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# **bedparse Documentation**

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Bedparse is a simple python module and CLI tool to perform common operations on BED files.

It offers the following functionality:

- BED format validation
- Filtering of transcripts based on annotations
- Joining of annotation files based on transcript names
- Conversion from GTF to BED format
- Conversion from UCSC to Ensembl chromosome names (and viceversa)
- Conversion from bed12 to bed6
- Promoter reporting
- Intron reporting
- CDS reporting
- UTR reporting



### 1.1 Installation

Bedparse is distributed on PyPI. To install, just run:

```
pip install bedparse
```

Alternatively, to install it from the Github repository:

```
pip install git+https://github.com/tleonardi/bedparse.git
```

### 1.2 Motivation

The BED (Browser Extensible Data) format is a plain text file format commonly used in bioinformatics to represent genomic features (e.g. genes, transcripts, peaks, regulatory regions, etc.). Each line in the file represents a genomic feature and consists of up to 12 tab-separated fields:

1. chromosome name
2. start coordinate in the chromosome
3. end coordinate in the chromosome
4. feature name
5. feature score
6. strand
7. thick start (conventionally the start codon for protein coding transcripts)
8. thick end (conventionally the stop codon for protein coding transcripts)
9. rgb color for visualisation in genome browsers
10. number of connected blocks (conventionally the number of exons)

11. comma separated list of blocks size
12. comma separated list of block starts relative to field 2 (i.e. genomic start of the feature)

One of the major advantages of the BED format over many of its alternatives is that each line includes all the information required to define an individual transcript. This characteristic allows to perform numerous operations on BED a file as part of unix pipes, for example using GNU awk.

For example, the following is a common approach to extract gene promoters (here defined as 500bp around the gene start):

```
awk 'BEGIN{OFS=FS="\t"}{print $1,$2-500,$3+500,$4,$5}' transcriptome.bed > promoters.  
↪bed
```

However, these one-liners can quickly get long and hard to read. For example, if we wanted to do the same as before but keeping the strand into considerations:

```
awk 'BEGIN{OFS=FS="\t"}{if($6=="+") {print $1,$2-500,$2+500,$4,$5}else{print $1,$3-500,  
↪$3+500,$4,$5}}' transcriptome.bed > promoters_stranded.bed
```

These and other more complex operations quickly get long to type and prone to errors and typos. Bedparse greatly simplifies the process:

```
bedparse promoter transcriptome.bed > promoters_stranded.bed
```

or:

```
bedparse promoter --unstranded transcriptome.bed > promoters.bed
```

Despite the simplicity of most of its operations, all functions in bedparse are thoroughly and rigorously tested through an automated test suit to ensure the accuracy and correctness of the results. Additionally, bedparse performs syntax validation checks on the input BED files and warns the user in case of malformed or unsupported formats.

Additionally, bedparse also provides two format conversion operations:

- gtf2bed allows converting Ensembl/Gencode Gene transfer format (GTF) files into bed format
- convertChr implements an internal dictionary that allows conversion of human and mouse chromosome names between the two most widely used formats, i.e. the Ensembl and the UCSC naming schemes.

## 1.3 Usage

```
usage: bedparse [-h] [--version]  
              {3pUTR,5pUTR,cds,promoter,introns,filter,join,gtf2bed,bed12tobed6,  
↪convertChr,validateFormat}  
              ...
```

Perform various simple operations on BED files.

positional arguments:

```
{3pUTR,5pUTR,cds,promoter,introns,filter,join,gtf2bed,bed12tobed6,convertChr,  
↪validateFormat}
```

	sub-command help
3pUTR	Prints the 3' of coding genes.
5pUTR	Prints the 5' of coding genes.
cds	Prints the CDS of coding genes.
promoter	Prints the promoters of transcripts.

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introns	Prints BED records corresponding to the introns of each transcript in the original file.
filter	Filters a BED file based on an annotation.
join	Joins a BED file with an annotation file using the BED name (col4) as the joining key.
gtf2bed	Converts a GTF file to BED12 format.
bed12tobed6	Converts a BED12 file to BED6 format
convertChr	Convert chromosome names between UCSC and Ensembl formats
validateFormat	Check whether the BED file adheres to the BED format specifications
optional arguments:	
-h, --help	show this help message and exit
--version, -v	show program's version number and exit

The basic syntax in the form: `bedparse sub-command [parameters]`.

For a list of all sub-commands and a brief explanation of what they do, use: `bedparse --help`

For a detailed explanation of each subcommand and a list of its parameters, use the `--help` option after the subcommand's name, e.g.: `bedparse promoter --help`

### 1.3.1 3'/5' UTRs

#### Usage

```
> bedparse 3pUTR --help
usage: bedparse 3pUTR [-h] [bedfile]
```

Report the 5' or 3' UTRs of each coding transcript in the BED file.

UTRs are defined as the region between transcript start/end and CDS start/end (the CDS is in turn defined as the region between `thickStart` and `thickEnd`).

Transcripts with an undefined CDS (i.e. with `thickStart` and `thickEnd` set to the same value) are not reported.

#### Examples

```
> cat transcripts.bed
chr1      167721988      167790819      ENST00000392121.
↪7        0            +            167722151      167787921      0            3            254,
↪167,3000,      0,43594,65831,

> bedparse 3pUTR transcripts.bed
chr1      167787921      167790819      ENST00000392121.
↪7        0            +            167787921      167787921      0            1            2898,
↪        0,
```

## 1.3.2 CDS

### Usage

```
> bedparse cds --help
usage: bedparse cds [-h] [--ignoreCDSonly] [bedfile]

Report the CDS of each coding transcript (i.e. transcripts with distinct
values of thickStart and thickEnd). Transcripts without CDS are not reported.

positional arguments:
  bedfile          Path to the BED file.

optional arguments:
  -h, --help      show this help message and exit
  --ignoreCDSonly Ignore transcripts that only consist of CDS.
```

### Examples

```
> cat transcripts.bed
chr1      167721988      167790819      ENST00000392121.
↪7        0              +              167722151      167787921      0              3              254,
↪167,3000,      0,43594,65831,

> bedparse cds transcripts.bed
chr1      167722151      167787921      ENST00000392121.
↪7        0              +              167722151      167787921      0              3              91,
↪167,102,      0,43431,65668,
```

---

## 1.3.3 Promoters

This command reports the promoter of each transcript in the input BED file. The promoter is defined as a fixed interval around the TSS.

### Usage

```
> bedparse promoter --help
usage: bedparse promoter [-h] [--up UP] [--down DOWN] [--unstranded] [bedfile]

Report the promoter of each transcript, defined as a fixed interval around its
start.

positional arguments:
  bedfile          Path to the BED file.

optional arguments:
  -h, --help      show this help message and exit
  --up UP         Get this many nt upstream of each feature.
  --down DOWN     Get this many nt downstream of each feature.
  --unstranded    Do not consider strands.
```

## Examples

```
> cat transcripts.bed
chr1      167721988      167790819      ENST00000392121.
↪7        0              +              167722151      167787921      0              3              254,
↪167,3000,      0,43594,65831,
```

```
> bedparse promoter transcripts.bed
chr1      167721488      167722488      ENST00000392121.7
```

```
> bedparse promoter --up 100 --down 100 transcripts.bed
chr1      167721888      167722088      ENST00000392121.7
```

## 1.3.4 Introns

Reports BED12 lines corresponding to the introns of each transcript. Unspliced transcripts are not reported.

### Usage

```
> bedparse introns --help
usage: bedparse introns [-h] [bedfile]

Report BED12 lines corresponding to the introns of each transcript. Unspliced
transcripts are not reported.

positional arguments:
  bedfile      Path to the BED file.

optional arguments:
  -h, --help  show this help message and exit
```

## Examples

```
> cat transcripts.bed
chr1      167721988      167790819      ENST00000392121.
↪7        0              +              167722151      167787921      0              3              254,
↪167,3000,      0,43594,65831,
```

```
> bedparse introns transcripts.bed
chr1      167722242      167787819      ENST00000392121.
↪7        0              +              167722242      167722242      0              2              43340,
↪22070,      0,43507,
```

## 1.3.5 Filter

Filters a BED file based on an annotation file. BED entries with a name (i.e. col4) that appears in the specified column of the annotation are printed to stdout. For efficiency reasons this command doesn't perform BED validation.

## Usage

```
> bedparse filter --help
usage: bedparse filter [-h] --annotation ANNOTATION [--column COLUMN]
                        [--inverse]
                        [bedfile]

Filters a BED file based on an annotation. BED entries with a name (i.e. col4)
that appears in the specified column of the annotation are printed to stdout.
For efficiency reasons this command doesn't perform BED validation.

positional arguments:
  bedfile                Path to the BED file.

optional arguments:
  -h, --help            show this help message and exit
  --annotation ANNOTATION, -a ANNOTATION
                        Path to the annotation file.
  --column COLUMN, -c COLUMN
                        Column of the annotation file (1-based, default=1).
  --inverse, -v         Only report BED entries absent from the annotation
                        file.
```

## Examples

```
> cat transcripts.bed
chr1      67092164      67231852      ENST00000371007.6      0      -
chr1      67092175      67127261      ENST00000371006.5      0      -
chr1      67092175      67127261      ENST00000475209.6      0      -
chr1      67092394      67134970      ENST00000371004.6      0      -
chr1      67092396      67127261      ENST00000621590.4      0      -
chr1      67092947      67134977      ENST00000544837.5      0      -
chr1      67093558      67231853      ENST00000448166.6      0      -
chr1      67096295      67134977      ENST00000603691.1      0      -
chr1      201283451     201332993     ENST00000263946.7      0      +
chr1      201283451     201332993     ENST00000367324.7      0      +

> cat filter.txt
GeneX      ENST00000263946.7      Other_field
GeneY      ENST00000367324.7      Another_field

> bedparse filter --annotation filter.txt --column 2 transcripts.bed
chr1      201283451     201332993     ENST00000263946.7      0      +
chr1      201283451     201332993     ENST00000367324.7      0      +
```

---

### 1.3.6 Join

Adds the content of an annotation file to a BED file as extra columns. The two files are joined by matching the BED Name field (column 4) with a user-specified field of the annotation file.

## Usage

```
> bedparse join --help
usage: bedparse join [-h] --annotation ANNOTATION [--column COLUMN]
      [--separator SEPARATOR] [--empty EMPTY] [--noUnmatched]
      [bedfile]

Adds the content of an annotation file to a BED file as extra columns. The two
files are joined by matching the BED Name field (column 4) with a user-
specified field of the annotation file.

positional arguments:
  bedfile                Path to the BED file.

optional arguments:
  -h, --help            show this help message and exit
  --annotation ANNOTATION, -a ANNOTATION
                        Path to the annotation file.
  --column COLUMN, -c COLUMN
                        Column of the annotation file (1-based, default=1).
  --separator SEPARATOR, -s SEPARATOR
                        Field separator for the annotation file (default tab)
  --empty EMPTY, -e EMPTY
                        String to append to empty records (default '.').
  --noUnmatched, -n    Do not print unmatched lines.
```

## Examples

```
> cat transcripts.bed
chr1      67092164      67231852      ENST00000371007.6      0      -
chr1      67092175      67127261      ENST00000371006.5      0      -
chr1      67092175      67127261      ENST00000475209.6      0      -
chr1      67092394      67134970      ENST00000371004.6      0      -
chr1      67092396      67127261      ENST00000621590.4      0      -
chr1      67092947      67134977      ENST00000544837.5      0      -
chr1      67093558      67231853      ENST00000448166.6      0      -
chr1      67096295      67134977      ENST00000603691.1      0      -
chr1      201283451      201332993      ENST00000263946.7      0      +
chr1      201283451      201332993      ENST00000367324.7      0      +
```

```
> cat annotation.txt
GeneX      ENST00000263946.7      Other_field
GeneY      ENST00000367324.7      Another_field
```

```
> bedparse join --column 2 --annotation annotation.txt transcripts.bed
chr1      67092164      67231852      ENST00000371007.6      0      -
↪      .
chr1      67092175      67127261      ENST00000371006.5      0      -
↪      .
chr1      67092175      67127261      ENST00000475209.6      0      -
↪      .
chr1      67092394      67134970      ENST00000371004.6      0      -
↪      .
chr1      67092396      67127261      ENST00000621590.4      0      -
↪      .
chr1      67092947      67134977      ENST00000544837.5      0      -
↪      .
```

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```

chr1      67093558      67231853      ENST00000448166.6      0      -
↪
chr1      67096295      67134977      ENST00000603691.1      0      -
↪
chr1      201283451      201332993      ENST00000263946.
↪7      0      +      GeneX      Other_field
chr1      201283451      201332993      ENST00000367324.
↪7      0      +      GeneY      Another_field

> bedparse join --column 2 --annotation annotation.txt --noUnmatched transcripts.bed
chr1      201283451      201332993      ENST00000263946.
↪7      0      +      GeneX      Other_field
chr1      201283451      201332993      ENST00000367324.
↪7      0      +      GeneY      Another_field

```

### 1.3.7 Convert GTF to BED

Converts a GTF file to BED12 format. This tool supports the Ensembl GTF format. The GTF file must contain 'transcript' and 'exon' features in field 3. If the GTF file also annotates 'CDS' 'start\_codon' or 'stop\_codon' these are used to annotate the thickStart and thickEnd in the BED file.

#### Usage

```

> bedparse gtf2bed --help
usage: bedparse gtf2bed [-h] [--extraFields EXTRAFIELDS]
                        [--filterKey FILTERKEY] [--filterType FILTERTYPE]
                        [gtf]

Converts a GTF file to BED12 format. This tool supports the Ensembl GTF
format. The GTF file must contain 'transcript' and 'exon' features in field 3.
If the GTF file also annotates 'CDS' 'start_codon' or 'stop_codon' these are
used to annotate the thickStart and thickEnd in the BED file.

positional arguments:
  gtf                Path to the GTF file.

optional arguments:
  -h, --help        show this help message and exit
  --extraFields EXTRAFIELDS
                    Comma separated list of extra GTF fields to be added
                    after col 12 (e.g. gene_id, gene_name).
  --filterKey FILTERKEY
                    GTF extra field on which to apply the filtering
  --filterType FILTERTYPE
                    Comma separated list of filterKey field values to
                    retain.

```

### 1.3.8 Convert BED12 to BED6

Convert the BED12 format into BED6 by reporting a separate line for each block of the original record.

## Usage

```
> bedparse bed12tobed6 --help
usage: bedparse bed12tobed6 [-h] [--appendExN] [--whichExon {all,first,last}]
                               [--keepIntrons]
                               [bedfile]

Convert the BED12 format into BED6 by reporting a separate line for each block
of the original record.

positional arguments:
  bedfile                Path to the GTF file.

optional arguments:
  -h, --help            show this help message and exit
  --appendExN           Appends the exon number to the transcript name.
  --whichExon {all,first,last}
                        Which exon to return. First and last respectively
                        report the first or last exon relative to the TSS
                        (i.e. taking strand into account).
  --keepIntrons        Add records for introns as well. Only allowed if
                        --whichExon all
```

## Examples

```
> cat transcripts.bed
chr1      67092164      67231852      ENST00000371007.6      0      -
↪      67093004      67127240      0      8      1440,187,70,113,158,92,86,
↪7,      0,3070,4087,23187,33587,35001,38977,139681,

> bedparse bed12tobed6 transcripts.bed
chr1      67092164      67093604      ENST00000371007.6      0      -
chr1      67095234      67095421      ENST00000371007.6      0      -
chr1      67096251      67096321      ENST00000371007.6      0      -
chr1      67115351      67115464      ENST00000371007.6      0      -
chr1      67125751      67125909      ENST00000371007.6      0      -
chr1      67127165      67127257      ENST00000371007.6      0      -
chr1      67131141      67131227      ENST00000371007.6      0      -
chr1      67231845      67231852      ENST00000371007.6      0      -
```

### 1.3.9 Convert chromosome names

Convert chromosome names between UCSC and Ensembl formats. The conversion supports the hg38 assembly up to patch 11 and the mm10 assembly up to patch 4. By default patches are not converted (because the UCSC genome browser does not support them), but can be enabled using the `-p` flag. When the BED file contains a chromosome that is not recognised, by default the program stops and throws an error. Alternatively, unrecognised chromosomes can be suppressed (`-s`) or artificially set to 'NA' (`-a`).

## Usage

```
> bedparse convertChr --help
usage: bedparse convertChr [-h] --assembly ASSEMBLY --target TARGET
                        [--allowMissing] [--suppressMissing] [--patches]
                        [bedfile]

Convert chromosome names between UCSC and Ensembl formats. The conversion
supports the hg38 assembly up to patch 11 and the mm10 assembly up to patch 4.
By default patches are not converted (because the UCSC genome browser does not
support them), but can be enabled using the -p flag. When the BED file
contains a chromosome that is not recognised, by default the program stops and
throws an error. Alternatively, unrecognised chromosomes can be suppressed
(-s) or artificially set to 'NA' (-a).

positional arguments:
  bedfile                Path to the BED file.

optional arguments:
  -h, --help            show this help message and exit
  --assembly ASSEMBLY  Assembly of the BED file (either hg38 or mm10).
  --target TARGET      Desired chromosome name convention (ucsc or ens).
  --allowMissing, -a   When a chromosome name can't be matched between USCS
                        and Ensembl set it to 'NA' (by default thrown as
                        error).
  --suppressMissing, -s When a chromosome name can't be matched between USCS
                        and Ensembl do not report it in the output (by default
                        throws an error).
  --patches, -p        Allows conversion of all patches up to p11 for hg38
                        and p4 for mm10. Without this option, if the BED file
                        contains contigs added by a patch the conversion
                        terminates with an error (unless the -a or -s flags
                        are present).
```

## Examples

```
> cat transcripts.bed
chr1      67092164      67231852      ENST00000371007.6      0      -
chr22_KI270928v1_alt      137191      137686      ENST00000630841.
↪1      0      -
chr1_KI270706v1_random      45985      46062      ENST00000611371.
↪2      0      +
chrM      3229      3304      ENST00000386347.1      0      +

> bedparse convertChr --assembly hg38 --target ens transcripts.bed
1      67092164      67231852      ENST00000371007.6      0      -
CHR_HSCHR22_3_CTG1      137191      137686      ENST00000630841.
↪1      0      -
KI270706.1      45985      46062      ENST00000611371.2      0      +
MT      3229      3304      ENST00000386347.1      0      +
```



### 1.3.10 Validate Format

Simply performs format validation on the input BED file. If any line doesn't adhere to the BED specifications the program reports an error and terminates. The `--fixSeparators` flag replaces fields separated by spaces into fields separated by a single tab. This is useful when writing a BED file by hand or when copy-pasting from a website.

#### Usage

```
usage: bedparse validateFormat [-h] [--fixSeparators] [bedfile]

Checks whether the BED file provided adheres to the BED format specifications.
Optionally, it can fix field speration errors.

positional arguments:
  bedfile                Path to the BED file.

optional arguments:
  -h, --help            show this help message and exit
  --fixSeparators, -f  If the fields are separated by multiple spaces (e.g.
                        when copy-pasting BED files), replace them into tabs.
```

#### Examples

```
> cat example.bed
  chr1 a213941196 213942363
  chr1 213942363 213943530
chr1 213943530          213944697

> bedparse validateFormat -f example.bed
chr1 213941196          213942363
chr1 213942363          213943530
chr1 213943530          213944697
```

## 1.4 Implementations notes

Internally, bedparse processes a bedfile line by line by instantiating objects of the bedline class. The bedline class implements an `init()` method that performs several checks on each field in order to ensure the correctness of the format, whereas the other methods of the class implement all the bedparse operations (see functionality).

## 1.5 bedparse.bedline module

**class** `bedparse.bedline.bedline` (*line=None*)

Bases: `object`

The bedline class defines an object that represents a single BED[3,4,6,12] line

**Parameters** `line` (*list*) – List where each element corresponds to one field of a BED file

**bed12tobed6** (*appendExN=False, whichExon='all'*)

Returns a list of bedlines (bed6) corresponding to the exons.

### Parameters

- **appendExN** (*bool*) – Appends the exon number to the transcript name
- **whichExon** (*str*) – Which exon to return. One of [“all”, “first”, “last”]. First and last respectively report the first or last exon relative to the TSS (i.e. taking strand into account).

**Returns** list of bedline objects, one per exon

**Return type** list

### Examples

```
>>> bl = bedline(["chr1", 100, 420, "Name", 0, "+", 210, 310, ".", 4, "20,20,
↳20,20,", "0,100,200,300,"])
>>> for i in bl.bed12tobed6(appendExN=True): print(i)
...
['chr1', 100, 120, 'Name_Exon001', 0, '+']
['chr1', 200, 220, 'Name_Exon002', 0, '+']
['chr1', 300, 320, 'Name_Exon003', 0, '+']
['chr1', 400, 420, 'Name_Exon004', 0, '+']
```

**cds** (*ignoreCDSonly=False*)

Return the CDS of a coding transcript. Transcripts without CDS are not reported

**Parameters** **ignoreCDSonly** (*bool*) – If True return None when the entire transcript is CDS

**Returns** The CDS as a bedline object

**Return type** *bedline*

### Examples

```
>>> bl = bedline(["chr1", 100, 500, "Tx1", 0, "+", 200, 300, ".", 1, "400,",
↳"0,"])
>>> print(bl.cds())
['chr1', 200, 300, 'Tx1', 0, '+', 200, 300, '.', 1, '100,', '0,']
```

**introns** ()

Returns a bedline object of the introns of a transcript

**Returns** The introns of the transcripts as a bedline object

**Return type** *bedline*

### Examples

```
>>> bl = bedline(["chr1", 100, 420, "Name", 0, "+", 210, 310, ".", 4, "20,20,
↳20,20,", "0,100,200,300,"])
>>> print(bl.introns())
['chr1', 120, 400, 'Name', 0, '+', 120, 120, '.', 3, '80,80,80,', '0,100,200,
↳']
>>> bl = bedline(["chr1", 100, 420, "Name", 0, "-", 210, 310, ".", 1, "320,",
↳"0,"])
>>> print(bl.introns())
None
```

**pprint** ()  
Prints a bedline object formatted as a python list

**print** (*end*='\n')  
Prints a bedline object

**Parameters** *end* – Line terminator character

**promoter** (*up*=500, *down*=500, *strand*=True)  
Returns the promoter of a bedline object

**Parameters**

- **up** (*int*) – Number of upstream bases
- **down** (*int*) – Number of downstream bases
- **strand** (*bool*) – If false strandedness is ignored

**Returns** The promoter as a bedline object

**Return type** *bedline*

## Examples

```
>>> bl = bedline(['chr1', 1000, 2000, 'Tx1', '0', '+'])
>>> print(bl.promoter())
['chr1', 500, 1500, 'Tx1']
```

**translateChr** (*assembly*, *target*, *suppress*=False, *ignore*=False, *patches*=False)  
Convert the chromosome name to Ensembl or UCSC

**Parameters**

- **assembly** (*str*) – Assembly of the BED file (either hg38 or mm10).
- **target** (*str*) – Desired chromosome name convention (ucsc or ens).
- **suppress** (*bool*) – When a chromosome name can't be matched between USCS and Ensembl set it to 'NA' (by default throws as error)
- **ignore** (*bool*) – When a chromosome name can't be matched between USCS and Ensembl do not report it in the output (by default throws an error)
- **patches** (*bool*) – Allows conversion of all patches up to p11 for hg38 and p4 for mm10. Without this option, if the BED file contains contigs added by a patch the conversion terminates with an error (unless the -a or -s flags are present)

**Returns** A bedline object with the converted chromosome

**Return type** *bedline*

## Examples

```
>>> bl = bedline(['chr1', 1000, 2000, 'Tx1', '0', '-'])
>>> print(bl.translateChr(assembly="hg38", target="ens"))
['1', 1000, 2000, 'Tx1', '0', '-']
>>> bl = bedline(['chr19_GL000209v2_alt', 1000, 2000, 'Tx1', '0', '-'])
>>> print(bl.translateChr(assembly="hg38", target="ens"))
['CHR_HSCHR19KIR_RP5_B_HAP_CTG3_1', 1000, 2000, 'Tx1', '0', '-']
```

**tx2genome** (*coord*, *stranded=False*)

Given a position in transcript coordinates returns the equivalent in genome coordinates. The transcript coordinates are considered without regard to strand, i.e. 0 is the leftmost position for both + and - strand transcripts, unless the *stranded* options is set to True.

**Parameters**

- **coord** (*int*) – Coordinate to convert from transcript-space to genome space
- **stranded** (*bool*) – If True use the rightmost base of negative strand transcripts as 0

**Returns** Coordinate in genome-space

**Return type** *int*

**Examples**

```
>>> bl = bedline(['chr1', 1000, 2000, 'Tx1', '0', '-'])
>>> bl.tx2genome(10)
1010
>>> bl.tx2genome(10, stranded=True)
1989
```

**utr** (*which=None*)

Returns the UTR of coding transcripts (i.e. those with a CDS)

**Parameters** **which** (*int*) – Which UTR to return: 3 for 3'UTR or 5 for 5' UTR

**Returns** The UTR as a bedline object

**Return type** *bedline*

**Examples**

```
>>> bl = bedline(["chr1", 100, 500, "Tx1", 0, "+", 200, 300, ".", 1, "400,",
↳ "0,"])
>>> print(bl.utr(which=5))
['chr1', 100, 200, 'Tx1', 0, '+', 100, 100, '.', 1, '100,', '0,']
```

## 1.6 Bedparse tutorial

Hi, thanks for your interest in bedparse!

The following is a short tutorial that will guide you through the functionality of bedparse. You can find the `example.bed` file in this repo under `docs/example.bed`. This file contains 6 human transcript models from Gencode. The first three are non-coding transcripts (i.e. field 7 and 8 contain the same coordinate), whereas the last three are coding (i.e. fields 7 and 8 indicate the `thickStart` and `thickEnd`, i.e. start and end of the CDS).

### 1.6.1 Extracting the promoters

The `bedparse promoter` command reports the promoter of each transcript, defined as a user specified interval around the TSS. For example, we can extract promoters consisting of 1000bp upstream and 500bp downstream of the CDS:

```
$ bedparse promoter --up 1000 --down 500 example.bed
chr1 10868 12368 ENST00000456328.2
chr1 11009 12509 ENST00000450305.2
chr1 29070 30570 ENST00000488147.1
chr1 922927 924427 ENST00000420190.6
chr1 924149 925649 ENST00000437963.5
chr1 924737 926237 ENST00000342066.7
```

Note how the TSS (and as a consequence the promoter) depends on the strand: for transcripts on the negative strand the TSS is the end coordinate, i.e. column 3. The `--unstranded` option allows you to override this behaviour and report promoters as an interval around column 2, thus disregarding the strand.

## 1.6.2 Extracting 5' or 3' UTRs

The UTRs are defined in a BED file as the region between the start (column 2) and the thickStart (column 7) for the 5' and between the thickEnd (column 8) and the end (column 3) for the 3'. These rules are reversed for transcripts on the - strand, and `bedparse` automatically takes care of this. Additionally, `bedparse` also handles correctly UTRs that span multiple exons: in these cases `bedparse` recomputes all exon starts and exon lengths as sets columns 11 and 12 accordingly.

```
$ bedparse 5pUTR example.bed
chr1 923927 924431 ENST00000420190.6 0 + 923927 923927 0
↳ 1 504, 0,
chr1 925149 925941 ENST00000437963.5 0 + 925149 925149 0
↳ 2 40,20, 0,772,
chr1 925737 925941 ENST00000342066.7 0 + 925737 925737 0
↳ 2 63,20, 0,184,
```

Clearly, as you can see from the output above, UTRs are only reported for coding transcripts.

## 1.6.3 Extracting the CDS

To extract the CDS of the coding transcripts in the BED file use the `bedparse cds` command:

```
$ bedparse cds example.bed
chr1 924431 939291 ENST00000420190.6 0 + 924431 939291 0
↳ 7 517,92,182,51,125,90,17, 0,1490,5723,6607,11340,14608,14843,
chr1 925941 935793 ENST00000437963.5 0 + 925941 935793 0
↳ 4 72,182,51,22, 0,4213,5097,9830,
chr1 925941 944153 ENST00000342066.7 0 + 925941 944153 0
↳ 13 72,182,51,125,90,186,163,116,79,500,125,111,246, 0,4213,5097,9830,
↳ 13098,13333,15202,16194,16468,16617,17311,17756,17966,
```

Note how non-coding transcripts are not reported (because by definition they don't have a CDS). Also, note how the number of exons (column 10) and exon lengths and starts (columns 11 and 12) have been readjusted to reflect the fact that the transcripts have "lost" the UTRs. To visualise this operation you can save the output of the command above to a new text file and upload it as a custom track in the UCSC genome browser: you'll see that the new transcripts only correspond to the thick portion of the original Gencode transcripts.

## 1.6.4 Extracting introns

In a BED file introns are implicitly defined as the genomic regions between exons. The `bedparse introns` command creates new "artificial" transcripts that correspond to the introns of the original transcripts:

```
$ bedparse introns example.bed
chr1 12227 13220 ENST00000456328.2 0 + 12227 12227 0
↳ 2 385,499, 0,494,
chr1 12057 13452 ENST00000450305.2 0 + 12057 12057 0
↳ 5 121,385,277,168,78, 0,170,640,995,1317,
chr1 14501 29533 ENST00000488147.1 0 - 14501 14501 0
↳ 10 503,757,659,92,177,237,172,206,6371,4642, 0,537,1446,2264,2554,2867,
↳3241,3560,3865,10390,
chr1 924948 939274 ENST00000420190.6 0 + 924948 924948 0
↳ 6 973,4141,702,4682,3143,145, 0,1065,5388,6141,10948,14181,
chr1 925189 935771 ENST00000437963.5 0 + 925189 925189 0
↳ 4 732,4141,702,4682, 0,824,5147,5900,
chr1 925800 943907 ENST00000342066.7 0 + 925800 925800 0
↳ 13 121,4141,702,4682,3143,145,1683,829,158,70,194,320,99, 0,213,4536,5289,
↳10096,13329,13660,15506,16451,16688,17258,17577,18008,
```

### 1.6.5 Convert BED12 to BED6

It's often convenient to convert a BED12 file into BED6, where each exon appears on its own line. This is easily done with `bedparse bed12tobed6`:

```
$ bedparse bed12tobed6 --appendExN example.bed
chr1 11868 12227 ENST00000456328.2_Exon001 0 +
chr1 12612 12721 ENST00000456328.2_Exon002 0 +
chr1 13220 14409 ENST00000456328.2_Exon003 0 +
chr1 12009 12057 ENST00000450305.2_Exon001 0 +
chr1 12178 12227 ENST00000450305.2_Exon002 0 +
chr1 12612 12697 ENST00000450305.2_Exon003 0 +
chr1 12974 13052 ENST00000450305.2_Exon004 0 +
chr1 13220 13374 ENST00000450305.2_Exon005 0 +
chr1 13452 13670 ENST00000450305.2_Exon006 0 +
chr1 14403 14501 ENST00000488147.1_Exon001 0 -
chr1 15004 15038 ENST00000488147.1_Exon002 0 -
chr1 15795 15947 ENST00000488147.1_Exon003 0 -
chr1 16606 16765 ENST00000488147.1_Exon004 0 -
chr1 16857 17055 ENST00000488147.1_Exon005 0 -
chr1 17232 17368 ENST00000488147.1_Exon006 0 -
chr1 17605 17742 ENST00000488147.1_Exon007 0 -
chr1 17914 18061 ENST00000488147.1_Exon008 0 -
chr1 18267 18366 ENST00000488147.1_Exon009 0 -
chr1 24737 24891 ENST00000488147.1_Exon010 0 -
chr1 29533 29570 ENST00000488147.1_Exon011 0 -
chr1 923927 924948 ENST00000420190.6_Exon001 0 +
chr1 925921 926013 ENST00000420190.6_Exon002 0 +
chr1 930154 930336 ENST00000420190.6_Exon003 0 +
chr1 931038 931089 ENST00000420190.6_Exon004 0 +
chr1 935771 935896 ENST00000420190.6_Exon005 0 +
chr1 939039 939129 ENST00000420190.6_Exon006 0 +
chr1 939274 939291 ENST00000420190.6_Exon007 0 +
chr1 925149 925189 ENST00000437963.5_Exon001 0 +
chr1 925921 926013 ENST00000437963.5_Exon002 0 +
chr1 930154 930336 ENST00000437963.5_Exon003 0 +
chr1 931038 931089 ENST00000437963.5_Exon004 0 +
chr1 935771 935793 ENST00000437963.5_Exon005 0 +
chr1 925737 925800 ENST00000342066.7_Exon001 0 +
chr1 925921 926013 ENST00000342066.7_Exon002 0 +
```

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chr1	930154	930336	ENST00000342066.7_Exon003	0	+
chr1	931038	931089	ENST00000342066.7_Exon004	0	+
chr1	935771	935896	ENST00000342066.7_Exon005	0	+
chr1	939039	939129	ENST00000342066.7_Exon006	0	+
chr1	939274	939460	ENST00000342066.7_Exon007	0	+
chr1	941143	941306	ENST00000342066.7_Exon008	0	+
chr1	942135	942251	ENST00000342066.7_Exon009	0	+
chr1	942409	942488	ENST00000342066.7_Exon010	0	+
chr1	942558	943058	ENST00000342066.7_Exon011	0	+
chr1	943252	943377	ENST00000342066.7_Exon012	0	+
chr1	943697	943808	ENST00000342066.7_Exon013	0	+
chr1	943907	944575	ENST00000342066.7_Exon014	0	+

The optional flag `--appendExN` adds ExonNNN to the end of each transcript name.

## 1.6.6 APIs

Bedparse can also be imported as a python module. The API documentation contains detailed information of the `bedline` class and its methods. The following is simple example of how to use it:

```
In [1]: from bedparse import bedline

In [2]: l = bedline(['chr1', 1000, 2000, 'Tx1', '0', '+'])

In [3]: prom = l.promoter()

In [4]: prom.print()
chr1    500    1500    Tx1

In [5]: prom.pprint()
['chr1', 500, 1500, 'Tx1']

In [6]: ens_prom = prom.translateChr(assembly="hg38", target="ens")

In [7]: ens_prom.print()
1      500    1500    Tx1
```





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