Using refGenome package

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1 Object types inside refGenome package

The central classes inside this package are `refGenome` derived (S4) classes. There is one class for Ensembl genomes `ensemblGenome` and one class for UCSC genomes `ucscGenome`. The objects basically contain annotation data in tables and the address of a folder (called "basedir").

The `ensemblExons` class centers on exon-intron-exon boundaries. The class also contains tabled annotation and a folder address ("basedir").

1.1 Creation of empty refGenome objects

Empty objects of `refGenome` derived classes can be created with `ensembleGenome()` or `ucscGenome()`. After creation of an empty object the first step usually is to set the basedir address:

```
> library(refGenome)
> beg<-ensemblGenome()
> basedir(beg)<-system.file("extdata",package="refGenome")
```

The "basedir" folder is intended to contain all data which is associated with the current annotation set, e.g. downloaded gtf files, saved object data, saved SQLite versions of the data and potenially sequence information. In order to fill an empty object, annotation data has to be imported from external files.

1.2 Importing annotation data

The basic importing mechanism for `refGenome` objects is to import a "gtf" file. Therefore, the "gtf" files have to be downloaded. The download source and mechanism is explained for `ensemblGenome` and `ucscGenome` separately. There are specialized mechanisms in order to provide additional information either from within the gtf file (ensembl) or via other external files (ucsc).

1.3 Saving and loading data

The data content of `refGenome` objects can be saved and re-loaded in several ways. One way is the `saveGenome` method where the content is written into a compressed ".RData" file. One alternative is to write the content into a SQLite database via `writeDB`.

2 Ensembl Genomes

The `ensemblGenome` class is specialized for managing annotation data for ensemble Genomes.

2.1 Download and import data

For ensemblGenome objects, gtf files can be downloaded from Ensemble servers. Therefore, go to

`http://www.ensembl.org/info/data/ftp/index.html`

and choose a file from the "Gene sets" column. They are labeled "GTF". For example Version 62 of human genomic annotation can be downloaded from

`ftp://ftp.ensembl.org/pub/release-62/gtf/homo_sapiens/Homo_sapiens.GRCh37.62.gtf.gz`

A copy of the obtained file should then be placed in the the "basedir" directory. With the appropriate setting of basedir, annotation data can be imported with:

```
> ens_gtf<-"hs.ensembl.62.small.gtf"
> read.gtf(beg,ens_gtf)

[read.gtf.refGenome] Reading file 'hs.ensembl.62.small.gtf'.
[read.gtf.refGenome] Parsing attributes.
[read.gtf.refGenome] Finished  135  rows and 424 gtfattributes lines.

> beg

Object of class 'ensemblGenome' with 135 rows and 11 columns.
   id seqid start   end feature score strand frame
25  1     1 11869 12227    exon     .      +     .
34  2     1 11872 12227    exon     .      +     .
41  3     1 11874 12227    exon     .      +     .
28  4     1 12010 12057    exon     .      +     .
29  5     1 12179 12227    exon     .      +     .
35  6     1 12190 12227     CDS     .      +     0
           gene_id   transcript_id         source
25 ENSG00000223972 ENST00000456328     pseudogene
34 ENSG00000249291 ENST00000515242 protein_coding
41 ENSG00000253101 ENST00000518655 protein_coding
28 ENSG00000223972 ENST00000450305     pseudogene
29 ENSG00000223972 ENST00000450305     pseudogene
35 ENSG00000249291 ENST00000515242 protein_coding
```

The top lines of the contained table are shown when the object is printed.

2.2 Attribute data in Ensembl Genome gtf files

In Ensembl gtf files there is additional data contained in the last column ("attributes"). Contained attribute types can be listet with "tableAttributeTypes". Specific attributes can be shifted into the main (gtf-) table by "moveAttributes":

```
> tableAttributeTypes(beg)

[tableAttributeTypes.refGenome] Row number in gtf-table: 135.

    exon_number       gene_name      protein_id
            135             135              19
transcript_name
            135

> moveAttributes(beg,c("gene_name","transcript_name","exon_number"))

[moveAttributes.ensemblGenome] Moving   135 'gene_name' attributes to 'gtf' table.
[moveAttributes.ensemblGenome] Moving   135 'transcript_name' attributes to 'gtf' table.
[moveAttributes.ensemblGenome] Moving   135 'exon_number' attributes to 'gtf' table.
[moveAttributes.ensemblGenome] Finished. Reduced attributes table size from 424 to 19 rows
```

3 UCSC Genomes

Downloading of annotation data for UCSC genomes is a bit more complicated than for Ensemble Genomes because additional data must be downloaded in separate files. The Homepage for UCSC browser can be found under:

`http://genome.ucsc.edu/`

In order to import UCSC annotation data into `refGenome` objects files containing the data have to be downloaded from the USCS Table Browser which can be found under:

`http://genome.ucsc.edu/cgi-bin/hgTables`

or by following the "Tablel Browser" link in the left panel on the homepage. On the Table Browser:

- Select genome, assembly and track (UCSC genes)
- Choose table (knownGene)
- Choose output format (GTF -gene transfer format for knownGene table)
- Insert a name for the output file
- Download the file (get output)

The basic table to be imported is "knownGene". The knownGene table has to be downloaded in GTF format (otherwise the read.gtf function will complain

about "wrong number of columns").

In order to extend the available information additionally the tables "kgXref", "knownToEnsembl" and "knownIsoforms" can be downloaded and imported. These tables come in plain "csv" format. Select "all fields from selected table" as output format.

Do not use "add custom tracks" or modify the tables elsewhere tracks because the importing functions will check for apropriate number of columns.

After downloading, all tables should be placed into a separate folder which we from now on call "basedir".`ucscGenome` objects keep a `basedir` as standard location for all writing and reading procedures.

```
> uc<-ucscGenome()
> basedir(uc)<-"/my/ucsc/basedir"
> read.gtf(uc,"ucsc_knownGene.gtf")
> addXref(uc,"kgXref.csv")
> addEnsembl(uc,"knownToEnsembl.csv")
> addIsoforms(uc,"ucsc_knownisoforms.csv")
```

3.1 Load stored data

Once, annotation data is imported and stored, `ucscGenome` objects can be restored with the `loadGenome` function which is shown below on example data:

```
> ucfile<-system.file("extdata", "hs.ucsc.small.RData", package="refGenome")
> uc<-loadGenome(ucfile)
> ensfile<-system.file("extdata", "hs.ensembl.62.small.RData", package="refGenome")
> ens<-loadGenome(ensfile)
```

4 Extracting data subsets

There are specialized functions for extracting data for multiple purposes.

4.1 Extracting data for sets of `seqid`'s

For preparation of `seqid` based extraction, the contained `seqid`'s can be tabled:

```
> tableSeqids(ens)
```

```
     1 GL000213.1
   111         24
```

Extraction of subsets based on `seqid` can be done with `extractSeqids`. The sequence id's for extraction are specified as regular expression:

```
> en1<-extractSeqids(ens,"^1$")
> en1
```

```
Object of class 'ensemblGenome' with 111 rows and 14 columns.
   id seqid start   end feature score strand frame
25  1     1 11869 12227    exon     .      +     .
34  2     1 11872 12227    exon     .      +     .
41  3     1 11874 12227    exon     .      +     .
28  4     1 12010 12057    exon     .      +     .
29  5     1 12179 12227    exon     .      +     .
35  6     1 12190 12227     CDS     .      +     0
           gene_id   transcript_id         source
25 ENSG00000223972 ENST00000456328     pseudogene
34 ENSG00000249291 ENST00000515242 protein_coding
41 ENSG00000253101 ENST00000518655 protein_coding
28 ENSG00000223972 ENST00000450305     pseudogene
29 ENSG00000223972 ENST00000450305     pseudogene
35 ENSG00000249291 ENST00000515242 protein_coding
    gene_name transcript_name exon_number
25    DDX11L1     DDX11L1-002           1
34 AL627309.2  AL627309.2-201           1
41   DDX11L11    DDX11L11-201           1
28    DDX11L1     DDX11L1-001           1
29    DDX11L1     DDX11L1-001           2
35 AL627309.2  AL627309.2-201           1
```

It looks cumbersome for single chromosomes but allows extraction of complex patterns.

4.2 Extracting primary assembly data

Usually the interesting part of the annotation data is the the primary assembly (where alternative haplotypes are excluded). Therefore functions which return the proper terms are supplied:

```
> ensPrimAssembly()

[1] "^([0-9]{1,2})$|^[XY]|MT$"

> ucPrimAssembly()

[1] "^chr[0-9XYM]{1,2}$"
```

Extraction of primary assembly `seqid`'s i is done by:

```
> enpa<-extractSeqids(ens,ensPrimAssembly())
> tableSeqids(enpa)

  1
111

> ucpa<-extractSeqids(uc,ucPrimAssembly())
> tableSeqids(ucpa)

chr1
   6
```

4.3 Extract features

Subsets defined by `features` can allso be tabled and extracted:

```
> tableFeatures(enpa)

        CDS        exon start_codon  stop_codon
          0           6           0           0

> enpf<-extractFeature(enpa,"exon")
> enpf

Object of class 'ensemblGenome' with 6 rows and 14 columns.
  gene_name seqid         source feature start   end score
1   DDX11L1  chr1 hg19_knownGene    exon 11874 12227     0
2   DDX11L1  chr1 hg19_knownGene    exon 11874 12227     0
3   DDX11L1  chr1 hg19_knownGene    exon 12613 12721     0
4   DDX11L1  chr1 hg19_knownGene    exon 12646 12697     0
5   DDX11L1  chr1 hg19_knownGene    exon 13221 14409     0
6   DDX11L1  chr1 hg19_knownGene    exon 13221 14409     0
  strand frame id    gene_id transcript_id         ensembl
1      +     .  1 uc001aaa.3    uc001aaa.3 ENST00000456328
2      +     .  4 uc010nxr.1    uc010nxr.1 ENST00000456328
3      +     .  2 uc001aaa.3    uc001aaa.3 ENST00000456328
4      +     .  5 uc010nxr.1    uc010nxr.1 ENST00000456328
5      +     .  3 uc001aaa.3    uc001aaa.3 ENST00000456328
6      +     .  6 uc010nxr.1    uc010nxr.1 ENST00000456328
  clusterId
1         1
2         1
3         1
4         1
5         1
6         1
```

4.4 Extract data for single genes and transcripts

There are some functions which extract objects that contain data for single genes (or transcripts). These functions provide a closer insight into specific regeions.

Objects which contain data for vectors of gene-names can be extracted with

```
> dxe<-extractByGeneName(enpa,"DDX11L1")
> dxu<-extractByGeneName(ucpa,"DDX11L1")
```

When gene-names did not match in the gtf-table of the object, a message including all names of not matching gene-names will be printed. When no gene-name matches, a message will be printed and the function returns `NULL`, which can be tested for later on.

From these extracts we can view the contained transcripts with the `tableTranscript.id` function:

```
> tableTranscript.id(enpa)

uc001aaa.3 uc010nxr.1
         3          3

> tableTranscript.id(ucpa)

uc001aaa.3 uc010nxr.1
         3          3
```

Data for interesting transcripts can be extracted by `extractTranscript`:

```
> extractTranscript(ens,"ENST00000456328")

Object of class 'ensemblGenome' with 3 rows and 14 columns.
    transcript_id id seqid start   end feature score strand
1 ENST00000456328  1     1 11869 12227    exon     .      +
2 ENST00000456328  9     1 12613 12721    exon     .      +
3 ENST00000456328 14     1 13221 14409    exon     .      +
  frame         gene_id     source gene_name
1     . ENSG00000223972 pseudogene   DDX11L1
2     . ENSG00000223972 pseudogene   DDX11L1
3     . ENSG00000223972 pseudogene   DDX11L1
  transcript_name exon_number
1     DDX11L1-002           1
2     DDX11L1-002           2
3     DDX11L1-002           3

> extractTranscript(uc,"uc010nxr.1")

Object of class 'ucscGenome' with 3 rows and 14 columns.
  transcript_id gene_name seqid         source feature
1    uc010nxr.1   DDX11L1  chr1 hg19_knownGene    exon
2    uc010nxr.1   DDX11L1  chr1 hg19_knownGene    exon
3    uc010nxr.1   DDX11L1  chr1 hg19_knownGene    exon
  start   end score strand frame id    gene_id
1 11874 12227     0      +     .  4 uc010nxr.1
2 12646 12697     0      +     .  5 uc010nxr.1
3 13221 14409     0      +     .  6 uc010nxr.1
         ensembl clusterId
1 ENST00000456328         1
2 ENST00000456328         1
3 ENST00000456328         1
```

5 Accumulate data for whole genes

The function `getGenePositions` accumulates position data for whole genes. Genes are grouped by `gene_name`. For both, *ensemblGenome* and *ucscGenome* the `gene_name` column is not present after the standard gtf-import. For *ensemblGenome*, `moveAttributes` must be used and for *ucscGenome*, `addXref` must be used. Respective warnings are given.

```
> gpe<-getGenePositions(ens)
> gpe
```

```
  id         gene_id  gene_name      seqid  start    end
2  2 ENSG00000223972    DDX11L1          1  11869  14409
7  7 ENSG00000249291 AL627309.2          1  11872  14412
8  8 ENSG00000253101   DDX11L11          1  11874  14409
3  3 ENSG00000227232     WASH7P          1  14363  29806
6  6 ENSG00000243485 MIR1302-10          1  29554  31109
1  1 ENSG00000221311 MIR1302-10          1  30366  30503
5  5 ENSG00000237613    FAM138A          1  34554  36081
4  4 ENSG00000237375 BX072566.1 GL000213.1 108007 139339
  strand
2      +
7      +
8      +
3      -
6      +
1      +
5      -
4      -
```

```
> gpu<-getGenePositions(ucpa)
> gpu
```

```
  id    gene_id gene_name seqid start   end strand
2  2 uc010nxr.1   DDX11L1  chr1 11874 14409      +
1  1 uc001aaa.3      <NA>  <NA> 11874 14409   <NA>
```

There is a slight difference between both results: The last column is `gene_id` for *ensemblGenome* and `clusterID` for *ucscGenome*. This is due to different information which is available for each.

6 Exon and splice-junction based views (only for Ensembl genomes)

6.1 Extract exon based table

Exon based view on annotation data can be obtained with `ensemblExons` which returns an object of class `ensemblExons`. Basically `ensemblExons` calls `extractFeature` for feature type "exon". Information about presence of cds start or end and start-codon or stop-codon is added.

```
> enex<-refExons(ens)
```

```
[refExons.refGenome] Extracting tables.
[refExons.refGenome] Adding 'CDS'.
[refExons.refGenome] Adding 'start_codon'.
[refExons.refGenome] Adding 'stop_codon'.
[refExons.refGenome] Finished.
```

```
> ucex<-refExons(uc)

[refExons.refGenome] Extracting tables.
[refExons.refGenome] Adding 'CDS'.
[refExons.refGenome] Adding 'start_codon'.
[refExons.refGenome] Adding 'stop_codon'.
[refExons.refGenome] Finished.
```

6.2 Extract splice-junction based views from `ensemblExons`

From `ensemblExons` information about adjacency of exons (which defines annotated splice-sites) can be obtained by putting exons with equal transcript_id and subsequent exon_number side by side.
The start and end positions of adjacent exons are renamed to lstart, lend and rstart and rend. The "l" prefix refers to the exon with lower start and end coordinates (i.e. left, lower exon_number). The "r" prefix refers to the exons with higher start and end coordinates (i.e. right, higher exon_number).
Setting `coding=TRUE` will restrict the result to exons for which `source` and `gene_biotype` equal "protein_coding".

```
> jens<-getSpliceTable(ens)
> jens

Object of class 'ensemblJunctions' with 92 rows and 12 columns.
  id      seqid lstart   lend rstart   rend         gene_id
1  1 GL000213.1 108007 108247 109884 110007 ENSG00000237375
2  2 GL000213.1 109884 110007 118422 118588 ENSG00000237375
3  3 GL000213.1 118422 118588 119629 119673 ENSG00000237375
4  4 GL000213.1 119629 119673 121073 121143 ENSG00000237375
5  5 GL000213.1 121073 121143 126648 126718 ENSG00000237375
6  6 GL000213.1 126648 126718 129228 129365 ENSG00000237375
   gene_name strand   transcript_id lexid rexid
1 BX072566.1      - ENST00000327822   112   115
2 BX072566.1      - ENST00000327822   115   117
3 BX072566.1      - ENST00000327822   117   119
4 BX072566.1      - ENST00000327822   119   121
5 BX072566.1      - ENST00000327822   121   123
6 BX072566.1      - ENST00000327822   123   125

> juc<-getSpliceTable(uc)
> juc

Object of class 'ucscJunctions' with 4 rows and 12 columns.
  id seqid lstart  lend rstart  rend    gene_id gene_name
1  1  chr1  11874 12227  12613 12721 uc001aaa.3   DDX11L1
2  2  chr1  12613 12721  13221 14409 uc001aaa.3   DDX11L1
3  3  chr1  11874 12227  12646 12697 uc010nxr.1   DDX11L1
4  4  chr1  12646 12697  13221 14409 uc010nxr.1   DDX11L1
  strand transcript_id lexid rexid
1      +    uc001aaa.3     1     2
2      +    uc001aaa.3     2     3
```

```
3      +    uc010nxr.1     4     5
4      +    uc010nxr.1     5     6
```

This generally leads to repeated occurrence of start and and positons when a splice-junction is contained in multiple transcripts. Additionally a handful splice-sites with multiple gene-id's are present.
The `unifyJunc` therefore calculates `nGenes` which represents the multiplicity of each gene-id at each splice-site and then selects a gene-id for which nGenes is maximal.
`unifyJuncs` adds a `uid` column to the basic `gtf` table which identifies each seqid, left-end, right-start combination uniquely. `unifyJuncs` also adds a new `ujs` table inside the contained environment.
`getUnifiedJuncs` takes the result of `unifyJuncs` and adds gene_name and strand information.

```
> ujens<-unifyJuncs(jens)
> ujuc<-unifyJuncs(juc)
> jeg<-getGenePositions(jens)
> jug<-getGenePositions(juc)
```

The result tables of `unifyJuncs` and `getGenePositions` are stored inside the internal environment of `ensemblJunctions`. From there, the results can easily be reproduced without recalculation. The tables are automatically included in `saveGenome` and `load.ensembl.juncs` mechanisms.

7 Overlapping

The overlap function is used to supply annotation for genomic ranges. The function takes two `data.frame`'s which contain query (qry) and reference (ref) ranges respectively. Each dataset will be identified by it's id.

The routine assumes that query and reference tables are ascending sorted by column 'start'. Otherwise the result will be incorrect (i.e. missing hits). The function assumes that there is no overlap between reference ranges. It will otherwise return only one, possibly arbitrary, hit per query range.

The function returns a `data.frame`. For each query range, there will be one row.

```
> qry<-data.frame(
+                   id=1:6,
+                   start=c(10,18,61,78,82,110),
+                   end=c(15,22,63,87,90,120))
> ref<-data.frame(
+                   id=1:5,
+                   start=c(20,40,60,80,100),
+                   end=c(25,45,65,85,105))
> overlap(qry,ref)

  overlap leftDiff rightDiff queryid refid
0      no        0         5       1     0
```

```
1       l       2       3       2       1
2       n       1       2       3       3
3       b       2       2       4       4
4       r       2       5       5       4
5      no       5       0       6       0
```

The query and reference record are identified by "queryid" and "refid". The type of overlap is encoded in the "overlap" column. The overlap encodings are explained as follows:

- **no**. The query range does not overlap with any reference ranges.
- **l** The query range overhangs the matching reference range on the left (lower coordinate) side.
- **n** The query range is completely contained within a reference range. There is no overhang.
- **b** The query range overhangs the matching reference range on both sides.
- **r** The query range overhangs the matching reference range on the right (higher coordinate) side.

The added "leftDiff" and "rightDiff" columns contain the distance between the query and reference range boundaries: leftDiff is the difference between the left (lower coordinate) margins and rightDiff is the difference between the right (higher coordinate) margins.