

HADEX: AN R PACKAGE AND WEB-SERVER FOR ANALYSIS OF DATA FROM HDX-MS EXPERIMENTS

DOMINIK CYSEWSKI, MICHAŁ BURDUKIEWICZ

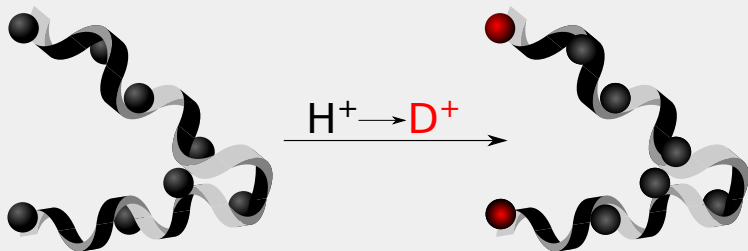
INSTITUTE OF BIOCHEMISTRY AND BIOPHYSICS, PAS

MI² DATA LAB, WARSAW UNIVERSITY OF SCIENCE AND TECHNOLOGY

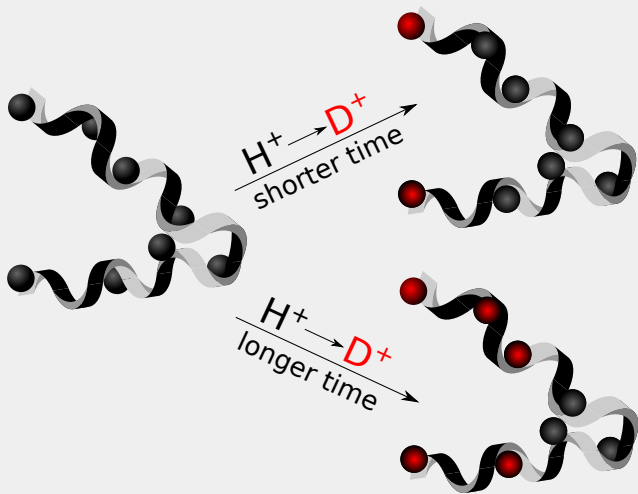
1 HDX-MS

2 HaDeX workflow

HDX-MS



After the incubation in heavy water (D_2O), the most exposed amide hydrogens of the protein backbone are being replaced by deuters.



The longer incubation time, the more protected hydrogens are being replaced by deuters.

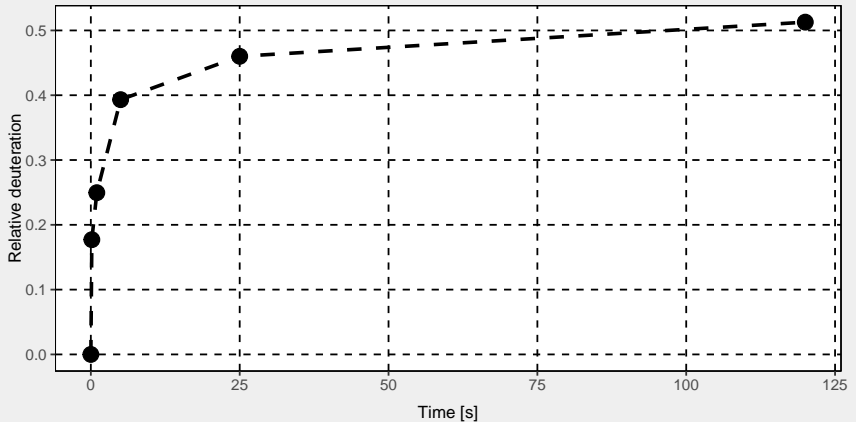
shorter incubation time



longer incubation time

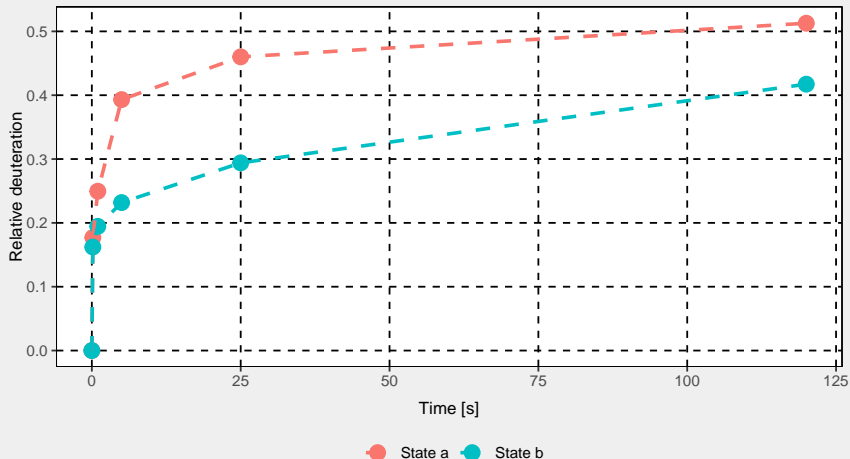


After the incubation, proteins are digested by a combination of proteases.



Thanks to the mass spectrometry, we are able to compute the m/z of each peptide depending on the duration of the incubation, thus we can compute a relative deuteration of a peptide.

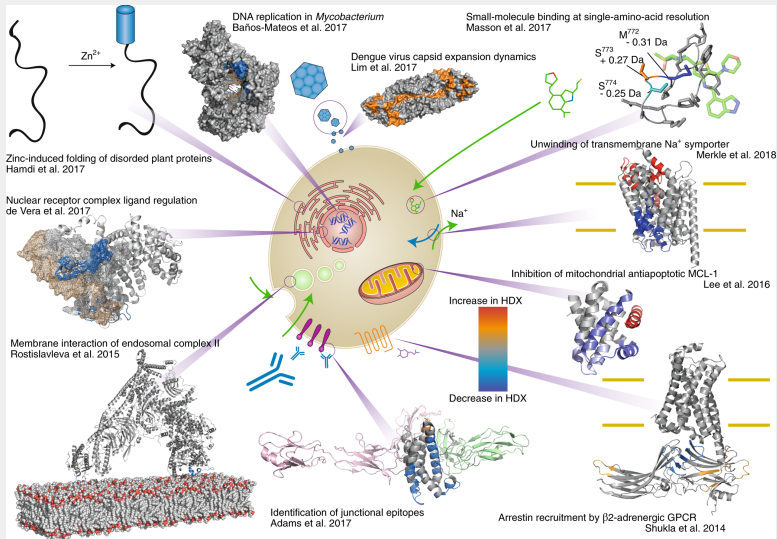
MULTI-STATE ANALYSIS



Peptides may come from proteins in different states, i.e. bounded by different cofactors.

During the analysis of the HDX-MS data we have to consider both time of the incubation and position of a peptide in a sequence.

APPLICATIONS



Source: Masson et al. (2019).

HADEX WORKFLOW

HaDeX:

- easy,
- comprehensive,
- reproducible.

Welcome!

Upload your file. Otherwise you will see example data.

Choose file:

Browse...

No file selected

File status:

Example file: KD_380T10_CD160_HVEM.csv

Please be aware that loading data (including example file) may take a while. Be patient.

Currently HaDeX supports files with only one protein.

Accepted file extensions: .csv, .xls, .xlsx.

In order for program to behave correctly, please make sure supplied file fulfills following requirements:

Show requirements

About

The HaDeX web server relies on functions from the **HaDeX R** package. For more information check our **documentation**. In case of any question or suggestion don't hesitate to contact us.

Authors

- Weronika Puchala
- Michał Burdukiewicz

Citation

Puchala W, Burdukiewicz M, Kistowski M, Dabrowska KA, Badaczewska-Dawid AE, Cysewski D and Dadlez M (2019). HaDeX: Analysis and Visualisation of Hydrogen/Deuterium Exchange Mass Spectrometry Data. R package version 1.0.

Funding

This work is supported by Foundation of Polish Science [TEAM TECH CORE FACILITY/2016-2/2 Mass Spectrometry of Biopharmaceuticals - improved methodologies for qualitative, quantitative and structural characterization of drugs, proteinaceous drug targets and diagnostic molecules].

Select parameters for the plot.

☐ Theoretical calculations

Choose values type:

- ☒ relative
☐ absolute

Comparison plot parameters:

Choose time parameters:

IN CHOSEN OUT
0001 1 1640

Choose states for comparison:

- ☒ CD160
☒ CD160_HVEM

Adjust colors

Woods plot parameters:

State 1 State 2
CD160 CD160_HVEM

Confidence limit 1:

98%

Confidence limit 2:

99%

Adjust plot:

Choose y range for comparison plot:

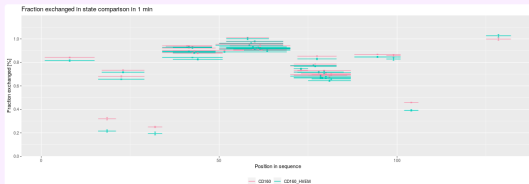


Choose y range for Woods plot:



Comparison plot

Data

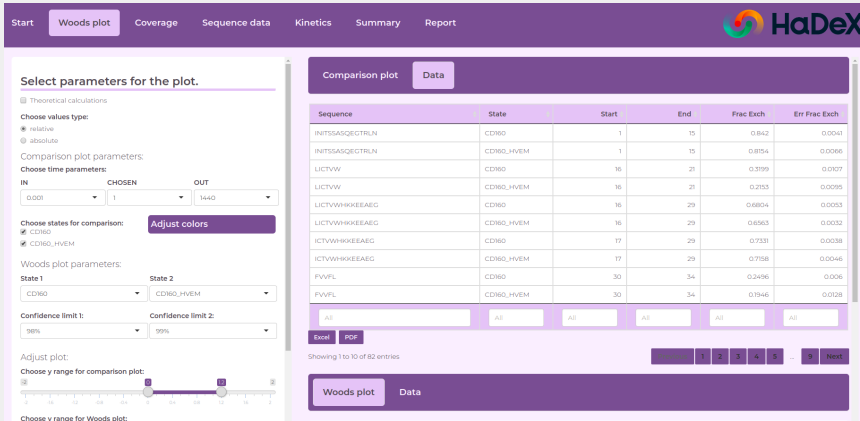


Save chart (.svg)

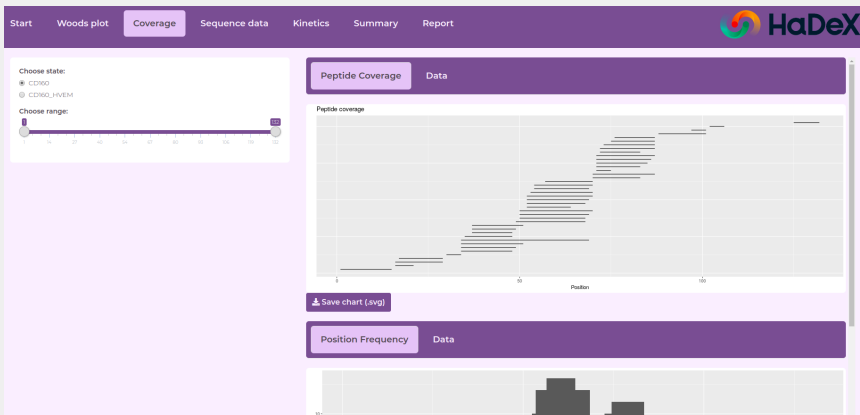
Woods plot

Data





Uncertainties derived by error propagation (Joint Committee for Guides in Metrology, 2008).



Protein name

db_CD160

Reconstructed sequence

INITSSASQEGTRLNLICTVWHKKEEAEGFVVFLLCKDRSGDCSPETSLKQLRLKRDPGIDGVGEISSQLMFTISQVTPLHSGTYQCCARSQKSGIRLQGHFFSILFxxxxxxxxxxxxxxxxxxFShNEGTL

Name	Value
Length	132
Coverage	86.36%
Cys	5

Hydro-

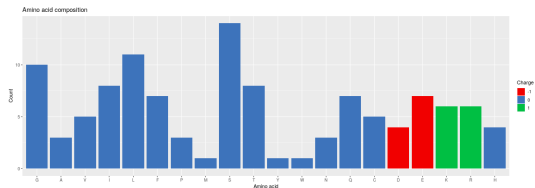
- ☒ Hydrophilic
☒ Hydrophobic

Correct sequence length:

132

If C-terminal of is not covered by peptides, enter its correct position.

Sequence length from the file is 132.



Save chart (.svg)

Select parameters for the plot.

☐ Theoretical calculations

Choose values type:

- ☒ relative
☐ absolute

Choose time parameters:

IN OUT
0.001 1440

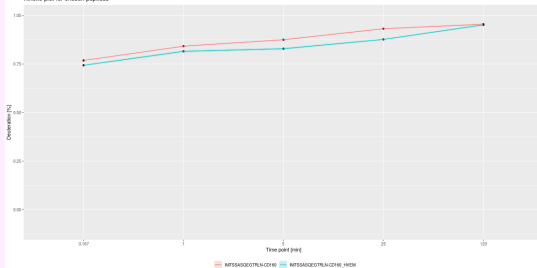
Choose peptide:

Sequence	State	Start	End
INITSSASQGETRLN	CD180	1	15
INITSSASQGETRLN	CD180_HVEM	1	15
LICTVW	CD180	16	21
LICTVW	CD180_HVEM	16	21
LICTVWHKKEEAEG	CD180	16	29
LICTVWHKKEEAEG	CD180_HVEM	16	29
ICTVWHKKEEAEG	CD180	17	29
ICTVWHKKEEAEG	CD180_HVEM	17	29
FVVFLL	CD180	30	34
FVVFLL	CD180_HVEM	30	34

Kinetic plot

Data

Kinetic plot for chosen peptides



Save chart (.svg)

Name	Value
HDX time course	8
Number of peptides	41
Sequence coverage	0.8636
Average peptide length	14.6829
Redundancy	4.5606
Replicates	4
Significant differences in HDX	0.98 0.99
All	All

[Excel](#) [PDF](#)

Showing 1 to 7 of 7 entries

This table was adapted from the HDX Data Summary (Masson, G.R., Burke, J.E., Ahn, N.C., Anand, G.S., Borchers, C., Brier, S., Bou-Assaf, G.M., Engen, J.R., Englander, S.W., Faber, J., et al. [2019]. Recommendations for performing, interpreting and reporting hydrogen deuterium exchange mass spectrometry (HDX-MS) experiments. Nat Methods 16, 595-602).

Choose items for report:

- | | |
|---|---|
| <input checked="" type="checkbox"/> Position Frequency | <input type="checkbox"/> Position Frequency Data |
| <input checked="" type="checkbox"/> Peptide Coverage | <input type="checkbox"/> Peptide Coverage Data |
| <input checked="" type="checkbox"/> Comparison Plot | <input type="checkbox"/> Comparison Plot Data |
| <input checked="" type="checkbox"/> Theoretical Comparison Plot | <input type="checkbox"/> Theoretical Comparison Plot Data |
| <input checked="" type="checkbox"/> Woods Plot | <input type="checkbox"/> Woods Plot Data |
| <input checked="" type="checkbox"/> Theoretical Woods Plot | <input type="checkbox"/> Theoretical Woods Plot Data |
| <input type="checkbox"/> Kinetic Plot | <input type="checkbox"/> Kinetic Plot Data |
| <input type="checkbox"/> Theoretical Kinetic Plot | <input type="checkbox"/> Theoretical Kinetic Plot Data |

Elements chosen for report have the same parameters as chosen in panel e.g. axis range and title. Adjust parameters for plots as needed in the report.

 Create report!

- a web-server (<http://mslab-ibb.pl/shiny/HaDeX/>),
- the R package
(<https://CRAN.R-project.org/package=HaDeX>),
- a standalone software
(<https://sourceforge.net/projects/HaDeX/>).

HaDeX developers:

- Weronika Puchala (main developer),
- Dominik Rafacz (frontend developer).

- Foundation of Polish Science TEAM TECH CORE FACILITY/2016-2/2 Mass Spectrometry of Biopharmaceuticals - improved methodologies for qualitative, quantitative and structural characterization of drugs, proteinaceous drug targets and diagnostic molecules.

MI² DATA LAB

MI² Data Lab (<https://mi2.mini.pw.edu.pl/>), Faculty of Mathematics and Information Science, Warsaw University of Technology.



Contact: michalburdukiewicz@gmail.com.

Presentation: <https://github.com/michbur/dpm>.

REFERENCES I

- Joint Committee for Guides in Metrology (2008). JCGM 100: Evaluation of Measurement Data - Guide to the Expression of Uncertainty in Measurement. Technical report, JCGM.
- Masson, G. R., Burke, J. E., Ahn, N. G., Anand, G. S., Borchers, C., Brier, S., Bou-Assaf, G. M., Engen, J. R., Englander, S. W., Faber, J., Garlish, R., Griffin, P. R., Gross, M. L., Guttman, M., Hamuro, Y., Heck, A. J. R., Houde, D., Iacob, R. E., Jørgensen, T. J. D., Kaltashov, I. A., Klinman, J. P., Konermann, L., Man, P., Mayne, L., Pascal, B. D., Reichmann, D., Skehel, M., Snijder, J., Strutzenberg, T. S., Underbakke, E. S., Wagner, C., Wales, T. E., Walters, B. T., Weis, D. D., Wilson, D. J., Wintrode, P. L., Zhang, Z., Zheng, J., Schriemer, D. C., and Rand, K. D. (2019). Recommendations for performing, interpreting and reporting hydrogen deuterium exchange mass spectrometry (HDX-MS) experiments. Nature Methods, 16(7):595–602.