# Evaluating the Addition of Supercharging Reagents During CESI-MS for Identification and Characterization of Biopharmaceuticals

*m/z* observed

218.1503

262.152

315.2031

347.1935

439.2193

236.6423

225.6136

280.6635

448.2772

457.2513

517.3084

255.1054

313.1451

263.6529

281.1845

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# **OVERVIEW**

**Purpose**: Previously, we reported on the integration of capillary electrophoresis (CE) with electrospray ionization (ESI) into one dynamic process which we have termed CESI. Our purpose of this study is to evaluate the effect of supercharging reagents on peptide ionization during CESI-MS.

Methods: Trypsin digested monoclonal antibody was separated by CE and identified by tandem MS. Supercharging reagents were added to the CE background electrolyte (BGE).

**Results**: Supercharging during CESI-MS showed a significant increase in the number of higher charge states attained by the peptides. In addition, for glycopeptides, the intensity of the precursor ions as well as fragment ions increased with supercharging.

# INTRODUCTION

- > The integration of CE and ESI into CESI is achieved by making the end of the separation of capillary porous to ion flow – this is achieved through an HF etching process.
- > This tip is then inserted into a metal needle and filled with a static conductive liquid to establish an electrical connection
- Analytes migrate according electrophoretic their to mobility inside the capillary and when it reaches the porous sprayer tip, it is electro-sprayed into MS.
- The ultra-low flow of the process, yields very high ionization efficiency and serves to reduce ion suppression.



### Schematic of CESI porous tip sprayer

- > Supercharging reagents have been demonstrated to enhance the multiple charging of proteins and peptides during ESI. This improves the mass measurement accuracy and fragmentation efficiency.
- Recently, Bonvin et al., have added the supercharging reagents to the sheath liquid interface during CE-MS to modulate charge state distribution of proteins.<sup>1</sup> But, the sheath flow dilutes the analytes migrating from the capillary as well as increasing the flow. Alternatively, supercharging reagents can be directly added to the BGE during CESI-MS, a ultra-low flow technique, to further maximize ionization.
- Palivizumab, a humanized therapeutic antibody, targeted against respiratory syncytial virus, is an ideal representation of IgG1 type monoclonal antibody. Tryptic peptides generated from this antibody were analyzed by CESI-MS and supercharging reagents were added to evaluate the efficiency of charge enhancement during CESI-MS.

# MATERIALS AND METHODS

**Sample preparation:** 100 µg of palivizumab was solubilized, reduced with DTT, alkylated with IAM, and digested overnight with trypsin. The sample was then dried down and resuspended in 100 µl of leading electrolyte (100 mM ammonium acetate at pH 4).

CESI-MS conditions: 50 nL of the sample was injected into the separation capillary and transient-isotachophoresis focuses the sample during electrophoretic separation. 10% acetic acid served as the BGE and a separation voltage of 20 kV at normal polarity was applied. For supercharging experiments, reagents such as *m*-NBA and sulfolane were added to the BGE.

CESI was connected on-line to a TripleTOF<sup>®</sup> 5600+ MS. Information dependent acquisition (IDA) mode consisting of 250 msec TOF MS survey scan and 50 msec IDA on the top 30 ions which exceed 150 cps was utilized for data acquisition. Rolling collision energy was used to induce fragmentation and the dynamic exclusion time was set to 5 sec. The total run time was set to 60 min. Auto-calibration was enabled once every 3 runs.

Data analysis: Data analysis was performed using AB SCIEX BioPharmaView<sup>™</sup> and ProteinPilot<sup>™</sup> software. Glycopeptides were extracted manually and MS/MS spectra were checked for diagnostic ions.

No.

2

3

4

5

6

14

15

Sequence

AK

SR

APK

EAK

EYK

VEPK

DTSK

VQWK

TISK

GQPR

**VDKR** 

SCDK

**ADYEK** 

**QPPGK** 

LTISK

# RESULTS

## I. Identification of small hydrophilic and large hydrophobic peptides

2.4e6 -

2.2e6 -

2.0e6 -

1.8e6 -

1.6e6 -

1.4e6 -

1.2e6 -

1.0e6 -

8.0e5 -

6.0e5 -

4.0e5 -

2.0e5

0.0e0 -

Extracted ion electropherogram (EIE) of tryptic peptides from palivizumab ranging from 2 to 5 amino acids. The table provides more information about each peptide in the EIE.

> Small hydrophilic peptides, which often lost in the void volume in RP-LC can be identified well by CESI-MS. Shown here are the peptides with 2-5 residues identified from both heavy and light chains (HC & LC) of palivizumab.





### II. Charge state distribution of peptides during supercharging

Antibody

chain

HC

HC

LC

LC

HC

HC

HC

HC

HC

HC

LC

HC

HC

Charge

1

- each instance.
- without any reagent.
- added.
- Addition of supercharaina reagents results in an increase in the average and maximum charges attained by the peptides.
- NBA was present.

EIE of 63-mer tryptic peptide without any reagent (A) and with 1% *m*-NBA (C). (B) and (D) show the mass spectrum without and with 1% *m*-NBA respectively.

> Electrophoretic separation is based on the charge-to-hydrodynamic volume ratio of the peptides and not based on hydrophilic / hydrophobic interactions with the separation capillary. Large hydrophobic peptides can also be identified equally well, resulting in the high sequence coverage obtainable by this technique.

| (sample 1)liv11, +TOF MS (250 - 3000): 701.3190 +/- 0.0025 Da<br>QGNVFSCSVMHEALHNHYTQK<br>25 26 27 28 29 30 31 32 33 34 35 36<br>Time, min<br>(sample 1)liv11, +TOF MS (250 - 3000): 896.3872 +/- 0.0025 Da | Sequence                                                 | <i>m/z</i><br>observed | Charge | No. of<br>amino<br>acids |
|-------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|----------------------------------------------------------|------------------------|--------|--------------------------|
| 25 26 27 28 29 30 31 32 33 34 35 36                                                                                                                                                                         | 420WQQGNVFSCSVM<br>HEALHNHYTQK442                        | 701.319                | 4      | 23                       |
| (sample 1)liv11, +TOF MS (250 - 3000): 937.4634 +/- 0.0025 Da                                                                                                                                               | <sup>84</sup> VTNMDPADTATYY<br>CAR <sup>99</sup>         | 896.3872               | 2      | 16                       |
| 25 26 27 28 29 30 31 32 33 34 35 36<br>Time, min<br>(sample 1)liv11, +TOF MS (250 - 3000): 848.7163 +/- 0.0025 Da                                                                                           | <sup>396</sup> TTPPVLDSDGSFF<br>LYSK <sup>412</sup>      | 937.4634               | 2      | 17                       |
| 25 26 27 28 29 30 31 32 33 34 35 36<br>Time, min<br>(sample 1)v11, +TOF MS (250 - 3000): 1178.1021 +/- 0.0025 Da                                                                                            | <sup>374</sup> GFYPSDIAVEWES<br>NGQPENNYK <sup>395</sup> | 848.7163               | 3      | 22                       |
| 25 26 27 28 29 30 31 32 33 34 35 36                                                                                                                                                                         | <sup>6</sup> ESGPALVKPTQTLTL<br>TCTFSGF <sup>27</sup>    | 1178.1021              | 2      | 22                       |
| rang, mm                                                                                                                                                                                                    |                                                          |                        |        |                          |

Extracted ion electropherogram (EIE) of peptides from palivizumab HC ranging from 16 to 23 amino acids. The table provides more information about each peptide in the EIE.

> Sulfolane and *m*-NBA were added to the BGE (10% acetic acid) at 1% concentration (v/v).

> Number of spectra obtained at 95% confidence was plotted as a function of charge states for

> With 1% sulfolane, the % of confident spectra at 3+ and 4+ charges increased compared to

 $\succ$  With 1% *m*-NBA, 50.4% of confident spectra were obtained from 3+ charge state.

> Small % of confident spectra at 5+ charges were present only when *m*-NBA or sulfolane was



> Relative intensity of the highest observed charge state also increases. For instance, in palivizumab HC, a 63 amino acid tryptic peptide without any missed cleavage had 5 fold more intensity with 1% *m*-NBA. > Consequently, this peptide was identified by MS/MS with a total of 22 b & y fragment ions only when m-



Mass/Charge, Da

**Precursor Charge Distribution** 

with 1% *m*-NBA

18.8%

Mass/Charge, Da

27.7%

### III. Glycopeptide identification and supercharging

- > Relative intensities of glycopeptides (G0F, G1F, & G2F) supercharging were compared.
- Addition of either reagent showed a significant increase in the relative abundance of all 3 glycopeptides at 3+ charge state.
- $\succ$  Increased intensity of glycopeptide precursor ions obtained from supercharging resulted in highly intense fragment ions during tandem MS. The intensity of [peptide+glycan] fragments increased by approximately 5 to 10 fold for all three glycopeptides.

Tandem MS spectra of G0F glycopeptide at 3+ charge with and without supercharging reagents. The inset shows the relative abundance of fragment ion at 1392 m/z.



In addition, 4+ charge state for all 3 glycopeptides were present only with addition of *m*-NBA. This is beneficial for glycopeptide analysis since it moves the high m/z glycopeptide precursor ions into the lower m/z region where it can be easily selected for MS/MS, without any limitation from the mass analyzer.

### CONCLUSIONS

- high sequence coverage for antibody identification.
- analysis.

### REFERENCES

Bonvin, G., Rudaz, S., Schappler, J. Analytica Chimica Acta 813 (2014) 97-105.

### TRADEMARKS/LICENSING

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from palivizumab with and without



> CESI-MS allowed efficient analysis of both short hydrophilic and long hydrophobic peptides, thereby providing

Charge state enhancement was observed for tryptic peptides during CESI-MS. Higher charge states fall in the lower m/z region and thus, can be fragmented without any limits for precursor selection during MS/MS.

Addition of supercharging reagents during CESI-MS enhances glycopeptide identification. The relative abundances of precursor and fragment ions significantly improved for different glycopeptides. Furthermore, greater S/N ratio, improved peak shape, and mass resolution can potentially benefit automated glycopeptide