Mass spectrometry of proteins, peptides and other analytes: principles and principal methods

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January 9, 2009

Objectives of the Lecture

1. Make ions
2. Separate/Analyze/Detect ions
3. What is mass resolution and mass accuracy?

What does a mass spectrometer do?

1. It measures mass better than any other technique.

2. It can give information about chemical structures.

What are mass measurements good for?

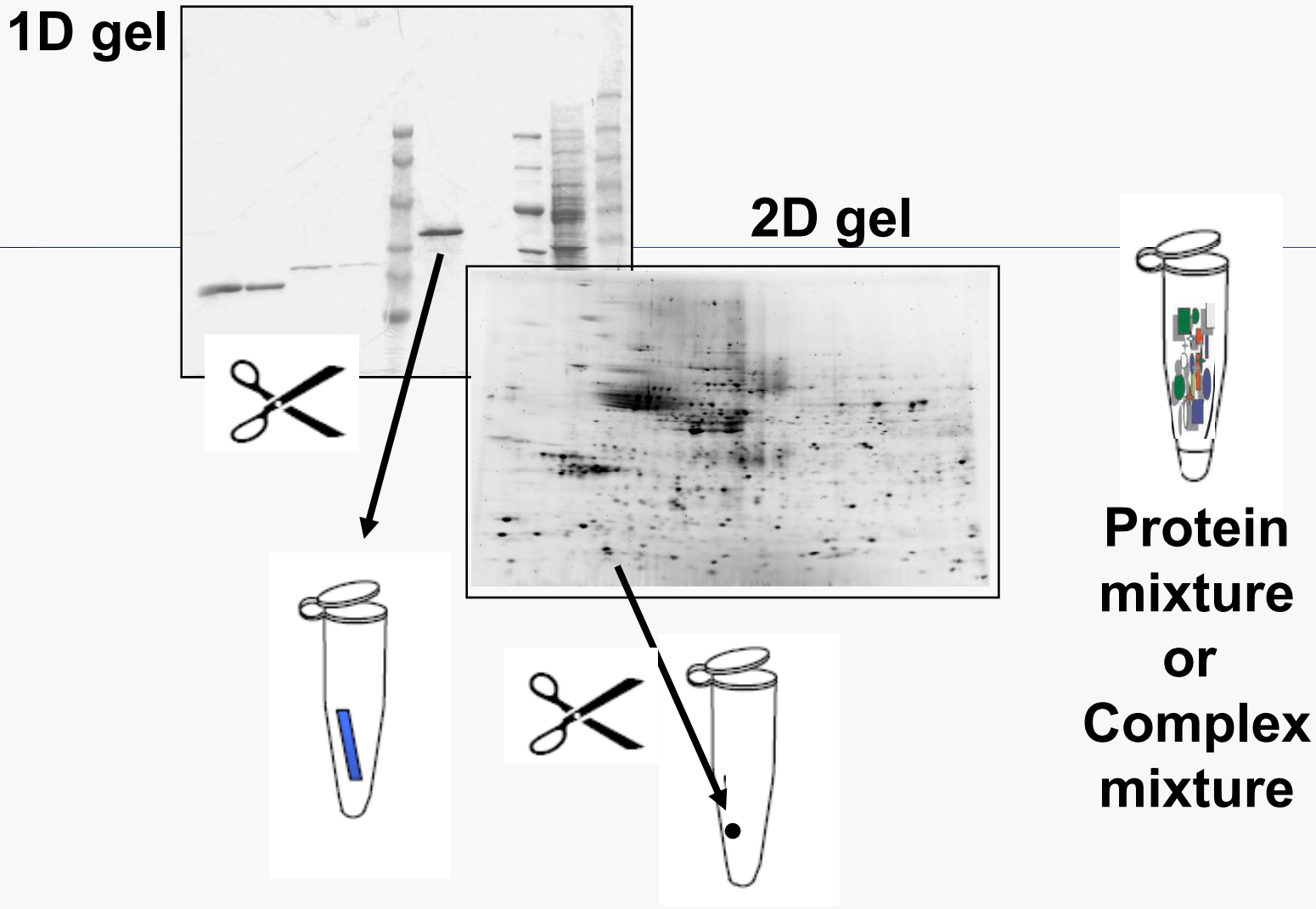
To identify, verify, and quantitate: metabolites, recombinant proteins, proteins isolated from natural sources, oligonucleotides, drug candidates, peptides, synthetic organic chemicals, polymers

Applications of Mass Spectrometry

- Pharmaceutical analysis
  - Bioavailability studies
  - Drug metabolism studies, pharmacokinetics
  - Characterization of potential drugs
  - Drug degradation product analysis

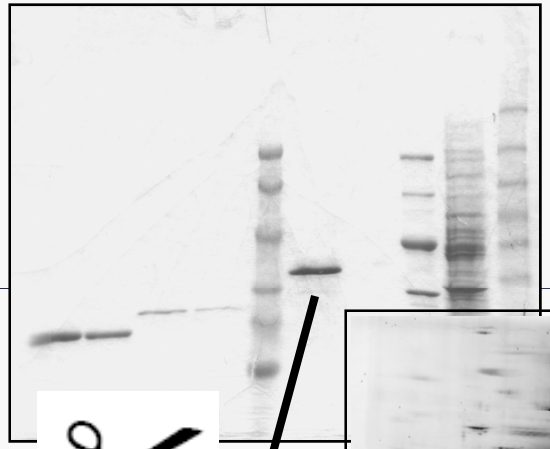
  - Screening of drug candidates
  - Identifying drug targets
- Biomolecule characterization
  - Proteins and peptides
  - Oligonucleotides
- Environmental analysis
  - Pesticides on foods
  - Soil and groundwater contamination
- Forensic analysis/clinical

MS of Proteins and Peptides

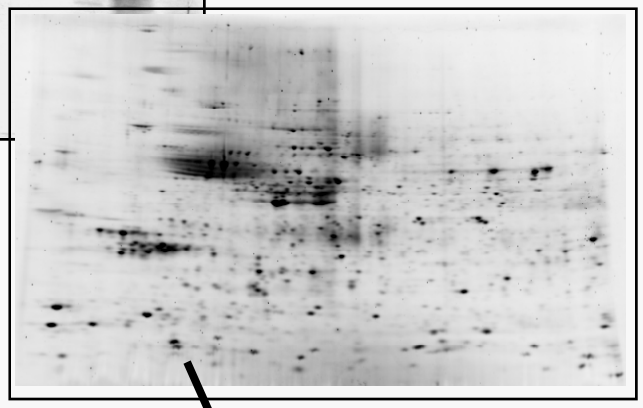


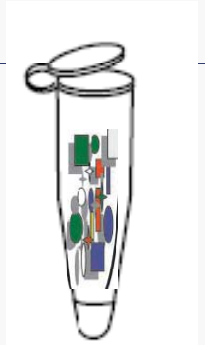
Put it in your machine and tell me the RIGHT answer

1D gel

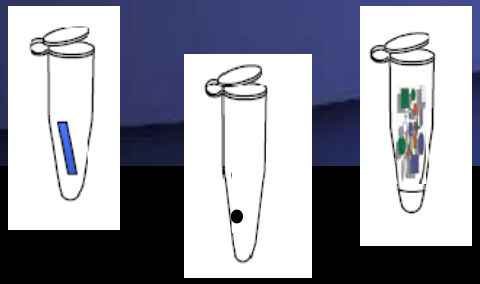


2D gel





Protein mixture or Complex mixture



Put it in

- We need Ions (+ or -)
- In the gas phase

Your machine

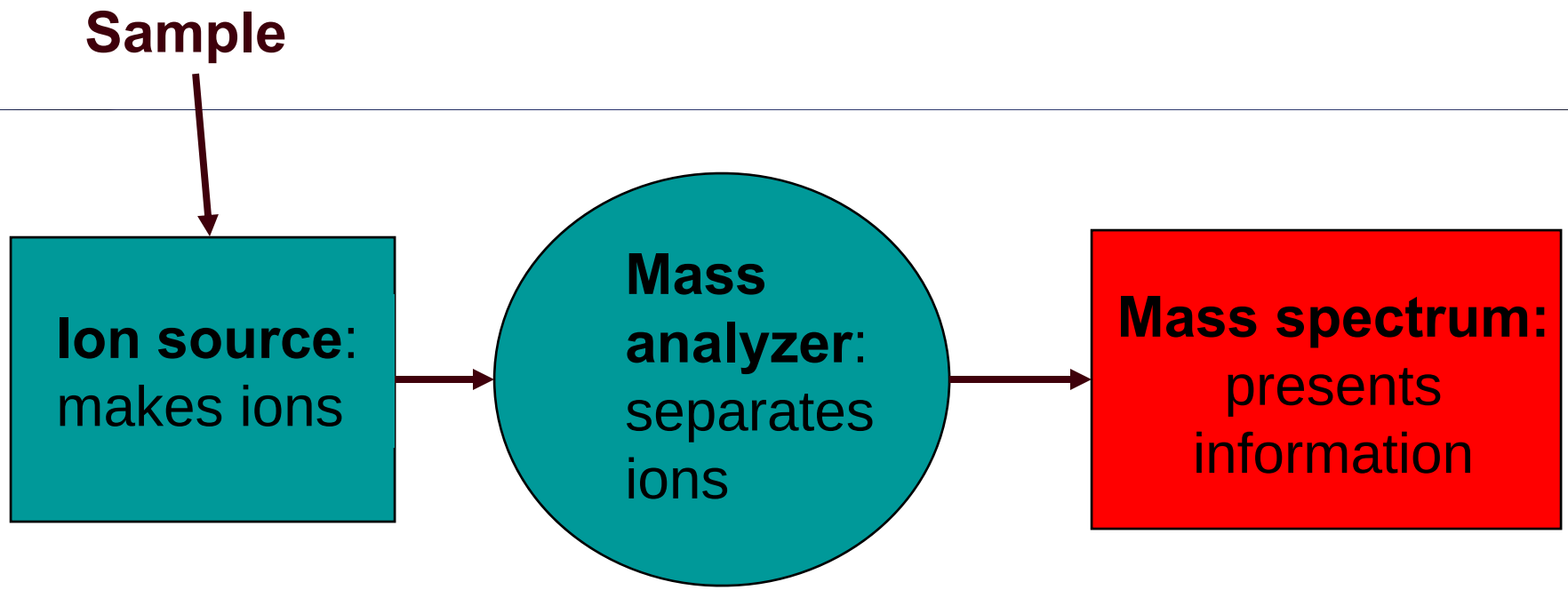
- Tof, Tof Tof
- Quadrupole, Ion trap
- FT-ICR, Orbitrap (high resolution)
- Hybrids

Tell me the <u>RIGHT</u> answer

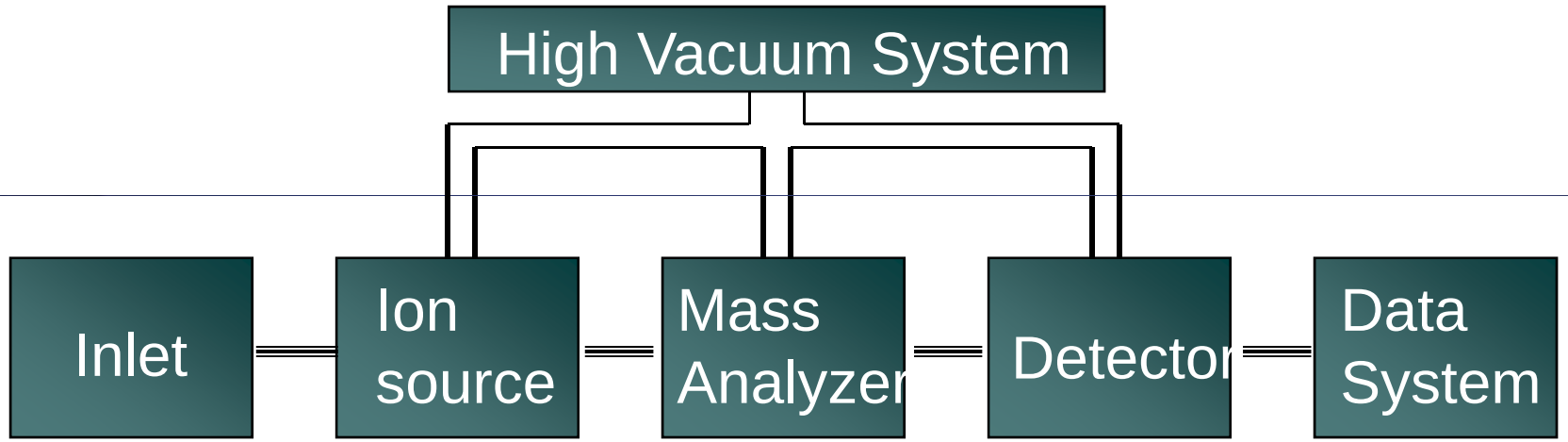
- How right is it?

  mass resolution and accuracy

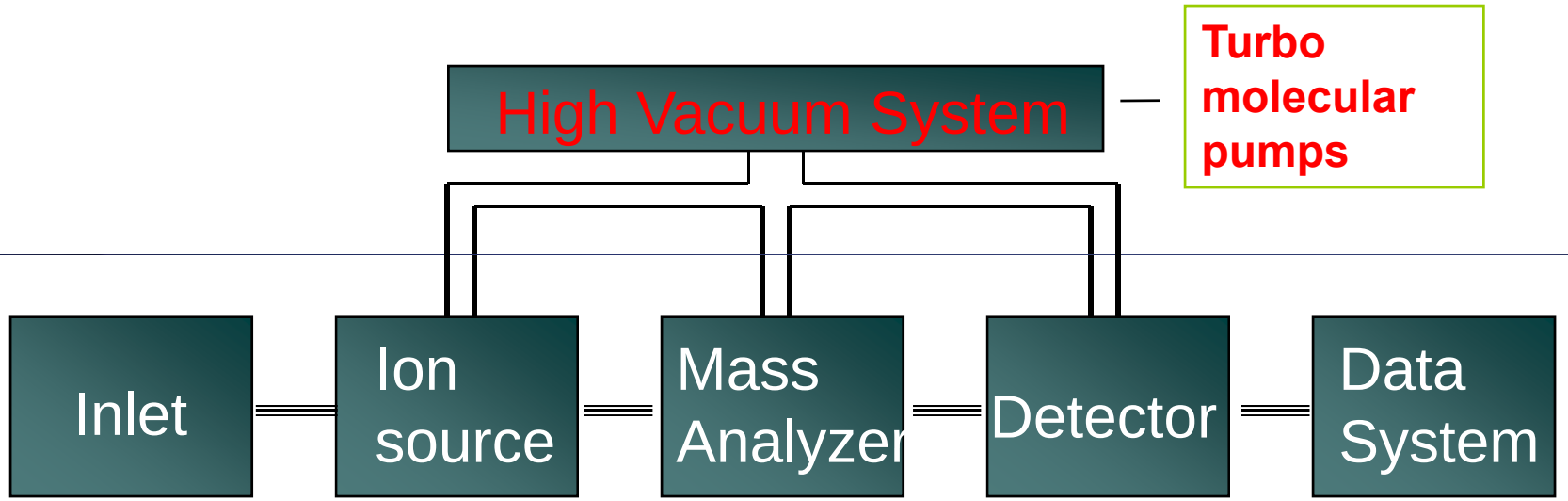
How does a mass spectrometer work?



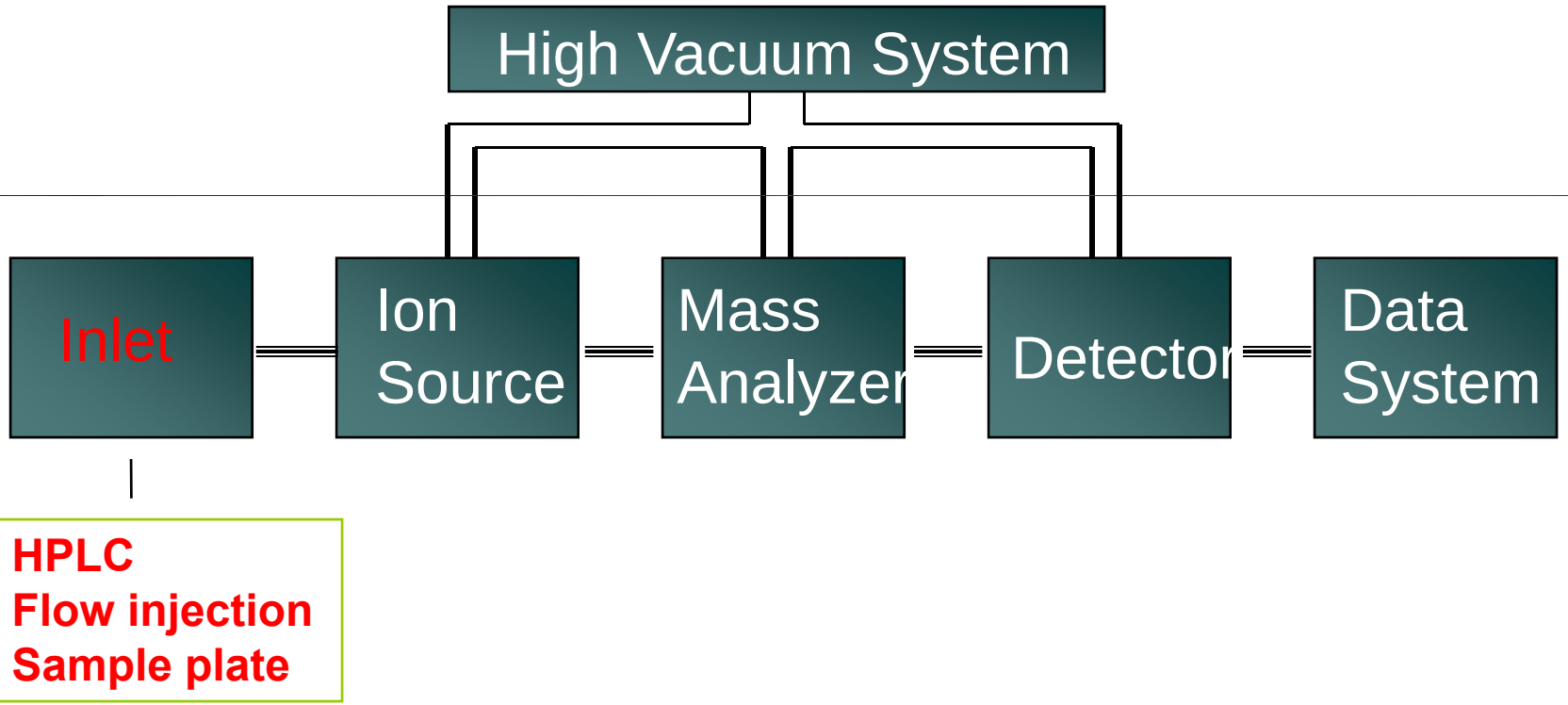
Mass Spectrometer Block Diagram



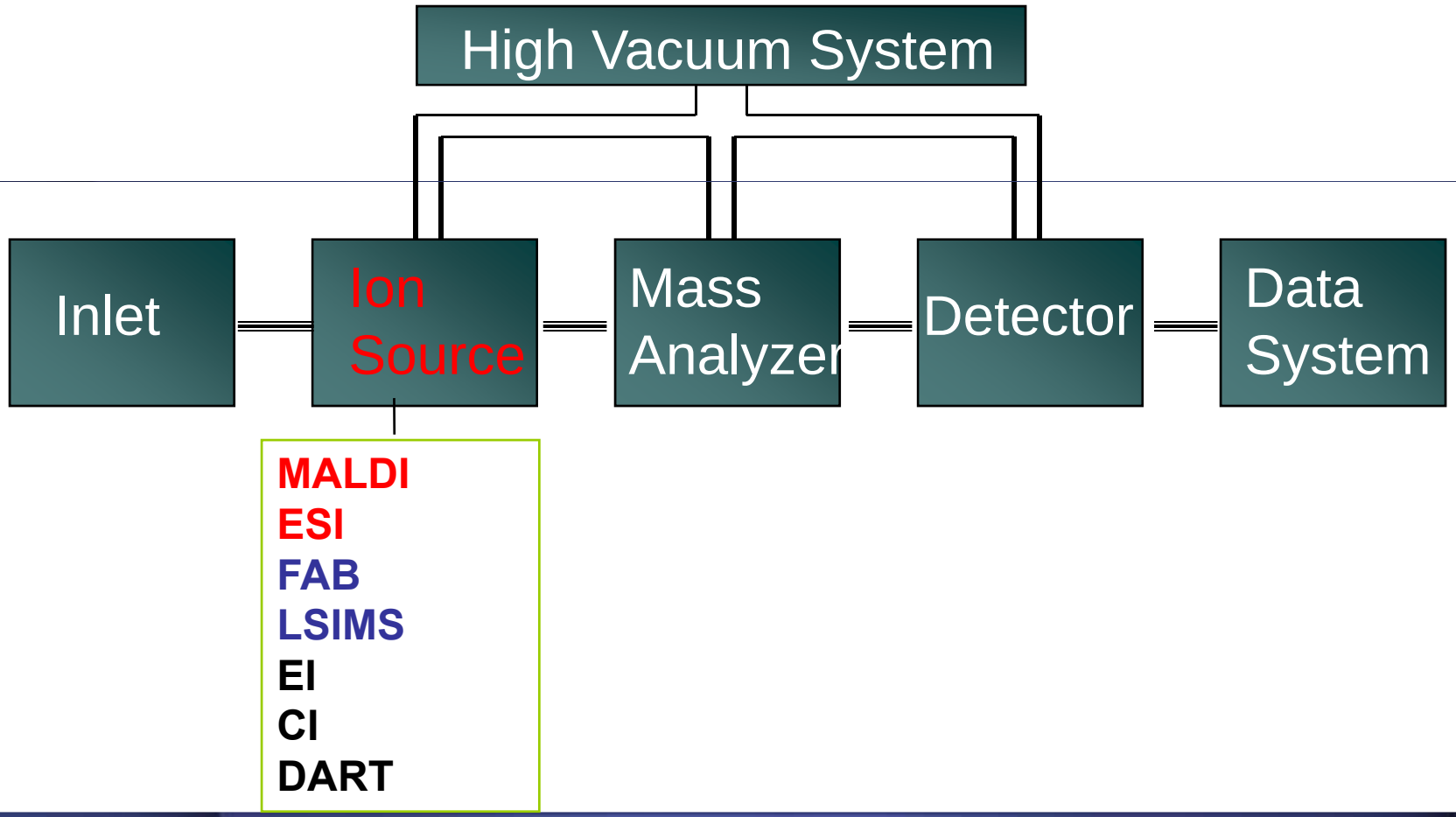
Mass Spectrometer Block Diagram



Sample Introduction

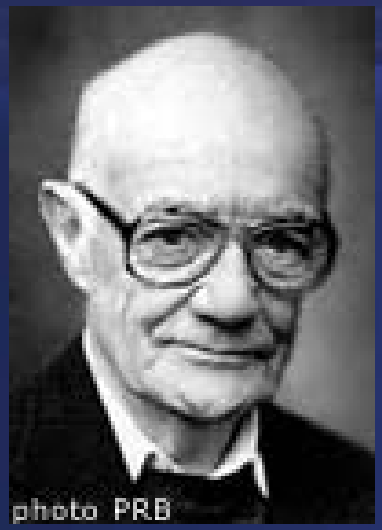


Ion Source


High Vacuum System
Inlet
Ion Source
Mass Analyzer
Detector
Data System
MALDI
ESI
FAB
LSIMS
EI
CI
DART

Nobel Prize in Chemistry- 2002

For getting proteins and peptides into the gas phase



John Fenn

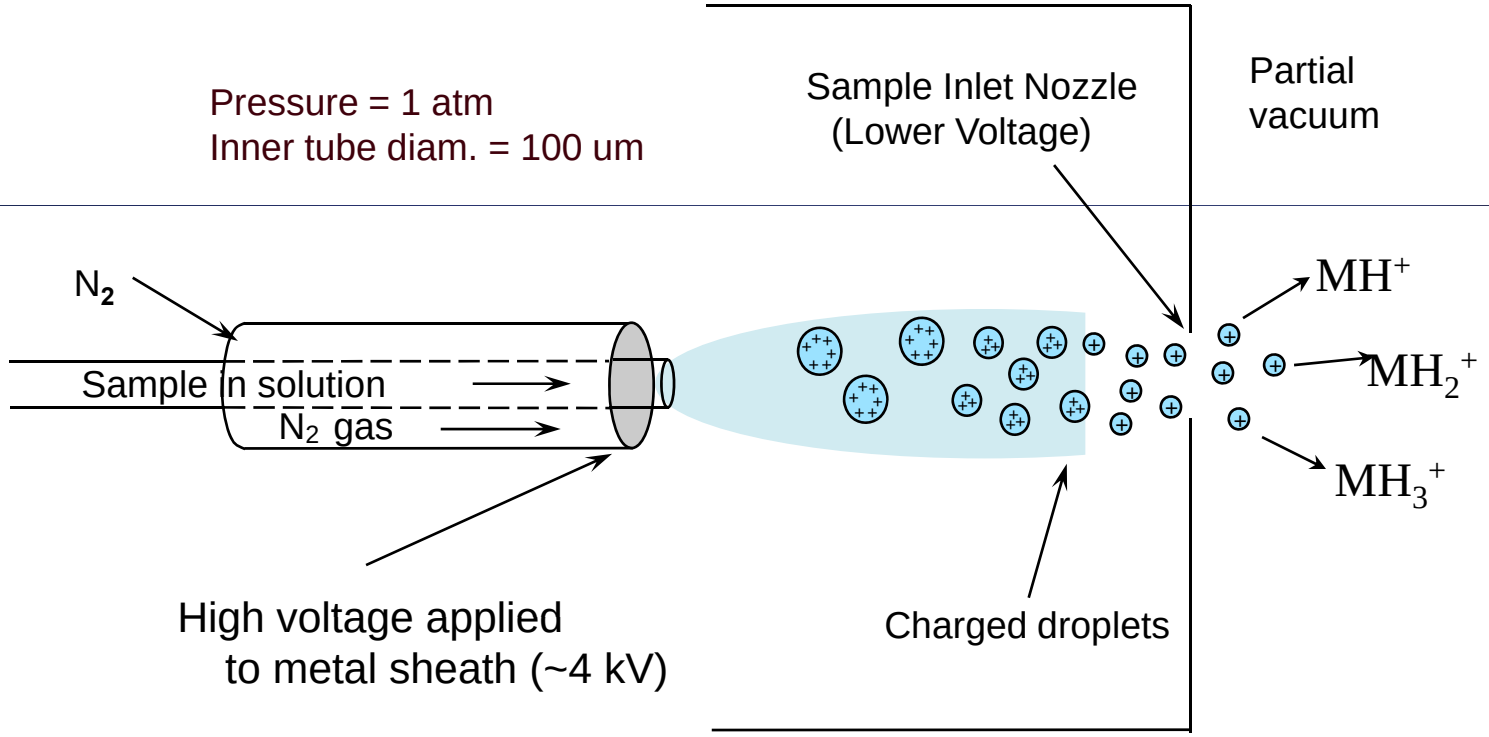
Koichi Tanaka

"*for the development of methods for identification and structure analyses of biological macromolecules*"
and
"*for their development of soft desorption ionisation methods for mass spectrometric analyses of biological macromolecules*"

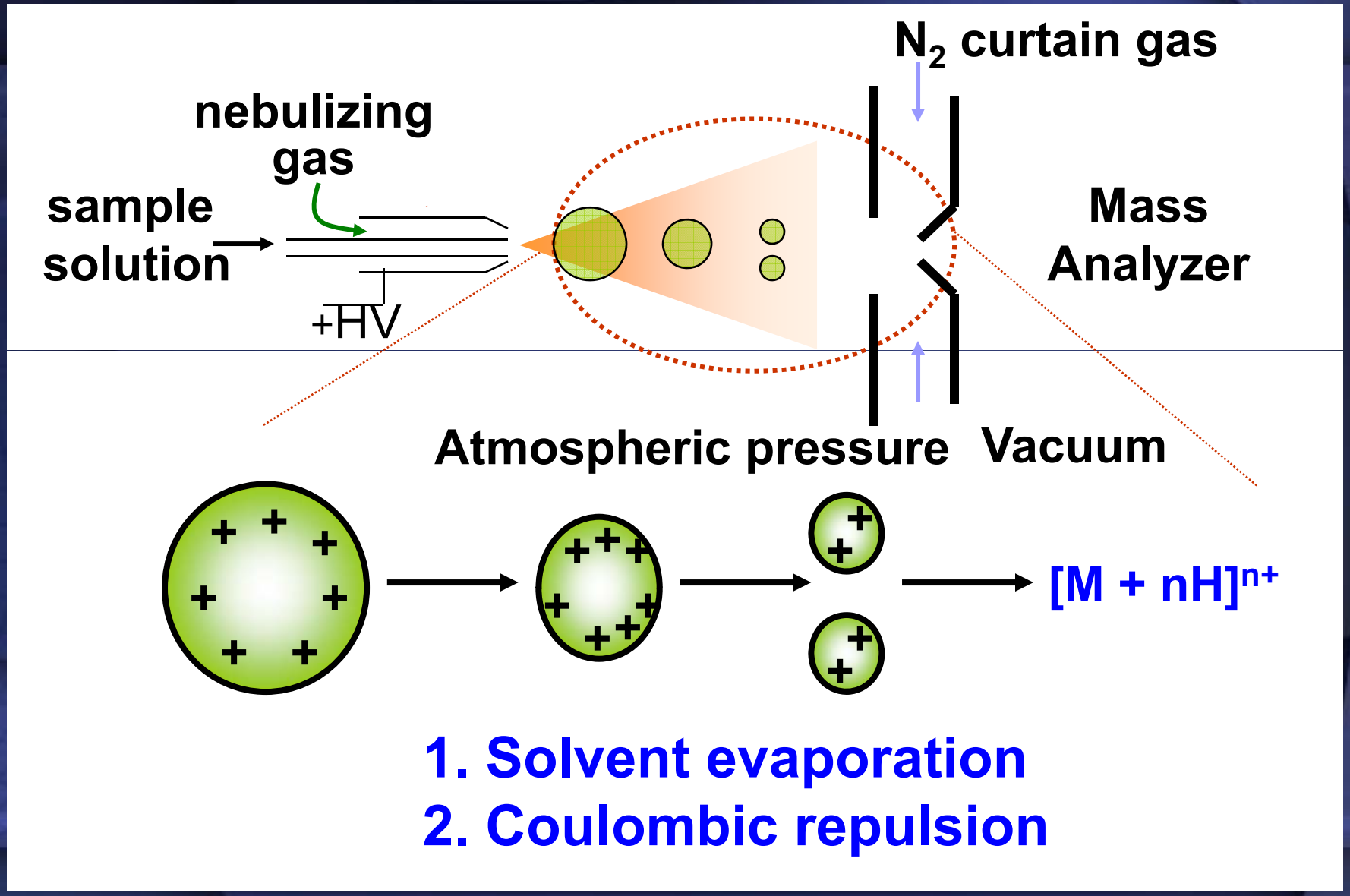
Ion Sources make ions from sample molecules

(Ions are easier to detect than neutral molecules.)

Electrospray ionization:

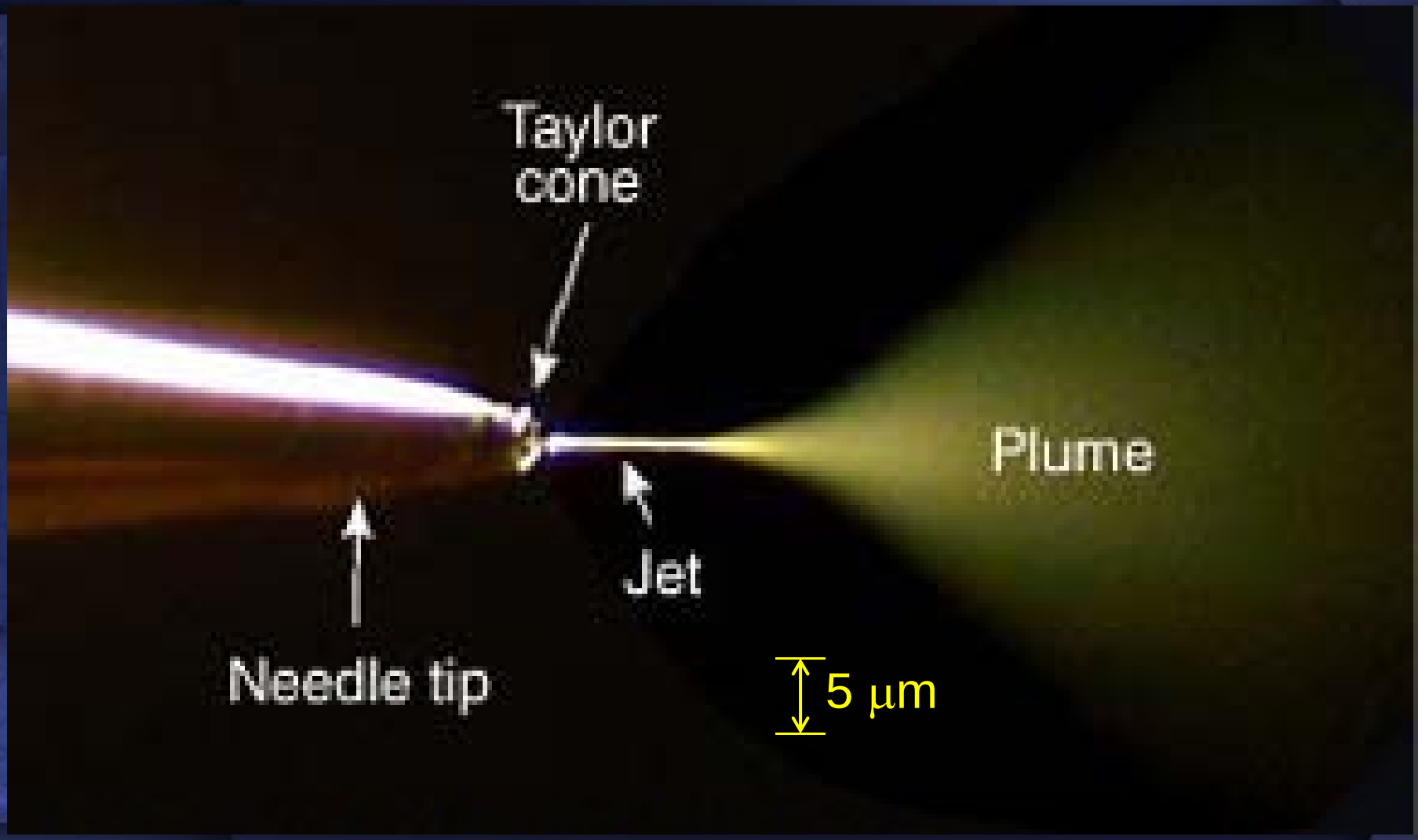


Electrospray Ionization (ESI)

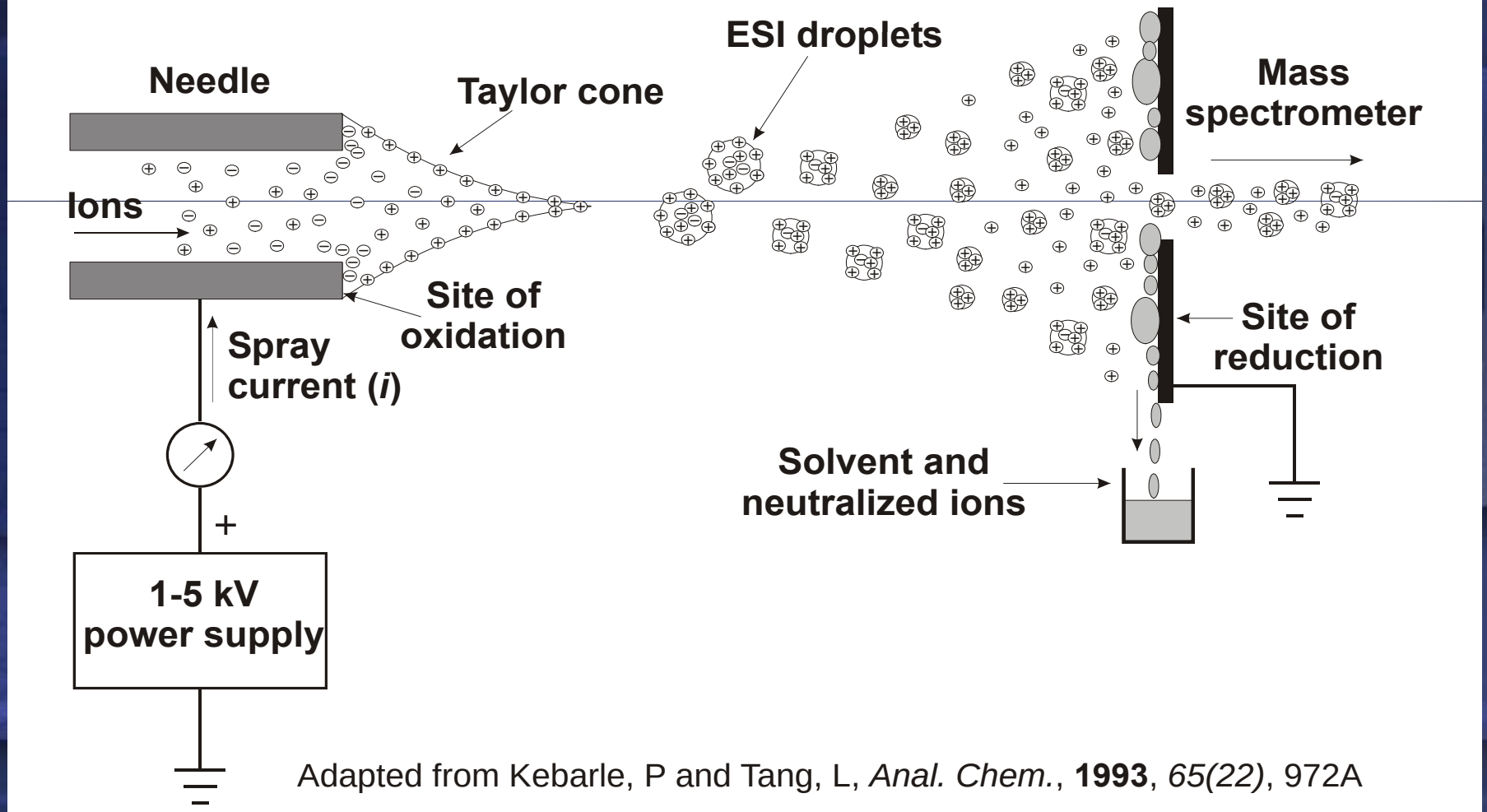


1. Solvent evaporation
2. Coulombic repulsion

Electrospray Ionization (ESI)

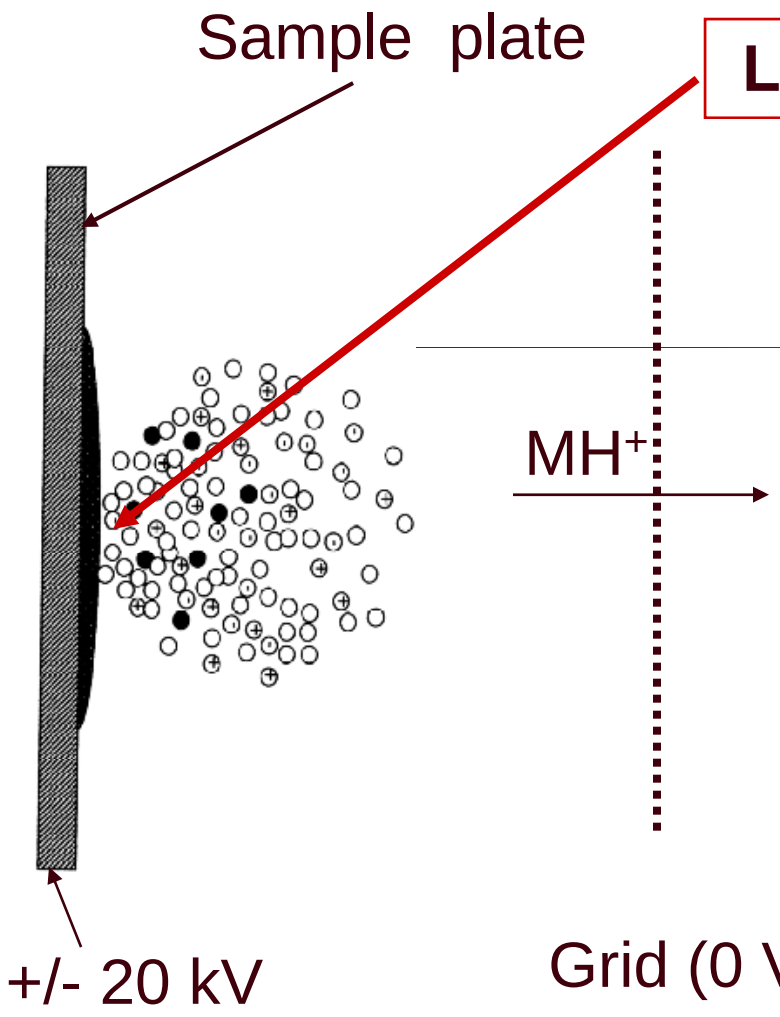


Electrospray Ionization (ESI) Process
(Positive Mode)



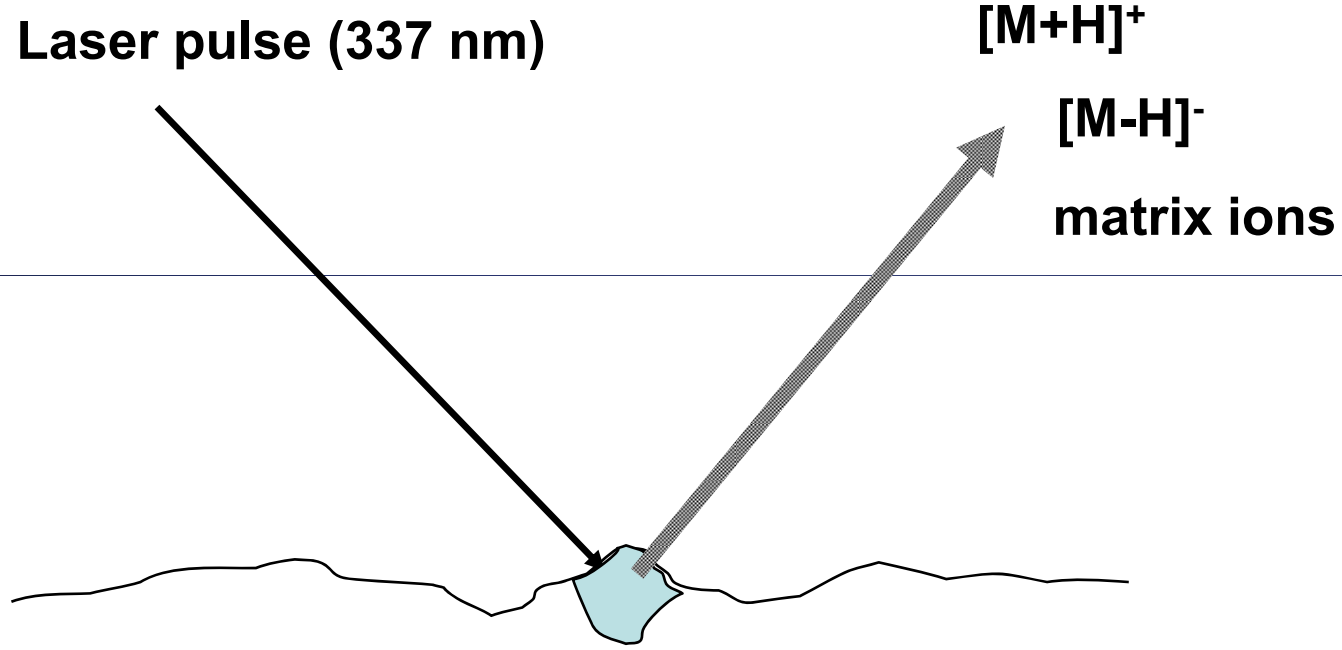
Adapted from Kebarle, P and Tang, L, *Anal. Chem.*, **1993**, *65(22)*, 972A

MALDI: Matrix Assisted Laser Desorption Ionization



1. Sample is mixed with matrix (X) and dried on plate.
2. Laser flash ionizes matrix molecules.
3. Sample molecules (M) are ionized by proton transfer: $XH^+ + M \rightarrow MH^+ + X$.

MALDI generation of ions
(Matrix-assisted laser desorption ionization)



Peptide/protein deposited on crystal surface

Sample mixed with a UV-absorbing matrix and is allowed to co-crystallize on the metal target.

Matrices for MALDI analysis

Peptides/proteins

- 3,5-dimethoxy-4-hydroxycinnamic acid (sinapinic acid)
- α-cyano-4-hydroxycinnamic acid (CHCA)
- 2,5-dihydroxybenzoic acid (DHB)
- 2-(4-hydroxyphenylazo)-benzoic acid (HABA)

Oligonucleotides

- 2-aminobenzoic acid
- 3-hydroxypicolinic acid (3-HPA)
- 2,4,6-trihydroxyacetophenone (THAP)

The choice of matrix depends greatly on the solute to be analyzed.

Put it in

- We need Ions (+ or -)
- In the gas phase

Your machine

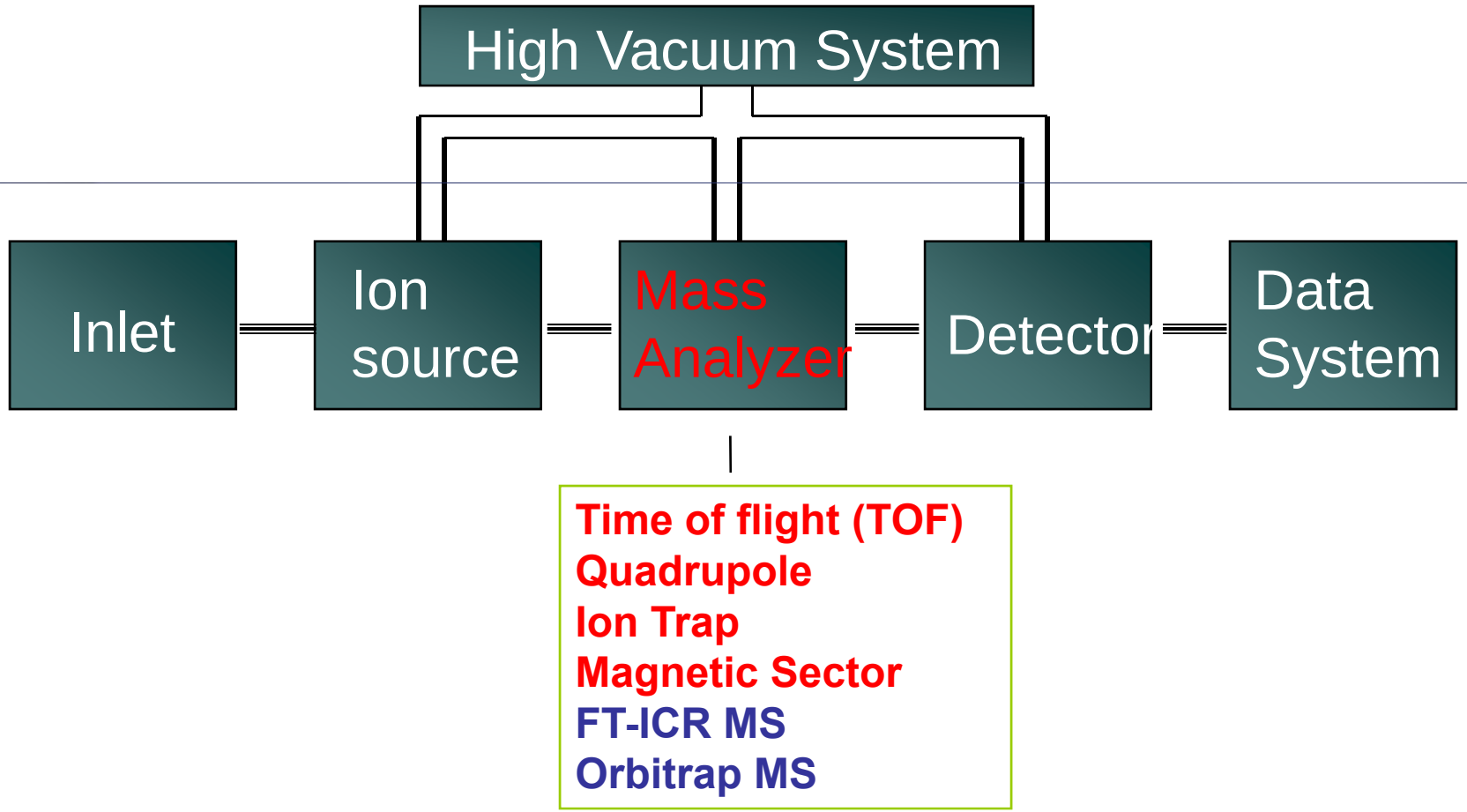
- Tof, Tof / Tof
- Quadrupole, Ion trap
- FT-ICR, Orbitrap (high resolution)
- Hybrids

Tell me the RIGHT answer

- How right is it?

  mass resolution and accuracy

Mass Analyzer



Mass analyzers separate ions based on their mass-to-charge ratio (m/z)

- Operate under high vacuum
(keeps ions from bumping into gas molecules)
- Actually measure mass-to-charge ratio of ions (m/z)

The importance of the mass-to-charge ratio is that according to classical electrodynamics two particles with the same mass-to-charge ratio move in the same path in a vacuum when subjected to the same electric and magnetic fields.

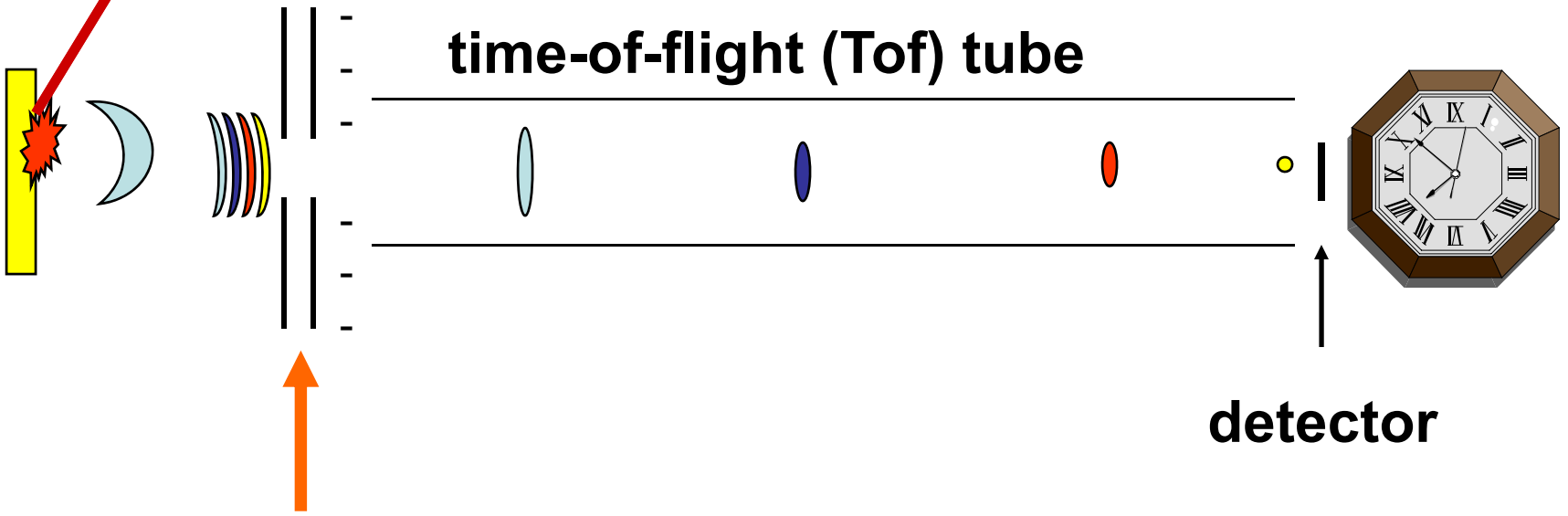
- $\mathbf{F}=m\mathrm{a}$ (Newton's second law of motion)
- $\mathbf{F}=q(\mathbf{E} + \mathrm{v} \times \mathrm{B})$ (Lorentz force Law)
- $(m/q)\mathrm{a} = \mathbf{E} + \mathrm{v} \times \mathbf{B}$

- Key specifications are resolution, mass measurement accuracy, and sensitivity.

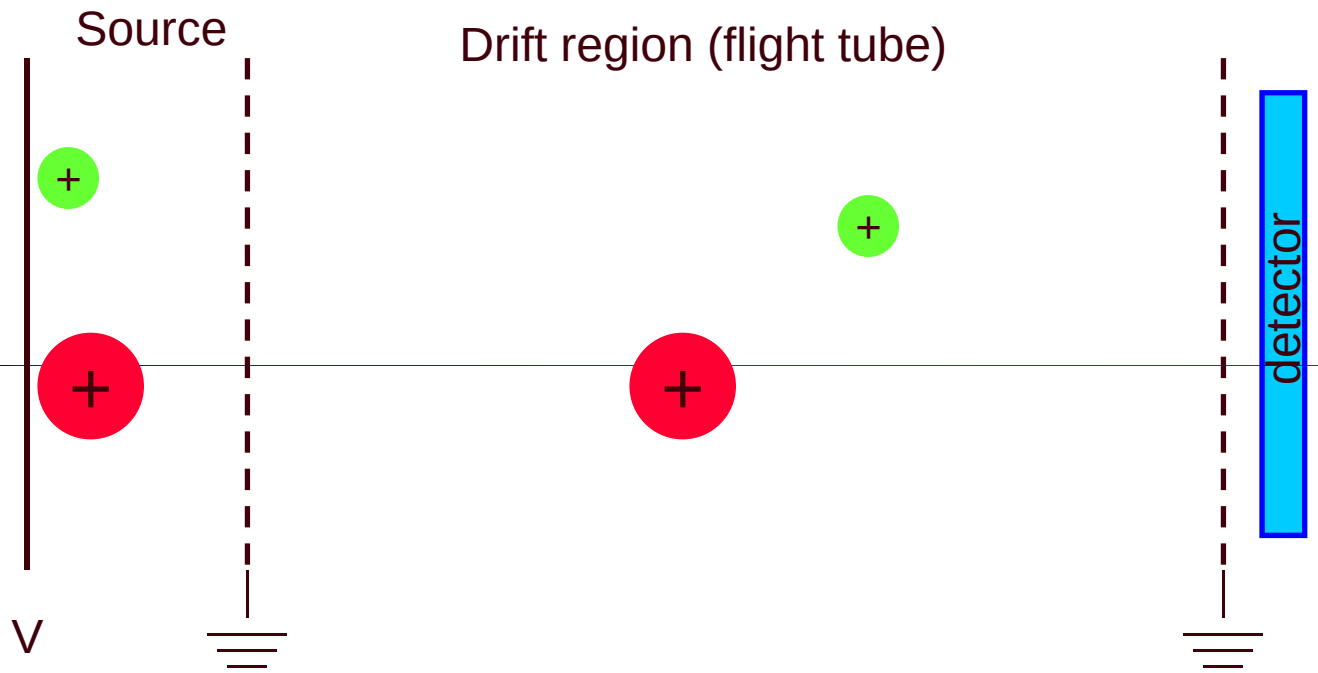
Time-of-flight (Tof) analyzer

MALDI

Laser; High Energy
Monochromatic Light Source

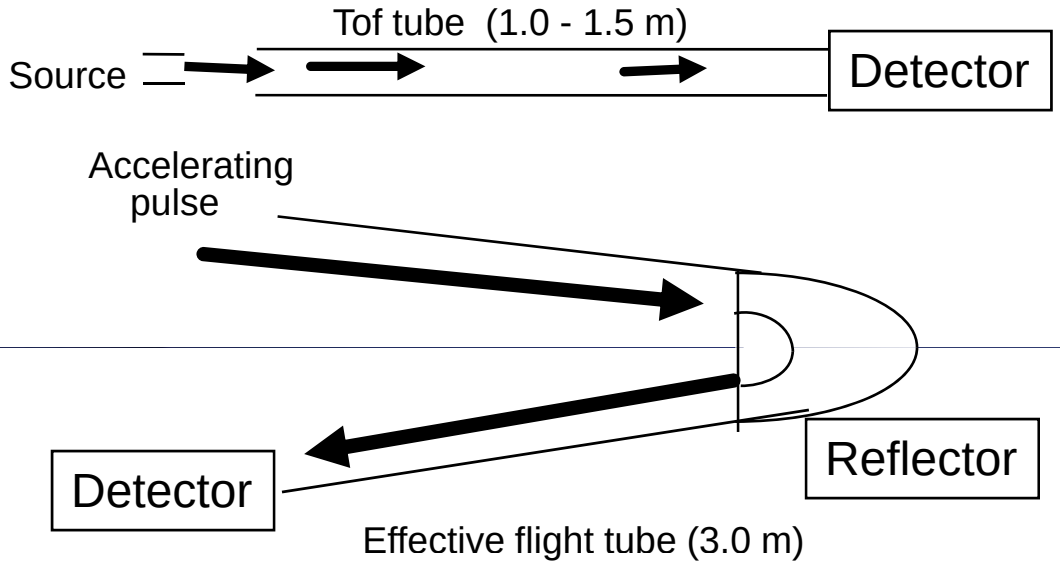


Time-of-flight (TOF) Mass Analyzer



- Ions are formed in pulses.
- The drift region is field free.
- Measures the time for ions to reach the detector.
- Small ions reach the detector before large ones.

Time-of-flight (Tof) analyzer



Resolution 2 x 10^4

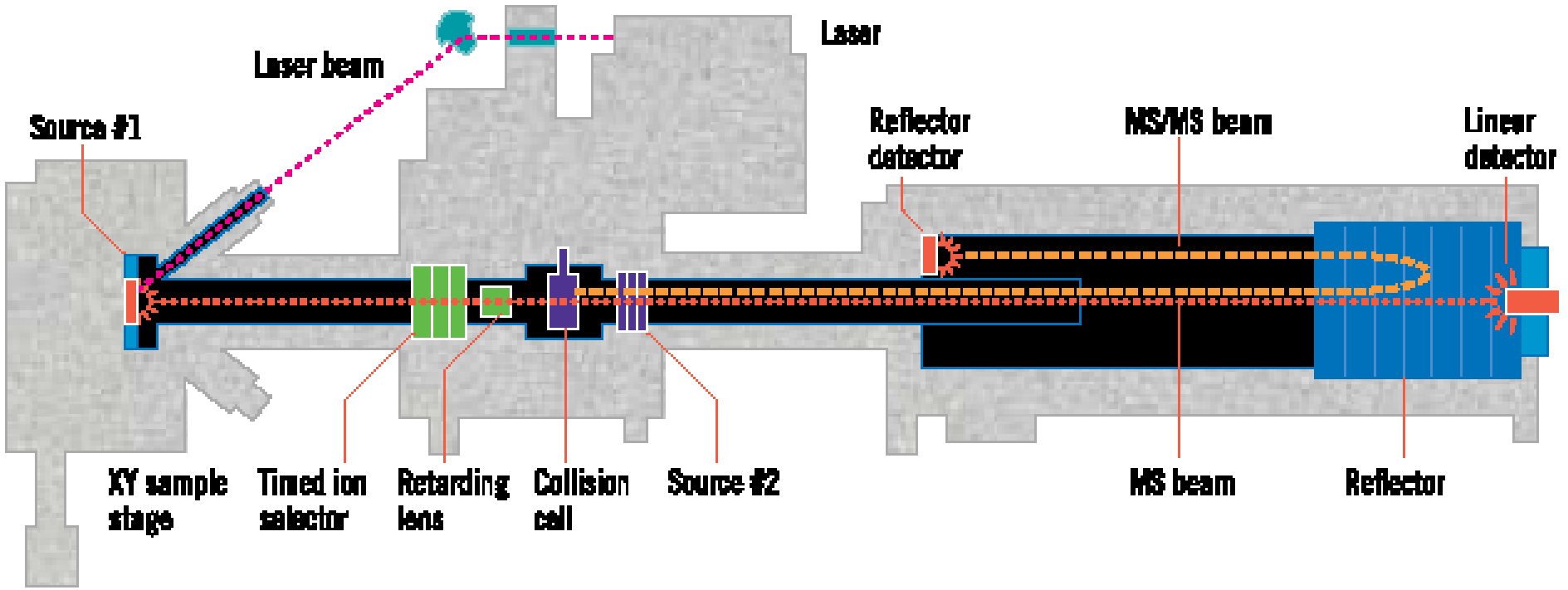
No upper limit of mass

Scan times ~ 1 μsec, good for LC-MSMS

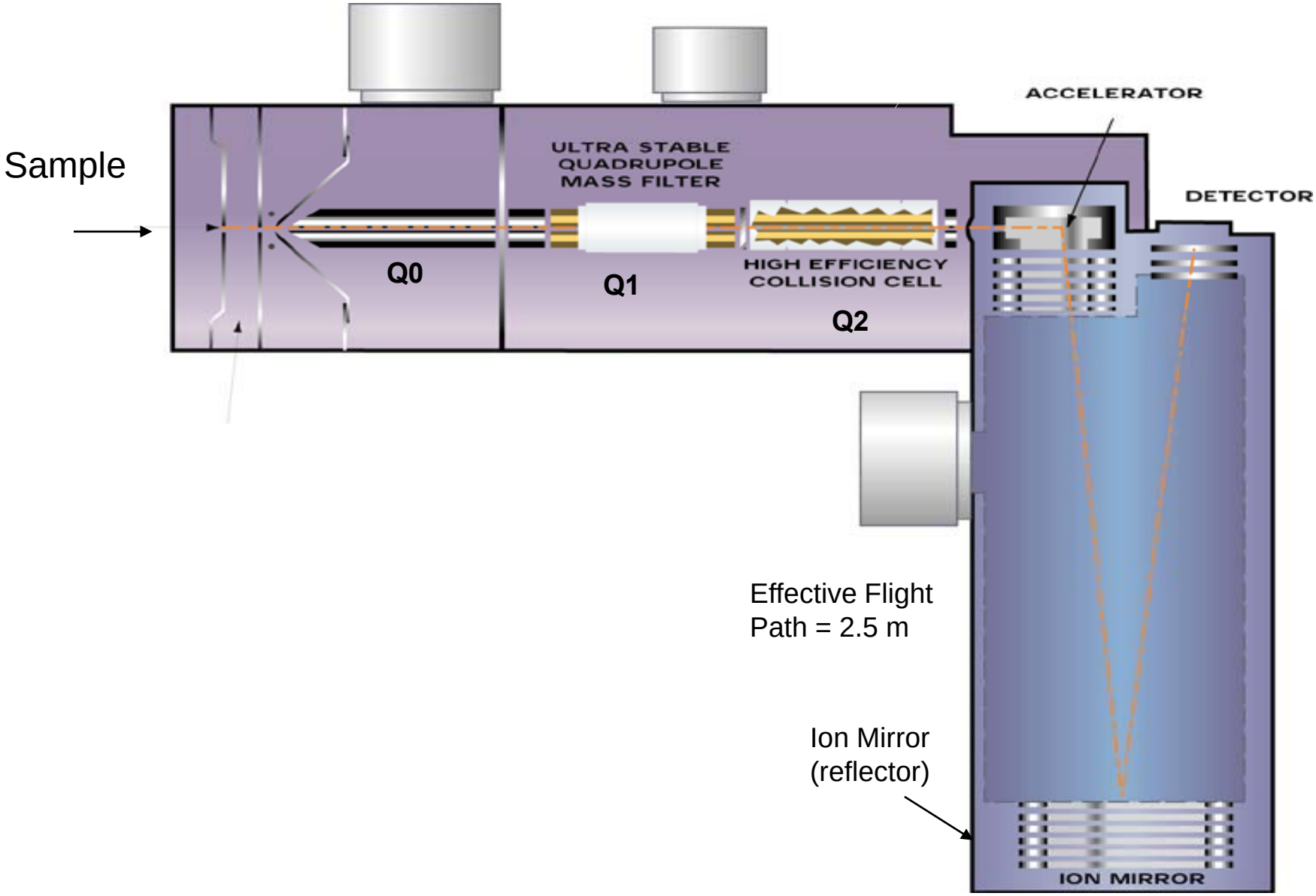
Ions are accelerated so that they have equal kinetic energy. The ions "drift" down a 1 - 1.5 meter tube before striking a photomultiplier detector. "time of flight" (t) depends on the mass of the ion (m), where $t = (m/2eV)^{1/2}*D$

V is the applied potential and D is the flight tube distance. For a given instrument, the flight time varies as the square root of the mass of the ion.

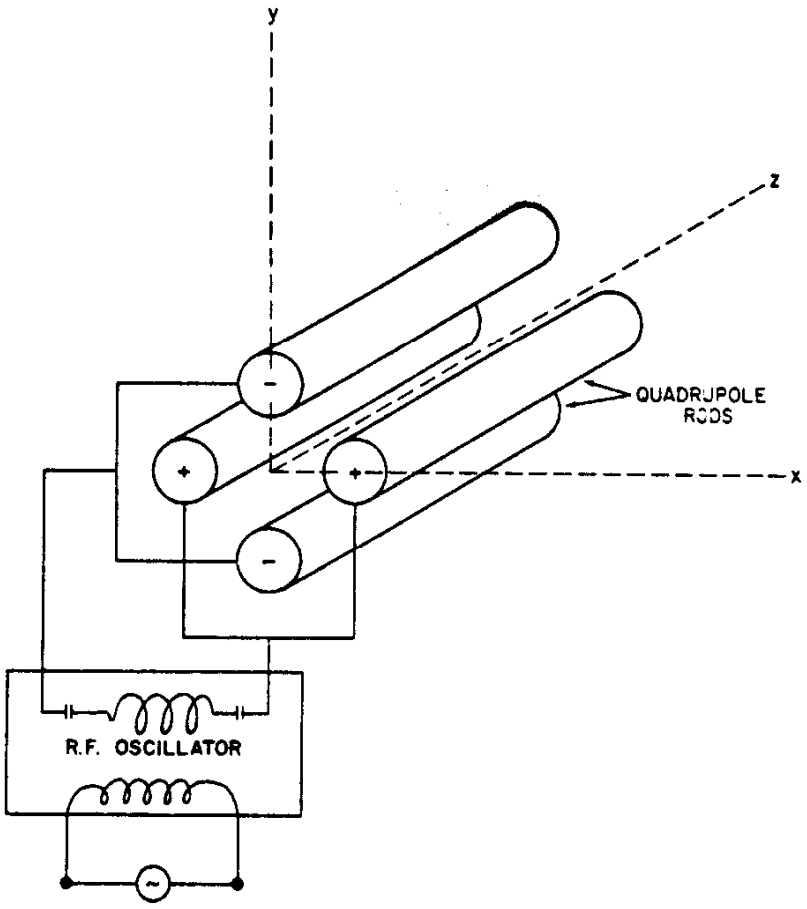
Principals of the MALDI-Tof/Tof 4700



QSTAR™ ESI QQ TOF or MALDI QQ TOF



Quadrupole Mass Analyzer



Uses a combination of RF and DC voltages to operate as a mass filter.

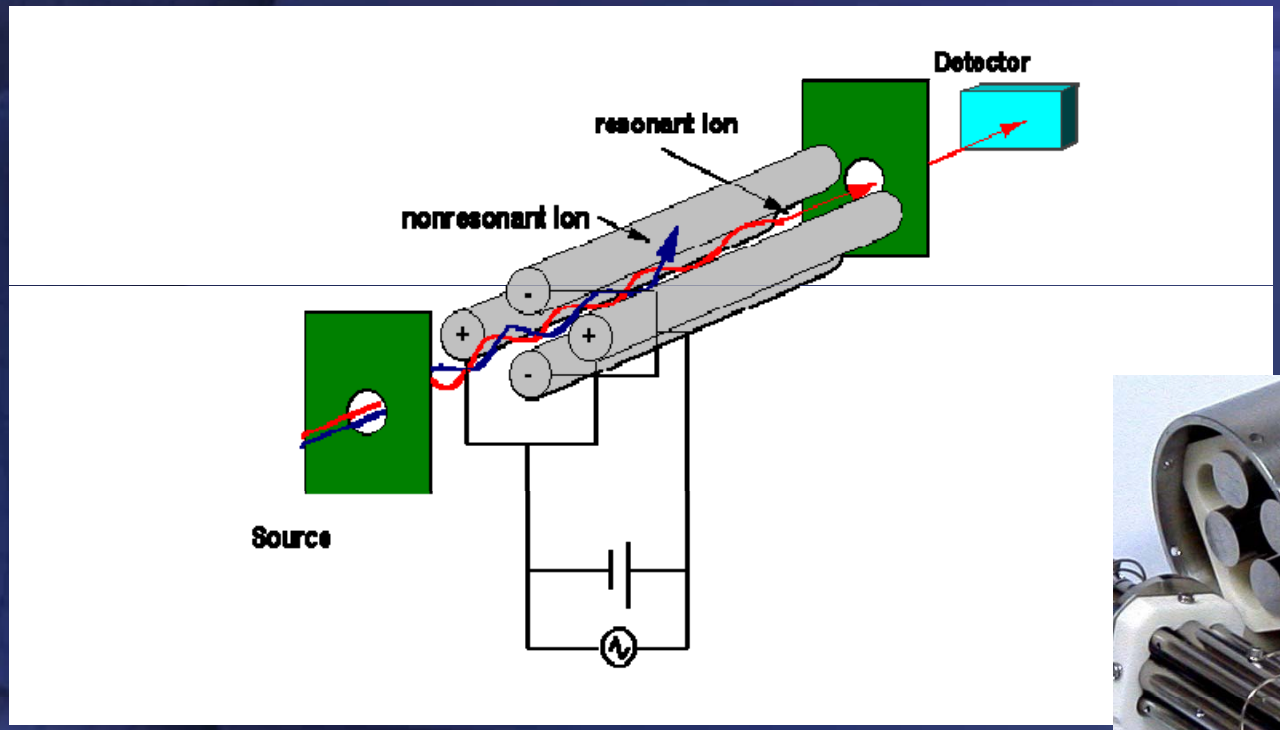
- Has four parallel metal rods.
- Lets one mass pass through at a time.
- Can scan through all masses or sit at one fixed mass.

Quadrupoles have variable ion transmission modes

mass scanning mode

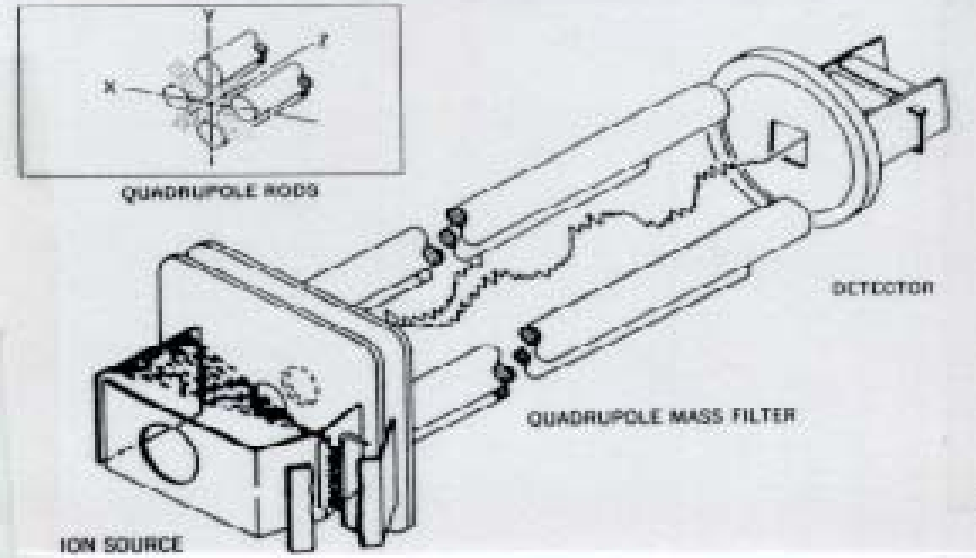
single mass transmission mode

Quadrupole mass filter / ion guide



Octapole

Quadrupole analyzer



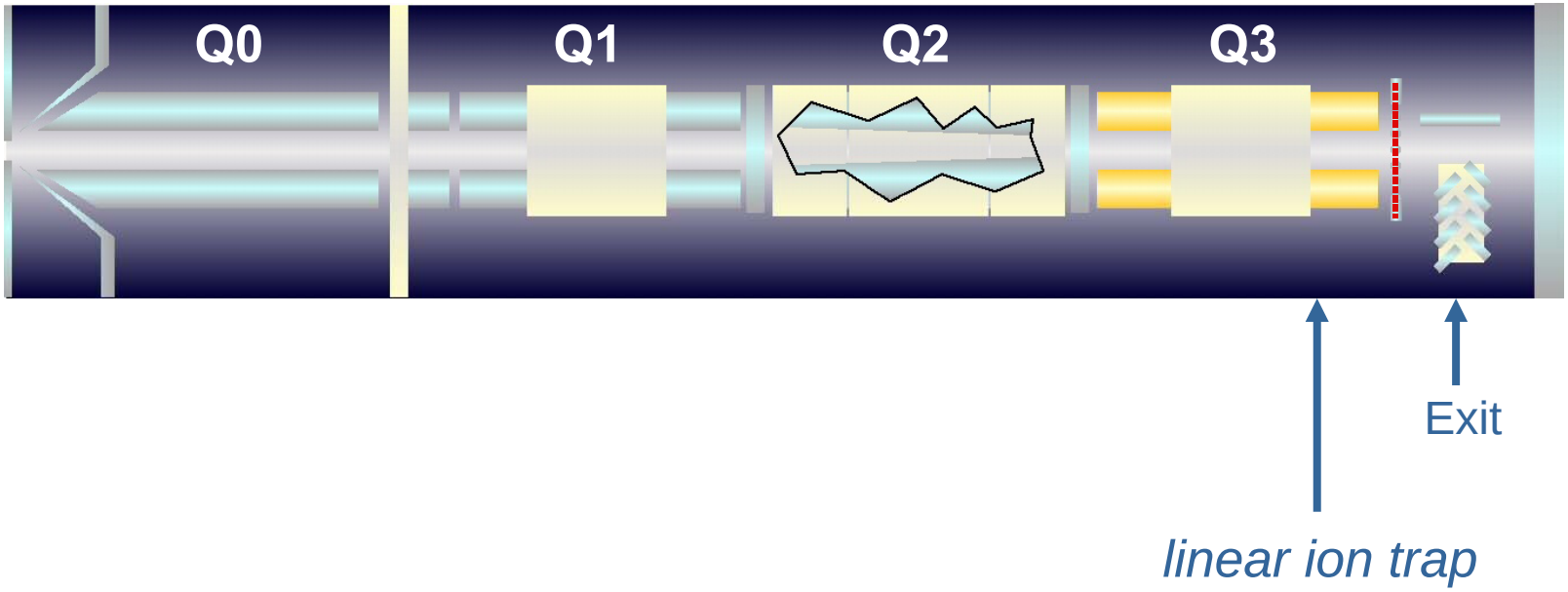
Mass resolution 2 x 10^3

Tolerant of relatively high pressure (10^{-4} torr)

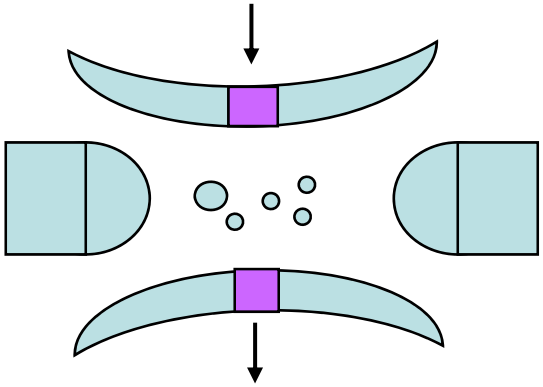
Upper limit for *m/z* is 3,000-4,000

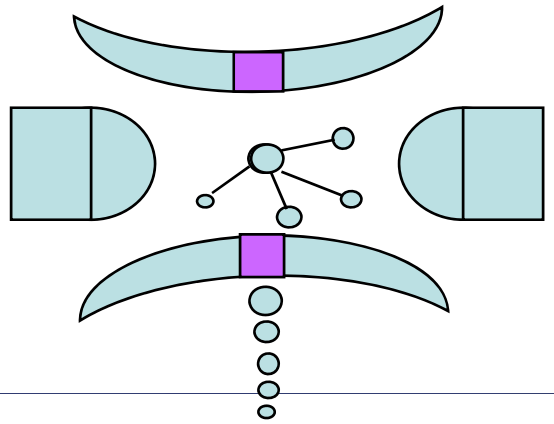
Ions are accelerated electrically (5-15V) and passed along the long central axis of four rods arranged symmetrically.
By applying combined DC and oscillating RF potentials, the ions drift along irregular flight paths along the rod axis. The DC/RF ratio is held constant and the absolute values of DC and RF are varied.
Only ions with a particular *m/z* value have stable trajectories for a given value of DC and RF.
If DC is set to 0, then all ions have stable trajectories. *A scan can be accomplished over a period of 10-1000 msec.*

QTRAP: Linear Ion Trap on a Triple Quadrupole



Ion Traps



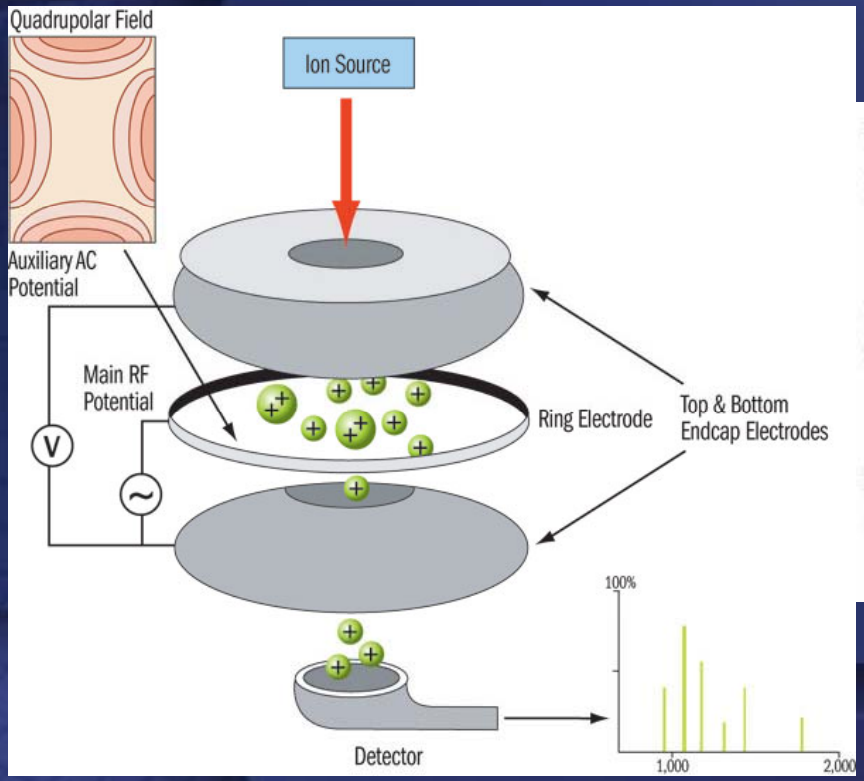


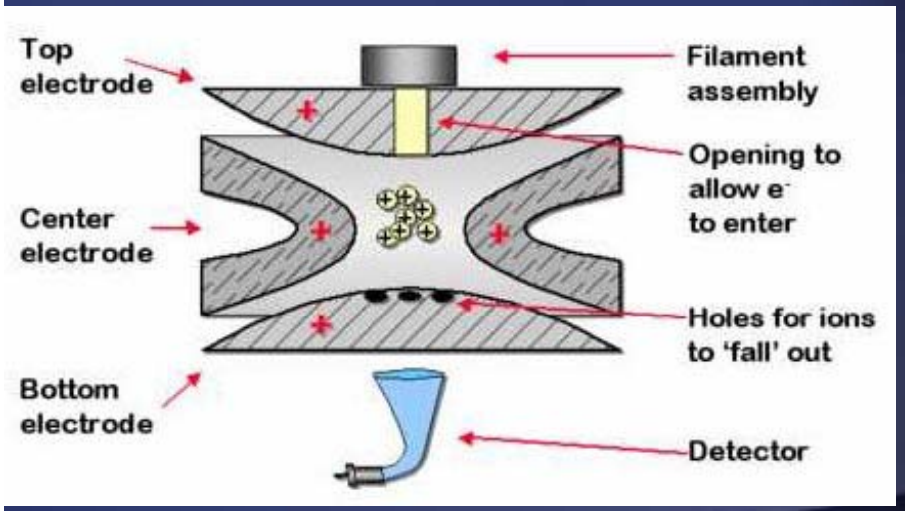
The ion trap is an energy well - ions with sufficient energy to enter the trap are retained by an energy barrier on the exit side of the trap. The advantage of the ion trap is that it accumulates selected ions prior to their analysis giving it high initial sensitivity (detection limit of approx. 20 fmol).

Ions are fragmented by collision with helium gas and their daughter ions analyzed within the trap. Selected daughter ions can undergo further fragmentation, thus allowing MS^n.

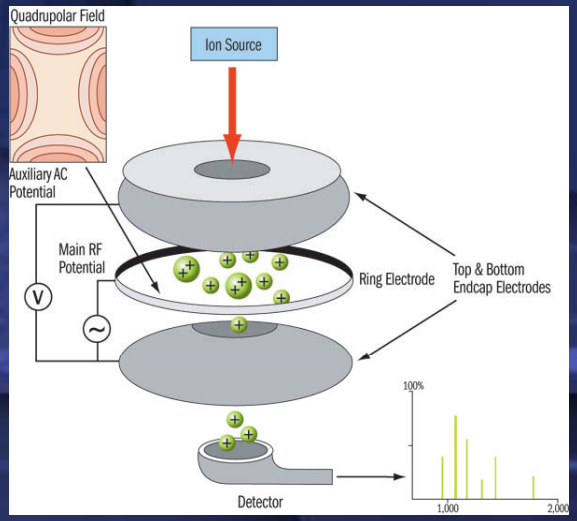
The ion trap has a high efficiency of transfer of fragment ions to the next stage of fragmentation (unlike the triple quadrupole instrument).

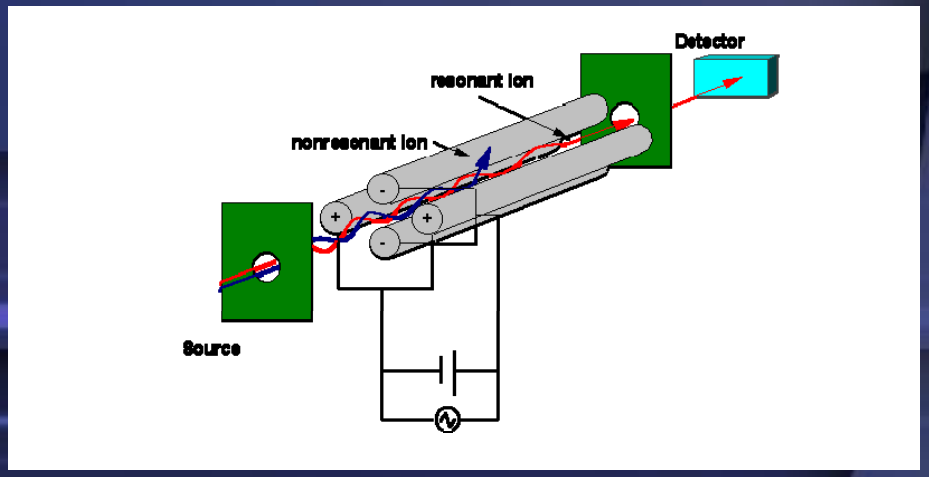
Expanded view of 3D ion trap



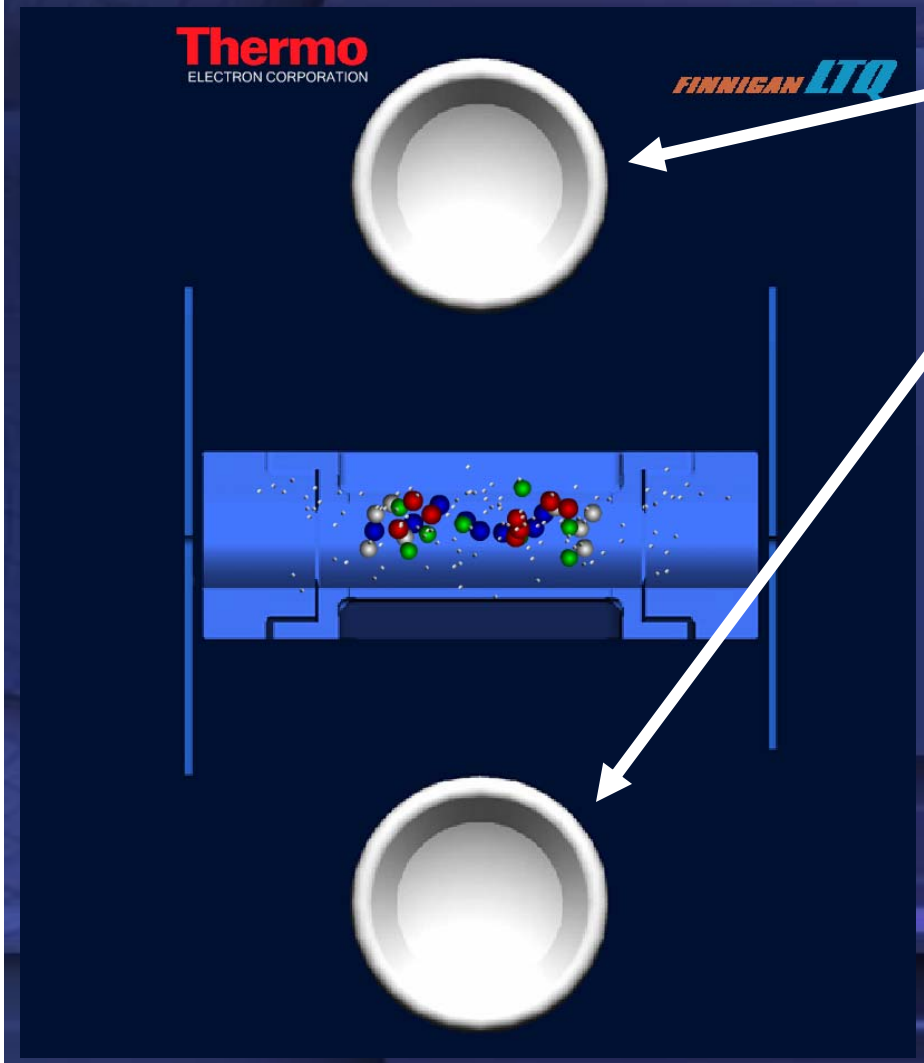


3D ion trap and 2D ion trap





2D ion trap detection



Conversion dynodes

Penning Trap (ICR cell)

Penning Trap (ICR cell)

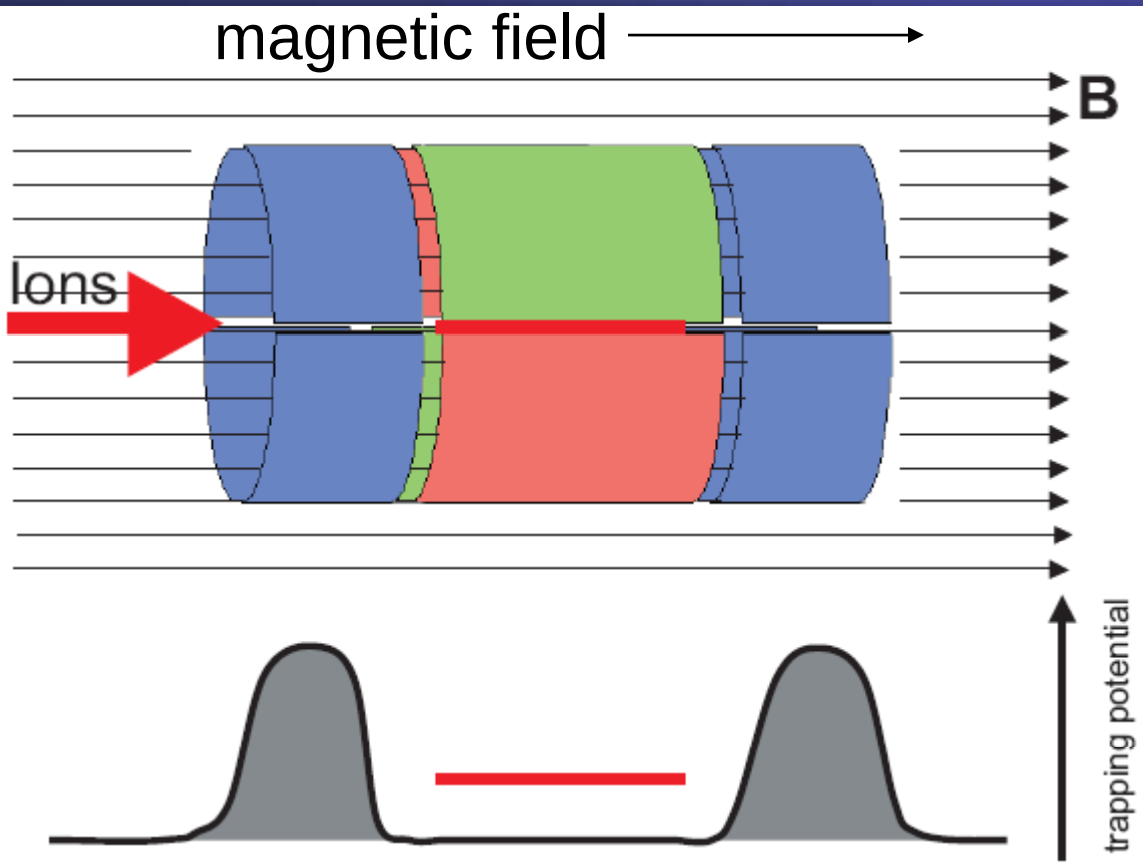
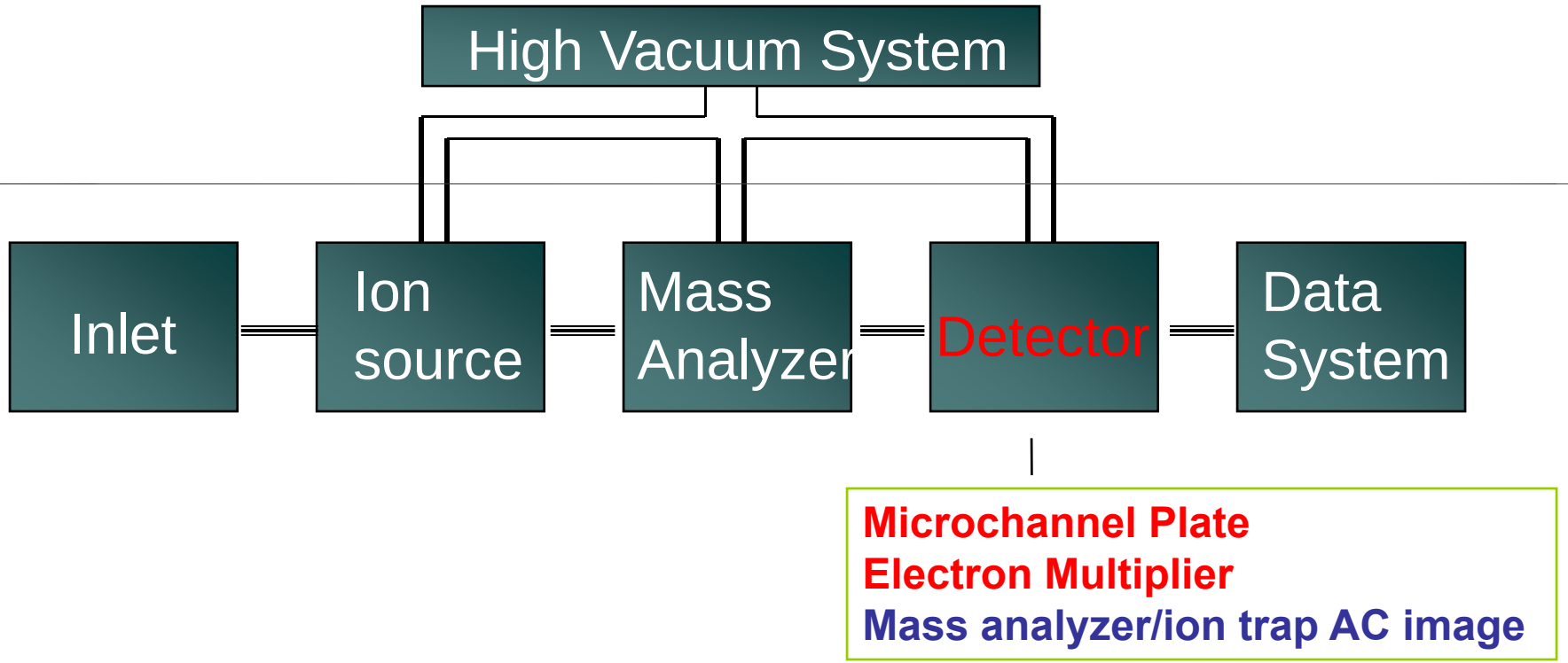


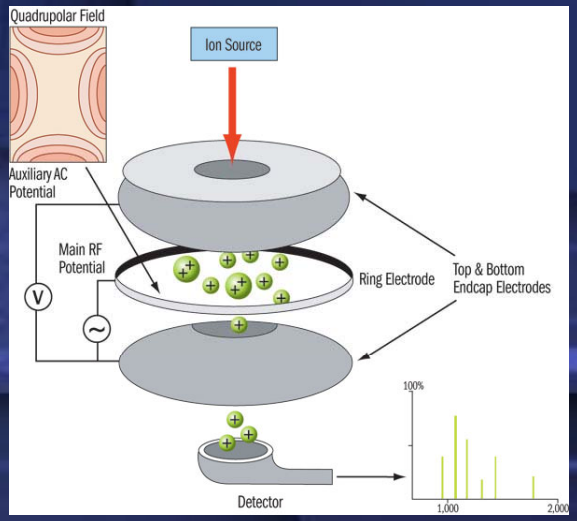
Figure 1-14. Operating principle of the Penning trap

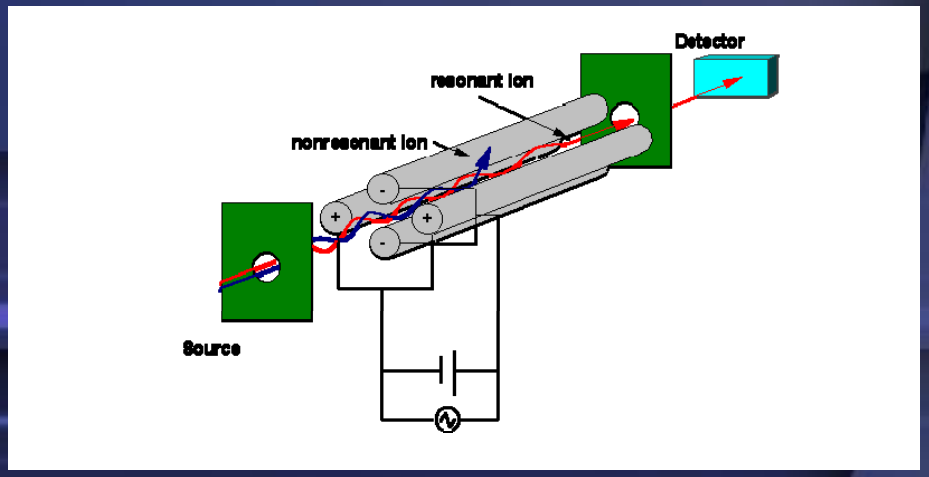
Put the trap in a high magnetic field Ion cyclotron resonance

Detector

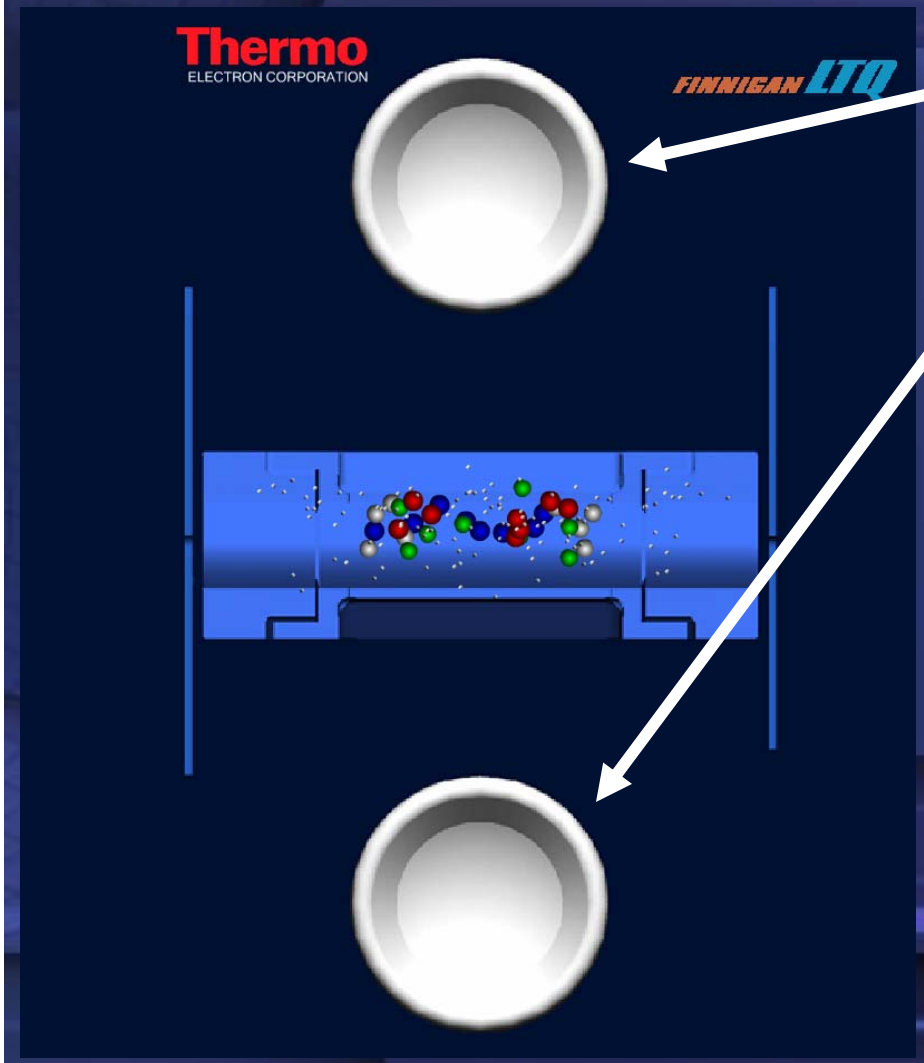

High Vacuum System
Inlet
Ion source
Mass Analyzer
Detector
Data System
Microchannel Plate
Electron Multiplier
Mass analyzer/ion trap AC image

3D ion trap and 2D ion trap



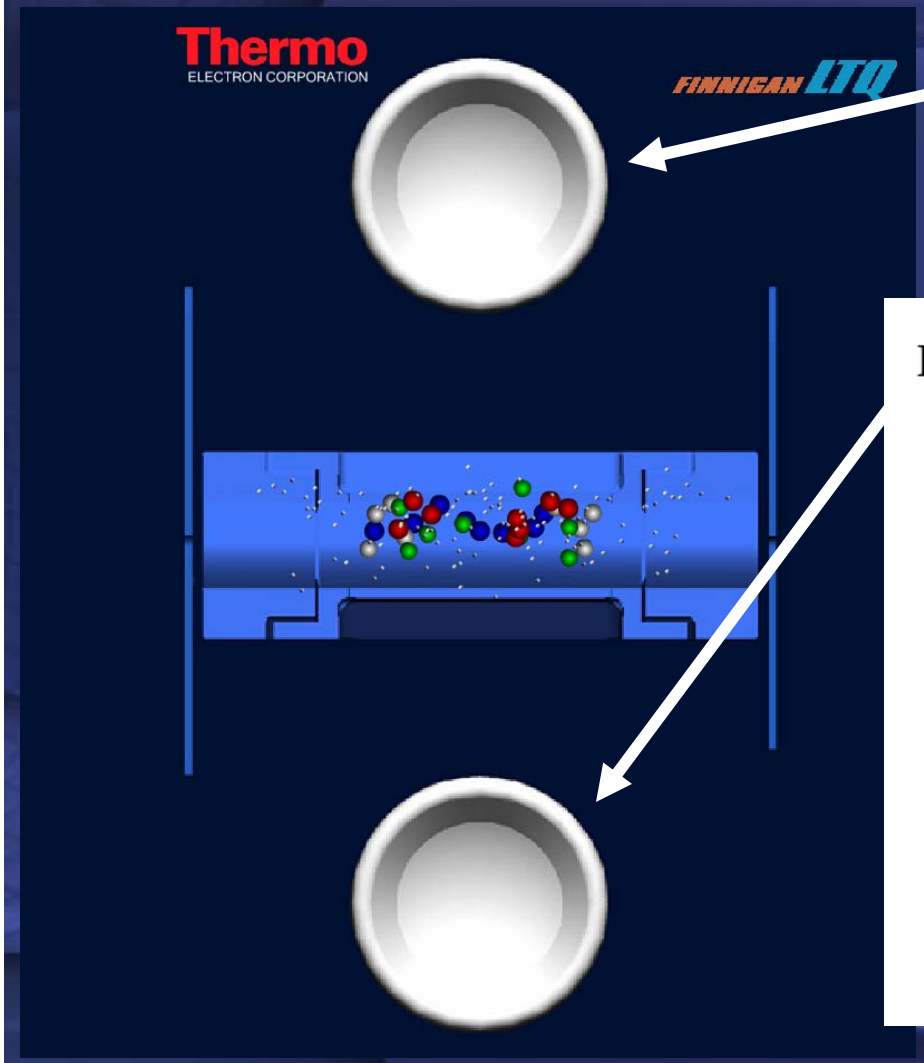


2D ion trap detection

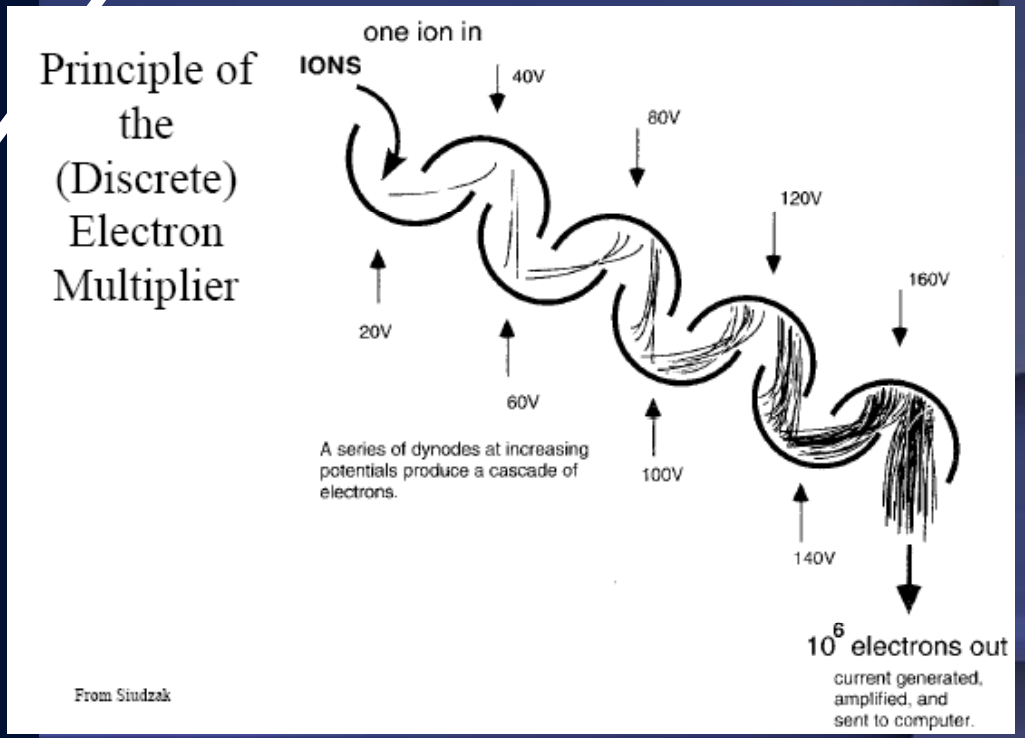


Conversion dynodes

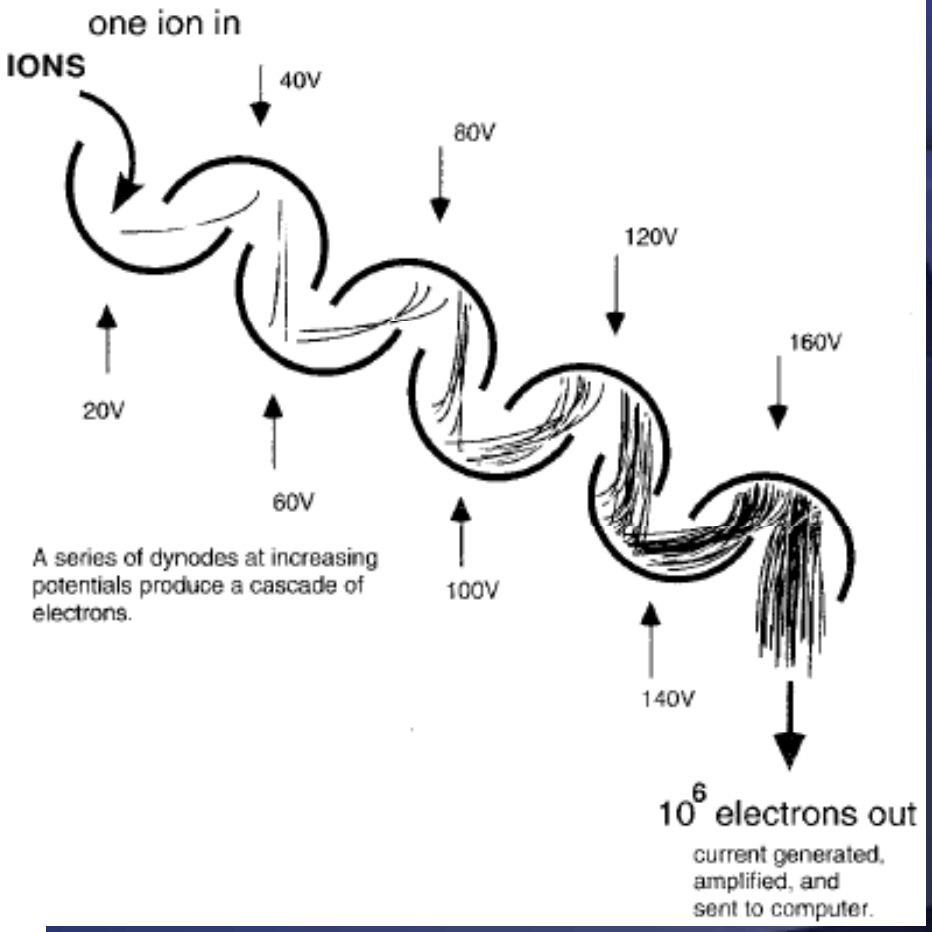
2D ion trap detection



Conversion dynodes (electron multipliers)

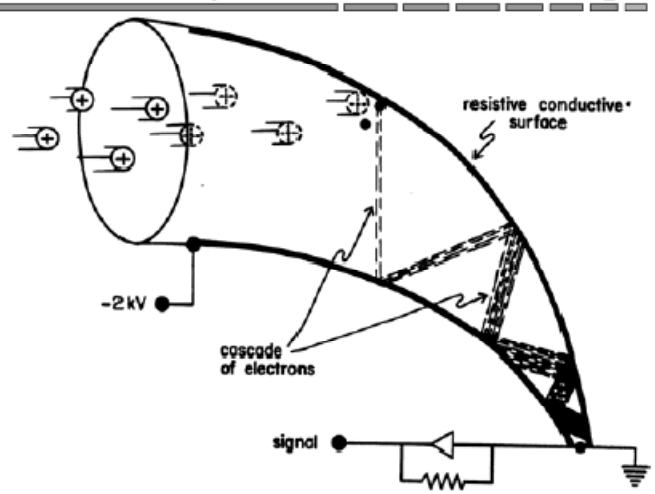


Principle of the (Discrete) Electron Multiplier



A series of dynodes at increasing potentials produce a cascade of electrons.

Continuous Dynode Electron Multiplier

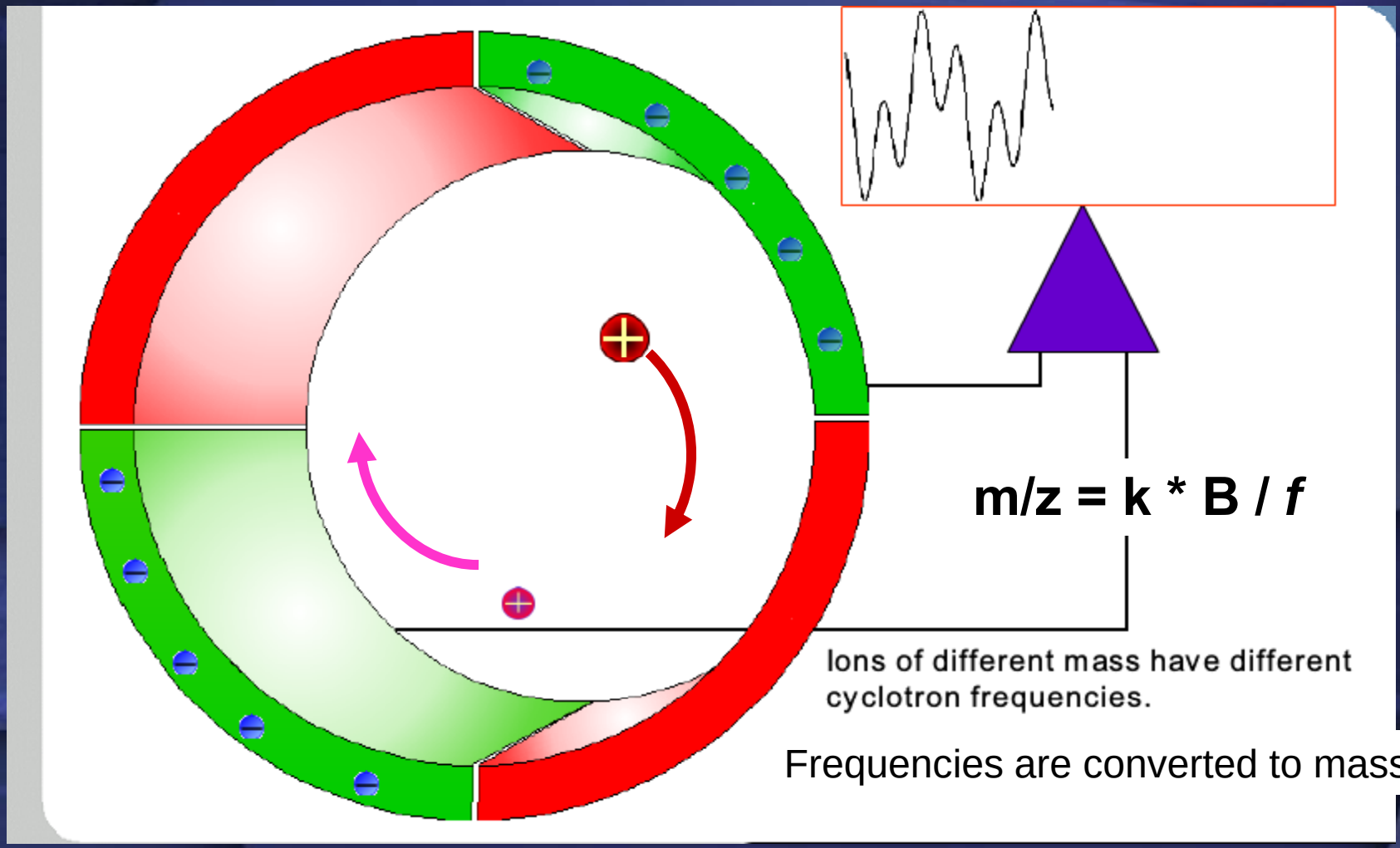


From Watson

FIG. 13.3. Conceptual diagram of a nonmagnetic electron multiplier; the field gradient along the resistive conductive internal surface of the cornucopia attracts the cascading electrons toward the preamplifier.

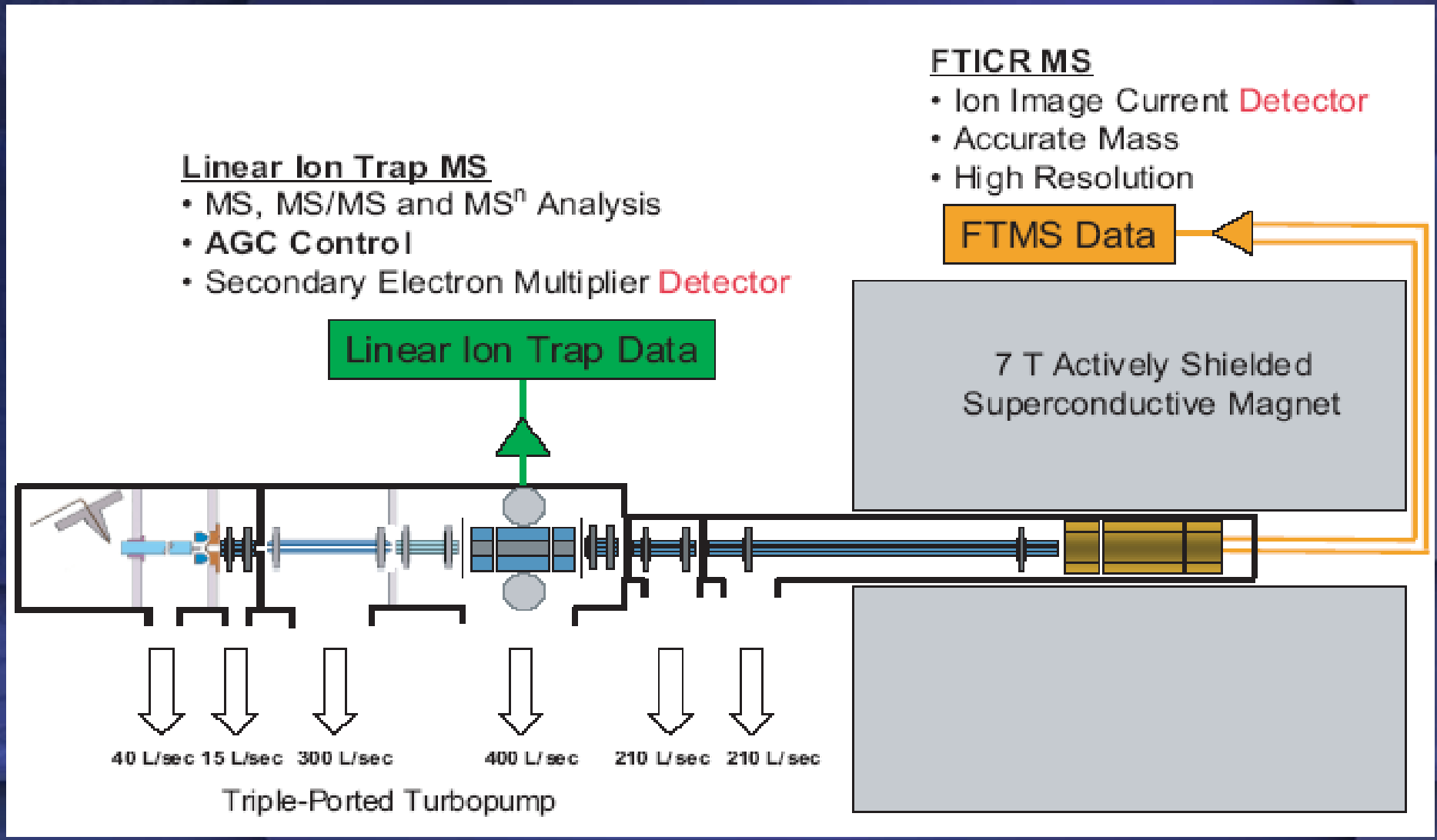
Detecting in the ion trap
Ion cyclotron resonance (ICR)

Fourier transform-
Ion cyclotron resonance
FT-ICR MS

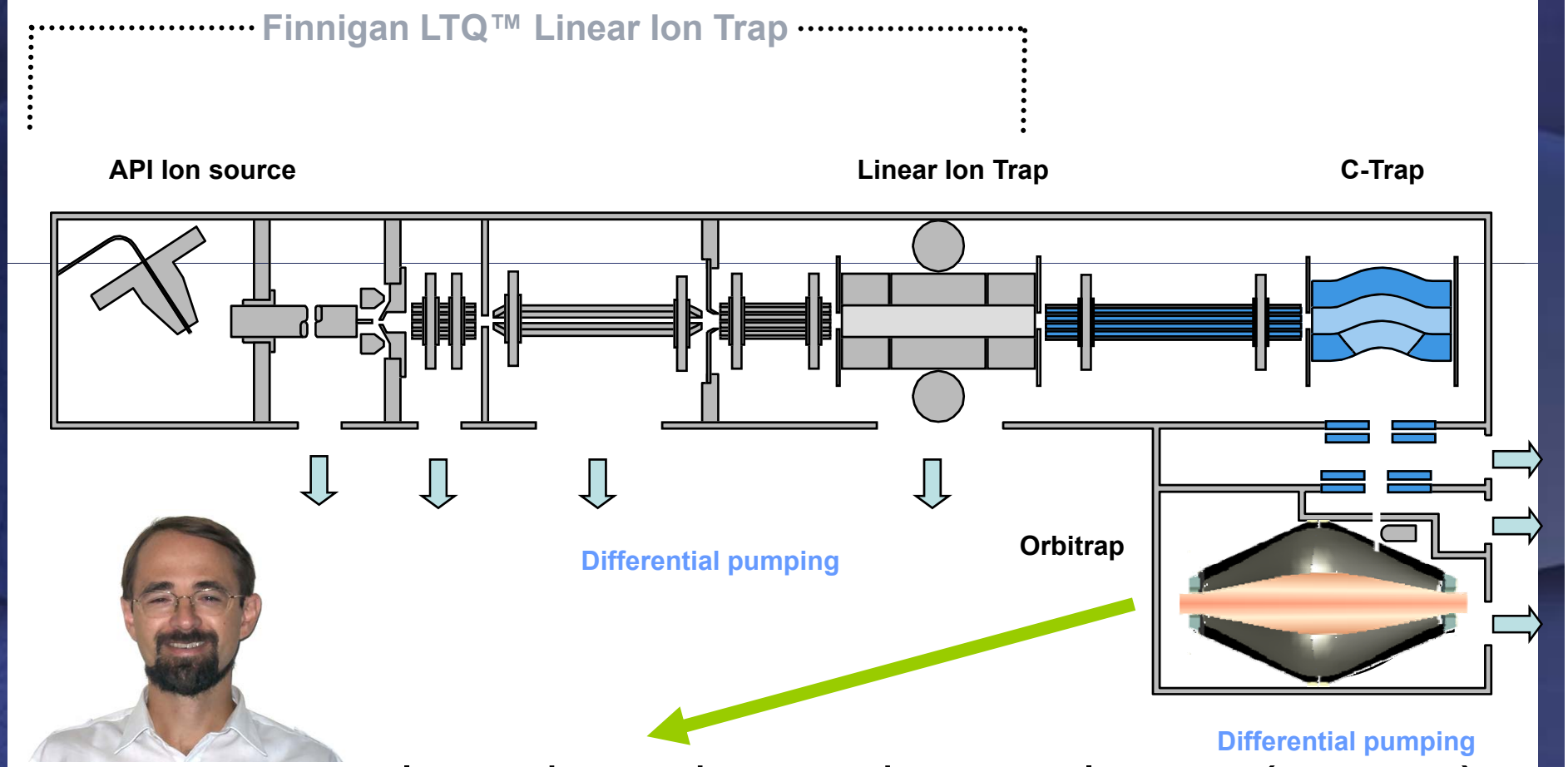


Frequencies are converted to masses.

ThermoFinnigan LTQ-FT



LTQ Orbitrap™ Hybrid Mass Spectrometer



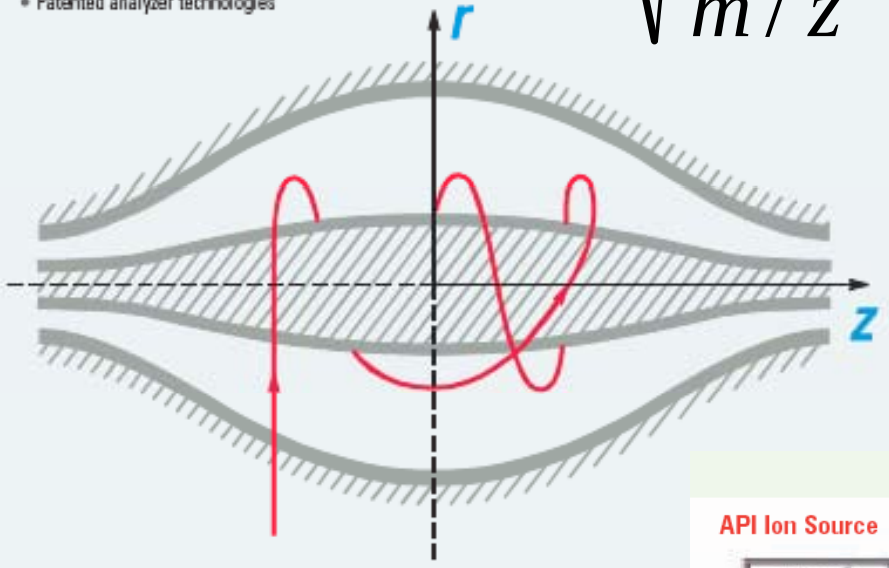
Inventor: Dr. Alexander Makarov, Thermo Electron (Bremen)

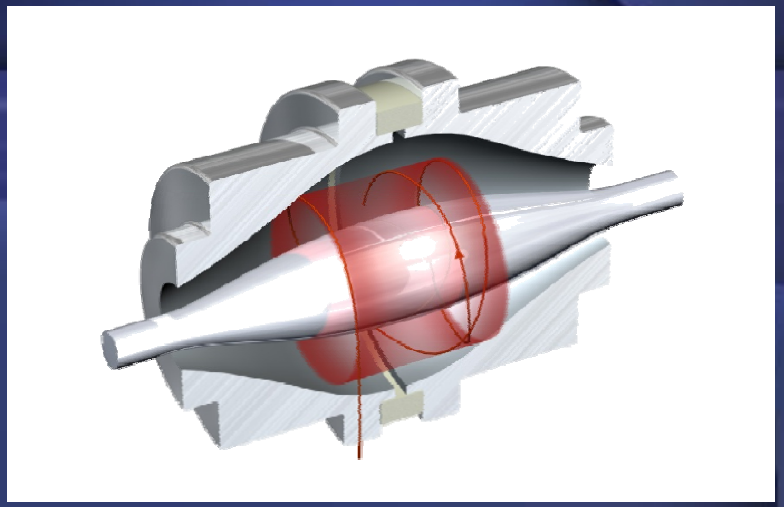
Orbitrap Mass Analyzer

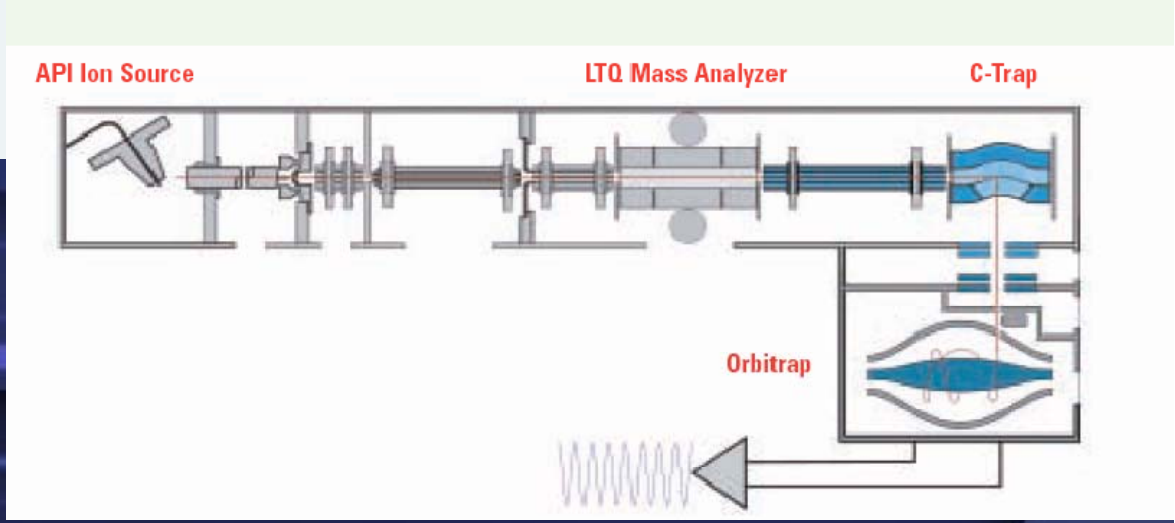
Orbitrap: A Breakthrough Electrostatic Ion Trap

- Highest ion trapping efficiencies
- Large ion capacity
- Stable and robust operation
- Patented analyzer technologies

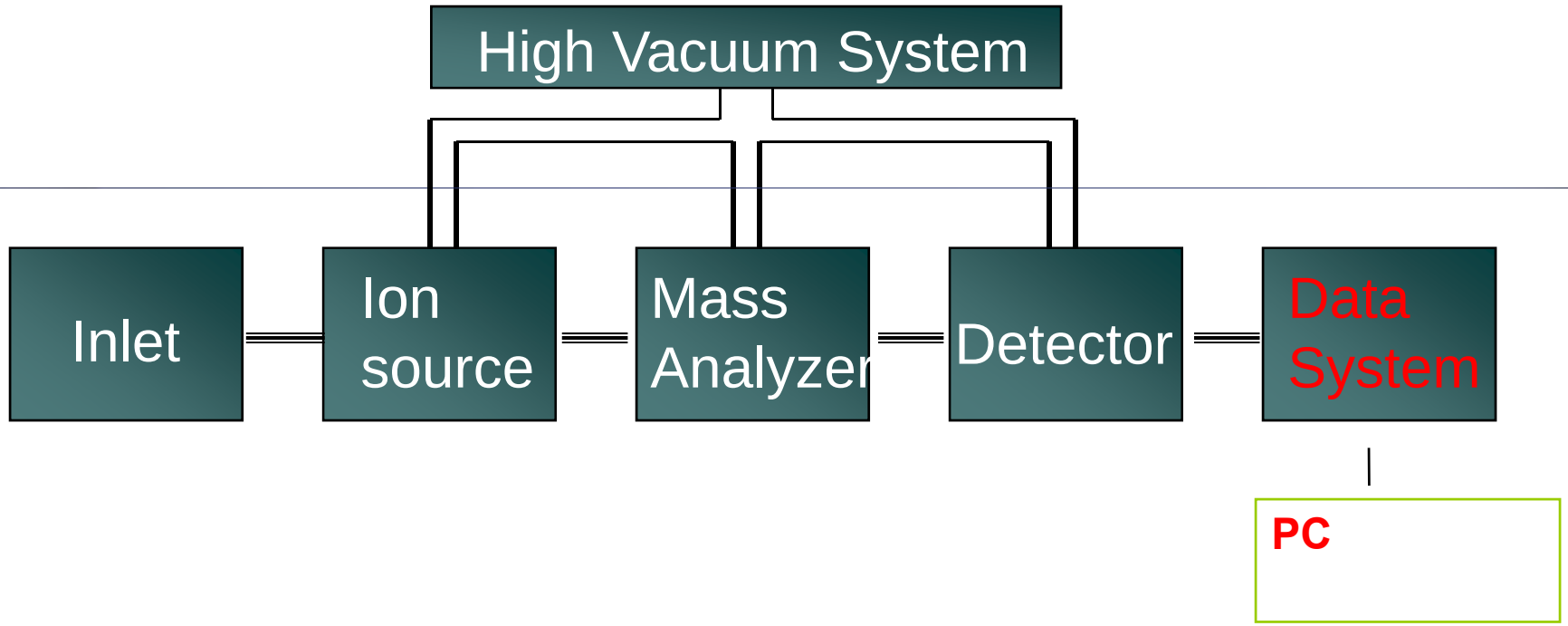
$$\omega = \sqrt{\frac{k}{m/z}}$$





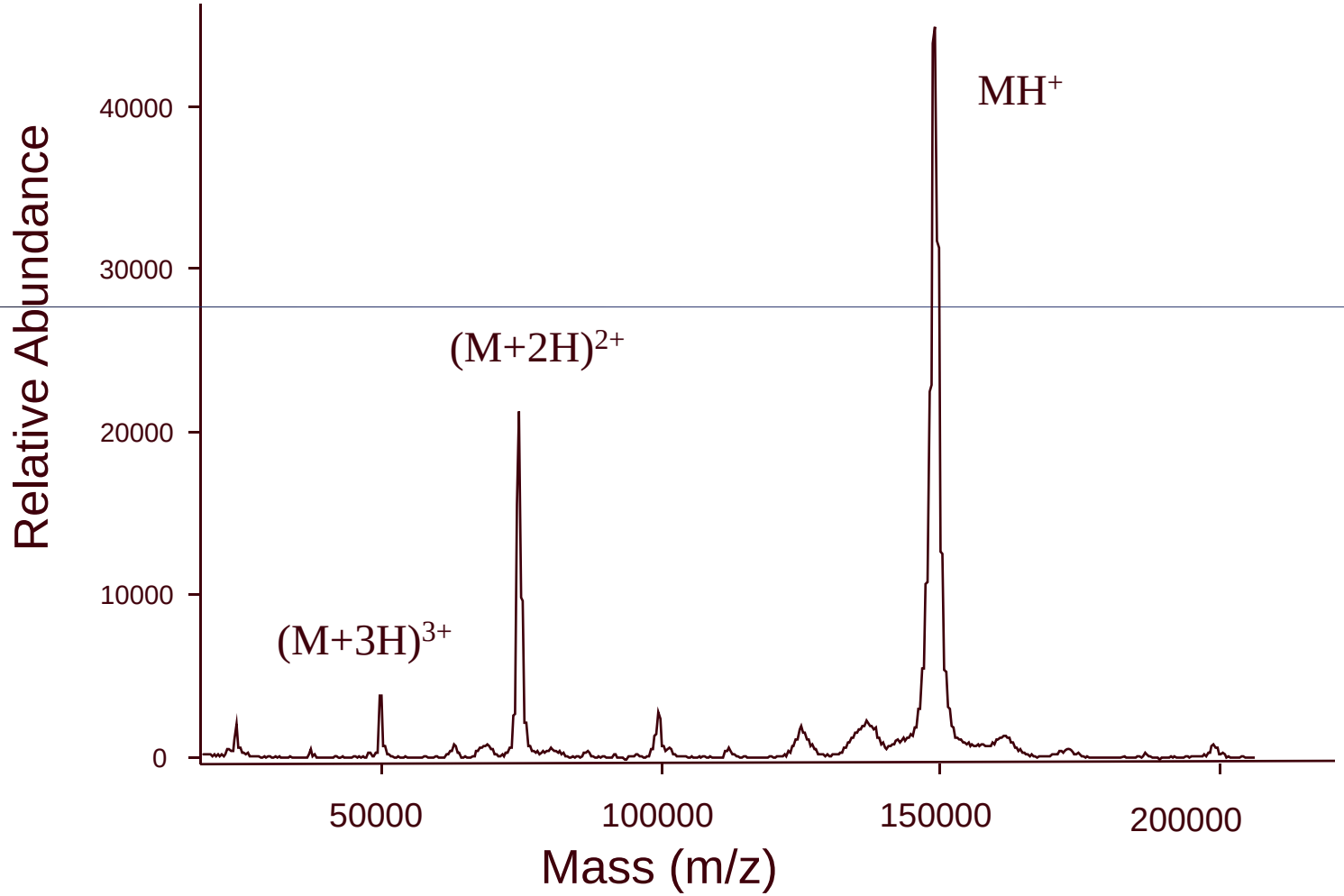


Data System

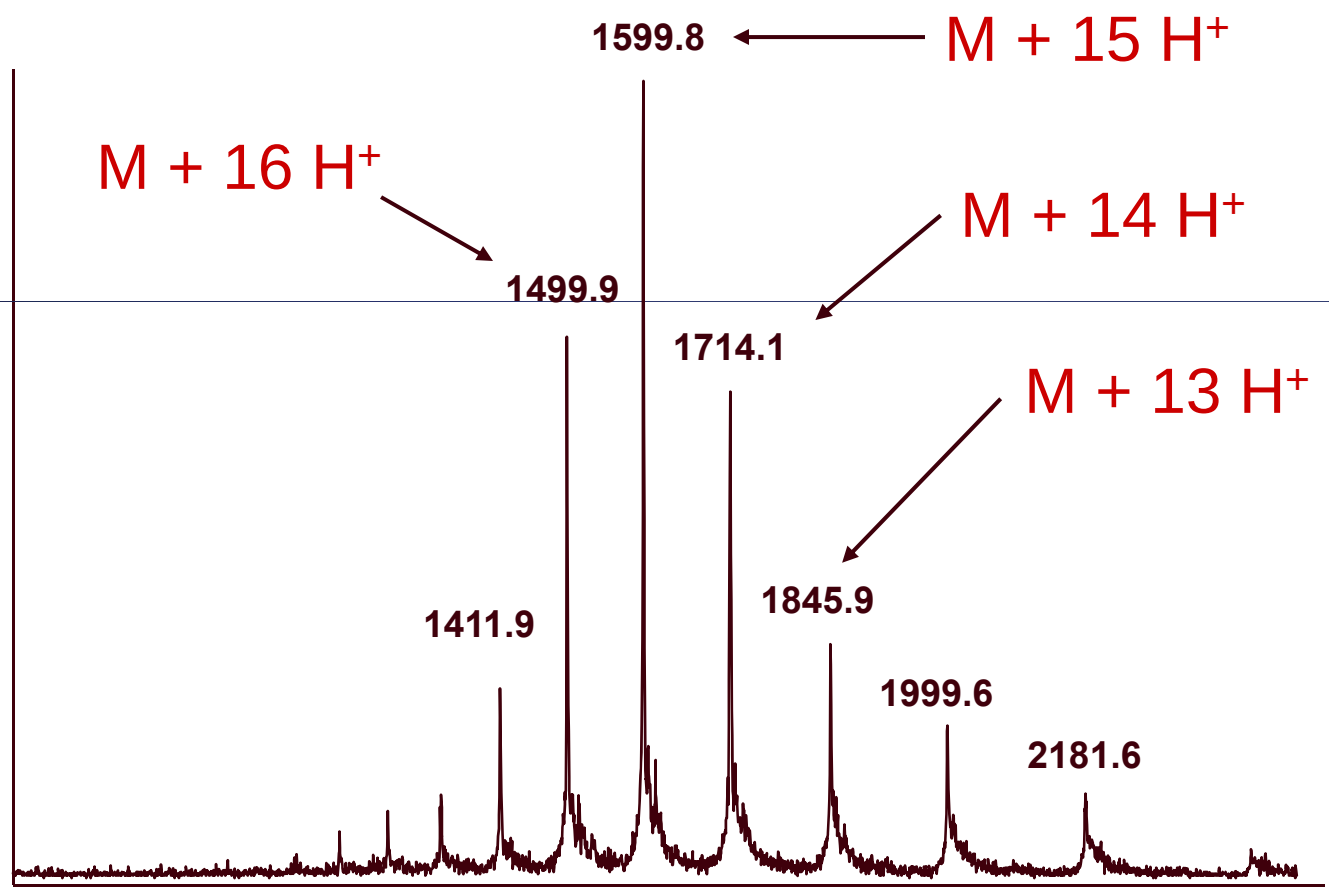

High Vacuum System
Inlet
Ion source
Mass Analyzer
Detector
Data System
PC

The mass spectrum shows the results

MALDI TOF spectrum of IgG



ESI Spectrum of Trypsinogen (MW 23983)



How do mass spectrometers get their names?

Types of ion sources:

- Electrospray (ESI)
- Matrix Assisted Laser Desorption Ionization (MALDI)

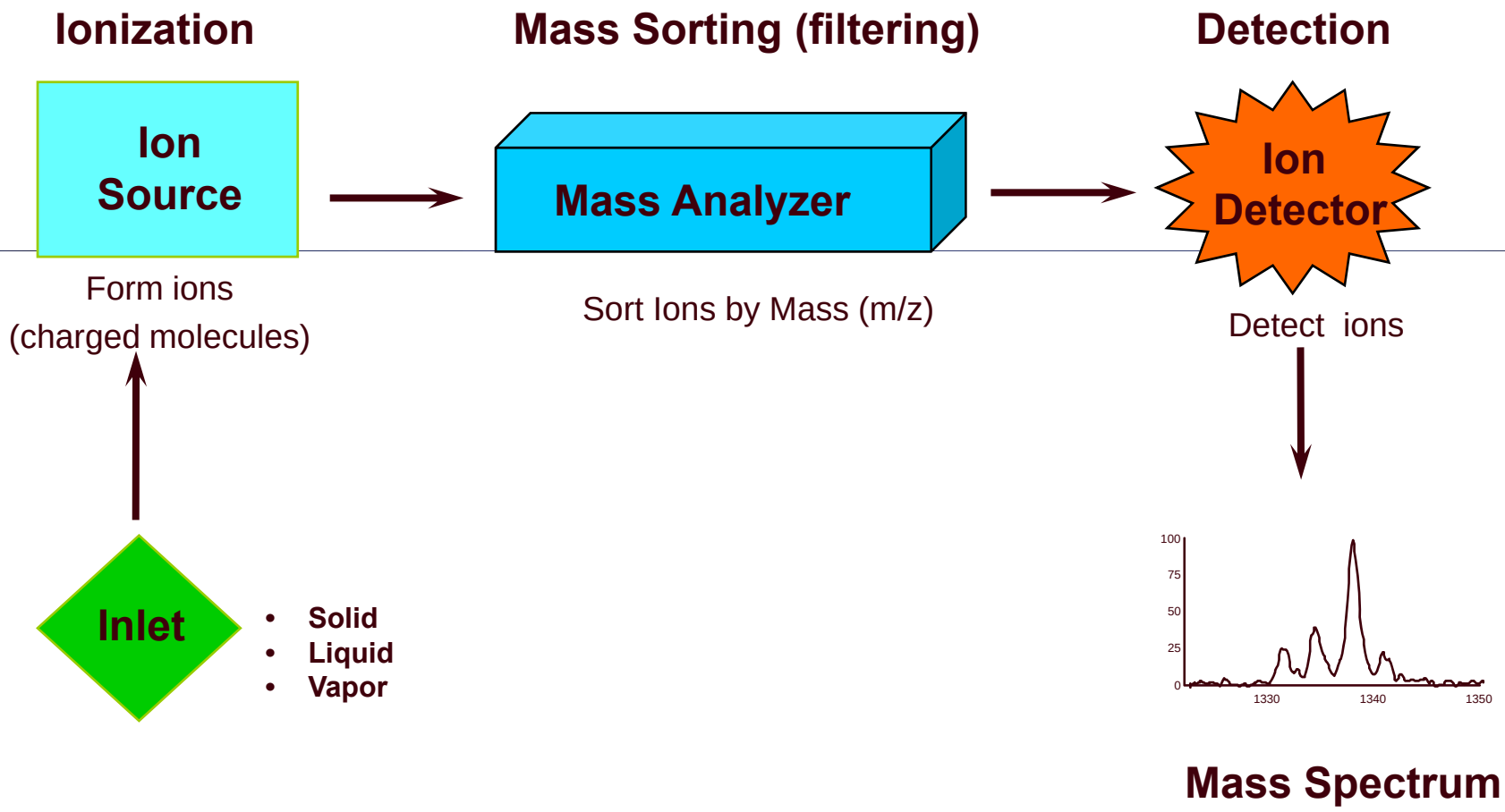
Types of mass analyzers:

- Quadrupole (Quad, Q)
- Ion Trap
- Time-of-Flight (TOF)

-Either source type can work with either analyzer type: “MALDI-TOF,” “ESI-Quad.”

-Analyzers can be combined to create “hybrid” instruments. ESI-QQQ, MALDI QQ TOF, Q Trap

Summary: acquiring a mass spectrum



Put it in

- We need Ions (+ or -)
- In the gas phase

Your machine

- Tof, Tof / Tof
- Quadrupole, Ion trap
- FT-ICR, Orbitrap (high resolution)
- Hybrids

Tell me the RIGHT answer

- How right is it?

  mass resolution and accuracy

How is mass defined?

Assigning numerical value to the intrinsic property of “mass” is based on using carbon-12, ^{12}C, as a reference point.

One unit of mass is defined as a Dalton (Da).

One Dalton is defined as 1/12 the mass of a single carbon-12 atom.

Thus, one ^{12}C atom has a mass of 12.0000 Da.

Isotopes

+Most elements have more than one stable isotope.

For example, most carbon atoms have a mass of 12 Da, but in nature, 1.1% of C atoms have an extra neutron, making their mass 13 Da.

+Why do we care?

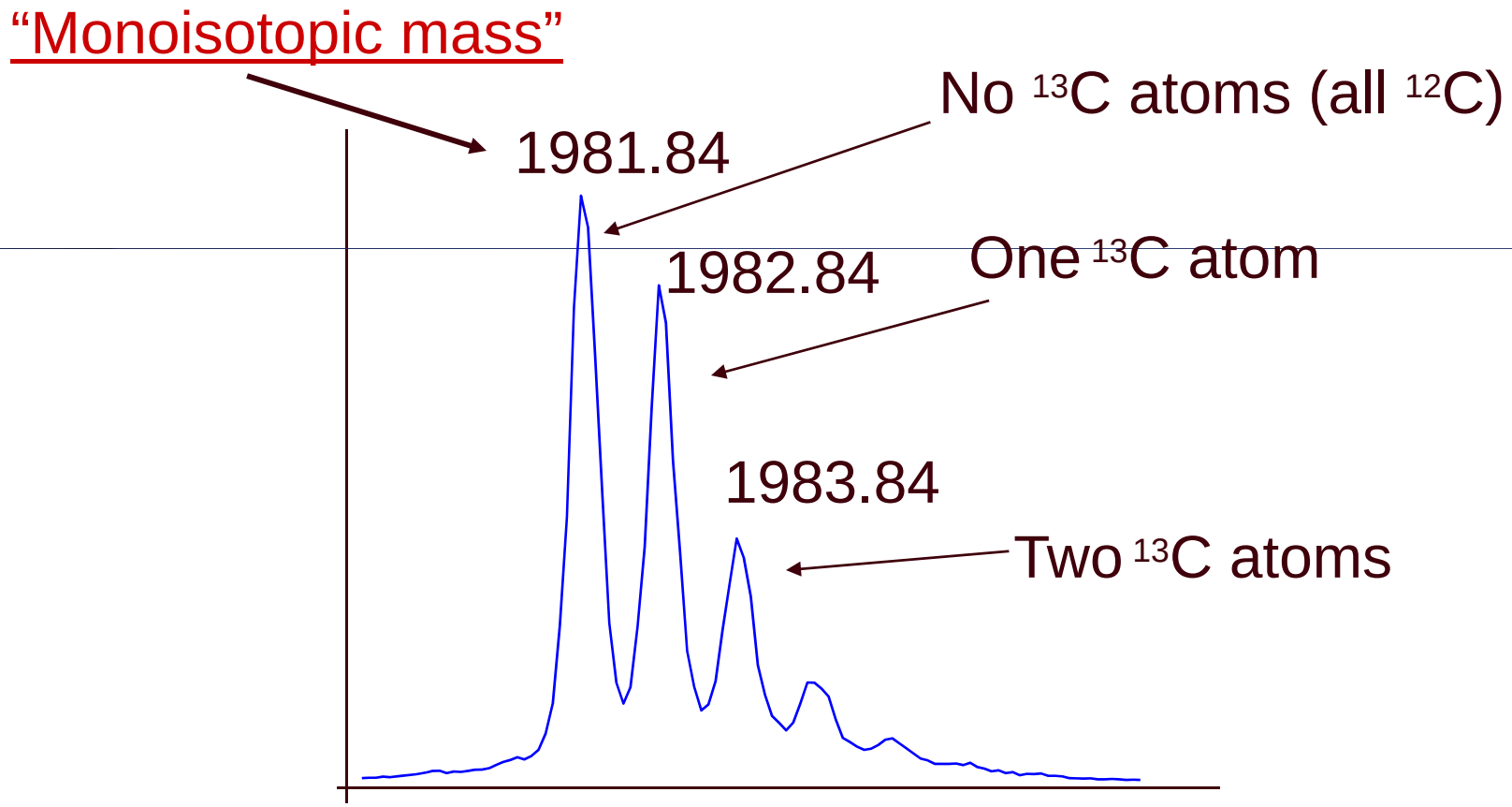
Mass spectrometers can “see” isotope peaks if their resolution is high enough.

If an MS instrument has resolution high enough to resolve these isotopes, better mass accuracy is achieved.

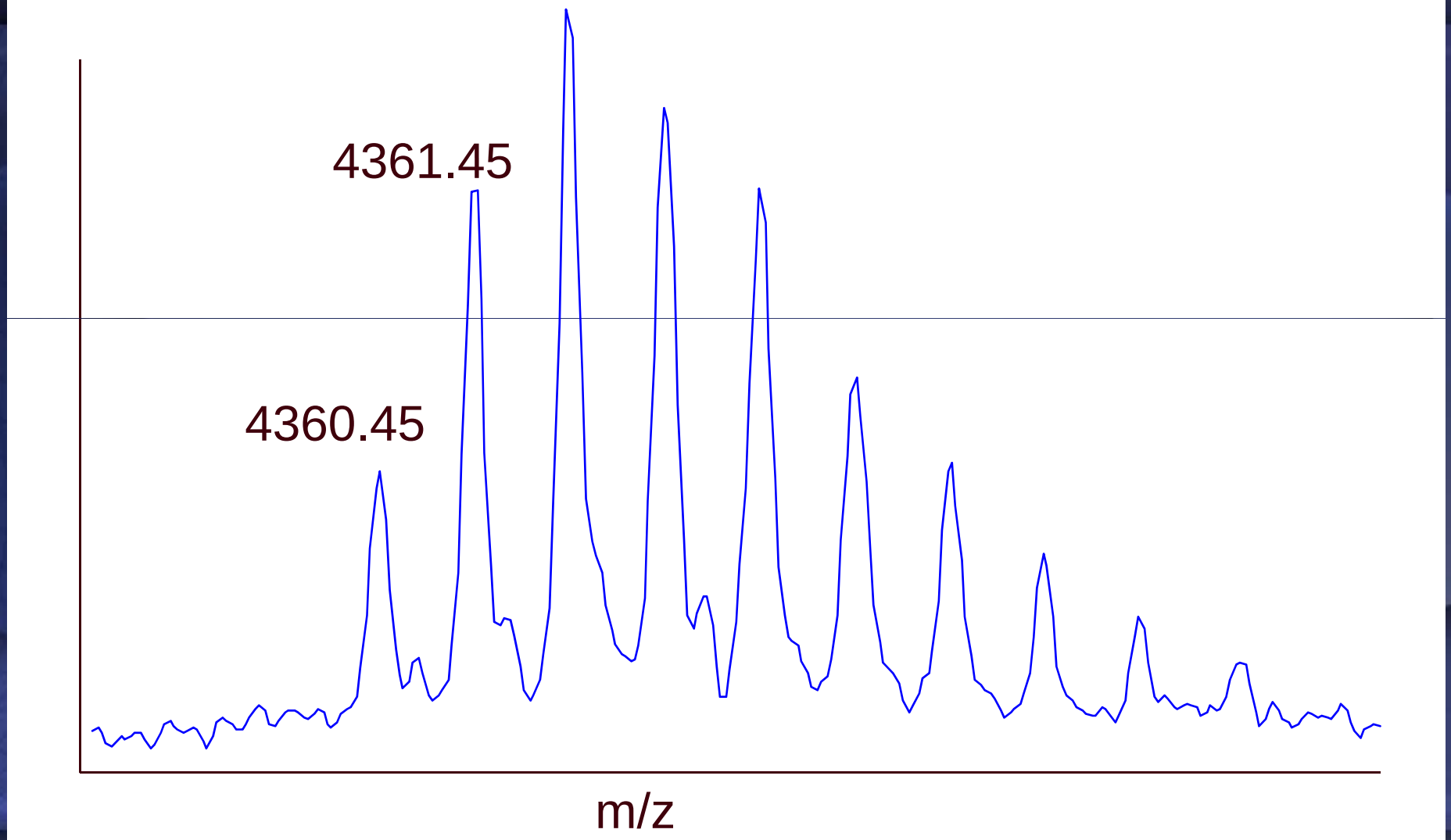
Stable isotopes of most abundant elements of peptides

Element	Mass	Abundance
H	1.0078 2.0141	99.985% 0.015
C	12.0000 13.0034	98.89 1.11
N	14.0031 15.0001	99.64 0.36
O	15.9949 16.9991 17.9992	99.76 0.04 0.20

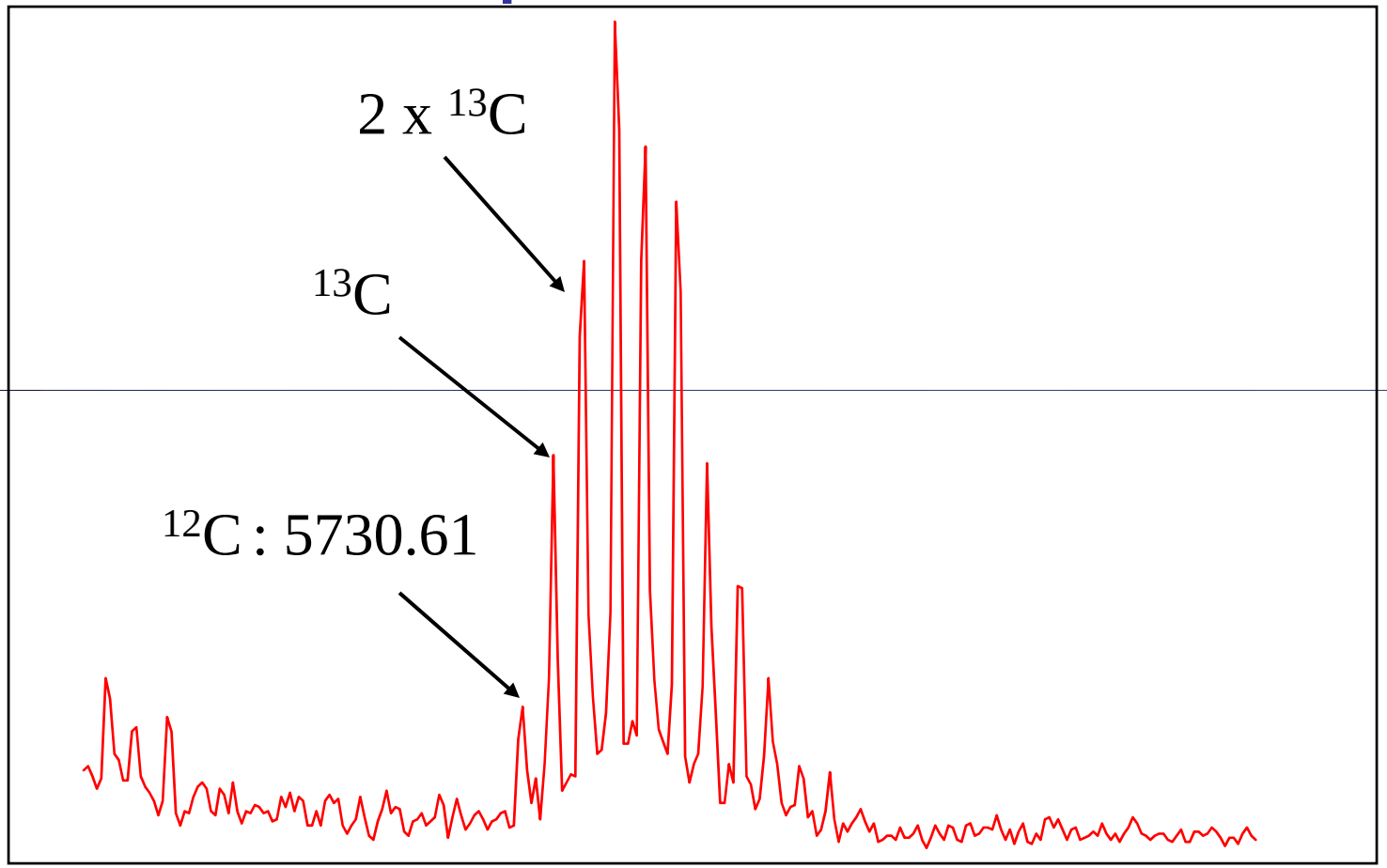
Mass spectrum of peptide with 94 C-atoms (19 amino acid residues)



Isotope pattern for a larger peptide (207 C-atoms)

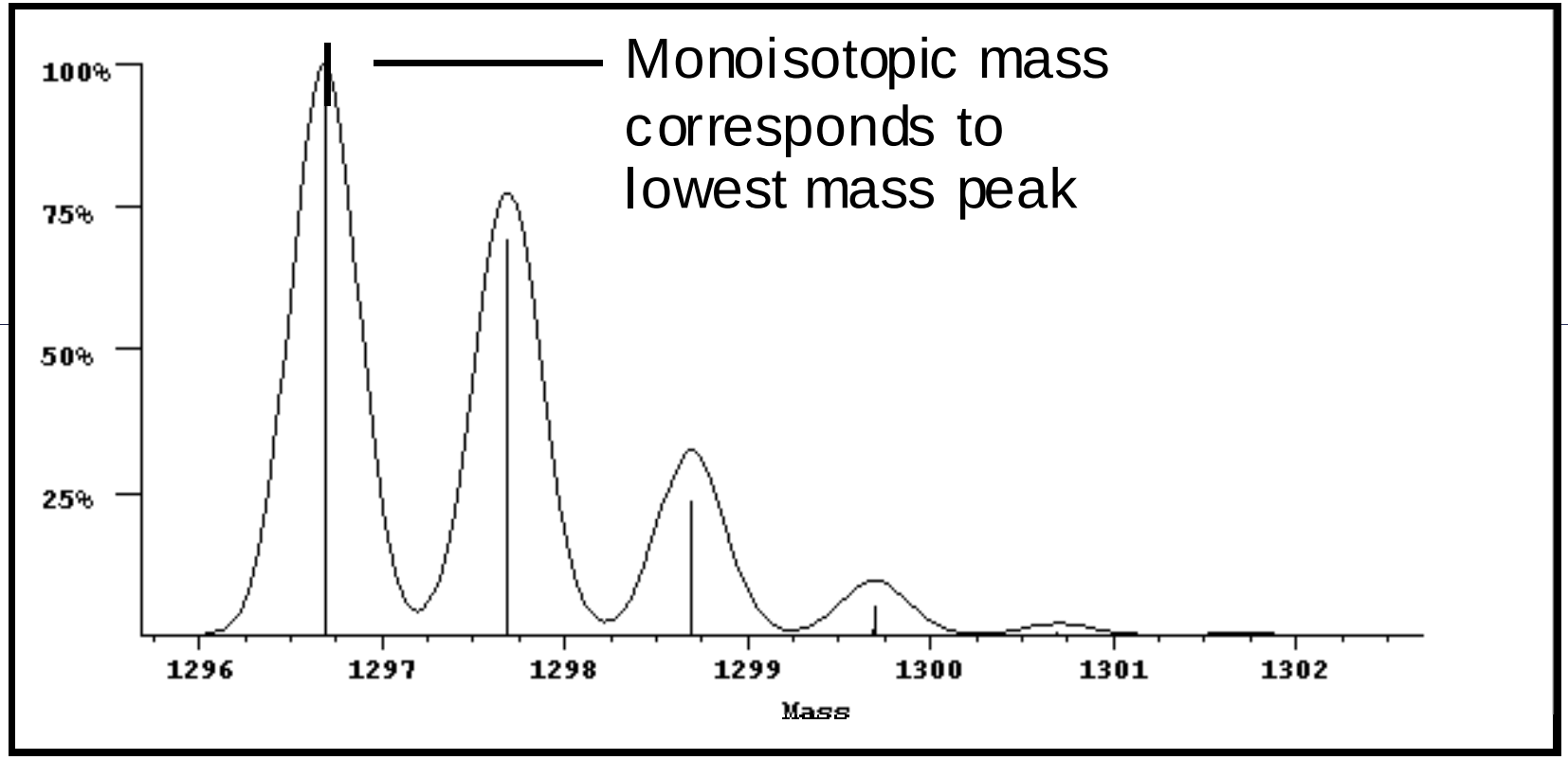


Mass spectrum of insulin



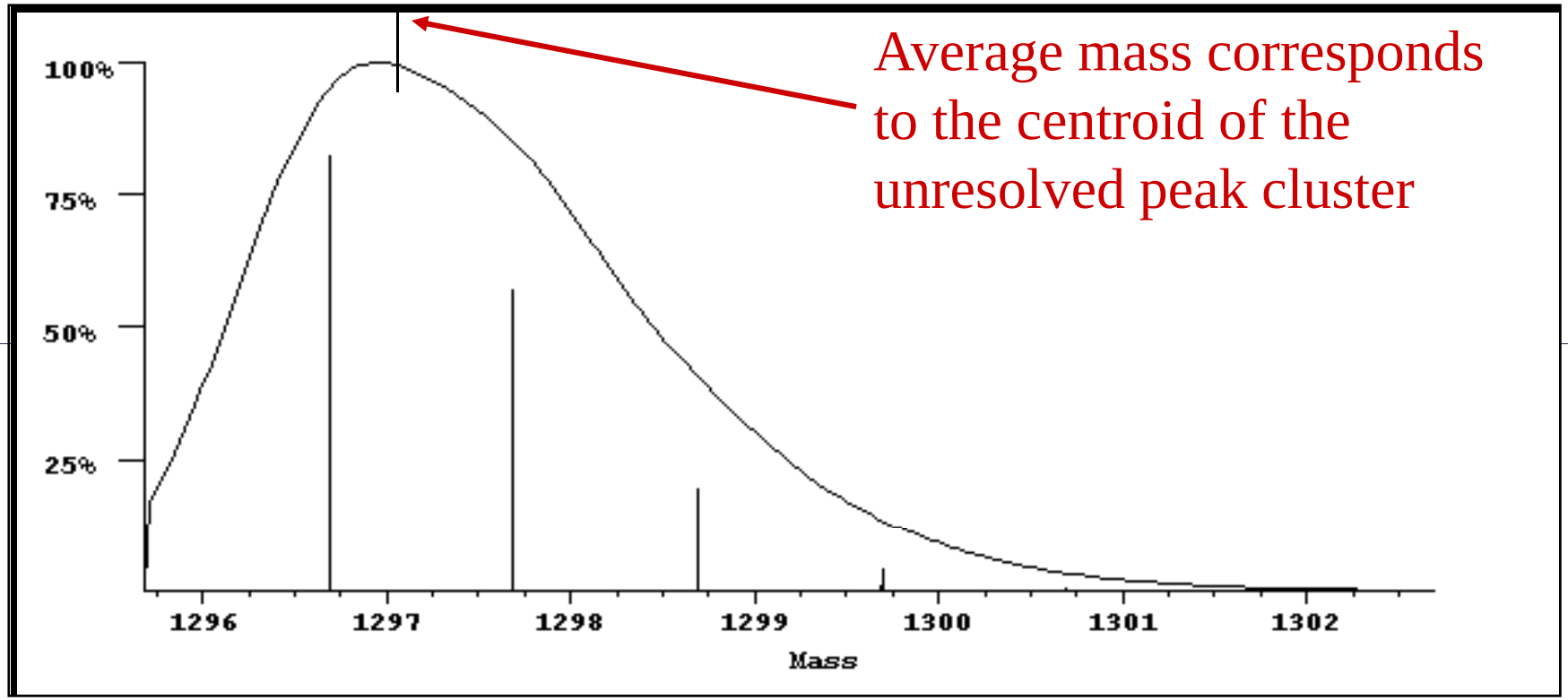
Insulin has 257 C-atoms. Above this mass, the monoisotopic peak is too small to be very useful, and the average mass is usually used.

Monoisotopic mass



When the isotopes are clearly resolved the **monoisotopic mass** is used as it is the most accurate measurement.

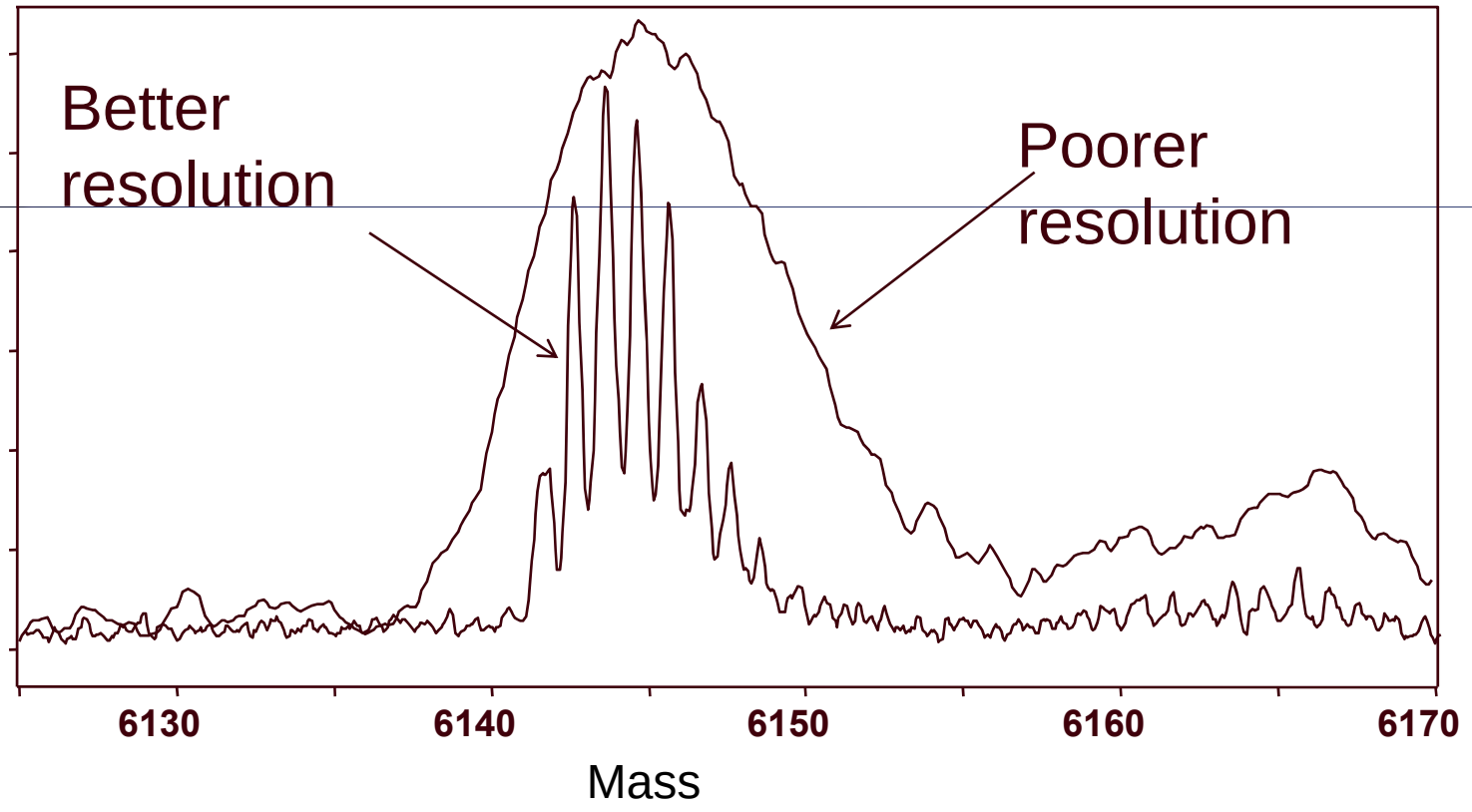
Average mass



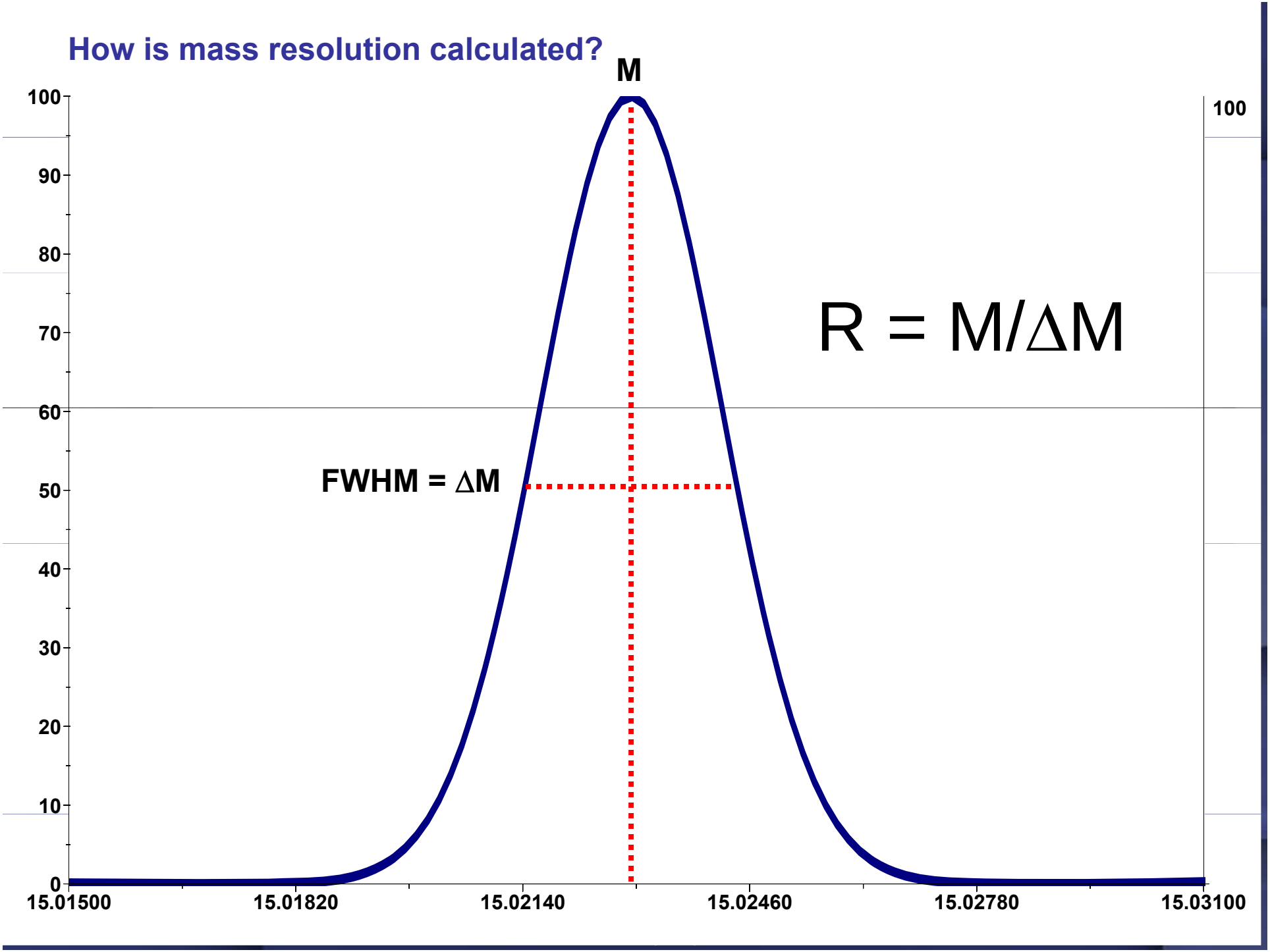
When the isotopes are not resolved, the centroid of the envelope corresponds to the weighted average of all the the isotope peaks in the cluster, which is the same as the average or chemical mass.

What if the resolution is not so good?

At lower resolution, the mass measured is the average mass.

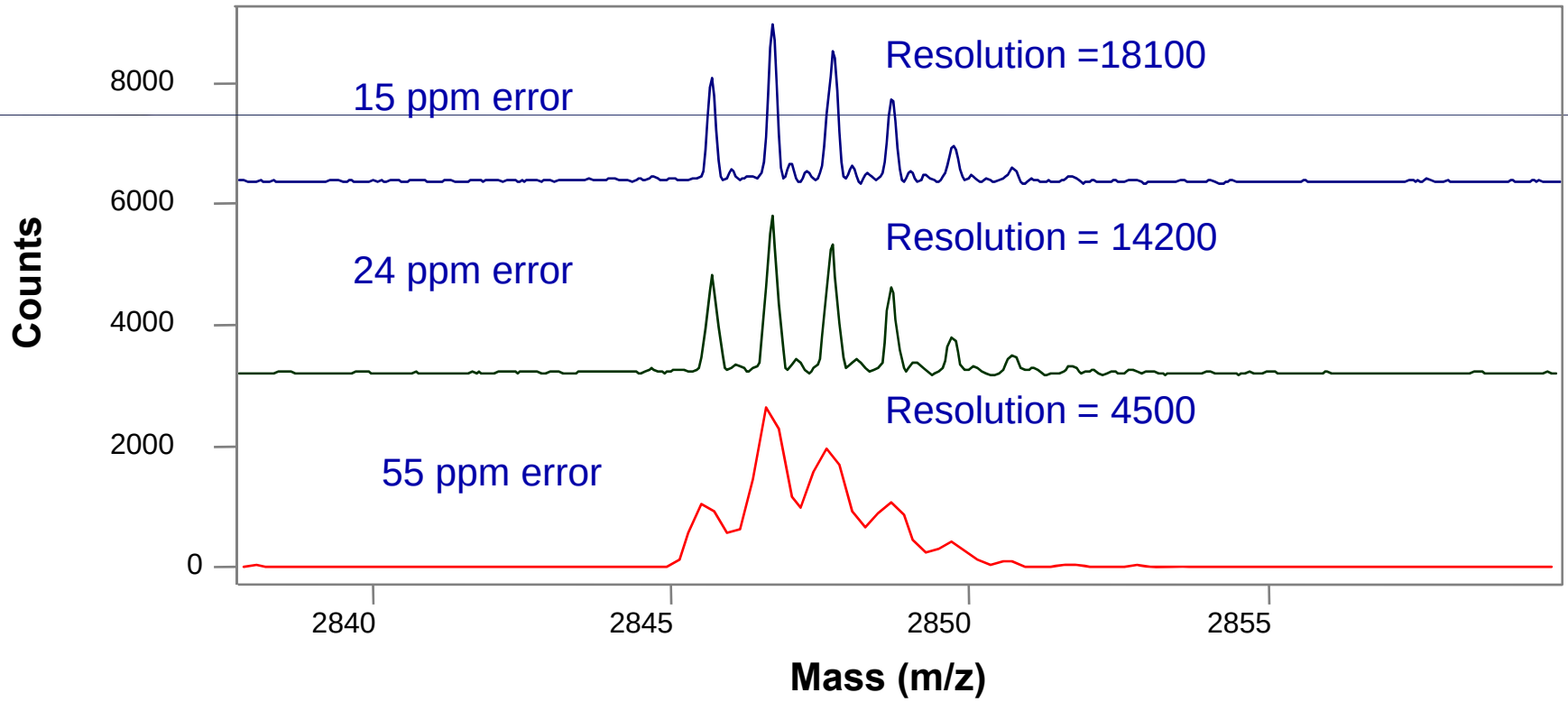


How is mass resolution calculated?



Mass measurement accuracy depends on resolution

High resolution means better mass accuracy



How do we achieve superior mass resolution?

Reflector TOF Mass Analyzer

Delayed Extraction on a MALDI source

Measure frequency

(FT-ICR MS, Orbitrap MS)

Important performance factors

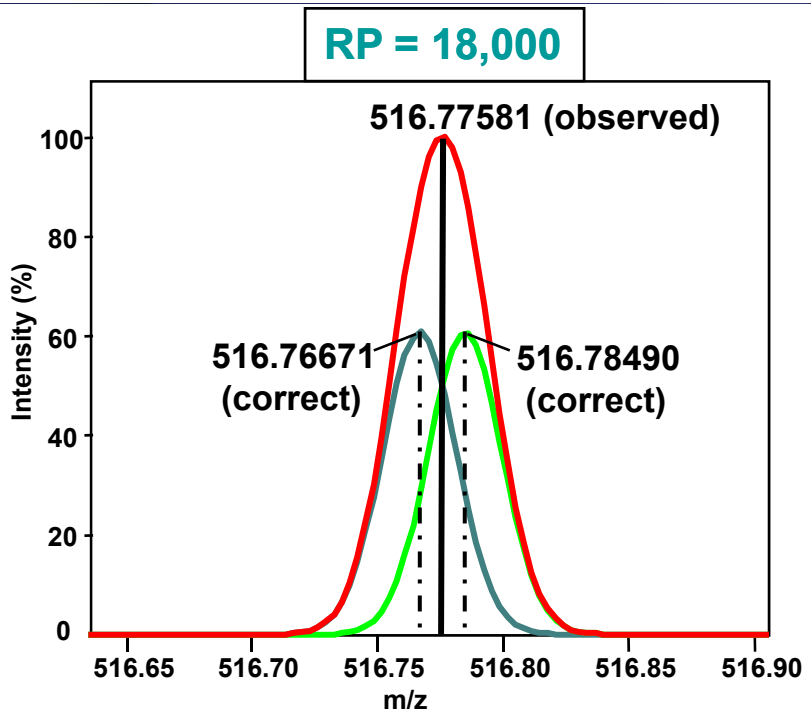
***Mass accuracy*: How accurate is the mass measurement?**

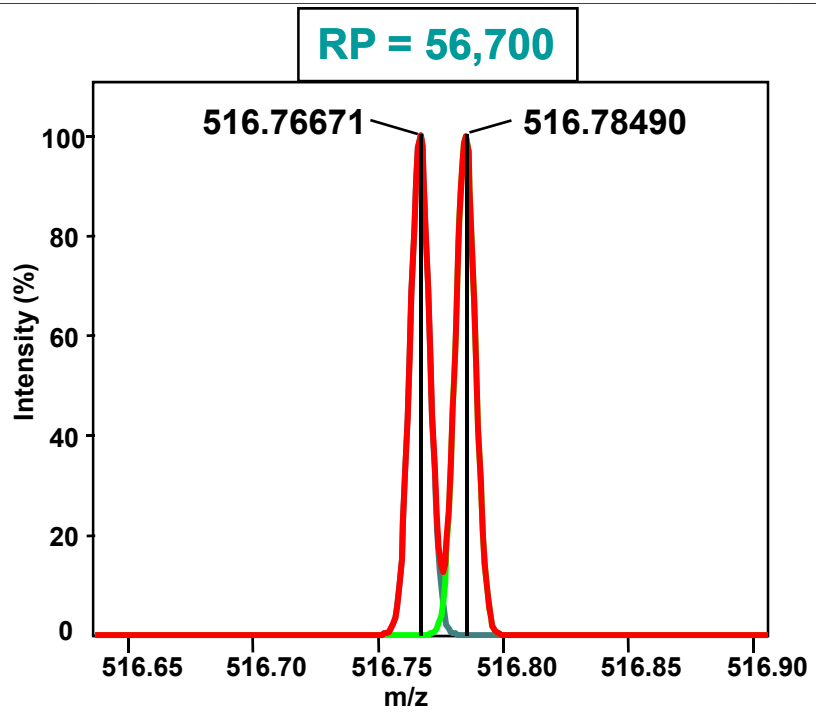
***Resolution:* How well separated are the peaks from each other?**

***Sensitivity:* How small an amount can be analyzed?**

Two peptides - same nominal mass - simulation

Peptide mixture:	**[Val^5]-Angiotensin II**	**Lys-des-Arg^9-Bradykinin**
Sequence:	**DRVYVHPF**	**KRPPGFSPF**
Formula:	**$C_{49}H_{69}N_{13}O_{12}$**	**$C_{50}H_{73}N_{13}O_{11}$**
Exact mass:	**$[M+2H]^{2+}$ = 516.76671**	**$[M+2H]^{2+}$ = 516.78490**
Δm (mmu):	**18.2 mmu**	



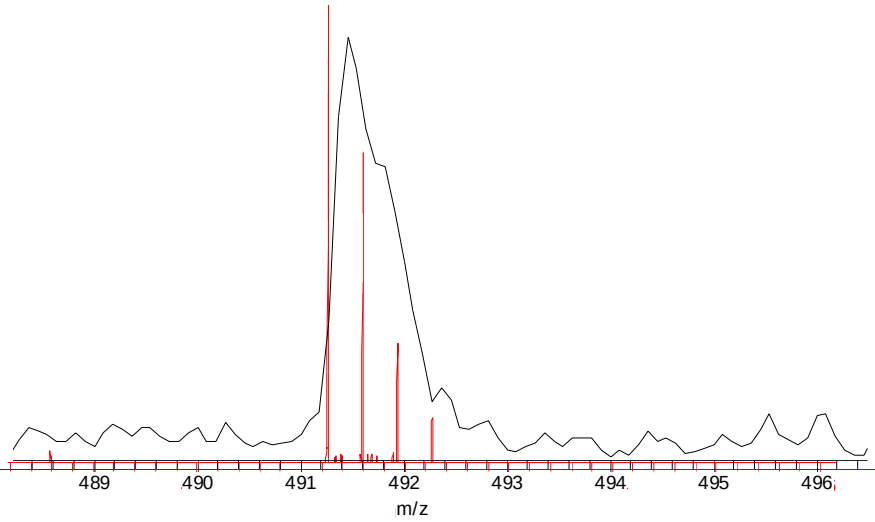


Is Mass Accuracy Important ?

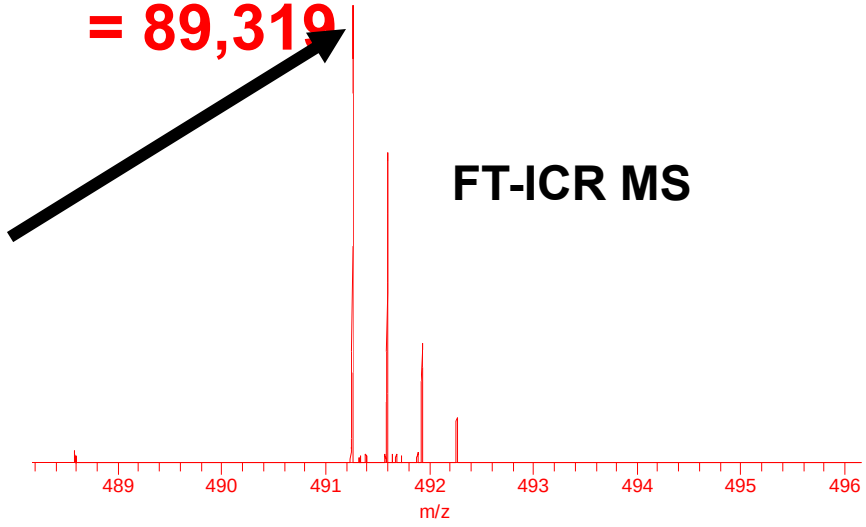
Results for error limit up to 5 ppm

	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}

Mass Resulution = m / Δm $_{50\%}$

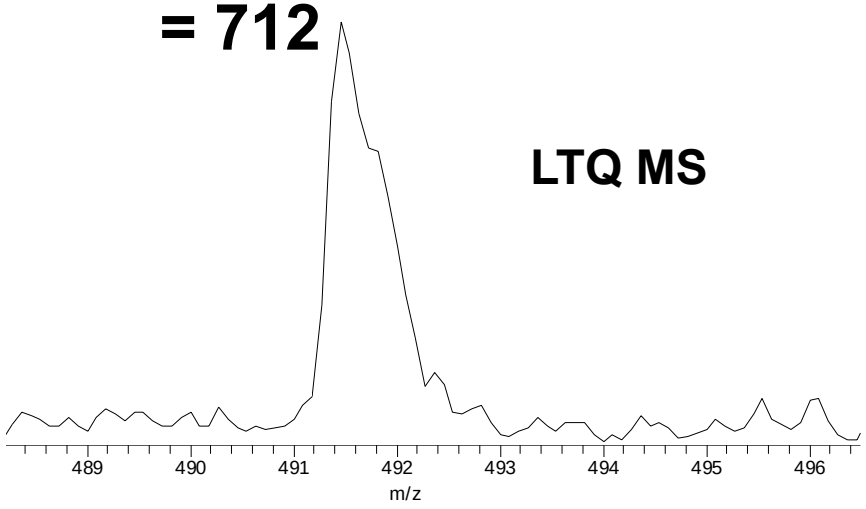


RP = 491.2594 / 0.0055 amu
= 89,319



FT-ICR MS

RP = 491.45 / 0.69 amu
= 712



LTQ MS

Objectives of the Lecture

1. Make ions
   - ESI, MALDI
2. Separate/Analyze/Detect ions
   - Tof, ion trap, quadrupole,
   - FT-ICR, Orbitrap
   - Electron multipliers
3. What is mass resolution and mass accuracy?

Slide Acknowledgements

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David Agard group

http://www.msg.ucsf.edu/agard/

(Univ. of Calif Sanfrancisco)