## **Lipids Analysis**

Stephen Barnes 3-5-15

## **Lipids**

- Lipids are mostly very hydrophobic
- Most are conjugates of fatty acids of a variety of chain lengths, which have different degrees of unsaturation, cis-trans isomers, and chiral centers
- The conjugating frame to which the fatty acids binds can be quite hydrophilic
- This results in a very wide (evergrowing) number of lipid species

## **Analysis of fatty acids**

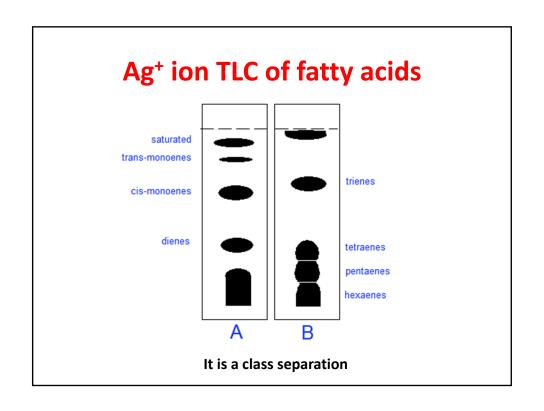
- Fractional crystallization
- Thin layer chromatography (TLC)
  - Argentation TLC (to separate according to number of double bonds)
- Gas liquid chromatography
  - Packed columns
  - Capillary columns
- LC-MS
- SWATH-MS
- Differential ion mobility
- DESI-MS

## **Fractional crystallization**

- Still used in industry
- Crystallization is used to determine whether adulteration of butter fat by other lower quality fats has occurred
- Unsaturated fats are more soluble at lower temperatures
  - Division into "stearins" and "oleins"
  - For fatty acids, make lead salts and cool in diethyl ether or ethanol – the saturated FAs crystallize out first

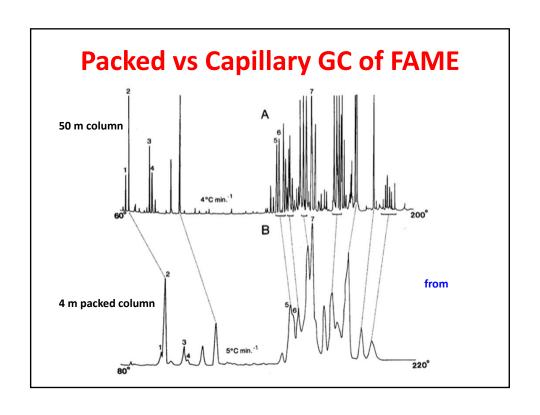
## Thin-layer chromatography

- Fatty acids or methylated fatty acids separated on alumina or silica gel TLC
- When  $AgNO_3$  is incorporated into the silica slurry before making the TLC plate, the observed separation is dependent on the degree of unsaturation ( $\pi$ -bonding)
  - Saturated
  - Mono-unsaturated
  - Di-unsaturated, etc.



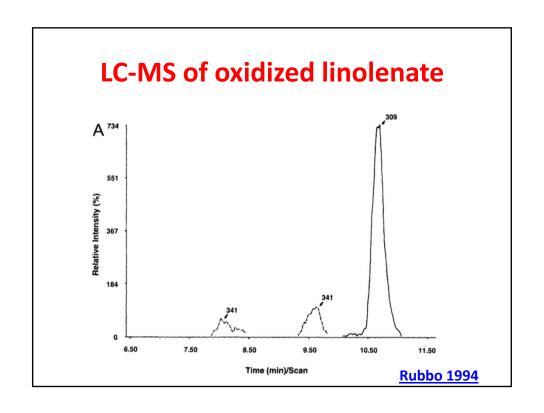
## **Gas-liquid chromatography**

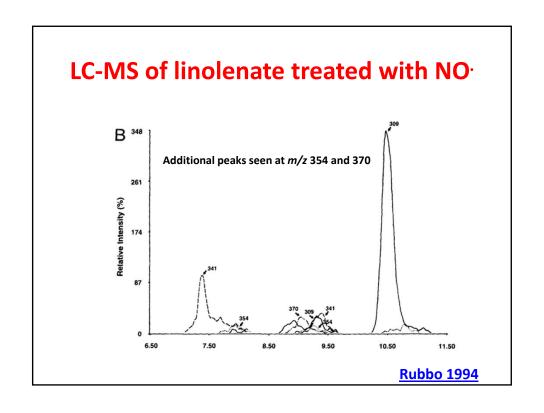
- 1952 Martin and James start GC by separating volatile fatty acids (C<sub>1</sub>-C<sub>6</sub>)
  - Quickly extended it to long chain FAs by methylating them
  - 5-6 feet x ¼ inch glass or stainless steel packed columns
- 1955 Patent for capillary, open tubular columns awarded
  - Did not enter commercial use until the mid-1970s

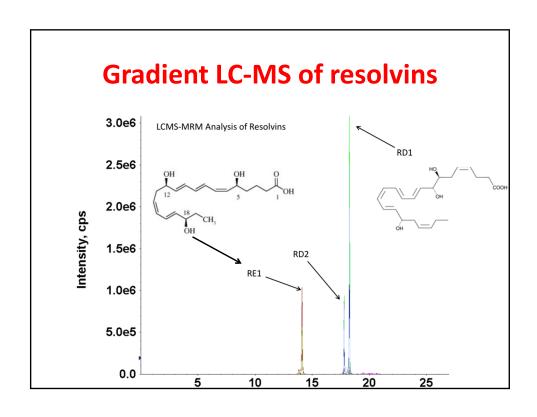


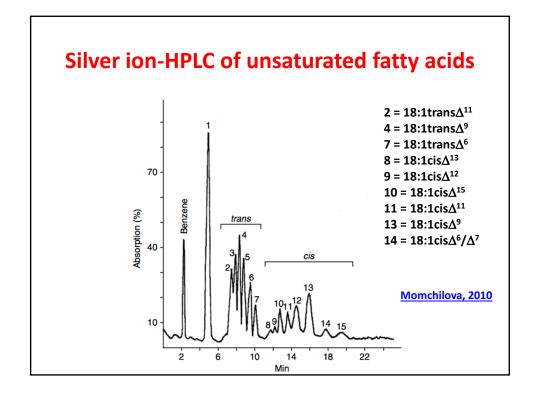
# (HP)LC

- Reverse-phase LC
  - Can be used for lipid class separation based on hydrophobicity
  - Again, Ag<sup>+</sup> can be introduced into the medium to enhance the separation of unsaturated fatty acids
  - Very difficult to detect lipids spectroscopically
  - LC-MS is the preferred method









## **Modern lipidomics**

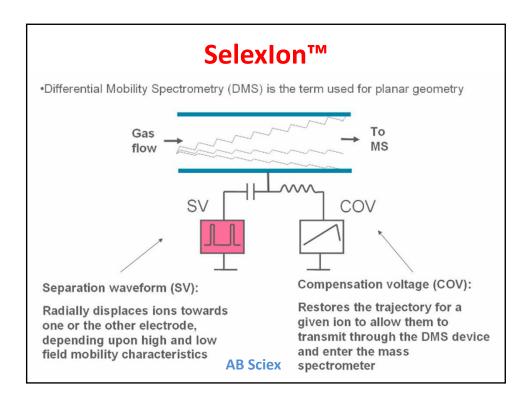
- Use of the SWATH-MS approach
- Preceded by total lipid extraction using a two-phase partition by adding CHCl<sub>3</sub>:MeOH
  - Bligh-Dyer and Folch extractions
    - Crucial to do so in an atmosphere of argon and in the presence of butylated hydroxytoluene to prevent oxidation

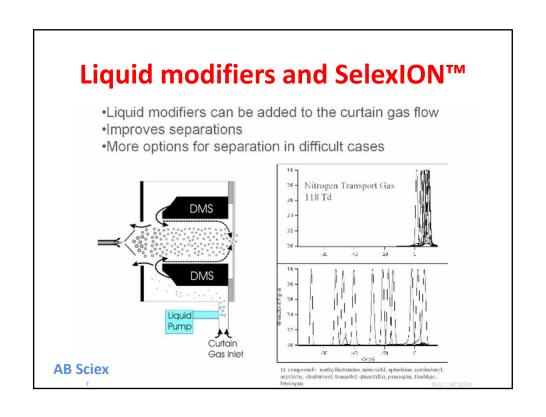
#### **SWATH-MS**

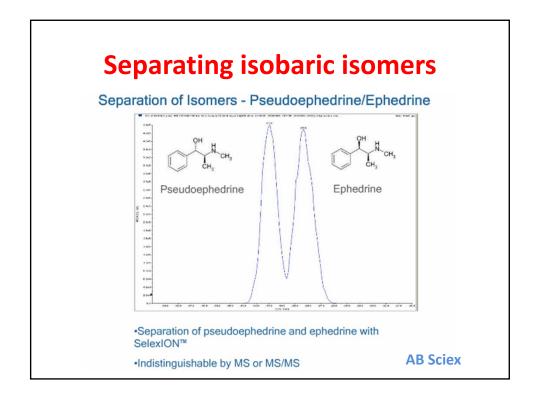
- Based on an infusion strategy on a 5600
  TripleTOF
  - lons are filtered 1.2 m/z at a time in the quadrupole over a m/z range of 200-1200
  - The filtered ions are collisionally dissociated and fragment ions analyzed by the TOF analyzer
  - MSMS spectra collected for 500 msec for each m/z, i.e., infusion for 500 sec (8.33 min)

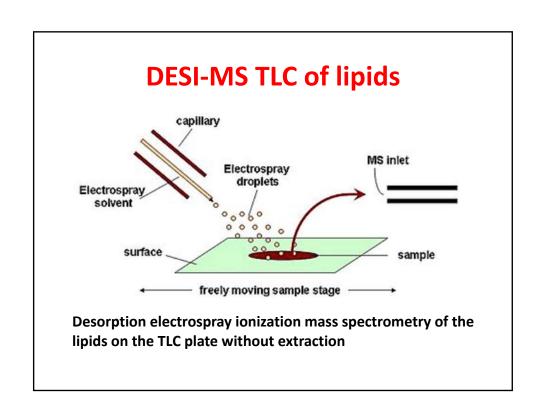
## Other MS methods for lipids

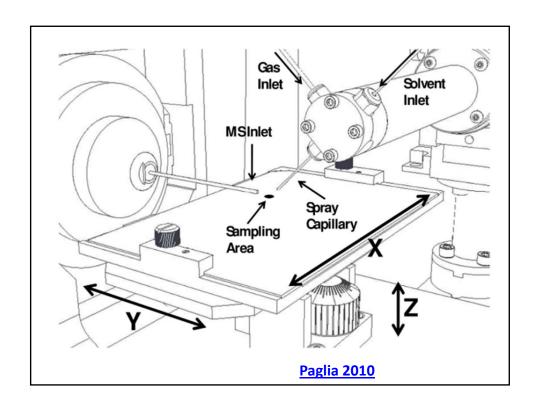
- SWATH-MS is comprehensive no stone unturned
- However, many lipids overlap in mass and there are also isomers with the same mass
- To observe more individual lipids, it is necessary to resolve lipids before analyzing them in the mass spectrometry
- Even then, isomers can be a problem
- A form of ion mobility may be the answer to this

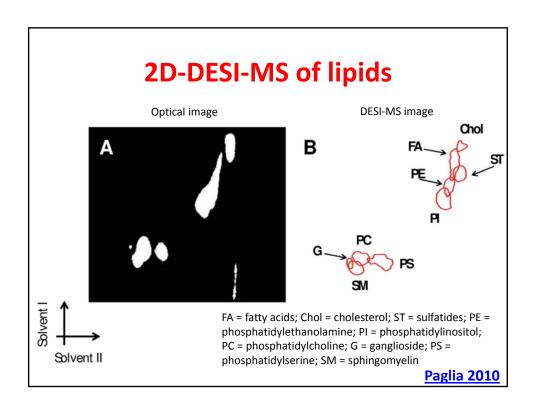












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