

**UAB**  
THE UNIVERSITY OF  
ALABAMA AT BIRMINGHAM  
Knowledge that will change your world

## Introduction to metabolomics research

Stephen Barnes, PhD  
University of Alabama at Birmingham  
[sbarnes@uab.edu](mailto:sbarnes@uab.edu)  
934-7117; MCLM 452\*

**T**argeted  
**M**etabolomics &  
**P**roteomics  
**L**aboratory

\*Will change in March to Bevill 709

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## Course goals

1. To understand the **vital** roles of small molecules/metabolites
  - To provide energy for the chemical and enzymatic processes of life
  - To provide the building blocks for the macromolecules (DNA, RNA, proteins, carbohydrates, lipids)
  - As co-factors
  - As signaling molecules
  - As biomarkers for disease

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## Course goals

### 2. To understand the **origins** of metabolites

- Produced by (human) cells
- Produced by **the things that we eat (the food-ome)**
  - Plants (wheat, corn)
  - Fruits (apples, oranges, strawberries)
  - Vegetables (rice, potatoes, broccoli, peas)
  - Dairy products, including fermented forms
  - Meat from other animals
  - Xenobiotics
- Produced by **microorganisms** in our bodies
- Therapeutics, smoking, household chemicals

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## Course goals

### 3. To understand that a metabolomics experiment is **high dimensional**

- i.e., it compares the intensities of hundreds, if not thousands, of distinct species
- Very important statistical consequences
- Cannot afford to do a robust experiment that fully satisfies theoretical statistical principles
- Very important to sit down with a statistician prior to executing an experiment

Dr. Hemant Tiwari

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## Course goals

### 4. To select the appropriate method for extracting/recovering metabolites

- **Metabolites encompass an enormous range of chemistries**
  - Gaseous (H<sub>2</sub>, H<sub>2</sub>S)
  - Volatile (butyric acid, acetone, skatole)
  - Hydrophilic (glucose)
  - Charged-positive/negative (amino acids, nucleotides, organic acids, amines)
  - Hydrophobic (lipids, steroids, hydrocarbons)
- **No single method suitable for all metabolites**

Dr. Prasain

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## Course goals

### 5. Selecting the analytical approach

- ***In situ* analysis**
  - Laser ablation of frozen tissue
  - Other desorption methods
  - Magic angle spinning NMR
  - Other spectroscopic methods
- **Extracted samples**
  - NMR
  - GC-MS (1- and 2D chromatography and MSMS)
  - LC-MS (1- and 2D chromatography and MSMS)
  - CE-MS
- **Targeted vs untargeted analysis**

Drs. Placzek and Barnes

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## Course goals

### 6. Analysis of the data

#### – Data alignment

- NMR methods
- LC-MS and GC-MS methods (XCMS; ADAP)

#### – Statistical evaluation

- Univariate and multivariate analysis (MetaboAnalyst)
- XCMSonline
- Peaks to Pathways (Metaboanalyst)

#### – Data visualization

- XCMSonline
- MZmine

Dr. Barnes

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## Course goals

### 7. Identifying metabolites

#### – Use of MS (absolute mass)

- METLIN
- Peaks to Pathways
- ChemSpider

#### – MSMS (fragmentation spectra)

- METLIN
- MS-DIAL

#### – Metabolite standards

#### – Importance of retention time

- Multiple column conditions

Dr. Barnes and Prasain

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## Course goals

### 8. Pathways and applications

- Peaks to Pathways/Metaboanalyst
- KEGG pathway mapping
- Applications to:
  - Adverse cardiovascular risk
  - Diabetes
  - Lens and kidney diseases
  - Cancer

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## What is “Metabolomics”?

- Metabolomics is like other types of –omics analysis (microarray, RNA-Seq, proteomics, etc.)
  - Offers a “comprehensive” view of all detectable chemicals (not just metabolites)
  - Can be applied to body fluids
    - Plasma/sera, urine, saliva, tears, fecal water, etc.
  - Also to tissues
    - Liver, lung, heart, kidney, brain, eyes, etc.
  - And to single cells
    - Human, rodent, yeast, bacteria, etc.

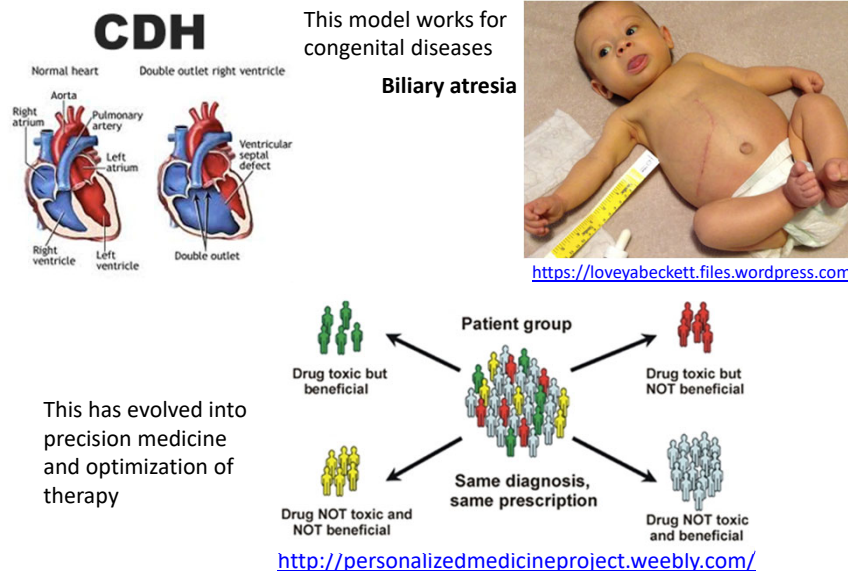
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## Defining who we are chemically

- Does an understanding of the functions of human genes define the chemical make up of our body fluids and tissues?
- How does metabolomics provide information on the circulating chemicals?
- Are the detected chemicals metabolites produced by human enzymes?
- So, what are we really exposed to? And does it make a difference?

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## A great deal of emphasis is being placed on the importance of DNA sequencing



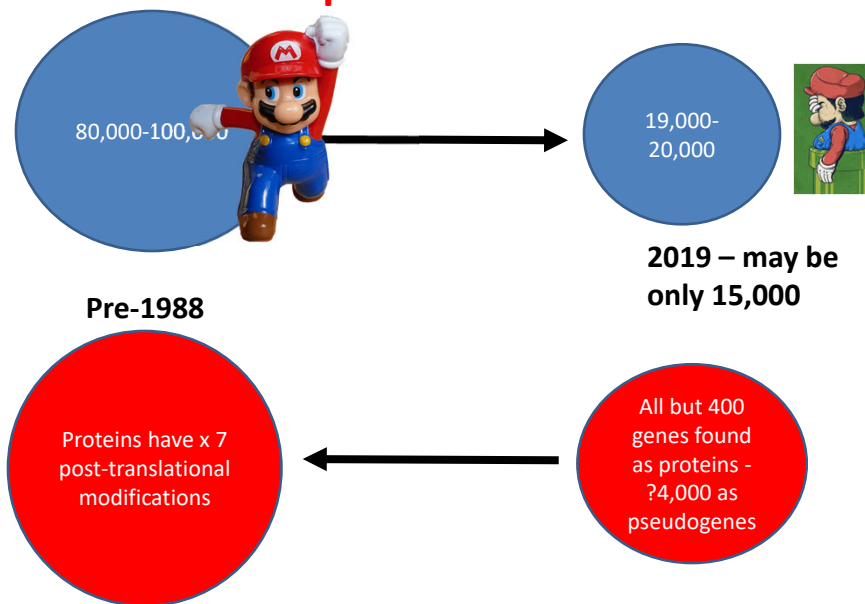
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## Metabolomics in the newborn

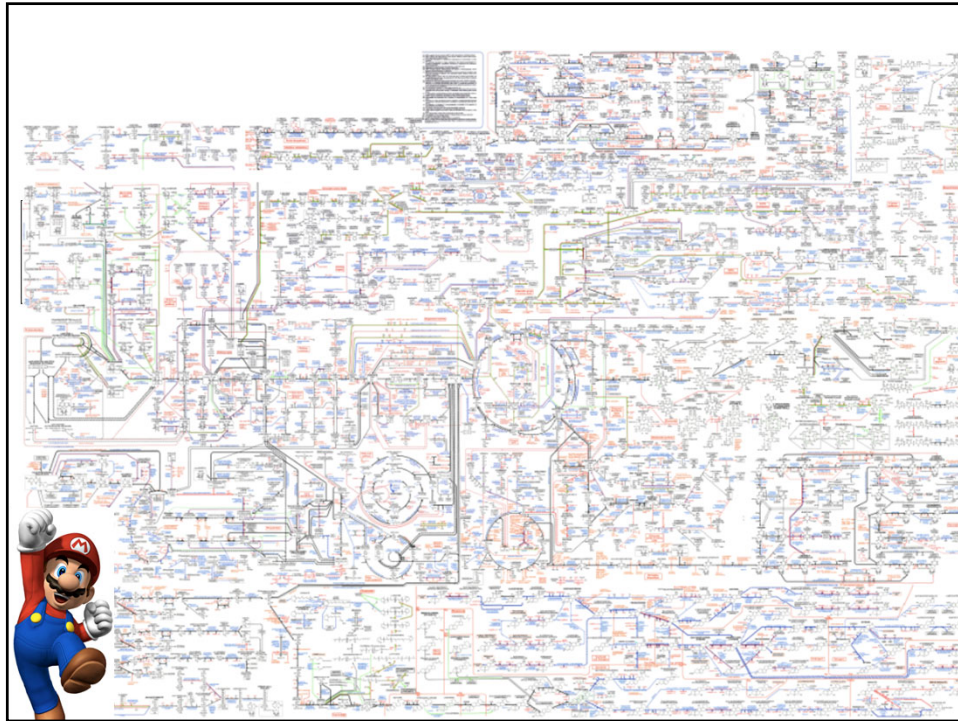
Dr. Dan Sharer

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### However, genes failed to meet expectations



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## The Undiagnosed Disease Network

Dr. Matthew Might



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## Where does the metabolome come from?

- It starts with what fixes CO<sub>2</sub> and N<sub>2</sub>



Trees convert CO<sub>2</sub> to organic compounds



Field of soybeans – they fix N<sub>2</sub> because of nitrogen-fixing bacteria in their root nodules

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## Plants have more genes than humans

- Why? Plants can't run away!!
- Instead, they have to practice chemical warfare to prevent attack by aphids and microorganisms
- Many plants are poisonous to us
- Understanding which plants were safe to eat, or were so if cooked, represented the rise of agriculture and civilization



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## Compounds in plants and fruits

- Carotenoids
- Many vitamins
- Polyphenols and anthocyanins
- Not made by human cells



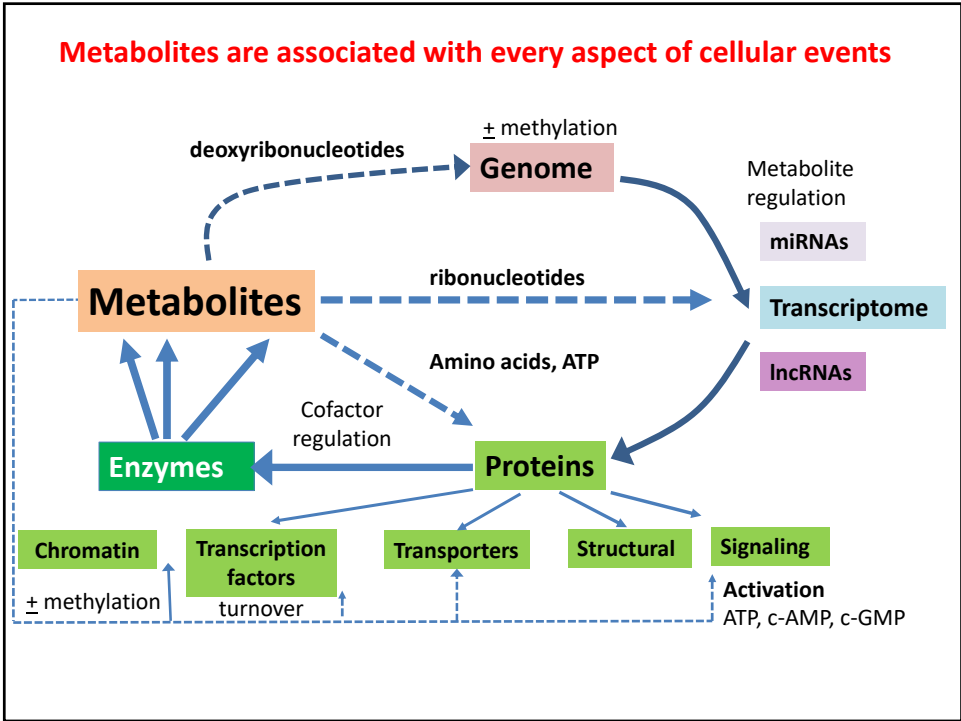
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## Other sources of body chemicals

- **The microbiomes**
  - Humans are not single organisms
  - Instead, we are super-organisms
  - The gut microbiome has 10 times the number of cells found in the rest of the (human) body
  - It makes novel compounds that are absorbed, enter the blood stream and tissues
- **Chemicals from the environment**
  - industrial contaminants, therapeutics, supplements
- **Interactions between the xenobiotics and the human enzyme systems**



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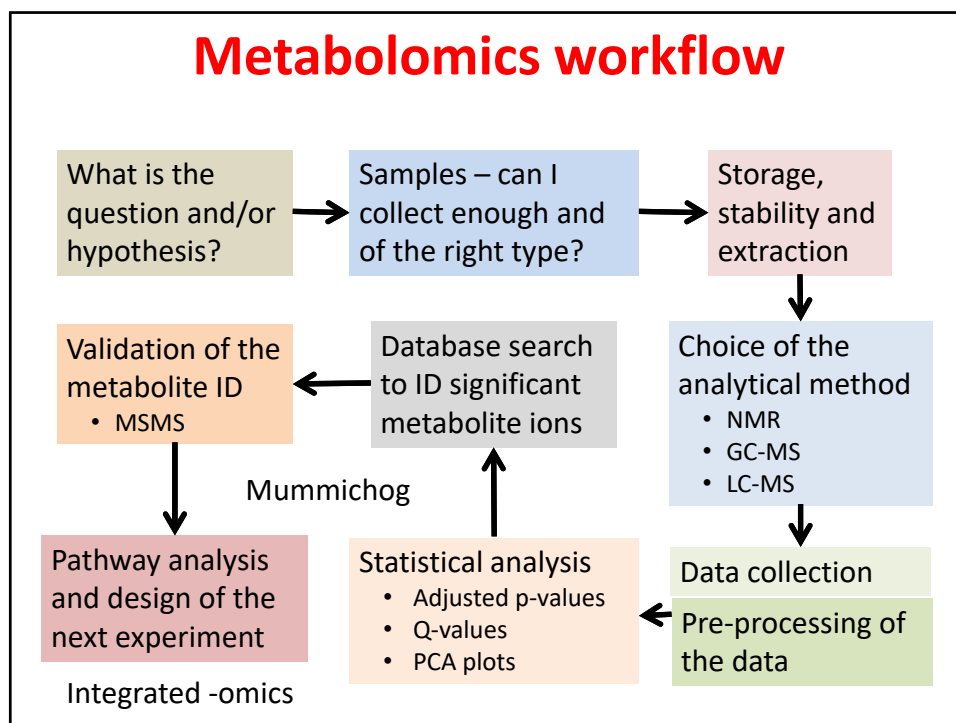
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## The metabolome is very complex!



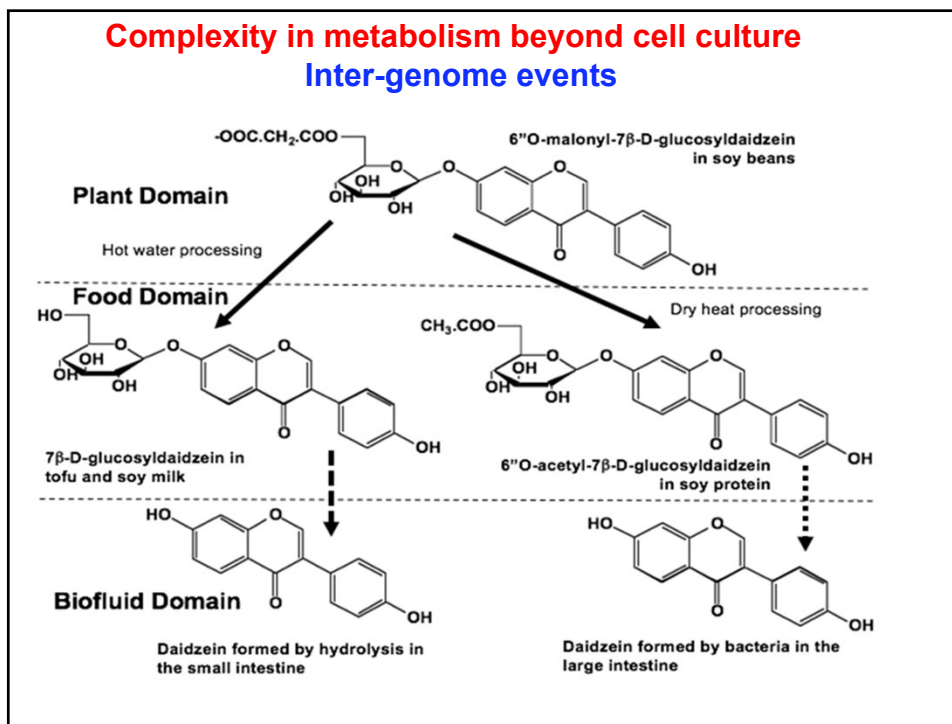
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## Metabolomics workflow



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**The Amazonian poison dart frog**

Their skin contains molecules like batrachotoxin which irreversibly poisons the Na<sup>+</sup>-channels

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## Two questions

**Why isn't the batrachotoxin a poison to the frog?**

ANSWER: The frog has mutations of three residues in the Na<sup>+</sup>-channel protein that prevent binding of the batrachotoxin

**Does the frog synthesize the toxin?**

ANSWER: It doesn't, it gets the toxin from what it eats – ants, beetles, etc.

So, it all depends on what you eat.

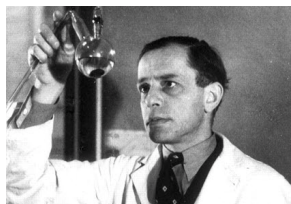
Dart frogs bred in captivity and fed a non-insect diet don't make batrachotoxin

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**Where did metabolomics come from?**

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## Transition of mass spectrometry to biology



Ralf Schoenheimer

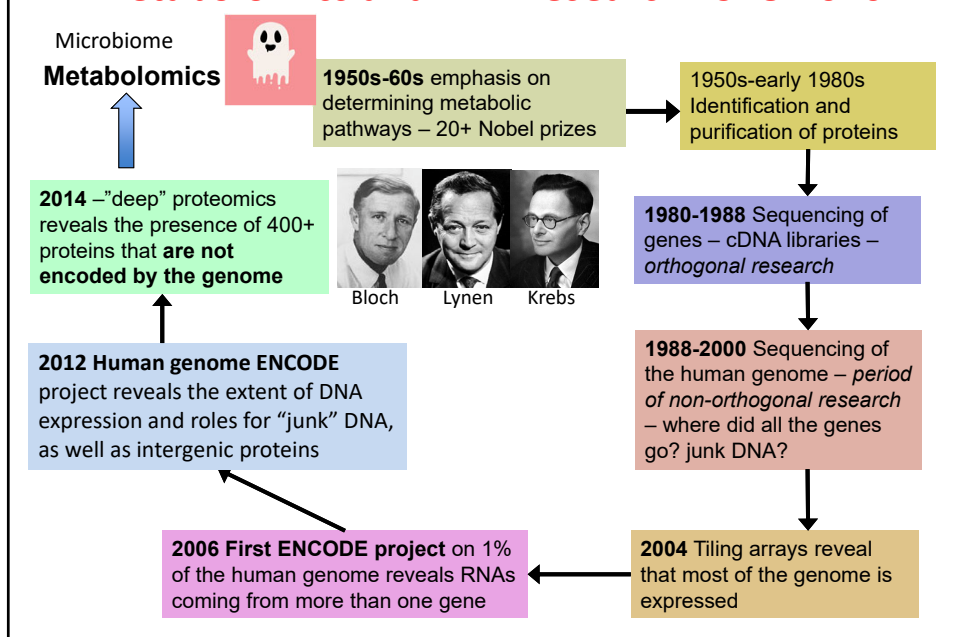


David Rittenberg

- While the politicians, tyrants, dictators and despots were salivating at the thought of developing nuclear weapons from unstable isotopes in the early part of the 20<sup>th</sup> Century, two scientists began the pursuit of the peaceful use of stable isotopes, initially deuterium (<sup>2</sup>H), and later carbon (<sup>13</sup>C) and nitrogen (<sup>15</sup>N), to study biochemical pathways
- Understanding the pathways of metabolism was born

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## Metabolomics and NIH Research 1948-2016



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## Progress in LC-MS

- Commercial HPLC appeared in the early 1970s to separate thermally stable and unstable molecules
- The challenge remained to find a way to get the unstable compounds into the gas phase
  - Applied to macromolecules (peptides, proteins) as well as metabolites
- Thermospray had some initial success
- **Electrospray ionization** and **chemical ionization** radically changed analysis, allowing compounds to go into the gas phase at atmospheric pressure and room temperature

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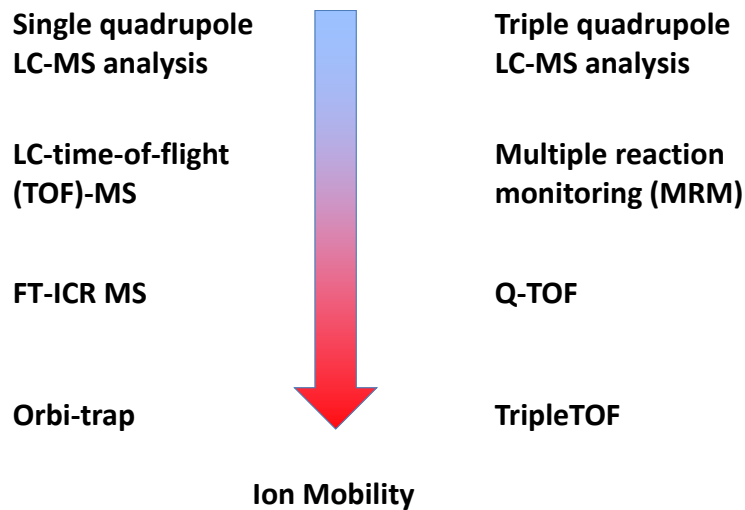
## LC-MS

- Suddenly, there were what appeared to be no limits (or very few) to what could be analyzed
- Unheard of, robust mass spectrometers came into play
  - “A reliable mass spectrometer” was considered in 1990 to be an oxymoron

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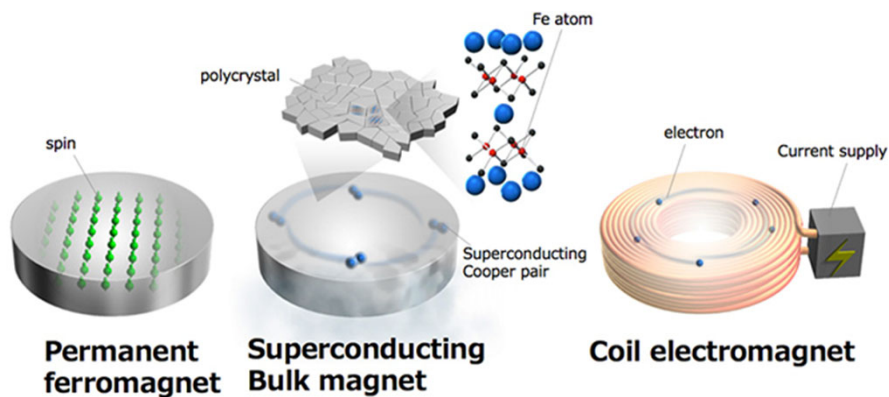


## Types of LC-MS analysis



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## NMR spectroscopy and metabolomics



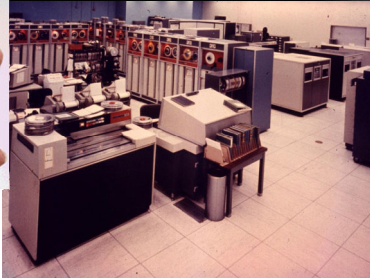
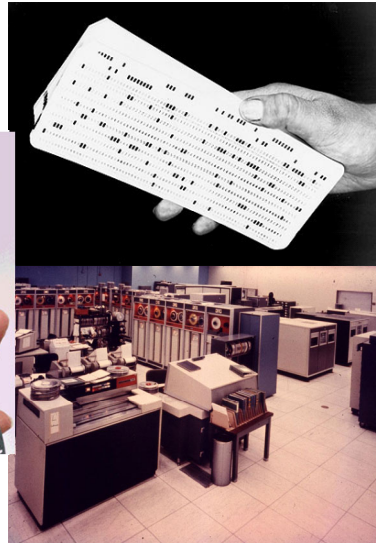
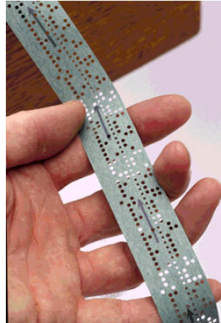
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NMR has had several critical development steps – Fourier Transform analysis of collected data, increase in field strength with superconducting magnets, micro-coil, cryogenic analysis, and hyperpolarization.

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## Changing times in Computing

- 1950 The Cambridge colleagues of Watson and Crick calculated the structure of DNA by putting data onto punched cards and taking them by train to London for analysis – and to the fog – the “cloud” in 1950s
- 1964 Seymour Cray develops the CDC 6600 (1 Mflops)
- 1967 I used paper tape to collect data from a radio gas chromatograph and then submitted them via a terminal reader to the CDC 6600 at the University of London



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## Today in Computing



### On my desk in 2019

- The Apple MacBook Air with 4 quad core Intel i7 processors
  - Operates at 2.0 GHz
  - Memory of 8 GB
    - Access 1.333 GHz
  - 512 GB Flash memory storage
  - 10 Gbs Thunderbolt I/O
- Also cost ~\$2,000




### Cheaha high-performance computing

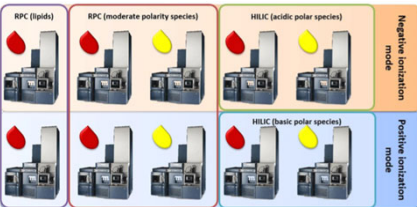
- Initially IBM Blue-Genie operating at 4.733 Tflop/s
- Replaced by Cheaha, in its current configuration it has 2800 conventional CPU cores and 6.6 PB raw storage
- It operates at 468 Tflop/s (max)

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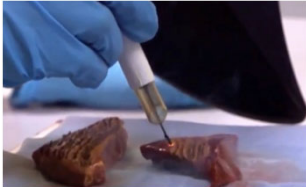
### MRC-NIHR National Phenome Centre



**600 MHz NMR instruments  
in surgical suite**



**Mass spectrometers (10 Q-TOFs) each  
dedicated to one assay format**



**lknife - revolutionizing surgery**

**This is Next-GEN precise medicine**

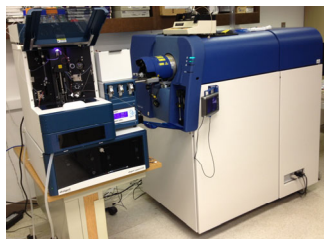
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### The UK National Phenome Center, LC-MS labs



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## UAB capabilities in metabolomics

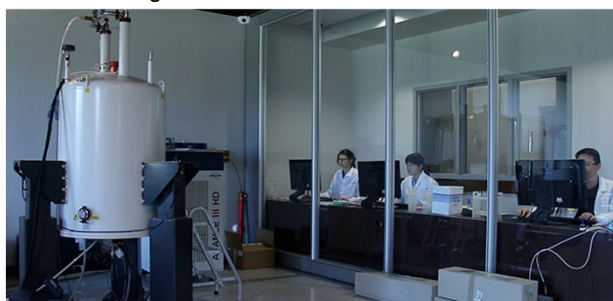


SCIEX 5600 TripleTOF  
with Eksigent nanoLC

TMPL mass spec lab  
MCLM 459/427  
Stephen Barnes, Director  
205-934-7117/3462



SCIEX 6500 Qtrap with SelexION



Central Alabama NMR facility  
Chemistry Bdg  
William Placzek, Director  
205-934-2465

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## Great challenges in metabolomics

- **The extent of the metabolome**
  - From gaseous hydrogen to earwax
- **Having complete databases**
  - METLIN has over 1 million metabolite records, but your problem always creates a need to have more
  - Improvement in the size of a MSMS database
- **Storing and processing TBs of data**
- **Standards and standard operating procedures**
- **Being able to do the analyses in real time**

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## NIH Common Fund Metabolomics Program

- **Metabolomics Workbench:** <http://www.metabolomicsworkbench.org/>
- **Regional Comprehensive Metabolomics Research Centers**
  - University of Michigan: <http://mrc2.umich.edu/index.php>
  - UC Davis Metabolomics Center: <http://metabolomics.ucdavis.edu/>
  - UNC-CH: <http://www.uncnri.org/wp-content/uploads/2016/12/NIHERCMRC.pdf>
  - SE Center for Integrated Metabolomics: <http://secim.ufl.edu/>
  - Resource Center for Stable Isotope Metabolomics: <http://bioinformatics.cesb.uky.edu/bin/view/RCSIRM/>
  - Mayo Clinic Metabolomics Resource: <http://www.mayo.edu/research/core-resources/metabolomics-resource-core/overview>
- **Other resources**
  - See this [link](#)