome_ranges))

## Calculating consensus regions for chromosome 1
## with a default region size of 30 bp (2 * extendingSize).
## Consensus regions are resized to include all genomic regions of
## included nucleosomes.
## Nucleosomes from at least 2 software must be present
## in a region to be retained as a consensus region.
chrList <- Seqinfo(c("chr1"), c(249250621), NA)
findConsensusPeakRegions(
  narrowPeaks = c(PING_nucleosome_ranges,
                  NORMAL_nucleosome_ranges,
                  NucPosSimulator_nucleosome_ranges),
  peaks = c(PING_nucleosome_positions,
            NORMAL_nucleosome_positions,
            NucPosSimulator_nucleosome_positions),
  chrInfo = chrList,
  extendingSize = 15,
  expandToFitPeakRegion = TRUE,
  shrinkToFitPeakRegion = TRUE,
  minNbrExp = 2,
  nbrThreads = 1)

```

---

NucPosSimulator\_nucleosome\_positions

*Nucleosome positions detected by the NucPosSimulator software using synthetic reads generated using a normal distribution. For demonstration purpose.*

---

## Description

Nucleosome positions detected by the NucPosSimulator software using synthetic reads generated using a normal distribution with a variance of 20 for regions chr1:10000-15000.

## Usage

```
data(NucPosSimulator_nucleosome_positions)
```

## Format

A GRanges containing one entry per detected nucleosome. The surrounding ranges associated to those nucleosomes are in the dataset NucPosSimulator\_nucleosome\_ranges.

## References

- Schöpfung R, Teif VB, Müller O, Weinberg C, Rippe K, and Wedemann G. 2013. Modeling nucleosome position distributions from experimental nucleosome positioning maps. *Bioinformatics* 29 (19): 2380-86.

## See Also

- [NucPosSimulator\\_nucleosome\\_ranges](#) the associate genomic ranges dataset.
- [findConsensusPeakRegions](#) for extracting regions sharing nucleosomes from more than one experiment.

## Examples

```
## Loading datasets
data(PING_nucleosome_positions)
data(PING_nucleosome_ranges)
data(NOrMAL_nucleosome_positions)
data(NOrMAL_nucleosome_ranges)
data(NucPosSimulator_nucleosome_positions)
data(NucPosSimulator_nucleosome_ranges)

## Assigning experiment name to each row of the dataset.
## Position and range datasets from the same software must
## have identical names.
names(PING_nucleosome_positions) <- rep("PING",
                                         length(PING_nucleosome_positions))
names(PING_nucleosome_ranges) <- rep("PING",
                                       length(PING_nucleosome_ranges))
names(NOrMAL_nucleosome_positions) <- rep("NOrMAL",
                                           length(NOrMAL_nucleosome_positions))
names(NOrMAL_nucleosome_ranges) <- rep("NOrMAL",
                                        length(NOrMAL_nucleosome_ranges))
names(NucPosSimulator_nucleosome_positions) <- rep("NucPosSimulator",
                                                    length(NucPosSimulator_nucleosome_positions))
names(NucPosSimulator_nucleosome_ranges) <- rep("NucPosSimulator",
                                                 length(NucPosSimulator_nucleosome_ranges))

## Calculating consensus regions for chromosome 1
## with a default region size of 50 bp (2 * extendingSize).
## The consensus regions are extended to include all genomic regions for
## all nucleosomes. However, if the consensus regions are larger than the
## genomic regions of the nucleosomes, the consensus regions are not
## shrunked.
## Nucleosomes from all software must be present in a region to
## be retained as a consensus region.
chrList <- Seqinfo(c("chr1"), c(249250621), NA)
findConsensusPeakRegions(
  narrowPeaks = c(PING_nucleosome_ranges,
                 NOrMAL_nucleosome_ranges,
                 NucPosSimulator_nucleosome_ranges),
  peaks = c(PING_nucleosome_positions,
```

```
        NORMAL_nucleosome_positions,  
        NucPosSimulator_nucleosome_positions),  
chrInfo = chrList,  
extendingSize = 25,  
expandToFitPeakRegion = TRUE,  
shrinkToFitPeakRegion = FALSE,  
minNbrExp = 3,  
nbrThreads = 1)
```

---

NucPosSimulator\_nucleosome\_ranges

*Ranges associated to nucleosomes detected by the NucPosSimulator software using synthetic reads generated using a normal distribution. For demonstration purpose.*

---

## Description

Ranges associated to nucleosomes detected by the NucPosSimulator software using synthetic reads generated using a normal distribution with a variance of 20 for regions chr1:10000-15000.

## Usage

```
data(NucPosSimulator_nucleosome_ranges)
```

## Format

A GRanges containing one entry per detected nucleosome. The ranges are surrounding the nucleosomes present in the dataset NucPosSimulator\_nucleosome\_positions. The genomic ranges have been obtained by adding 73 bps on each side of the detected positions.

## References

- Schönpfing R, Teif VB, Müller O, Weinberg C, Rippe K, and Wedemann G. 2013. Modeling nucleosome position distributions from experimental nucleosome positioning maps. *Bioinformatics* 29 (19): 2380-86.

## See Also

- [NucPosSimulator\\_nucleosome\\_positions](#) the associate genomic positions dataset.
- [findConsensusPeakRegions](#) for extracting regions sharing nucleosomes from more than one experiment.

**Examples**

```
## Loading datasets
data(PING_nucleosome_positions)
data(PING_nucleosome_ranges)
data(NOrMAL_nucleosome_positions)
data(NOrMAL_nucleosome_ranges)
data(NucPosSimulator_nucleosome_positions)
data(NucPosSimulator_nucleosome_ranges)

## Assigning experiment name to each row of the dataset.
## Position and range datasets from the same software must
## have identical names.
names(PING_nucleosome_positions) <- rep("PING",
                                         length(PING_nucleosome_positions))
names(PING_nucleosome_ranges) <- rep("PING",
                                      length(PING_nucleosome_ranges))
names(NOrMAL_nucleosome_positions) <- rep("NOrMAL",
                                           length(NOrMAL_nucleosome_positions))
names(NOrMAL_nucleosome_ranges) <- rep("NOrMAL",
                                        length(NOrMAL_nucleosome_ranges))
names(NucPosSimulator_nucleosome_positions) <- rep("NucPosSimulator",
                                                    length(NucPosSimulator_nucleosome_positions))
names(NucPosSimulator_nucleosome_ranges) <- rep("NucPosSimulator",
                                                length(NucPosSimulator_nucleosome_ranges))

## Calculating consensus regions for chromosome 1
## with a default region size of 60 bp (2 * extendingSize).
## Consensus regions are resized to include all genomic regions of
## included nucleosomes.
## Nucleosomes from at least 2 software must be present
## in a region to be retained as a consensus region.
chrList <- Seqinfo(c("chr1"), c(249250621), NA)
findConsensusPeakRegions(
  narrowPeaks = c(PING_nucleosome_ranges,
                 NOrMAL_nucleosome_ranges,
                 NucPosSimulator_nucleosome_ranges),
  peaks = c(PING_nucleosome_positions,
            NOrMAL_nucleosome_positions,
            NucPosSimulator_nucleosome_positions),
  chrInfo = chrList,
  extendingSize = 30,
  expandToFitPeakRegion = TRUE,
  shrinkToFitPeakRegion = TRUE,
  minNbrExp = 2,
  nbrThreads = 1)
```

---

PING\_nucleosome\_positions

*Nucleosome positions detected by the PING software using synthetic reads generated using a normal distribution. For demonstration purpose.*

---

## Description

Nucleosome positions detected by the PING software using synthetic reads generated using a normal distribution with a variance of 20 for regions chr1:10000-15000.

## Usage

```
data(PING_nucleosome_positions)
```

## Format

A GRanges containing one entry per detected nucleosome. The surrounding ranges associated to those nucleosomes are in the dataset PING\_nucleosome\_positions.

## References

- Sangsoon W, Zhang X, Sauteraud R, Robert F and Gottardo R. 2013. PING 2.0: An R/Bioconductor package for nucleosome positioning using next-generation sequencing data. *Bioinformatics* 29 (16): 2049-50.

## See Also

- [PING\\_nucleosome\\_ranges](#) the associate genomic ranges dataset.
- [findConsensusPeakRegions](#) for extracting regions sharing nucleosomes from more than one experiment.

## Examples

```
## Loading datasets
data(PING_nucleosome_positions)
data(PING_nucleosome_ranges)
data(NOrMAL_nucleosome_positions)
data(NOrMAL_nucleosome_ranges)
data(NucPosSimulator_nucleosome_positions)
data(NucPosSimulator_nucleosome_ranges)

## Assigning experiment name to each row of the dataset.
## Position and range datasets from the same software must
## have identical names.
names(PING_nucleosome_positions) <- rep("PING",
                                         length(PING_nucleosome_positions))
names(PING_nucleosome_ranges) <- rep("PING",
                                       length(PING_nucleosome_ranges))
names(NOrMAL_nucleosome_positions) <- rep("NOrMAL",
```

```

                                length(NOrMAL_nucleosome_positions))
names(NOrMAL_nucleosome_ranges) <- rep("NOrMAL",
                                length(NOrMAL_nucleosome_ranges))
names(NucPosSimulator_nucleosome_positions) <-rep("NucPosSimulator",
                                length(NucPosSimulator_nucleosome_positions))
names(NucPosSimulator_nucleosome_ranges) <- rep("NucPosSimulator",
                                length(NucPosSimulator_nucleosome_ranges))

## Calculating consensus regions for chromosome 1
## with a default region size of 20 bp (2 * extendingSize).
## The consensus regions are not resized to fit genomic ranges of the
## included nucleosomes.
## Nucleosomes from at least 2 software must be present in a region to
## be retained as a consensus region.
chrList <- Seqinfo(c("chr1"), c(249250621), NA)
findConsensusPeakRegions(
  narrowPeaks = c(PING_nucleosome_ranges,
                  NOrMAL_nucleosome_ranges,
                  NucPosSimulator_nucleosome_ranges),
  peaks = c(PING_nucleosome_positions,
            NOrMAL_nucleosome_positions,
            NucPosSimulator_nucleosome_positions),

  chrInfo = chrList,
  extendingSize = 10,
  expandToFitPeakRegion = FALSE,
  shrinkToFitPeakRegion = FALSE,
  minNbrExp = 3,
  nbrThreads = 1)

```

---

PING\_nucleosome\_ranges

*Ranges associated to nucleosomes detected by the PING software using synthetic reads generated using a normal distribution. For demonstration purpose.*

---

## Description

Ranges associated to nucleosomes detected by the PING software using synthetic reads generated using a normal distribution with a variance of 20 for regions chr1:10000-15000.

## Usage

```
data(PING_nucleosome_ranges)
```

## Format

A GRanges containing one entry per detected nucleosome. The ranges are surrounding the nucleosomes present in the dataset PING\_nucleosome\_positions. The genomic ranges have been obtained by adding 73 bps on both sides of the detected positions.

## References

- Sangsoo W, Zhang X, Sauteraud R, Robert F and Gottardo R. 2013. PING 2.0: An R/Bioconductor package for nucleosome positioning using next-generation sequencing data. *Bioinformatics* 29 (16): 2049-50.

## See Also

- [PING\\_nucleosome\\_positions](#) the associate genomic positions dataset.
- [findConsensusPeakRegions](#) for extracting regions sharing nucleosomes from more than one experiment.

## Examples

```
## Loading datasets
data(PING_nucleosome_positions)
data(PING_nucleosome_ranges)
data(NOrMAL_nucleosome_positions)
data(NOrMAL_nucleosome_ranges)
data(NucPosSimulator_nucleosome_positions)
data(NucPosSimulator_nucleosome_ranges)

## Assigning experiment name to each row of the dataset.
## Position and range datasets from the same software must
## have identical names.
names(PING_nucleosome_positions) <- rep("PING",
                                         length(PING_nucleosome_positions))
names(PING_nucleosome_ranges) <- rep("PING",
                                       length(PING_nucleosome_ranges))
names(NOrMAL_nucleosome_positions) <- rep("NOrMAL",
                                           length(NOrMAL_nucleosome_positions))
names(NOrMAL_nucleosome_ranges) <- rep("NOrMAL",
                                        length(NOrMAL_nucleosome_ranges))
names(NucPosSimulator_nucleosome_positions) <- rep("NucPosSimulator",
                                                    length(NucPosSimulator_nucleosome_positions))
names(NucPosSimulator_nucleosome_ranges) <- rep("NucPosSimulator",
                                                length(NucPosSimulator_nucleosome_ranges))

## Calculating consensus regions for chromosome 1
## with a default region size of 20 bp (2 * extendingSize).
## which is not extended to include all genomic regions for the closest
## peak to the median position of all peaks included in the region (for
## each experiment).
## Nucleosomes from at least 2 software must be present in a region to
## be retained as a consensus region.
chrList <- Seqinfo(c("chr1"), c(249250621), NA)
findConsensusPeakRegions(
  narrowPeaks = c(PING_nucleosome_ranges,
                 NOrMAL_nucleosome_ranges,
                 NucPosSimulator_nucleosome_ranges),
  peaks = c(PING_nucleosome_positions,
            NOrMAL_nucleosome_positions,
```



```
                                NucPosSimulator_nucleosome_positions),
chrInfo = chrList,
extendingSize = 10,
expandToFitPeakRegion = FALSE,
shrinkToFitPeakRegion = FALSE,
minNbrExp = 2,
nbrThreads = 1)
```

---

readNarrowPeakFile      *Extract narrow regions and peaks from a narrowPeak file*

---

### Description

Read a narrowPeak file and extract the narrow regions and/or the peaks, as specified by used. The narrowPeak file must fit the UCSC specifications. See <https://genome.ucsc.edu/FAQ/FAQformat.html#format12> for more details. The file can have one or many header lines. However, the total number of header lines must be inferior to 250 lines.

### Usage

```
readNarrowPeakFile(file_path, extractRegions = TRUE, extractPeaks = TRUE)
```

### Arguments

file_path	the name of the file.
extractRegions	a logical indicating if the narrow regions must be extracted. If TRUE, a GRanges containing the narrow regions will be returned. Otherwise, NULL is returned. Default = TRUE.
extractPeaks	a logical indicating if the peaks must be extracted. If TRUE, a GRanges containing the peaks will be returned. Otherwise, NULL is returned. Default = TRUE.

### Value

a list containing 2 entries:

- narrowPeak a GRanges containing the narrow regions extracted from the file. NULL when not needed by user.
- peak a GRanges containing the peaks extracted from the file. NULL when not

### Author(s)

Astrid Deschênes

**Examples**

```
## Set file information
test_narrowPeak <- system.file("extdata",
                              "A549_FOSL2_ENCSR000BQ0_MZW_part_chr_1_and_12.narrowPeak",
                              package = "consensusSeeker")

## Read file to extract peaks and regions
data <- readNarrowPeakFile(test_narrowPeak, extractRegions = TRUE,
                          extractPeaks = TRUE)

## To access peak data (GRanges format)
head(data$peak)

## To access region data (GRanges format)
head(data$narrowPeak)
```

---

refineRegion	<i>Refine the selected region by using an iterative process.</i>
--------------	--

---

**Description**

Find regions sharing the same features for a minimum number of experiments using called peaks of signal enrichment based on pooled, normalized data (mainly coming from narrowPeak files). The analysis is limited to one chromosome. The peaks and narrow peaks are used to identify the consensus regions. The minimum number of experiments that must have at least one peak in a region so that it is retained as a consensus region is specified by user, as well as the size of mining regions.

**Usage**

```
refineRegion(peaks, setPeaks, extendingSize, region_width, currentPeak)
```

**Arguments**

peaks	a GRanges containing all peaks from the current analysis sorted by position.
setPeaks	a GRanges containing the peaks present in the current selected region.
extendingSize	a numeric value indicating the size of padding at each side of the peaks median position to create the consensus region. The minimum size of the consensus region will be equal to twice the value of the extendingSize parameter. The size of the extendingSize must be a positive integer.
region_width	a numeric value indicating the size of the region which is equivalent to twice the extendingSize.
currentPeak	a GRanges with 1 range containing the information about the current peak used as the starting point for the selected region.

**Value**

an object of class "commonFeatures".

*refineRegion*

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**Author(s)**

Astrid Deschênes

# Index

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