



Top-Down Proteomics Workshop at ASMS 2025

Software and Data Analysis Strategies for Getting the Most Out of Your Data

Presiders:

Corinne Lutomski (Robinson Lab / Oxford University)

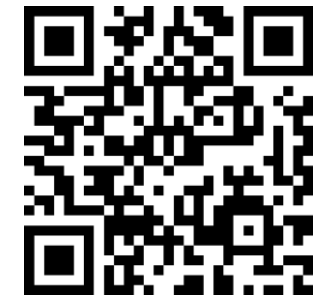
Fanny Caroline Liu (Bleiholder Lab / Florida State University)

Please access Slido Q&A

73rd Annual ASMS Meeting

Baltimore, MD

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Code: 5417047

Top-Down Proteomics Workshop at ASMS 2025

Software and Data Analysis Strategies for Getting the Most Out of Your Data



Xiaowen Liu
TopPIC



Boris Krichel
MASH



Ken Durbin
ProSight Suite



Kyowon Jeong
FLASH



Jack Bennett
precisIION



Mariangela Kosmopoulou
OmniScape



Michael Shortreed
Proteoform Suite

Please access Slido Q&A



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Intro survey



Do you currently use Top-Down (TDP) approach in your research?

Yes



No, but I would like to



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Is TDP software an obstacle in your work?

Yes



89%

No



11%



What is the largest obstacle you encounter with TDP software?

Choosing the right software for my application



Spectral deconvolution



Evaluating the quality of the proteoform identification



Other/not listed



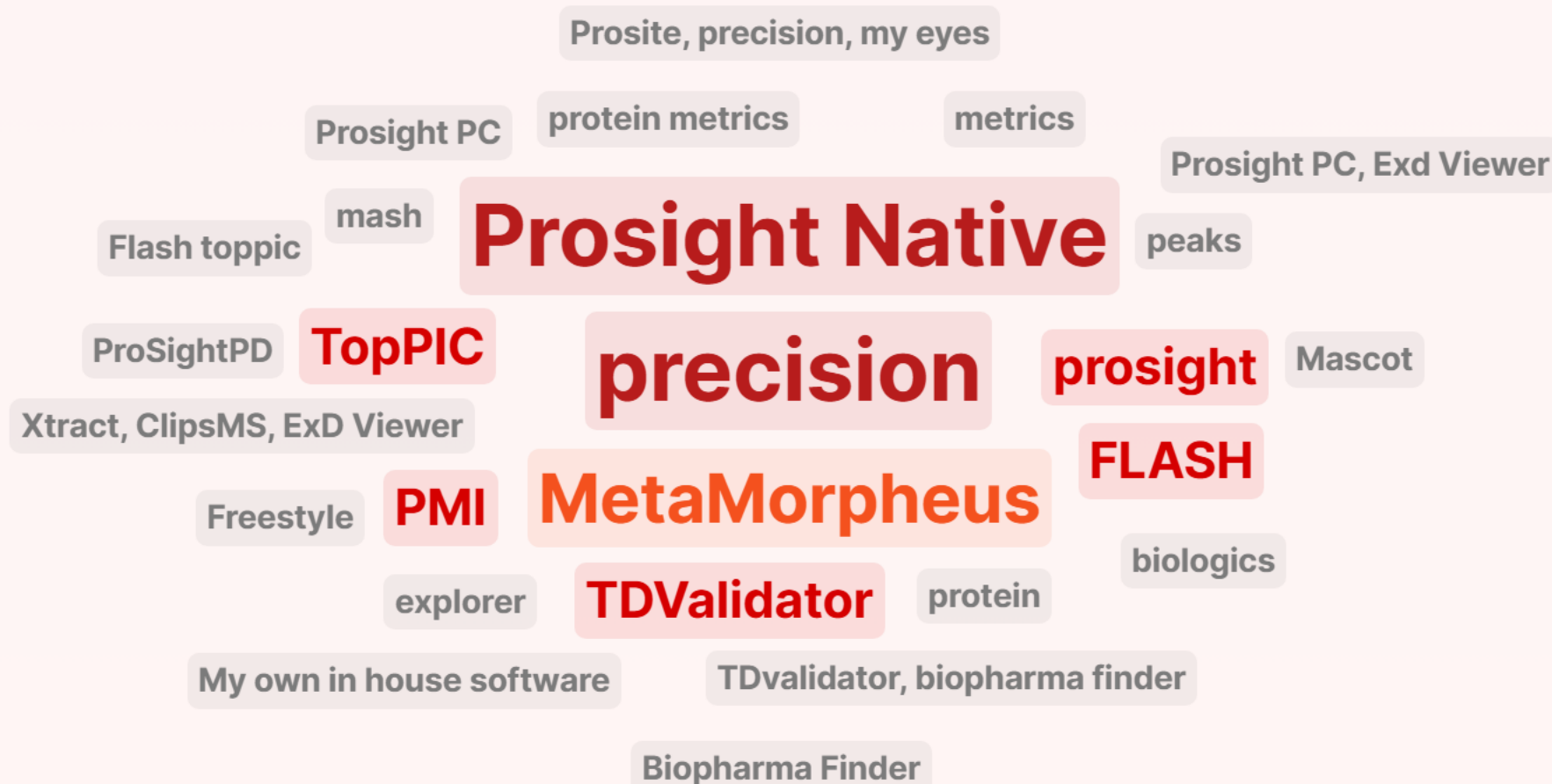
Database searching



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What software do you use?



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What would you like TDP software to be able to do but currently can't?

Internal fragments are a problem

Better validation (Retention/Migration Times, deconvolution, etc.) and classification.

Give me reliable monoisotopic masses for both precursors and fragments; I often get masses that are clearly 1-2 Da or even more off, especially for larger masses! And there's no good way to minimize false positives while not making huge concessions for false negatives!

Untargeted analysis of top down proteomics

Manually inspect annotations; Integrated data visualization



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What would you like TDP software to be able to do but currently can't?

Correct for monoisotopic mis assignment instead of assigning as hydrogen atom exchange

Identify ligand binding sites and variable modifications

Data interpretation and visualization

Get reliable label-free quantification

Accurate deconvolution

Be more user friendly !!!! Always very complicated



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What would you like TDP software to be able to do but currently can't?

be more user friendly :::: Always very complicated

database searching and mapping like bottom up proteomics

Assign more peaks based on empirical observations of fragmentation propensity in tdms rather than adopting bottom up approaches

Glyco

Label raw mass spectra, be more reliable in fragment assignment and protein ID

Automatic fragment peak annotation



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What topic are you most interested in discussing in this workshop?

Evaluating the quality of proteoform IDs



Database searching



Which software is right for my application?



What makes data high quality and how to obtain it



Spectral deconvolution



Something else (fill in blank below)





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Do you have any suggestions for the topic of the next TDP workshop?

When we have BUP and TDP data of one sample how can we relate them together to find specific PTMs shared between them??

How to obtain high-quality proteoform identification in complex TDP datasets

End-to-End validation of identification quality

Proteoform quantitation for function and mechanism

Sample preparation and separation of intact proteins

What are the best softwares for database searching and relative



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Do you have any suggestions for the topic of the next TDP workshop?

Proteoform quantitation for function and mechanism

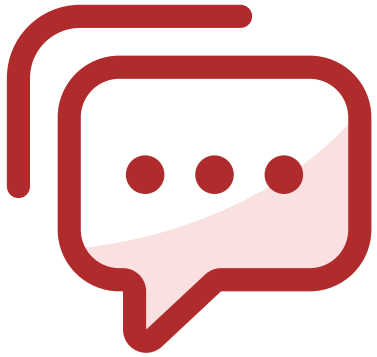
Sample preparation and separation of intact proteins

What are the best softwares for database searching and relative quantitation?

Analysing complex samples natively

No


Lc-ms tips



Audience Q&A




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 Anonymous

3


How can we analyze modifications other than PTM in different tools? for identification and quantification

 Anonymous

2

Do the softwares analyze protein information including adducts or metal complex in the MS1 and/or MS2 data?

Latest question


 Anonymous

2

How do you normalize ???




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 Anonymous

1

Which software suites have support for ion mobility?


 Anonymous

1

How do you approach proteoform quantification

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Latest question


 Anonymous

2

How do you normalize ???




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 Anonymous

0

What does each developer think their software is most well suited for?


 Anonymous

0

Can you please specify the difference between Top-Down MS and Top-Down Proteomics? And which Tool is suitable for TD MS and TDP?

[↑ Back to top](#)

Latest question

 Anonymous

2

How do you normalize ???



Anonymous

0

Can you please specify the difference between Top-Down MS and Top-Down Proteomics? And which Tool is suitable for TD MS and TDP?

Anonymous

0

Can I quantitatively analyze proteins with and without glycosylation using intact or top down method?

↑ Back to top

Latest question

Anonymous

2

How do you normalize ???

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Thank you for participating in this work

**See you in the Top-Down Proteomics Workshop
at ASMS 2026**