

Peptide ion fragmentation in mass spectrometry

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4-7117

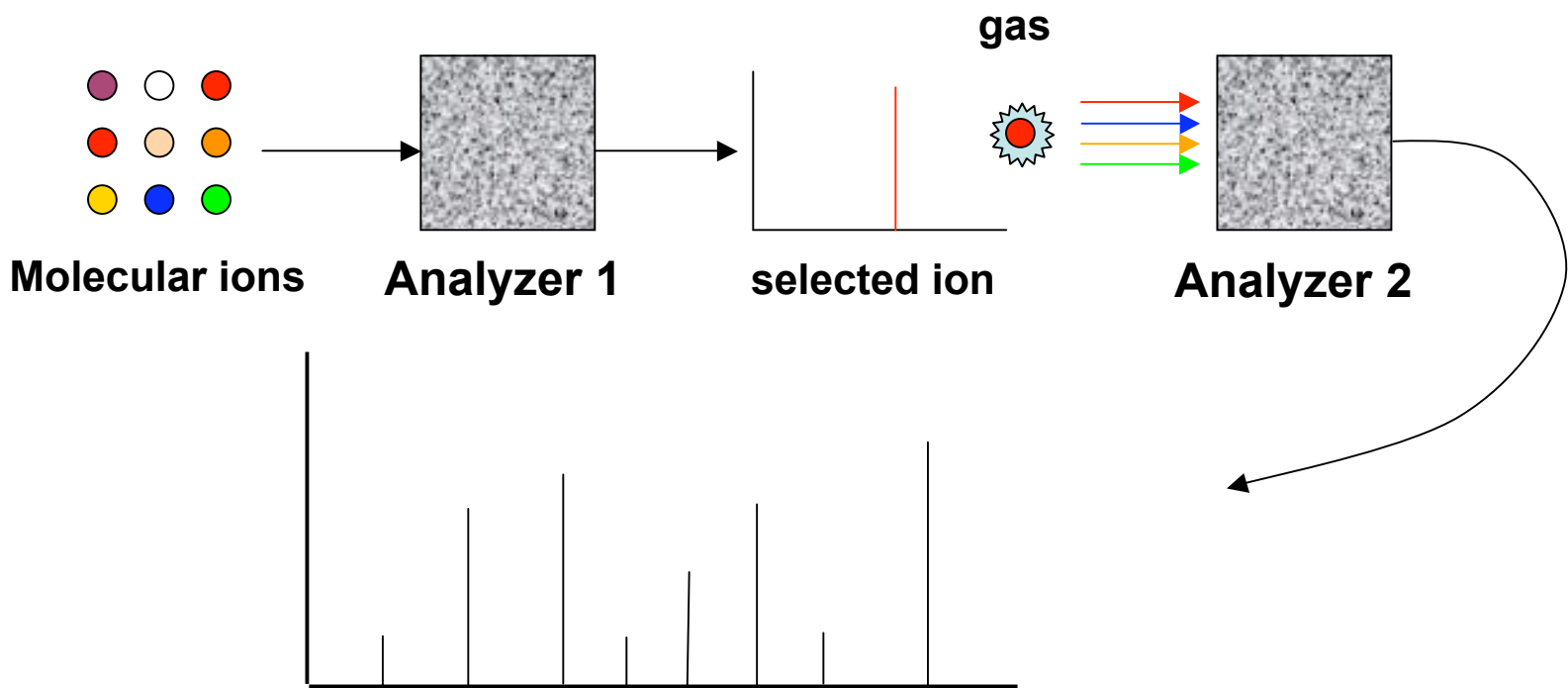
Lecture goals

- **Value of fragmentation in determining structure**
- **How peptides fragment**
 - Interpreting the tandem mass spectrum
- **Automating identification of peptides from their fragment ions**
 - pros and cons
- **Controlling fragmentation**
 - Choice of ionization and fragmentation methods

Why ion fragmentation provides useful information

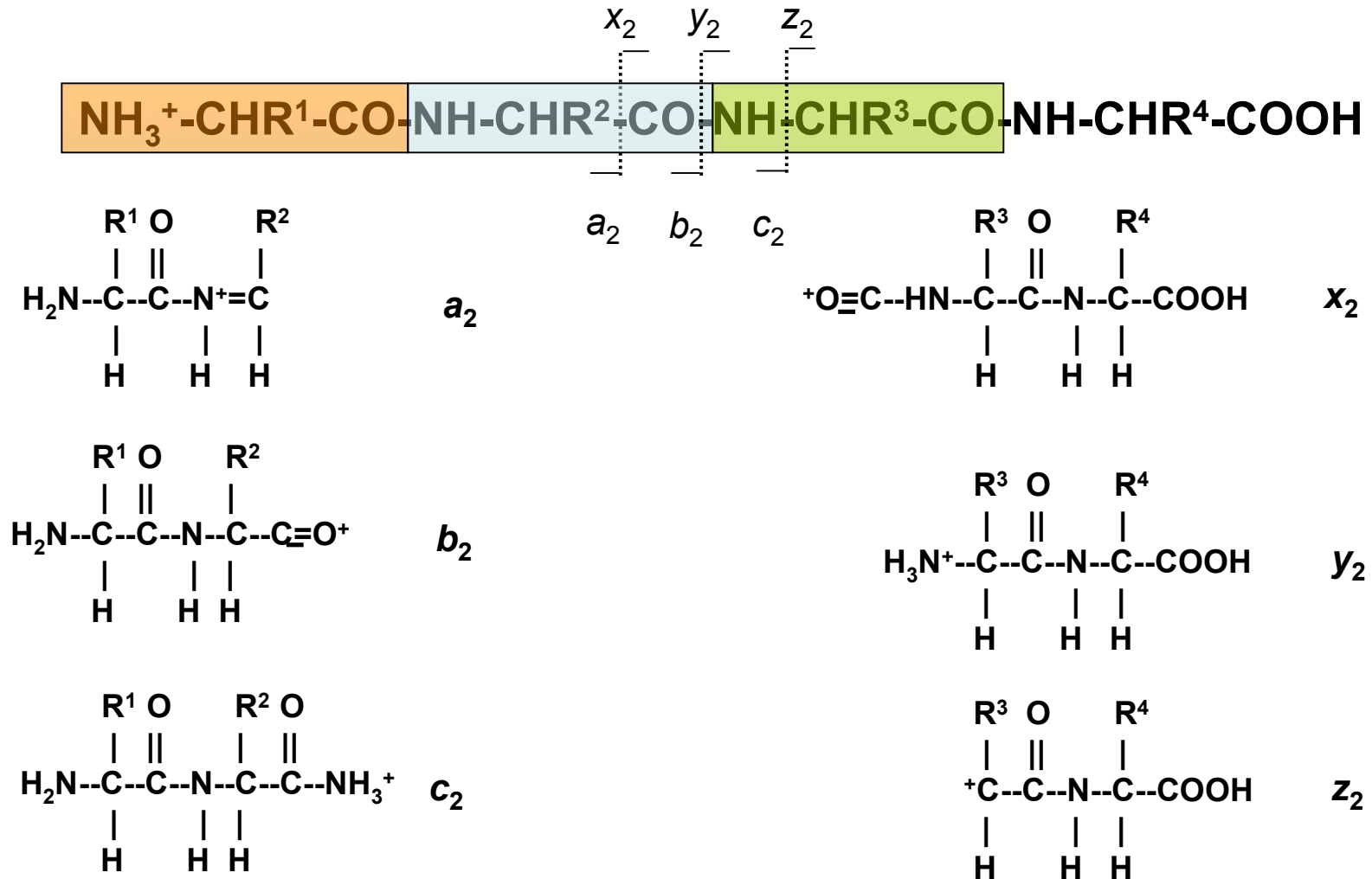
- Compounds can have the same empirical formula, i.e., the same molecular weight or m/z , but be different chemically.
- Breaking them into parts (fragmenting them) helps to identify what they are.
- Each of the following peptides gives rise to the same $[M+2H]^{2+}$ ion
 - $\text{NH}_2\text{VFAQHLK-COOH}$ $\text{NH}_2\text{VAFQHLK-COOH}$
 - $\text{NH}_2\text{VFQHALK-COOH}$ $\text{NH}_2\text{VHLAFQK-COOH}$
- In proteomics we want to distinguish these peptides

What is MS-MS (tandem mass spectrometry)?



Collision-induced dissociation

Fragmenting a peptide



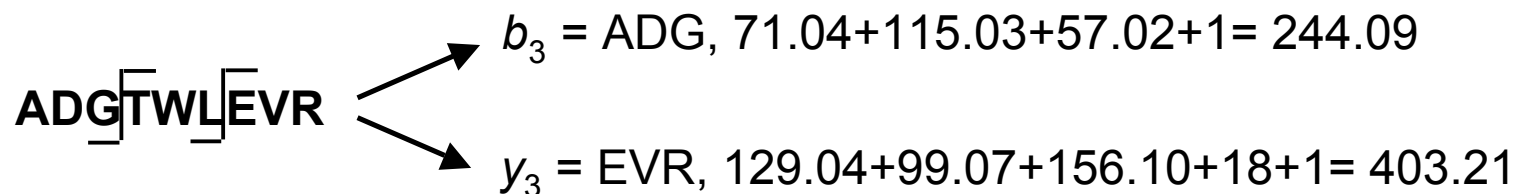
Adapted from http://www.matrixscience.com/help/fragmentation_help.html

Calculating expected *b*- and *y*-ion fragments

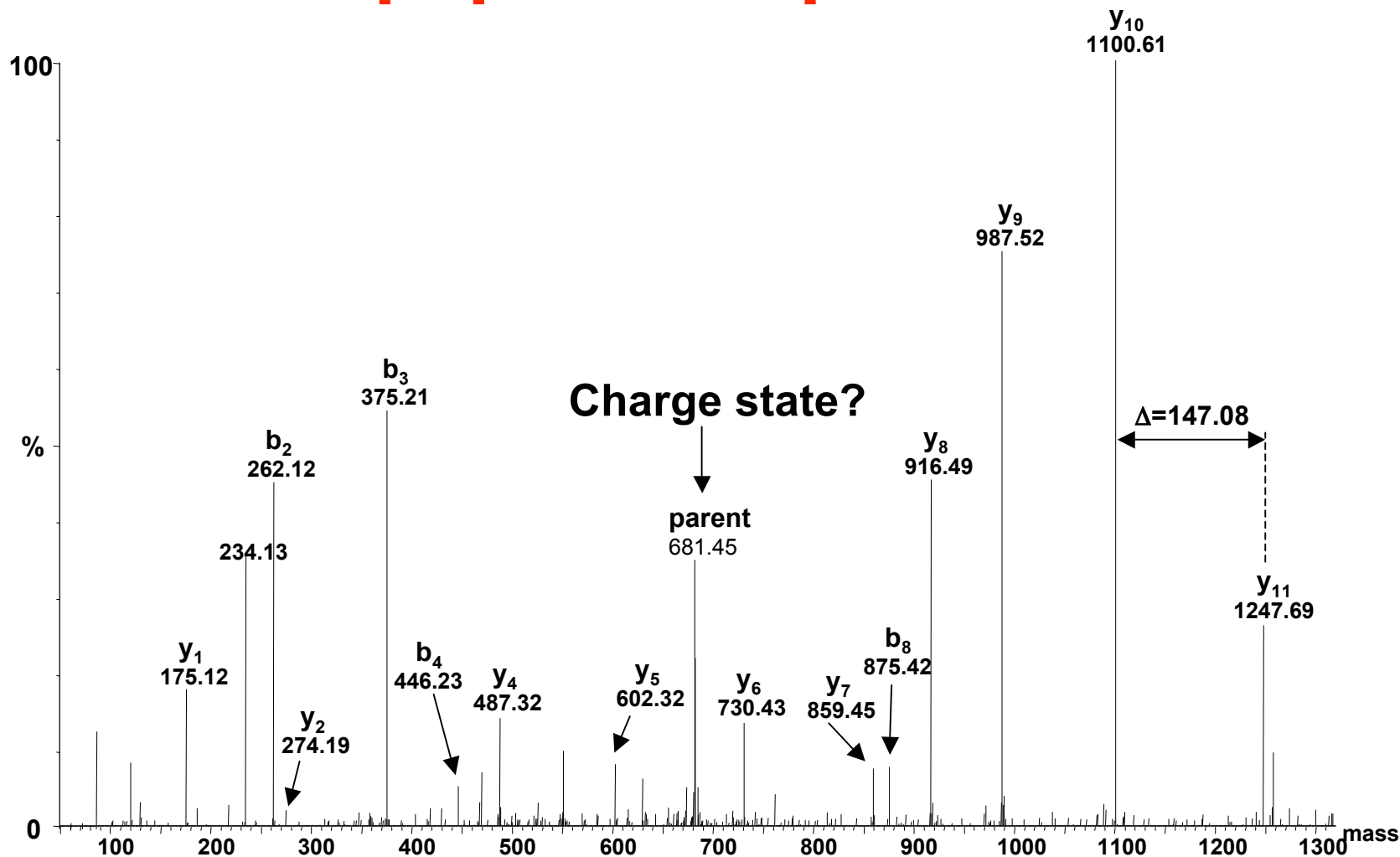
Alanine	71.037	Leucine	113.084
Arginine	156.101	Lysine	128.094
Asparagine	114.043	Methionine	131.040
Aspartic acid	115.027	Phenylalanine	147.068
Cysteine	103.009	Proline	97.053
Glutamic acid	129.043	Serine	87.032
Glutamine	128.058	Threonine	101.048
Glycine	57.021	Tryptophan	186.079
Histidine	137.059	Tyrosine	163.063
Isoleucine	113.084	Valine	99.068

b_n = [residue masses + 1] - these come from the N-terminus

y_n = [residue masses + H₂O + 1] = these come from the C-terminus



Identification of daughter ions and peptide sequence



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What's in a peptide MSMS spectrum?

- In most cases, some, but rarely all, of the theoretic *b*- and *y*-ions are observed
- Besides *b*- and *y*-ions, other types of fragmentation can occur to form a_n and x_n ions, as well as also losing CO, NH₃ and H₂O groups
- Internal cleavage reactions can occur at acidic (Asp - Glu) residue sites

Identifying a peptide by de novo sequencing

- Take the partial sequence that can be identified manually and submit it to PROWL (<http://prowl.rockefeller.edu/>) - click on PROTEININFO and enter sequence - select all species
- Use suggested sequences to fill in the gaps and then check all theoretical ions using MS-Product at <http://prospector.ucsf.edu/prospector/4.27.1/cgi-bin/msform.cgi?form=msproduct>

Laboratory of Mass Spectrometry and Gaseous Ion Chemistry

PROWL

- > ProFound
- > ProteinInfo
- > PeptideMap
- > PepFrag
- > X! Tandem
- > X! Hunter
- > GPMDB
- > PROWL
- > Chait Lab



The Rockefeller University
1230 York Avenue,
New York, NY 10021
(212) 327-8000



National Center for

PROTEININFO

Advanced Sequence Search

Analyze Amino Acid Sequence

Select a database:

Enter keywords:

Enter sequence:

Select Search Category:

All Categories

- | <input type="radio"/> Bacteria | <input type="radio"/> Eukaryota | <input type="radio"/> Viruses |
|---|---|---|
| <input type="radio"/> Firmicutes <ul style="list-style-type: none"> <input type="radio"/> Bacillus subtilis <input type="radio"/> Mycoplasma <input type="radio"/> Other Firmicutes | <input type="radio"/> Dictyostelium discoideum <ul style="list-style-type: none"> <input type="radio"/> Fungi <ul style="list-style-type: none"> <input type="radio"/> Pneumocystis carinii <input type="radio"/> Saccharomyces cerevisiae <input type="radio"/> Schizosaccharomyces pombe <input type="radio"/> Other Fungi <input type="radio"/> Metazoa <ul style="list-style-type: none"> <input type="radio"/> Caenorhabditis elegans <input type="radio"/> Chordata <ul style="list-style-type: none"> <input type="radio"/> Fugu rubripes <input type="radio"/> Danio rerio <input type="radio"/> Mammalia <ul style="list-style-type: none"> <input type="radio"/> Primates <ul style="list-style-type: none"> <input type="radio"/> Homo sapiens <input type="radio"/> Other primates <input type="radio"/> Rodentia <ul style="list-style-type: none"> <input type="radio"/> Mus musculus | <input type="radio"/> Hepatitis C V <ul style="list-style-type: none"> <input type="radio"/> Other Viruses |
| <input type="radio"/> Proteobacteria <ul style="list-style-type: none"> <input type="radio"/> Enterobacteria <ul style="list-style-type: none"> <input type="radio"/> Escherichia coli <input type="radio"/> Other Enterobacteria <input type="radio"/> Other Proteobacteria | | |
| <input type="radio"/> Other Bacteria <ul style="list-style-type: none"> | | |

MS-Product

Peptide Sequence

N term **C term**

Enter Sequence in Capital letters except:
 | m - Met-ox | h - Homoserine lactone | U - Selenocysteine |
 | s,t,y - Phosphorylated S,T,Y | u, v, w, x - user specified amino acids |

**Constant
 Mods**

Acetyl (K)
 Amino (Y)
 Asn->Succinimide (N)
 Carbamidomethyl (C)

User Specified AA Elem Comp (u)

User Specified AA Elem Comp (v)

User Specified AA Elem Comp (w)

User Specified AA Elem Comp (x)

Use instrument specific defaults to override ion types below

AA Composition	ions N-term sequence	ions C-term sequence	ions Internal Fragment-ions	Ladder sequencing ions
i <input checked="" type="checkbox"/> m <input type="checkbox"/> a <input checked="" type="checkbox"/> b <input checked="" type="checkbox"/> c <input type="checkbox"/> x <input type="checkbox"/> y <input checked="" type="checkbox"/> Y <input type="checkbox"/> z <input type="checkbox"/>			internal <input checked="" type="checkbox"/>	N-term <input type="checkbox"/> C-term <input type="checkbox"/>
Satellite Sequence Ions (side-chain loss)	Neutral-loss Sequence Ions		Peeling Sequence Ions	
d <input type="checkbox"/> v <input type="checkbox"/> w <input type="checkbox"/>	-H ₂ O <input checked="" type="checkbox"/> S, T, E, DR, K, Q, N	-NH ₃ <input checked="" type="checkbox"/> (S, T, Y + PO ₄)	-H ₃ PO ₄ <input type="checkbox"/> s, t, y	-SOCH ₄ <input type="checkbox"/> m
			Multiple Losses <input type="checkbox"/>	b+H ₂ O <input checked="" type="checkbox"/> R, H, K

Induce Fragmentation

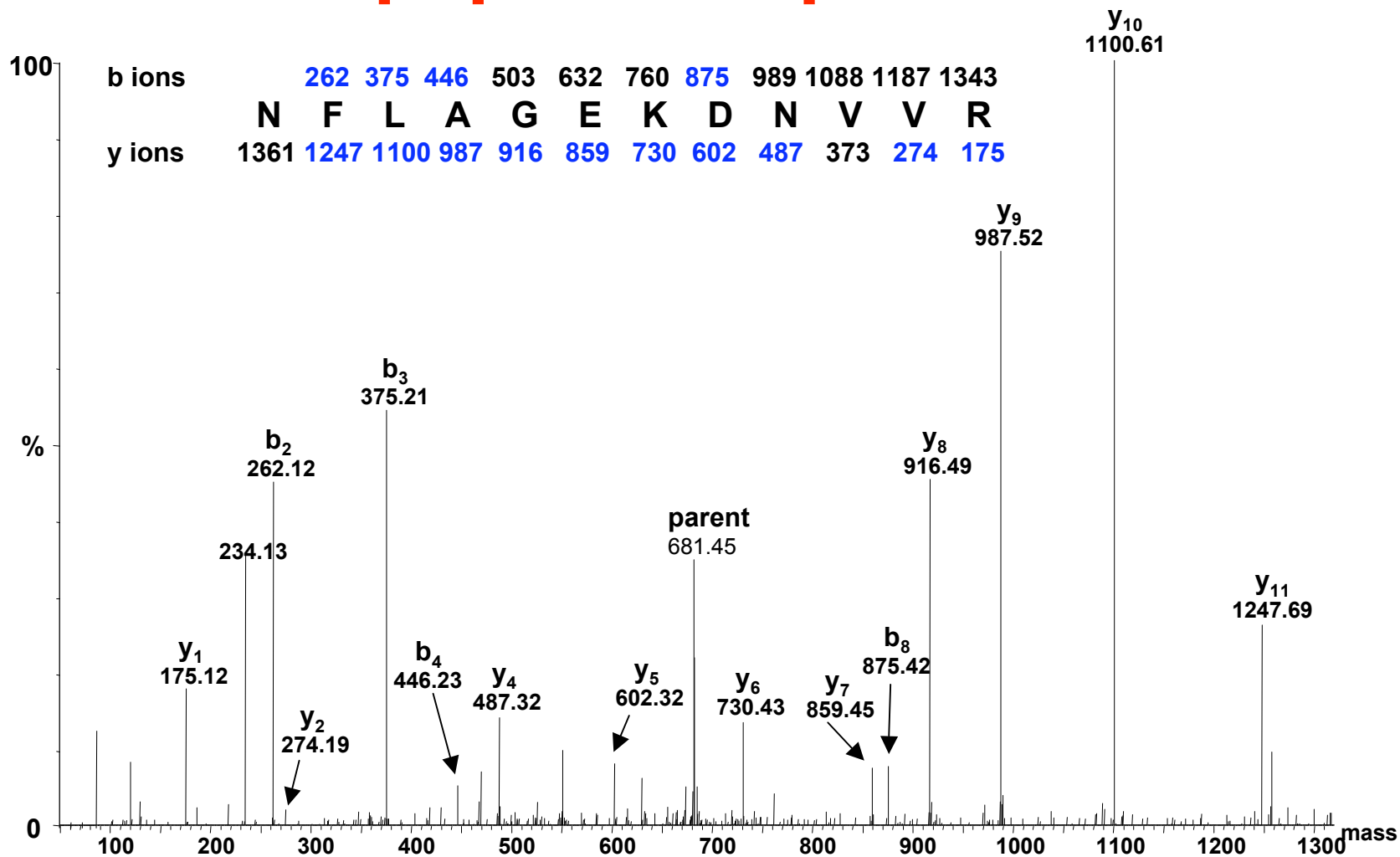
Display Graph Max. Charge Output Type Hits to file Name

Other ions observed in CID peptide fragmentation

Immonium and Related Ions

	87.06	120.08	86.10	---	---	102.05	84.08 101.11 129.10	88.04	87.06	72.08	72.08	70.07 87.09 100.09 112.09
N-terminal ions												
a-NH ₃ ions	---	217.10	330.18	401.22	458.24	587.28	715.38	830.40	944.45	1043.52	1142.58	---
a ions	---	234.12	347.21	418.24	475.27	604.31	732.40	847.43	961.47	1060.54	1159.61	---
b-NH ₃ ions	---	245.09	358.18	429.21	486.23	615.28	743.37	858.40	972.44	1071.51	1170.58	---
b-H ₂ O ions	---	---	---	---	---	614.29	742.39	857.42	971.46	1070.53	1169.59	---
b ions	---	262.12	375.20	446.24	503.26	632.30	760.40	875.43	989.47	1088.54	1187.61	---
	1	2	3	4	5	6	7	8	9	10	11	12
H -	N	F	L	A	G	E	K	D	N	V	V	R
y ions	---	1247.67	1100.61	987.52	916.48	859.46	730.42	602.33	487.30	373.26	274.19	175.12
y-NH ₃ ions	---	1230.65	1083.58	970.50	899.46	842.44	713.39	585.30	470.27	356.23	257.16	158.09
y-H ₂ O ions	---	1229.66	1082.60	969.51	898.47	841.45	712.41	584.32-	---	---	--	---

Identification of daughter ions and peptide sequence



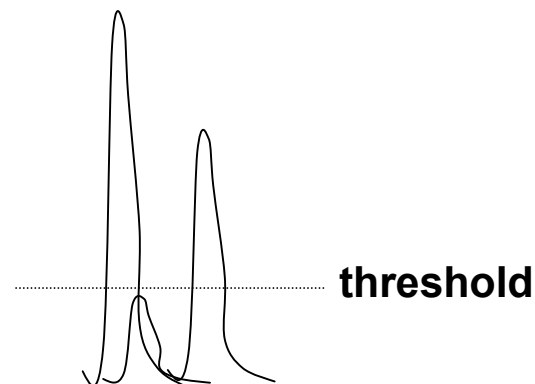
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Towards automated MSMS sequencing

- The 2D-LC-ESI-MSMS method (MuDPIT) generates 50,000+ MSMS spectra for each sample
- If it takes 15 min to hand interpret one MS-MS spectrum, then it would take 12,500 hours to complete the analysis. For someone working 8 hours/day and a five-day week, this would be about 6 years!
- Using SEQUEST and MASCOT, methods were developed to use computer-driven approaches to analyze MSMS data

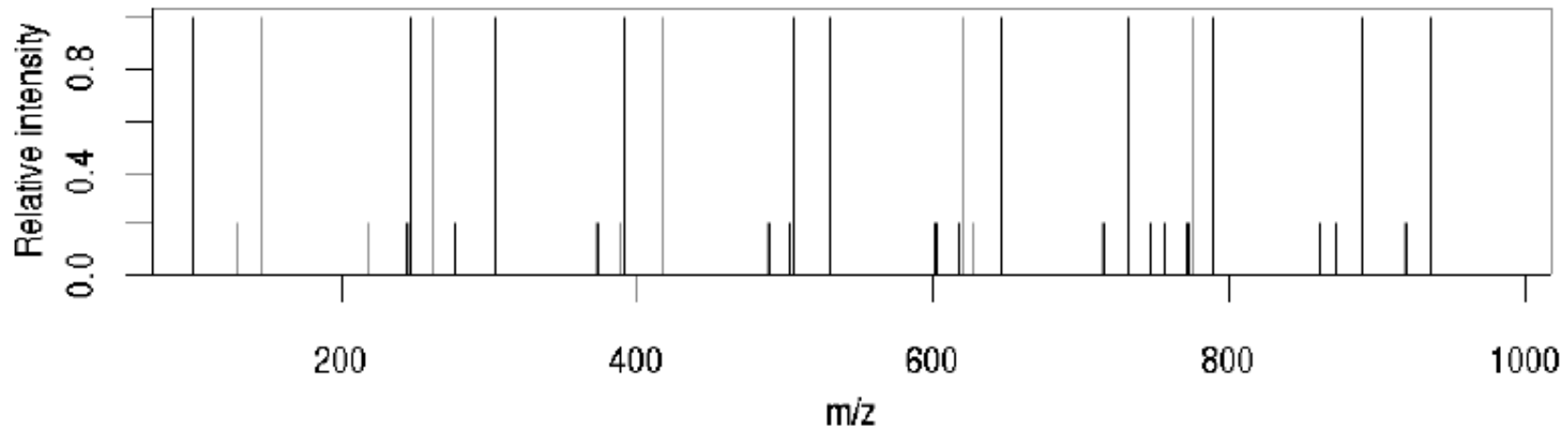
Issues in MS-MS experiment

- **At any one moment, several peptides may be co-eluting**
- **Data-dependent operation:**
 - **The most intense peptide molecular ion is selected first (must exceed an initial threshold value)**
 - **A 2-3 Da window is used (to maximize the signal)**
 - **The ion must be in 2⁺ or 3⁺ state**
 - **Since the ion trap scan of the fragment ions takes ~ 1 sec, only the most intense ions will be measured**
 - **However, can use an exclusion list on a subsequent run to study minor ions**



The SEQUEST approach

- Each observed MSMS spectrum has a corresponding molecular ion $[M+nH]^{n+}$. For ion trap data, ions are selected from the known or virtual proteome that are within 1 Da. These are then “fragmented” *in silico* to produce *b*- and *y*-ions and less abundant fragment ions.



- The cross correlations of the observed MSMS spectrum to each of the virtual MSMS spectra are calculated. The peptides are scored and the one having the highest score is deemed to be “identified”.

What has SEQUEST provided to proteomics?

- Initially, it seemed an awful lot! Typically, the “identified” proteins covered most of known biochemistry, so they satisfied everybody
- But the method obviously has limitations. There is redundancy - each protein yields multiple peptides
- The number of unique proteins is much less than the observed peptides
- **Critically, it was missing controls**

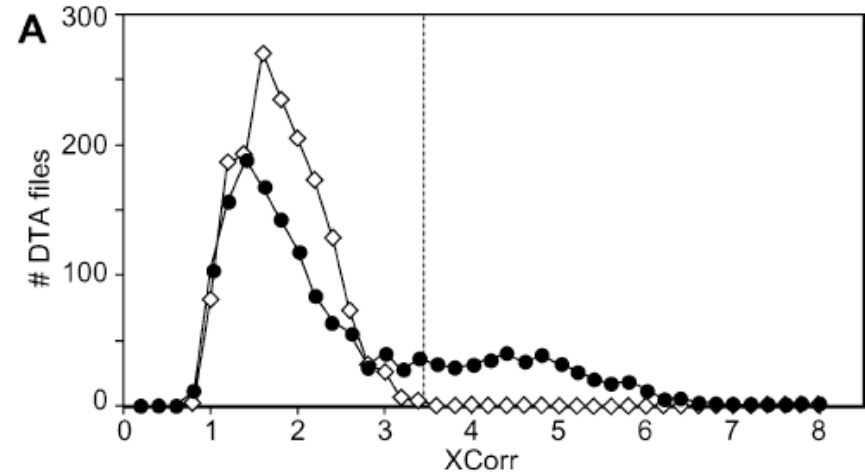
SEQUEST sequencing

- **Use of SEQUEST requires considerable computing power - if there are 500 possible peptides to compare, then examination of 50,000+ spectra would require 25 million correlations**
- **Data analysis is typically carried out using computer clusters to accelerate the analysis**

More haste, less speed?

- Post analysis, the masses of the peptides triggering MS-MS are used to create a set of virtual peptides with masses within ± 1 Da
- Predicted MS-MS are compared to the observed and the best fit is reported as a hit
- The abundance of these hits are plotted in the figure as closed circles

Resing et al. (2004)



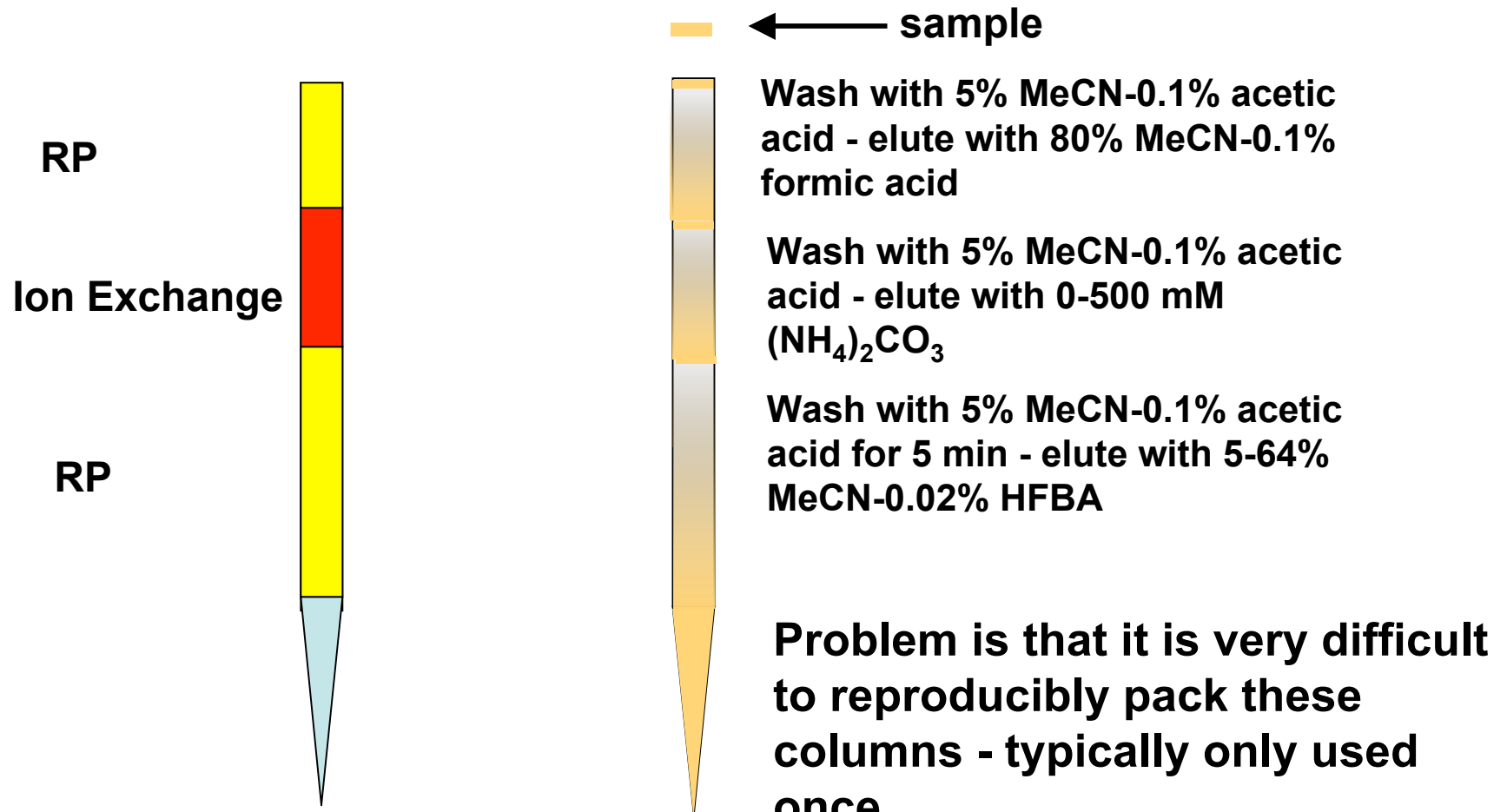
However, if the sequences of the peptides within ± 1 Da are reversed *in silico* and their predicted MS-MS compared to the observed spectra, a similar histogram is obtained (open circles), but without the right side tail

A forced fit to a set of data will always come up with a match, but not necessarily the truth

How to improve MUDPIT

- **Reproducible column engineering**
 - Tandem columns, each built to separate, but high specifications
 - Columns on a chip
- **More careful selection of the parent ion**
 - Accurate measurement of the peptide's mass will eliminate many false peptides
 - Accurate measurement of peptide fragments' masses
- **Greater stringency in assessing score cutoff**

Engineering of a MuDPIT column



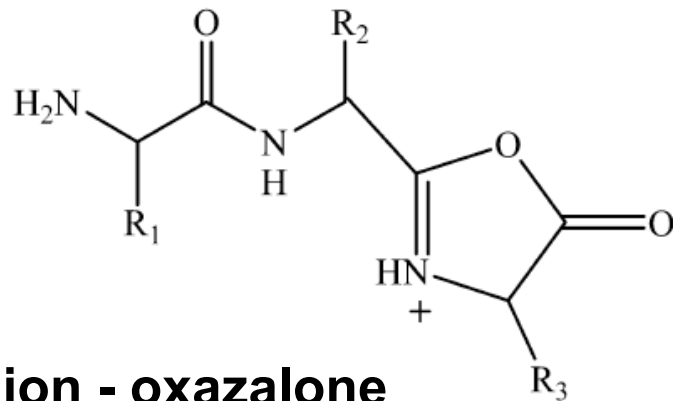
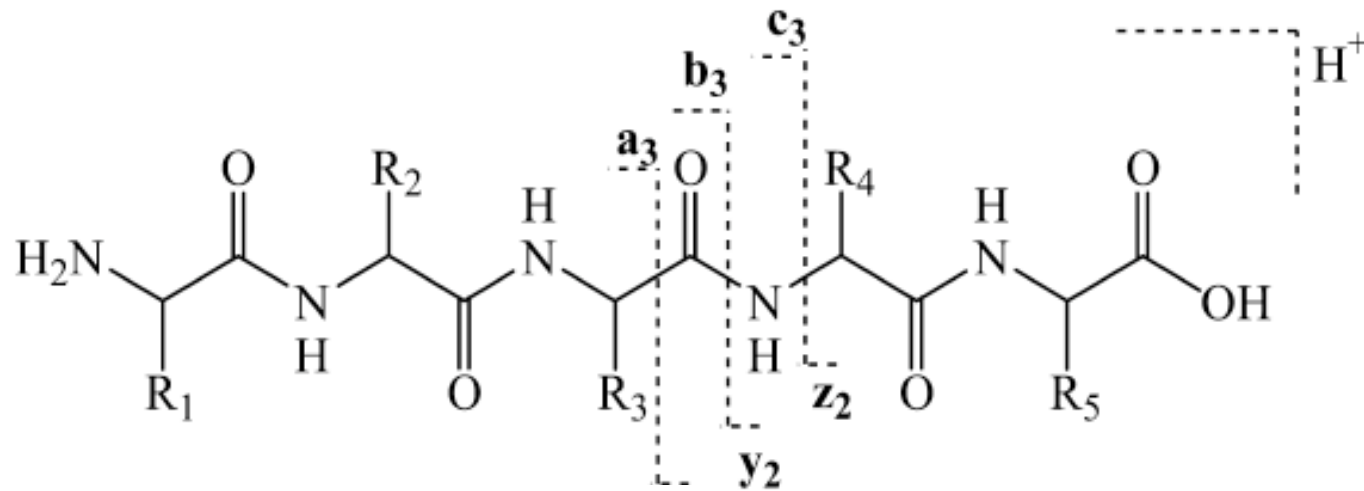
Packing right to end of the tip - no dead volume

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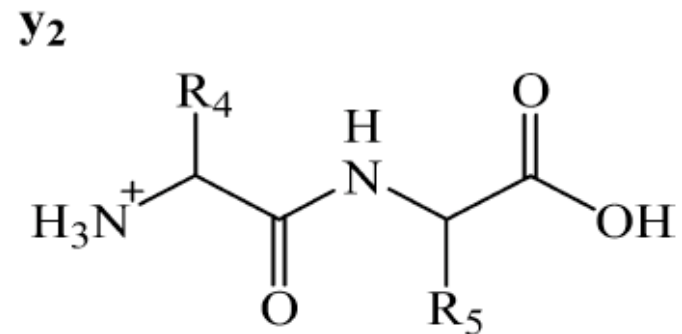
Mass accuracy limits the ions to consider

	Theoretical Mass	Delta [ppm]	Delta [mmu]	RDB	Composition
1 ppm (4)	516.76671	0.0	0.0	21.0	C ₄₉ H ₇₁ O ₁₂ N ₁₃
	516.76647	0.5	0.2	15.0	C ₄₉ H ₇₉ O ₁₁ N ₉ S ₂
	516.76638	0.6	0.3	12.0	C ₄₁ H ₇₅ O ₁₄ N ₁₅ S ₁
	516.76705	-0.7	-0.3	11.5	C ₄₃ H ₇₇ O ₁₅ N ₁₂ S ₁
2 ppm (10)	516.76604	1.3	0.7	16.0	C ₄₈ H ₇₅ O ₁₆ N ₉
	516.76738	-1.3	-0.7	20.5	C ₅₁ H ₇₃ O ₁₃ N ₁₀
	516.76604	1.3	0.7	21.5	C ₄₇ H ₆₉ O ₁₁ N ₁₆
	516.76580	1.8	0.9	15.5	C ₄₇ H ₇₇ O ₁₀ N ₁₂ S ₂
	516.76772	-2.0	-1.0	16.5	C ₄₄ H ₇₃ O ₁₁ N ₁₆ S ₁
	516.76773	-2.0	-1.0	11.0	C ₄₅ H ₇₉ O ₁₆ N ₉ S ₁
5 ppm (23)	516.76805	-2.6	-1.3	25.5	C ₅₂ H ₆₉ O ₉ N ₁₄
	516.76537	2.6	1.3	16.5	C ₄₆ H ₇₃ O ₁₅ N ₁₂
	516.76807	-2.6	-1.4	7.0	C ₃₈ H ₇₉ O ₁₄ N ₁₅ S ₂
	516.76513	3.0	1.6	10.5	C ₄₆ H ₈₁ O ₁₄ N ₈ S ₂
	516.76513	3.1	1.6	16.0	C ₄₅ H ₇₅ O ₉ N ₁₅ S ₂
	516.76839	-3.3	-1.7	16.0	C ₄₆ H ₇₅ O ₁₂ N ₁₃ S ₁
	516.76479	3.7	1.9	20.0	C ₅₂ H ₇₅ O ₁₁ N ₉ S ₁
	516.76872	-3.9	-2.0	25.0	C ₅₄ H ₇₁ O ₁₀ N ₁₁
	516.76470	3.9	2.0	17.0	C ₄₄ H ₇₁ O ₁₄ N ₁₅
	516.76874	-3.9	-2.0	6.5	C ₄₀ H ₈₁ O ₁₅ N ₁₂ S ₂
	516.76446	4.3	2.2	11.0	C ₄₄ H ₇₉ O ₁₃ N ₁₁ S ₂
	516.76897	-4.4	-2.3	12.5	C ₄₀ H ₇₃ O ₁₆ N ₁₆
	516.76907	-4.6	-2.4	15.5	C ₄₈ H ₇₇ O ₁₃ N ₁₀ S ₁

Let's take a closer look at fragmentation



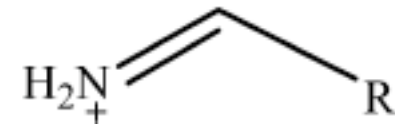
b_3 ion - oxazolone
no b_1 ion



Other amino acid fragment ions

m/z values of common immonium ions

Immonium ion (<i>m/z</i>)	Amino acid residue	Major (M) or minor (m) peak
60.04	S	M
70.07	R or P	M
72.08	V	M
73.00	R	m
74.06	T	M
84.08	K or Q	M
86.1	I or L	M
87.09	N or R	M
88.04	D	M
100.09	R	m
101.11	K or Q	M
102.06	E	M
104.05	M	M
110.07	H	M
112.09	R	M
120.08	F	M
126.06	P	M
129.1	K or Q	m
136.08	Y	M
138.07	H	m
159.09	W	M



Detecting posttranslational modifications (PTMs) by MS

- **A key issue is that the energy of ionization or the collisional process should not exceed the dissociational energy of the PTM**
- **MALDI-TOF MS with a N₂ laser causes fragmentation of a nitrated tyrosine residue**
 - Use ESI to make the molecular ion
 - Go to another laser wavelength (YAG laser at 355 nm or IR)
- **O-glucosyl and phospho groups fragment more easily than the peptide to which they are attached**
 - Use electron capture dissociation

Types of fragmentation (1)

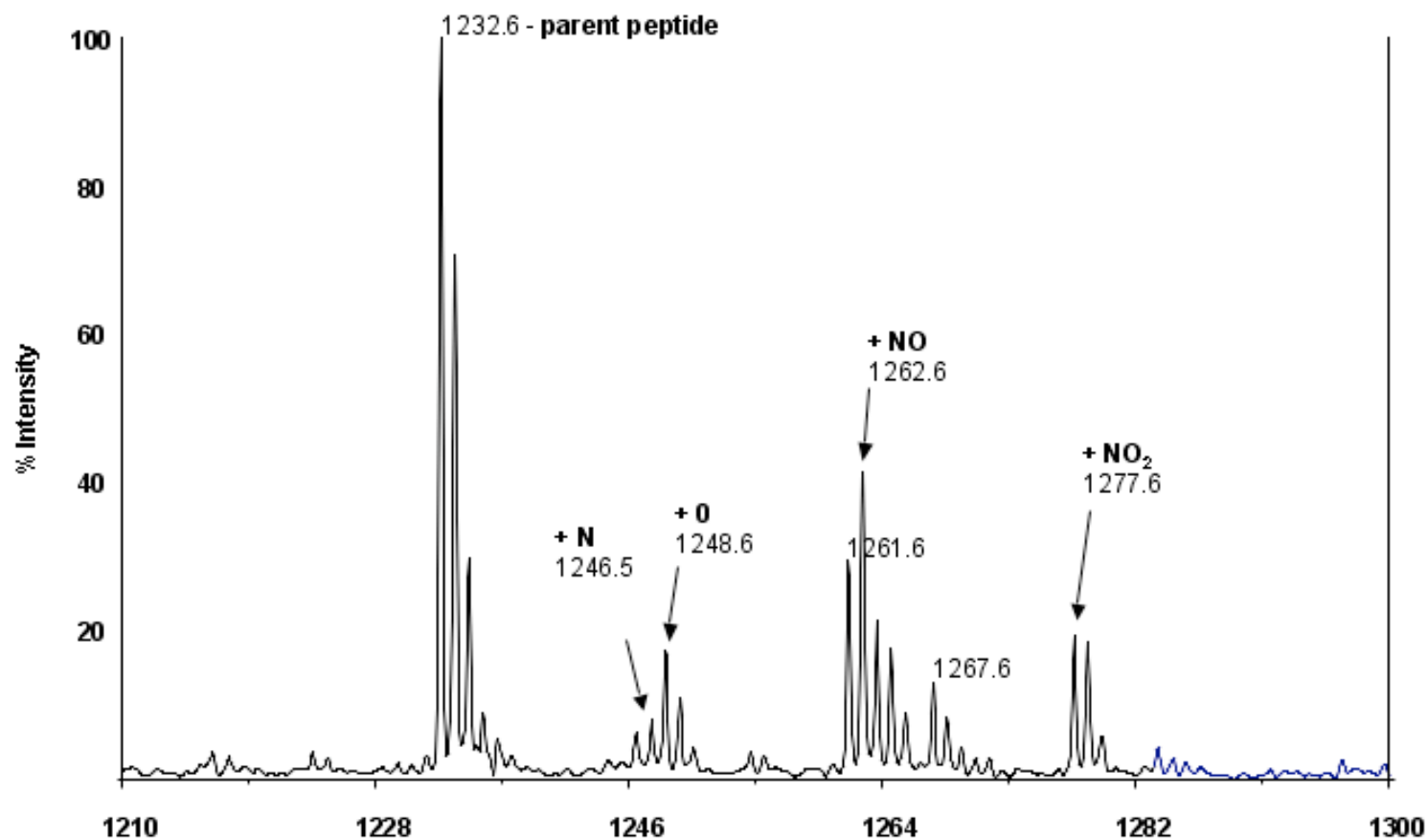
- **Collision-induced dissociation (CID)**
 - Also called CAD (collision-activated dissociation)
 - Multiply charged peptide ions are isolated by an m/z based filter
 - Selected ions are accelerated into a field of inert gas (He, N₂, Ar, Xe) at moderate pressure
 - The energy gained in collision events increases vibrational and stretching modes of the peptide backbone (and anything attached to it!)
 - The increased motion of the energized peptide causes breaks that occur typically at the peptide bond
 - Side chain groups can also be broken, some times more easily than the peptide chain

Types of fragmentation (2)

IRMPD

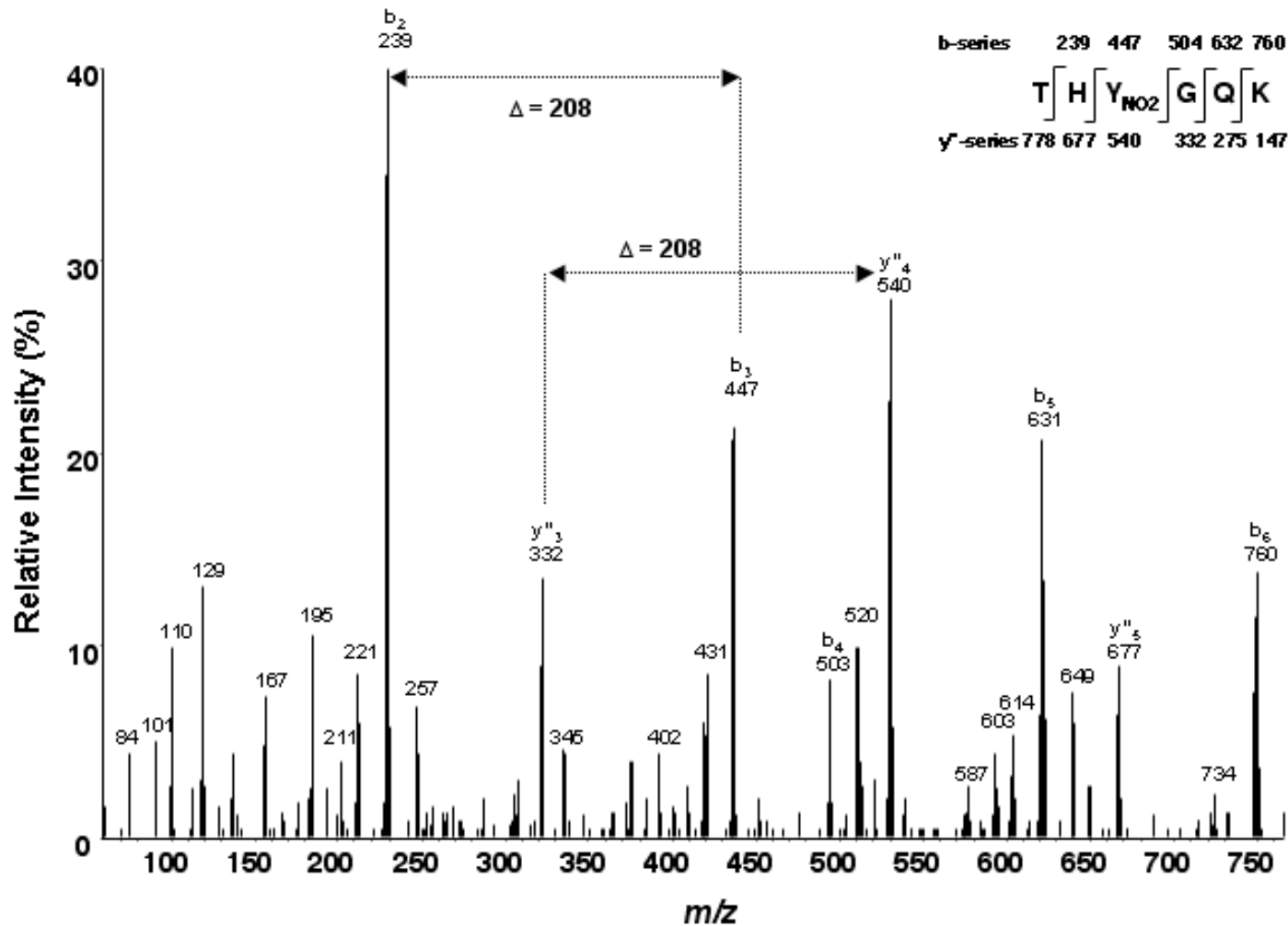
- **InfraRed Multi-Photon Dissociation**
 - Used in FT-ICR instruments where a vacuum better than 1×10^{-10} torr is necessary for the analysis of peptide ions
 - The infra-red radiation is delivered by an IR laser operating at 10.6 microns
 - No gas is involved
 - In this case, the fragmentation is induced in the ICR cell
 - Effects are essentially equivalent to CID

Fragmentation of nitrated peptides in MALDI-TOF experiment



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ESI-tandem MS of a nitrated peptide



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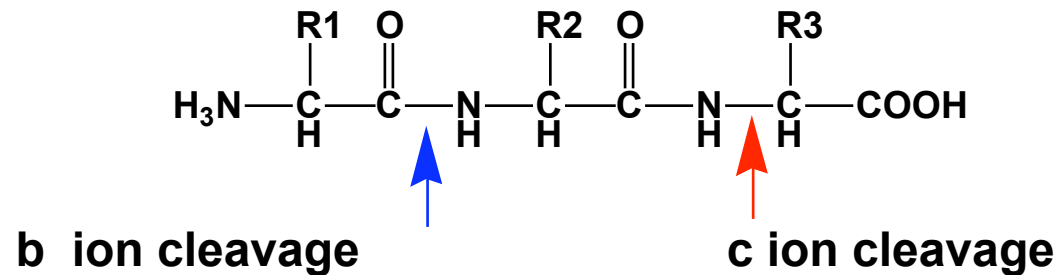
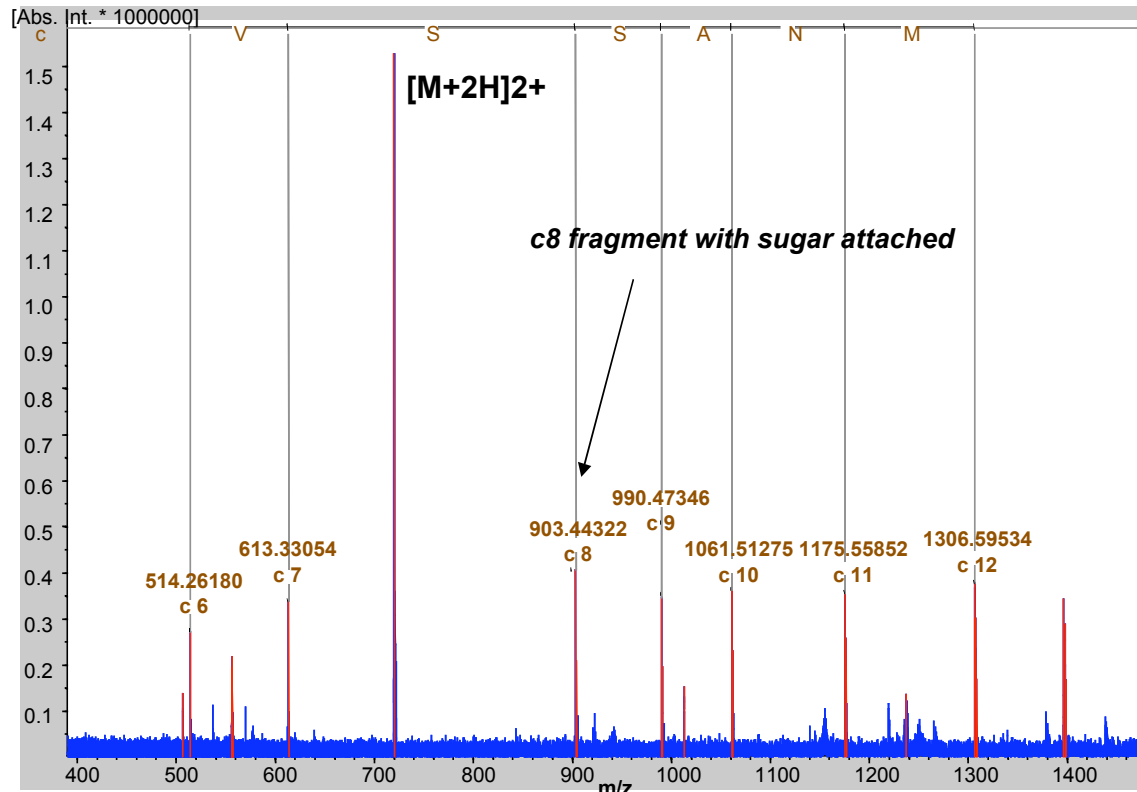
Types of fragmentation (3)

ECD

- **Electron Capture Dissociation**
 - Used in an ICR cell of an FT-MS instrument
 - Low energy electrons interact with the multiply charged peptide and are absorbed
 - They disturb bonding of the peptide backbone and cleave it without altering the side chain
 - Yields c- and z-ions
 - MS-MS spectra often very clean
 - In conjunction with an IR laser, ECD can fragment whole proteins (top-down)

Sequencing O-GlcNAc peptides by ECD FT-ICR-MS

Casein kinase II - AGGSTPVSSANMSG



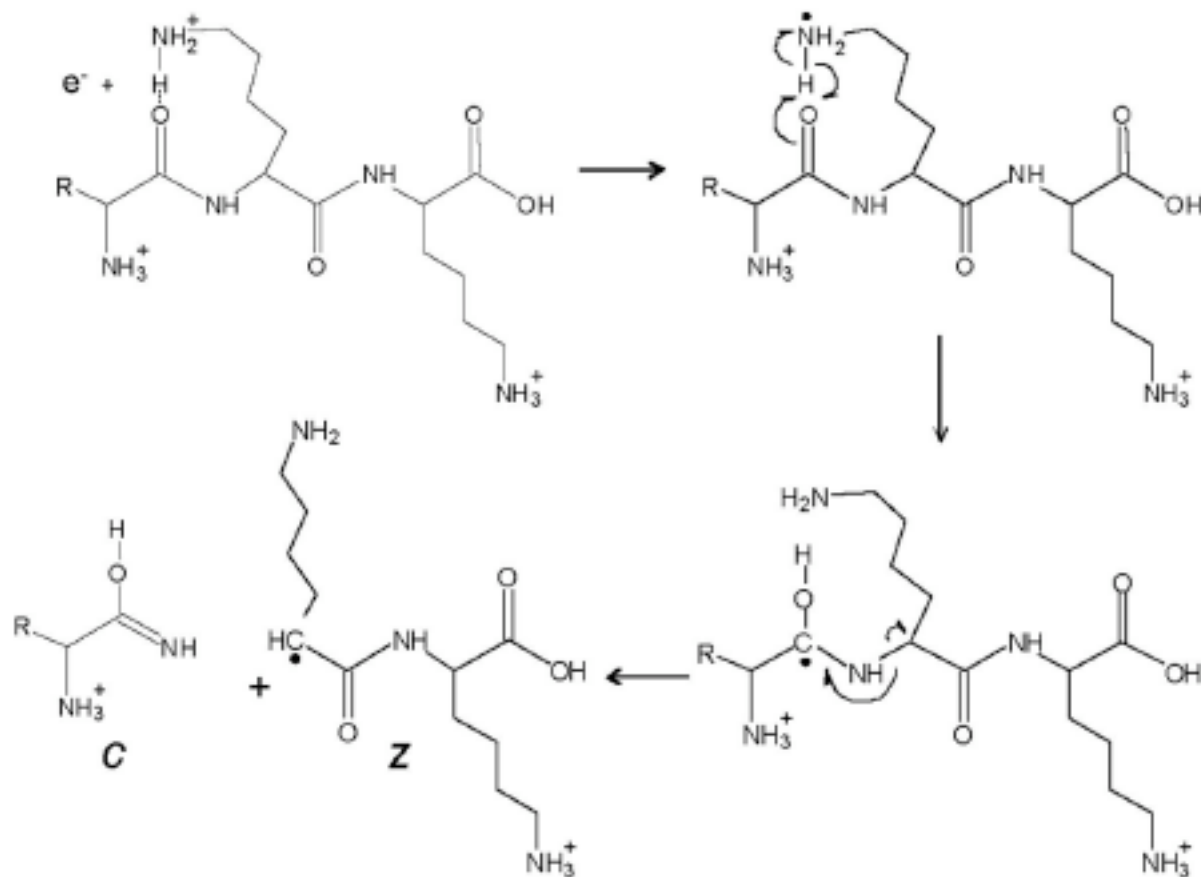
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Types of fragmentation (4)

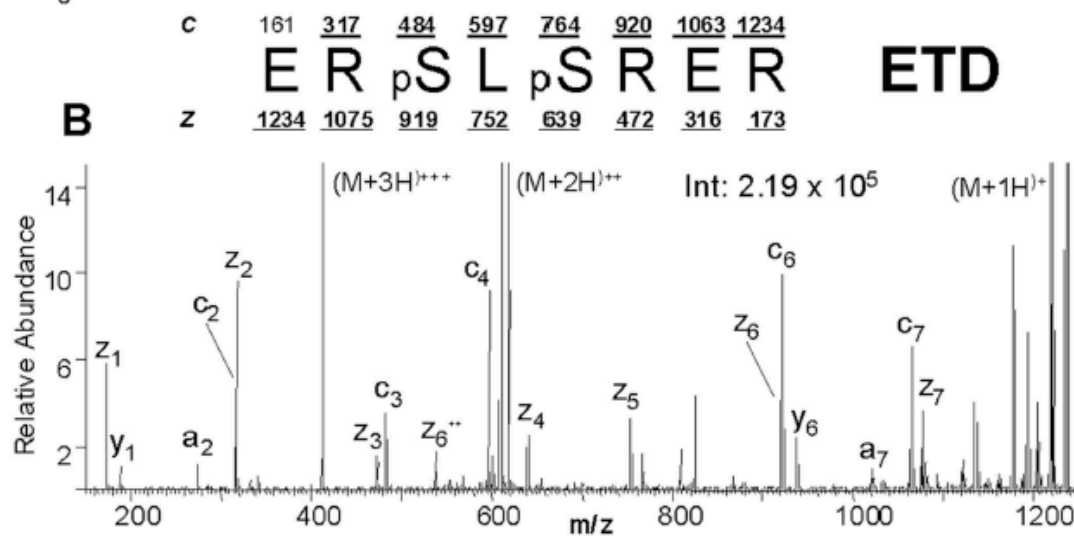
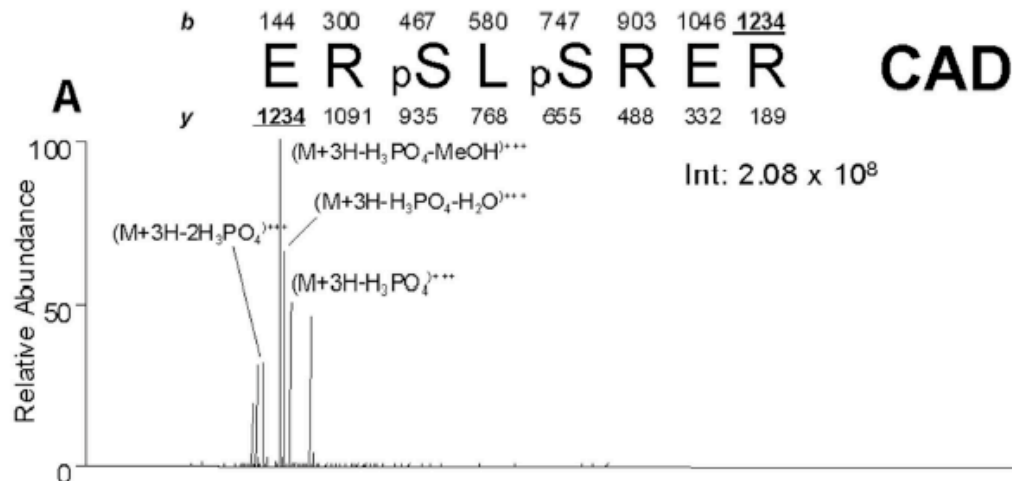
ETD

- **Electron Transfer Dissociation**
 - The electron is provided by an electron donating chemical species, a radical anion (azobenzene, fluoranthene) directly infused as a reagent gas, or from their precursors introduced by ESI - 9-anthracenecarboxylic acid, 2-fluoro-5-iodobenzoic acid, and 2-(fluoranthene-8-carbonyl)benzoic acid)

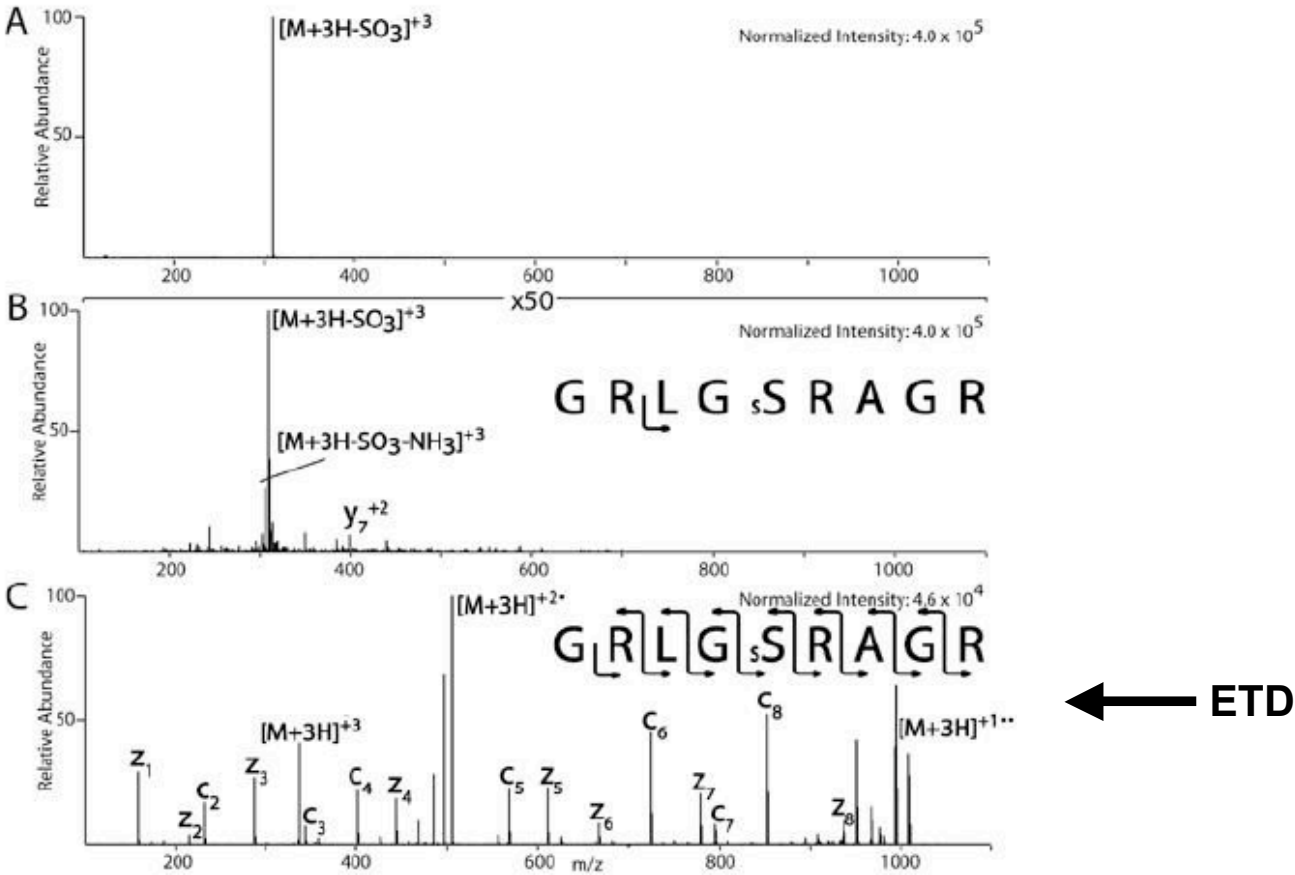
Electron transfer dissociation



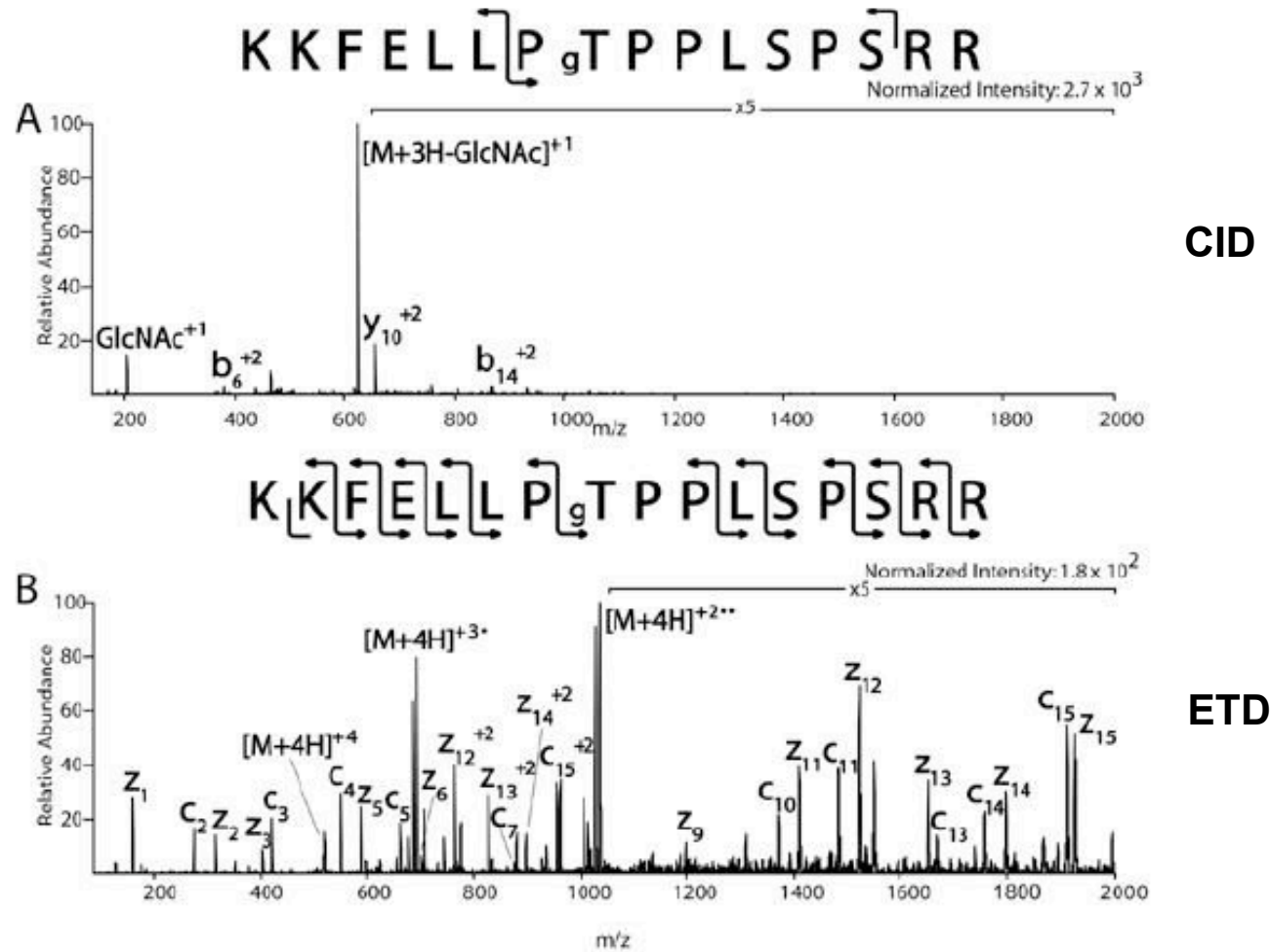
CAD versus ETD for phosphopeptide



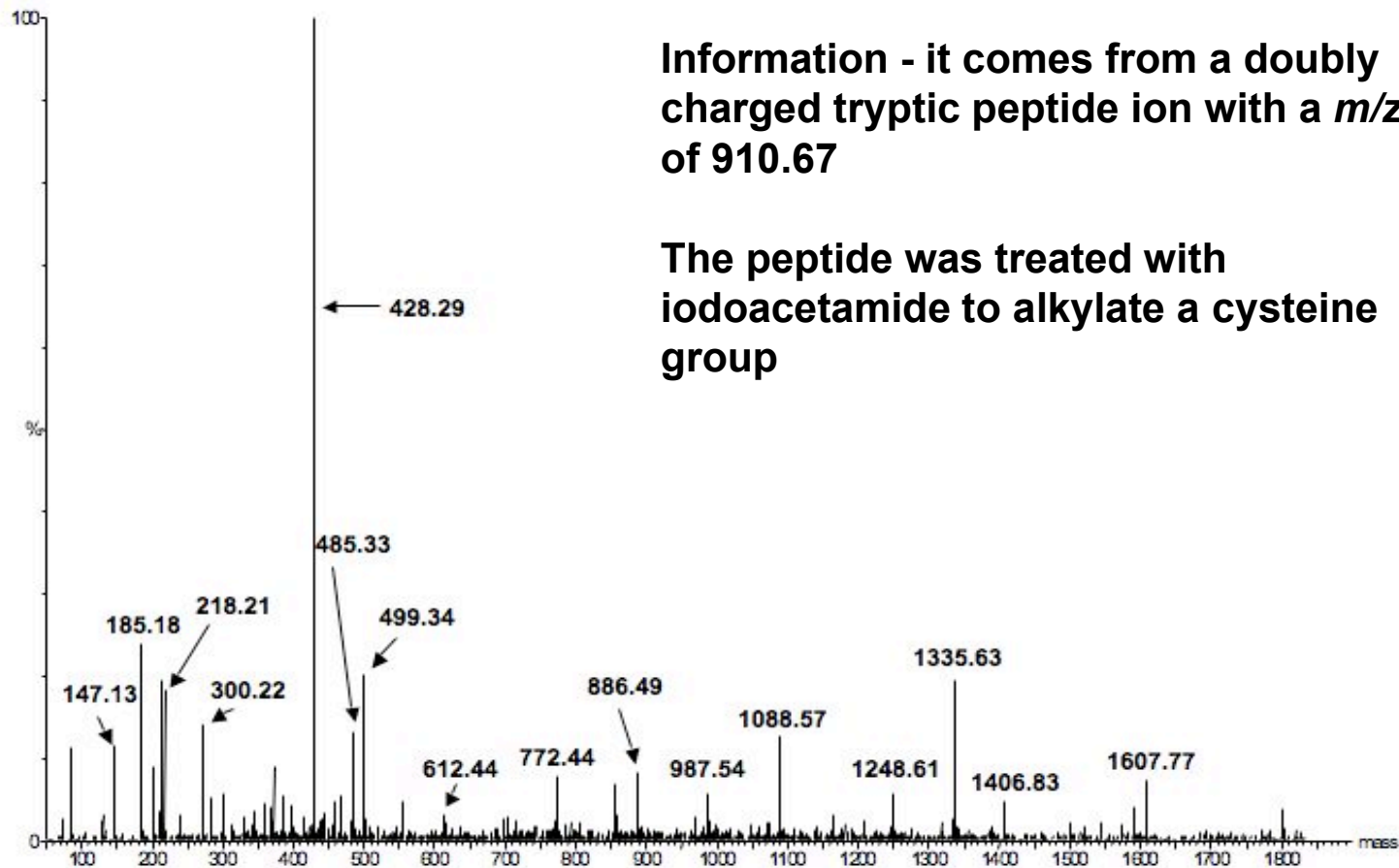
ETD better for sulfonated peptides



ETD and O-glycosylation

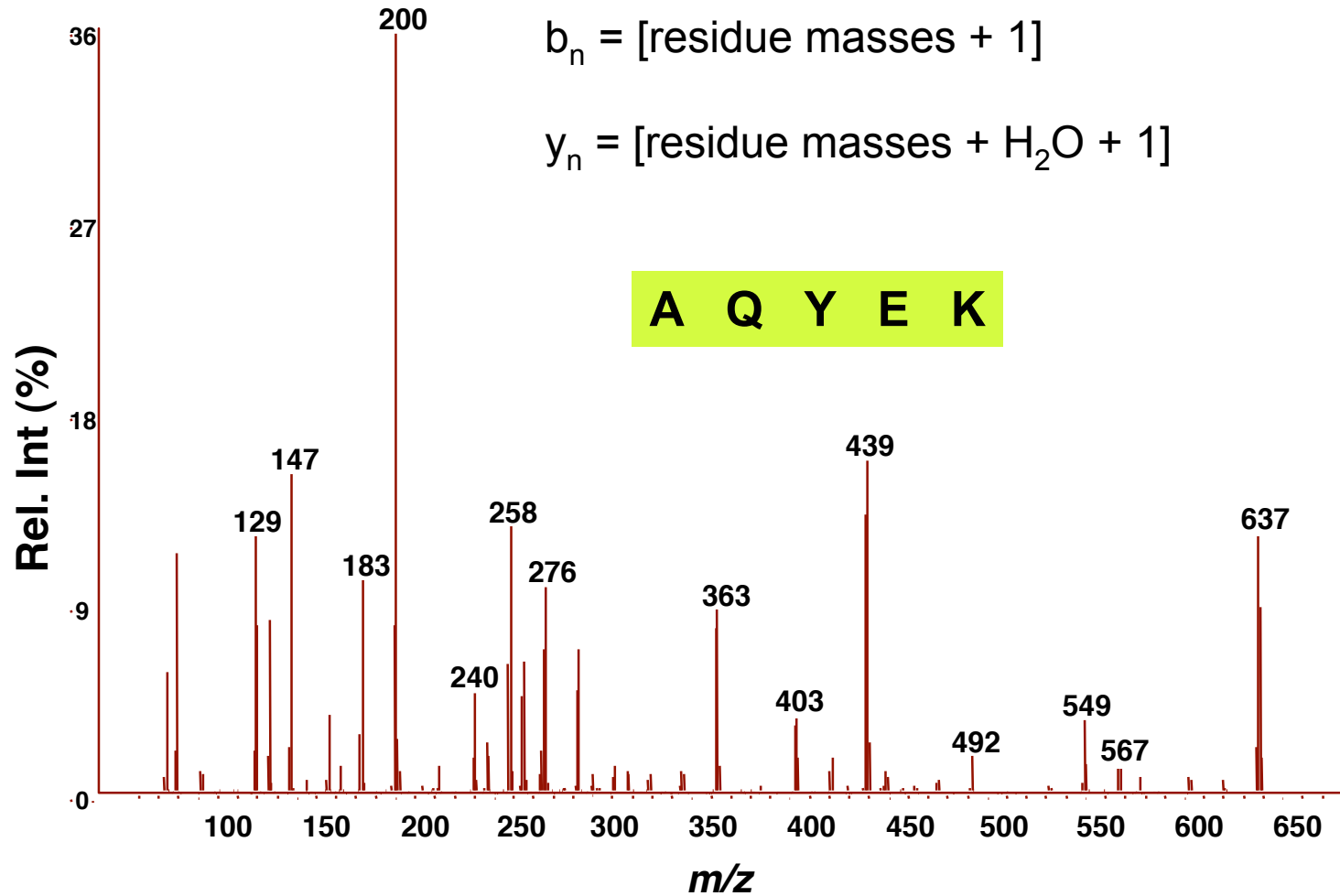


A tandem mass spectrum - what is the peptide? (Homework due Jan 29)



Fragment ions of a small 5-mer peptide

Homework - write down the masses of the b and y ions



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