

In-silico cleavage of polypeptides using the *cleaver* package

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Contents

1	Introduction	1
2	Simple Usage	1
3	Insulin & Somatostatin Example	4
4	Isotopic Distribution Of Tryptic Digested Insulin	5
5	Session Information	6

1 Introduction

Most proteomics experiments need protein (peptide) separation and cleavage procedures before these molecules could be analyzed or identified by mass spectrometry or other analytical tools.

cleaver allows in-silico cleavage of polypeptide sequences to e.g. create theoretical mass spectrometry data.

The cleavage rules are taken from the [ExPASy PeptideCutter tool](#)⁴.

2 Simple Usage

Loading the *cleaver* package:

In-silico cleavage of polypeptides using the *cleaver* package

```
> library("cleaver")
```

Getting help and list all available cleavage rules:

```
> help("cleave")
```

Cleaving of *Gastric juice peptide 1 (P01358)* using *Trypsin*:

```
> ## cleave it
> cleave("LAAGKVEDSD", enzym="trypsin")

$LAAGKVEDSD
[1] "LAAGK" "VEDSD"

> ## get the cleavage ranges
> cleavageRanges("LAAGKVEDSD", enzym="trypsin")

$LAAGKVEDSD
      start end
[1,]      1  5
[2,]      6 10

> ## get only cleavage sites
> cleavageSites("LAAGKVEDSD", enzym="trypsin")

$LAAGKVEDSD
[1] 5
```

Sometimes cleavage is not perfect and the enzyme miss some cleavage positions:

```
> ## miss one cleavage position
> cleave("LAAGKVEDSD", enzym="trypsin", missedCleavages=1)

$LAAGKVEDSD
[1] "LAAGKVEDSD"

> cleavageRanges("LAAGKVEDSD", enzym="trypsin", missedCleavages=1)

$LAAGKVEDSD
      start end
[1,]      1 10

> ## miss zero or one cleavage positions
> cleave("LAAGKVEDSD", enzym="trypsin", missedCleavages=0:1)

$LAAGKVEDSD
[1] "LAAGK"      "VEDSD"      "LAAGKVEDSD"
```

In-silico cleavage of polypeptides using the *cleaver* package

```
> cleavageRanges("LAAGKVEDSD", enzym="trypsin", missedCleavages=0:1)

$LAAGKVEDSD
      start end
[1,]      1  5
[2,]      6 10
[3,]      1 10
```

Combine *cleaver* and the *Biostrings* R package⁵:

```
> ## create AAStringSet object
> p <- AAStringSet(c(gaju="LAAGKVEDSD", pnm="AGEPKLDAGV"))
>
> ## cleave it
> cleave(p, enzym="trypsin")

AAStringSetList of length 2
[["gaju"]] LAAGK VEDSD
[["pnm"]] AGEPK LDAGV

> cleavageRanges(p, enzym="trypsin")

IRangesList of length 2
$gaju
IRanges object with 2 ranges and 0 metadata columns:
      start      end      width
  <integer> <integer> <integer>
[1]        1        5         5
[2]        6       10         5

$pnm
IRanges object with 2 ranges and 0 metadata columns:
      start      end      width
  <integer> <integer> <integer>
[1]        1        5         5
[2]        6       10         5

> cleavageSites(p, enzym="trypsin")

$gaju
[1] 5

$pnm
[1] 5
```

3 Insulin & Somatostatin Example

Downloading *Insulin* (P01308) and *Somatostatin* (P61278) sequences from the UniProt⁶ database using the *UniProt.ws* R package¹.

```
> ## load UniProt.ws library
> library("UniProt.ws")
>
> ## select species Homo sapiens
> UniProt.ws <- UniProt.ws(taxId=9606)
>
> ## download sequences of Insulin/Somatostatin
> s <- select(UniProt.ws, keys=c("P01308", "P61278"), columns=c("SEQUENCE"))

Getting extra data for P01308, P61278

'select()' returned 1:1 mapping between keys and columns

> ## fetch only sequences
> sequences <- setNames(s$SEQUENCE, s$UNIPROTKB)
>
> ## remove whitespaces
> sequences <- gsub(pattern="[:space:]", replacement="", x=sequences)
```

Cleaving using *Pepsin*:

```
> cleave(sequences, enzym="pepsin")

$P01308
  [1] "MA"          "L"          "WMRLLP"     "LL"
  [5] "A"           "WGPDPAAA"   "F"          "VNQH"
  [9] "CGSH"        "VEA"        "Y"          "VCGERG"
 [13] "FF"          "YTPKTRREAED" "QVGQVE"     "GGGPGAGS"
 [17] "LQP"         "LA"         "EGS"        "QKRGIVEQCCTSICS"
 [21] "YQ"          "ENYCN"

$P61278
  [1] "ML"          "SCRL"       "QCA"
  [4] "L"           "AA"         "SIV"
  [7] "A"           "GCVTGAPSDPRL" "RQ"
 [10] "FL"          "QKS"        "LAAAAGKQEL"
 [13] "AKY"         "AE"         "SEPNQTENDA"
 [16] "LEPED"       "SQAAEQDEMRL" "EL"
 [19] "QRSANSNPAMAPRERKAGCKN" "FF"         "WKT"
```

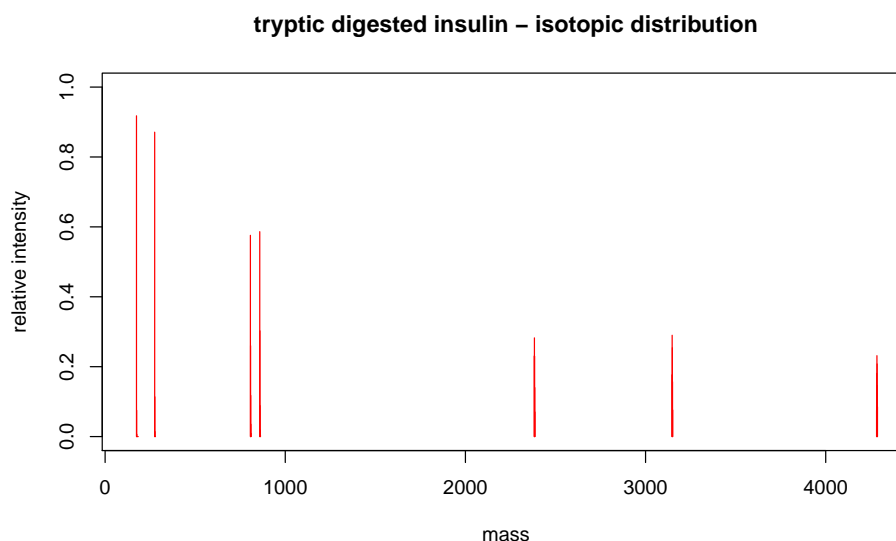
```
[22] "FTSC"
```

4 Isotopic Distribution Of Tryptic Digested Insulin

A common use case of in-silico cleavage is the calculation of the isotopic distribution of peptides (which were enzymatic digested in the in-vitro experimental workflow). Here the *BRAIN* R package^{2;3} is used to calculate the isotopic distribution of *cleaver*'s output. (please note: it is only a toy example, e.g. the relation of intensity values between peptides isn't correct).

```
> ## load BRAIN library
> library("BRAIN")
>
> ## cleave insulin
> cleavedInsulin <- cleave(sequences[1], enzym="trypsin")[[1]]
>
> ## create empty plot area
> plot(NA, xlim=c(150, 4300), ylim=c(0, 1),
+      xlab="mass", ylab="relative intensity",
+      main="tryptic digested insulin - isotopic distribution")
>
> ## loop through peptides
> for (i in seq(along=cleavedInsulin)) {
+   ## count C, H, N, O, S atoms in current peptide
+   atoms <- BRAIN::getAtomsFromSeq(cleavedInsulin[[i]])
+   ## calculate isotopic distribution
+   d <- useBRAIN(atoms)
+   ## draw peaks
+   lines(d$masses, d$isoDistr, type="h", col=2)
+ }
```

In-silico cleavage of polypeptides using the *cleaver* package



5 Session Information

- R version 3.5.1 Patched (2018-07-12 r74967),
x86_64-apple-darwin15.6.0
- Locale: C/en_US.UTF-8/en_US.UTF-8/C/en_US.UTF-8/en_US.UTF-8
- Running under: OS X El Capitan 10.11.6
- Matrix products: default
- BLAS:
/Library/Frameworks/R.framework/Versions/3.5/Resources/lib/libRblas.0.dylib
- LAPACK:
/Library/Frameworks/R.framework/Versions/3.5/Resources/lib/libRlapack.dylib
- Base packages: base, datasets, grDevices, graphics, methods, parallel,
stats, stats4, utils
- Other packages: BRAIN 1.28.0, BiocGenerics 0.28.0, Biostrings 2.50.0,
IRanges 2.16.0, PolynomF 1.0-2, RCurl 1.95-4.11, RSQLite 2.1.1,
S4Vectors 0.20.0, UniProt.ws 2.22.0, XVector 0.22.0, bitops 1.0-6,
cleaver 1.20.0, knitr 1.20, lattice 0.20-35
- Loaded via a namespace (and not attached): AnnotationDbi 1.44.0,
Biobase 2.42.0, BiocFileCache 1.6.0, BiocManager 1.30.3,
BiocStyle 2.10.0, DBI 1.0.0, R6 2.3.0, Rcpp 0.12.19, assertthat 0.2.0,
backports 1.1.2, bindr 0.1.1, bindrcpp 0.2.2, bit 1.1-14, bit64 0.9-7,

In-silico cleavage of polypeptides using the *cleaver* package

blob 1.1.1, compiler 3.5.1, crayon 1.3.4, curl 3.2, dbplyr 1.2.2, digest 0.6.18, dplyr 0.7.7, evaluate 0.12, glue 1.3.0, grid 3.5.1, highr 0.7, htmltools 0.3.6, httr 1.3.1, magrittr 1.5, memoise 1.1.0, pillar 1.3.0, pkgconfig 2.0.2, purrr 0.2.5, rappdirs 0.3.1, rlang 0.3.0.1, rmarkdown 1.10, rprojroot 1.3-2, stringi 1.2.4, stringr 1.3.1, tibble 1.4.2, tidyselect 0.2.5, tools 3.5.1, yaml 2.2.0, zlibbioc 1.28.0

References

- [1] Marc Carlson. *UniProt.ws: R Interface to UniProt Web Services*. R package version 2.0.0.
- [2] Jürgen Claesen, Piotr Dittwald, Tomasz Burzykowski, and Dirk Valkenburg. An efficient method to calculate the aggregated isotopic distribution and exact center-masses. *Journal of The American Society for Mass Spectrometry*, 23(4):753–763, 2012.
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- [4] Elisabeth Gasteiger, Christine Hoogland, Alexandre Gattiker, S'everine Duvaud, Marc R. Wilkins, Ron D. Appel, and Amos Bairoch. Protein identification and analysis tools on the expasy server. In John M. Walker, editor, *The Proteomics Protocols Handbook*, pages 571–607. Humana Press, 2005. ISBN 978-1-58829-343-5. doi: 10.1385/1-59259-890-0:571. URL <http://dx.doi.org/10.1385/1-59259-890-0%3A571>.
- [5] H. Pages, P. Abouyoun, R. Gentleman, and S. DebRoy. *Biostrings: String objects representing biological sequences, and matching algorithms*. R package version 2.28.0.
- [6] The UniProt Consortium. Reorganizing the protein space at the universal protein resource (uniprot). *Nucleic Acids Research*, 40(D1):D71–D75, 2012. doi: 10.1093/nar/gkr981. URL <http://nar.oxfordjournals.org/content/40/D1/D71.abstract>.