

# Package ‘synapterdata’

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**Type** Package

**Title** Data accompanying the synapter package

**Version** 1.0.0

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**Description** Data independant acquisition of UPS1 protein mix in an *E. coli* background obtained on a Waters Synapt G2 instrument.

**Depends** R (>= 2.10), synapter (>= 0.99.6)

**License** GPL-2

**biocViews** ExperimentData, MassSpectrometry, MassSpectrometryData,Proteomics

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synapterdata-package *Data accompanying the synapter package*

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## Description

Data independant acquisition of UPS1 protein mix in an *E. coli* background obtained on a Waters Synapt G2 instrument.

## Details

See the synapter package for details.

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getHDMSeFinalPeptide	PLGS csv data and fasta files
getMaster	Get _master_ HDMSe data
ups25a	'Synapter' spiked-in example data.

## Author(s)

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## References

Shliaha P.V., Gatto L., Bond N.J. and Lilley K.S. Synapter: Improving qualitative and quantitative performance for label free proteomics, in prep.

Shliaha, P.V., Bond N.J., Gatto L. and Lilley K.S. The Effects of Ion Mobility Separation on Data Independent Acquisition in Proteomics Studies., in prep.

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getHDMSeFinalPeptide    *PLGS csv data and fasta files*

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## Description

The PLGS HDMSe final peptide, MSe final peptide and MSe Pep3D output files are provided as gzipped csv files and their respective full paths can be obtained with getHDMSeFinalPeptide, getMSeFinalPeptide and getMSePep3D. These can then be used directly in the respective synpater functions and methods, as read.csv automatically uncompressed the files.

The fasta database file is also available in as a gunzip archive. Fasta file are however not automatically handled in gzipped format. getFasta first decompresses the file in a temporary directory and returns the full path to that uncompressed file.

## Usage

```
getHDMSeFinalPeptide()
getMSeFinalPeptide()
getMSePep3D()
getFasta()
```

## Examples

```
getHDMSeFinalPeptide()
```

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getMaster	<i>Get master HDMSe data</i>
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**Description**

TODO A concise (1-5 lines) description of what the function does.

**Usage**

```
getMaster()  
loadMaster()
```

**Details**

TODO If necessary, more details than the description above

**Author(s)**

Laurent Gatto <lg390@cam.ac.uk>

**References**

Bond N. J., Shliaha P.V., Gatto L. and Lilley K.S., in prep.

See the synapter vignette from the synapter package, available with ysynapterGuide() for a description of the underlying concepts and detailed description of the pipeline.

**See Also**

[ups25a](#) and [getHDMSeFinalPeptide](#)

**Examples**

```
loadMaster()  
master
```

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ups25a	<i>Synapter spiked-in example data.</i>
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**Description**

Objects of class Synapter, implmented in the synapter package. The 6 instances represent triplicate run of the Universal Proteomics Standard (UPS1) 48 protein mix in an E. coli background, spiked in at 25 and 50 femtomoles.

**Usage**

```
data(ups25c)
```

**Details**

Each instance has been created with the `synergise` function. The respective MSe final peptide and MSe Pep3D final are also provided in the package (see [getMSeFinalPeptide](#) and [getMSePep3D](#)). The identification peptides is a master HDMS file (see [getMaster](#)). The code generating the instances is available in the `synergise.R` R file, in the `scripts` package directory.

**Source**

Bond N. J., Shliaha P.V., Gatto L. and Lilley K.S., in prep.

**References**

See the `synapter` vignette from the `synapter` package, available with `ysnapterGuide()` for a description of the underlying concepts and detailed description of the pipeline.

**Examples**

```
library(synapter)
data(ups25a)
performance(ups25a)
```

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