

## Introduction to metabolomics research

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

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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; MS-DIAL)

#### – 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 (IROA kit)

#### – 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
- Integration with other –Omics
- Machine learning/Artificial Intelligence

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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.

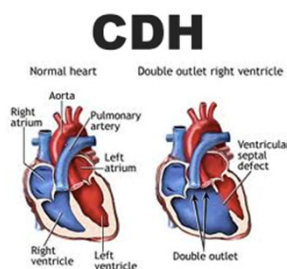
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## Defining who we are chemically

- 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 has been placed on the importance of DNA sequencing



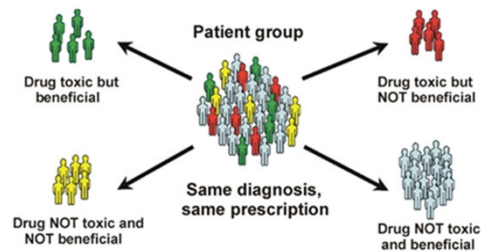
This model works for congenital diseases

**Biliary atresia**



<https://loveyabeckett.files.wordpress.com>

This has evolved into precision medicine and optimization of therapy



<http://personalizedmedicineproject.weebly.com/>

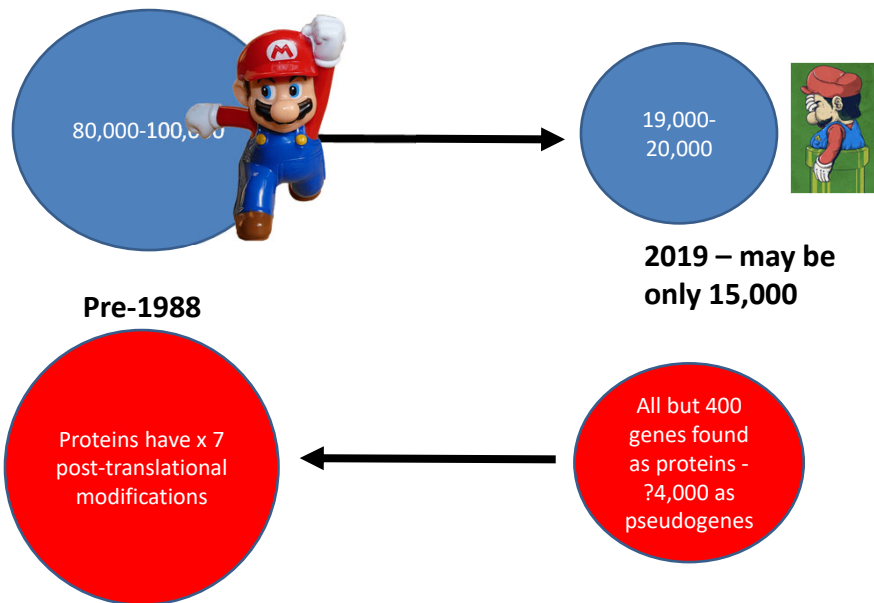
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## Metabolomics in the newborn

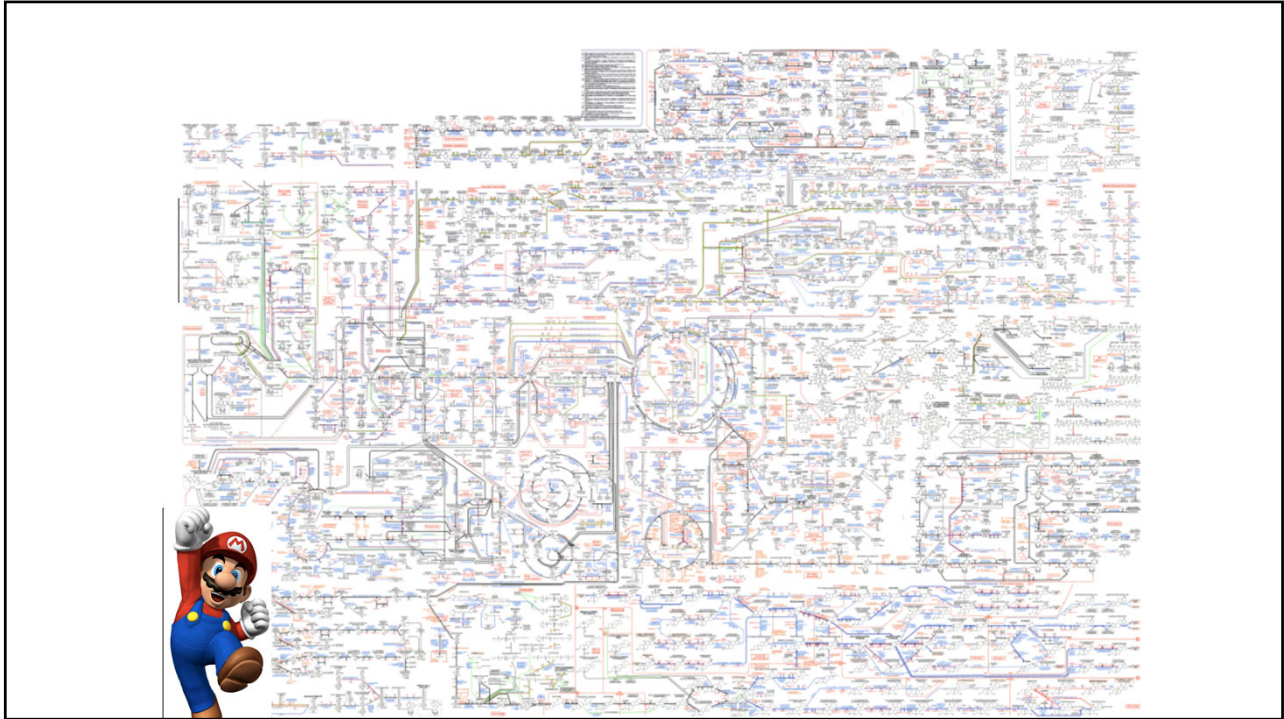
Dr. Dan Sharer

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### However, genes failed to meet full expectations



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## The Precision Medicine Institute

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



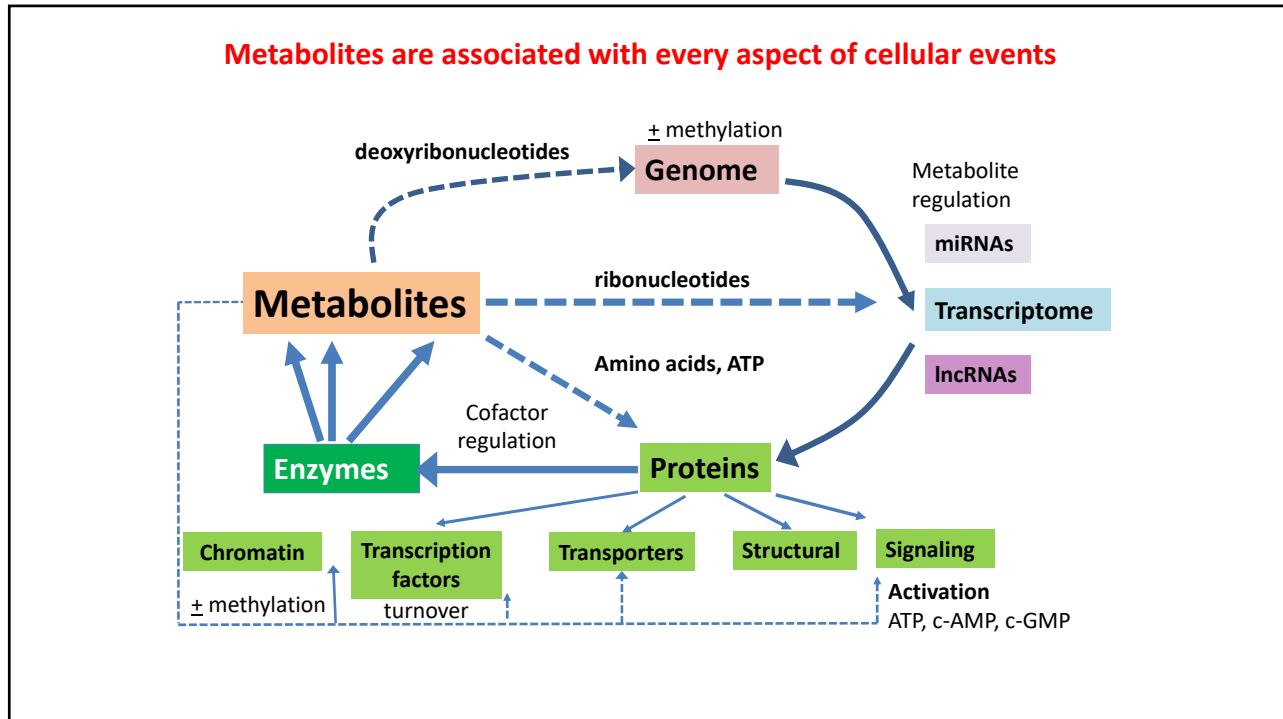
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## Other sources of body chemicals

- **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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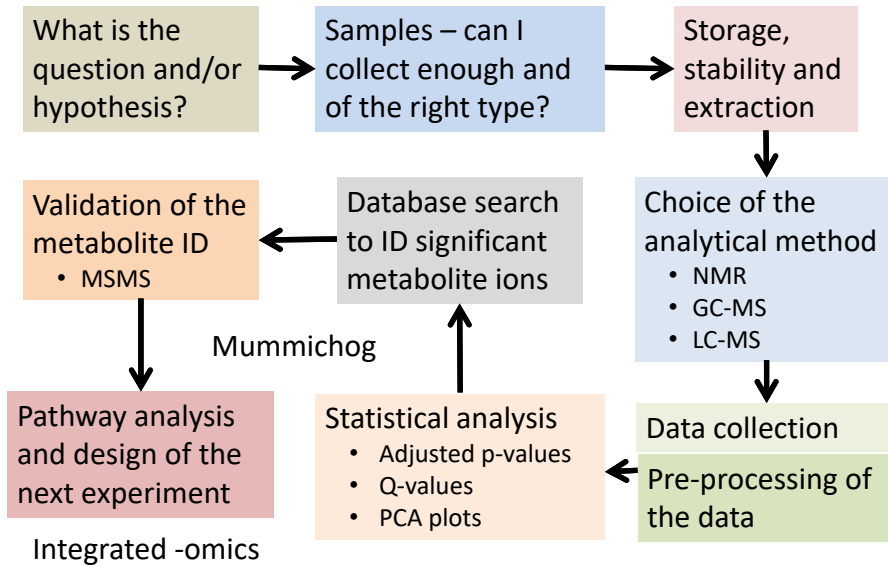
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# The metabolome is very complex!



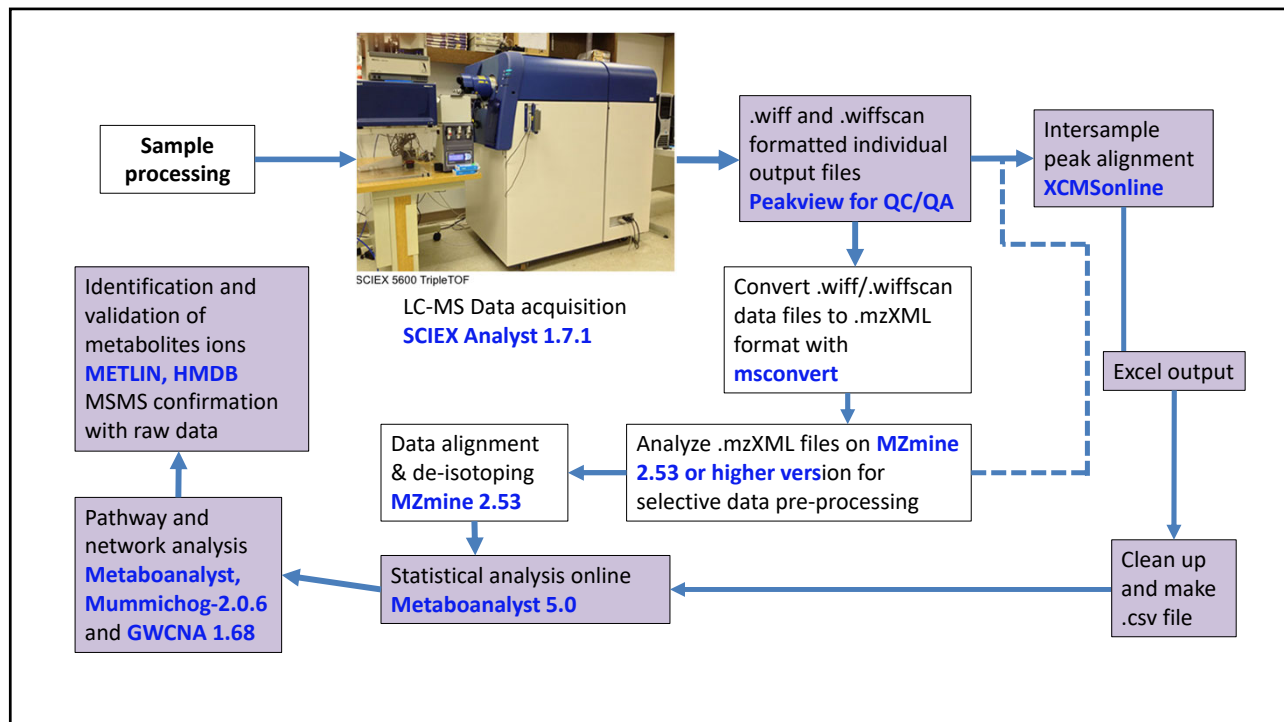
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## Metabolomics workflow

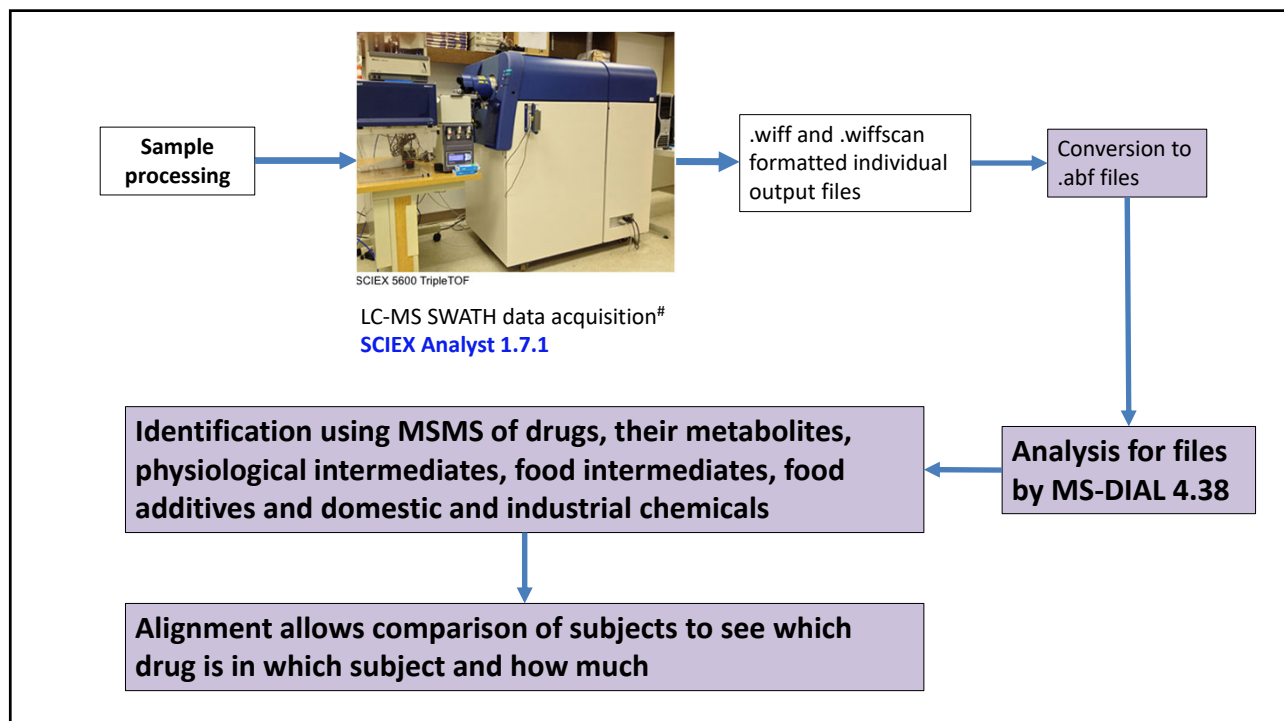


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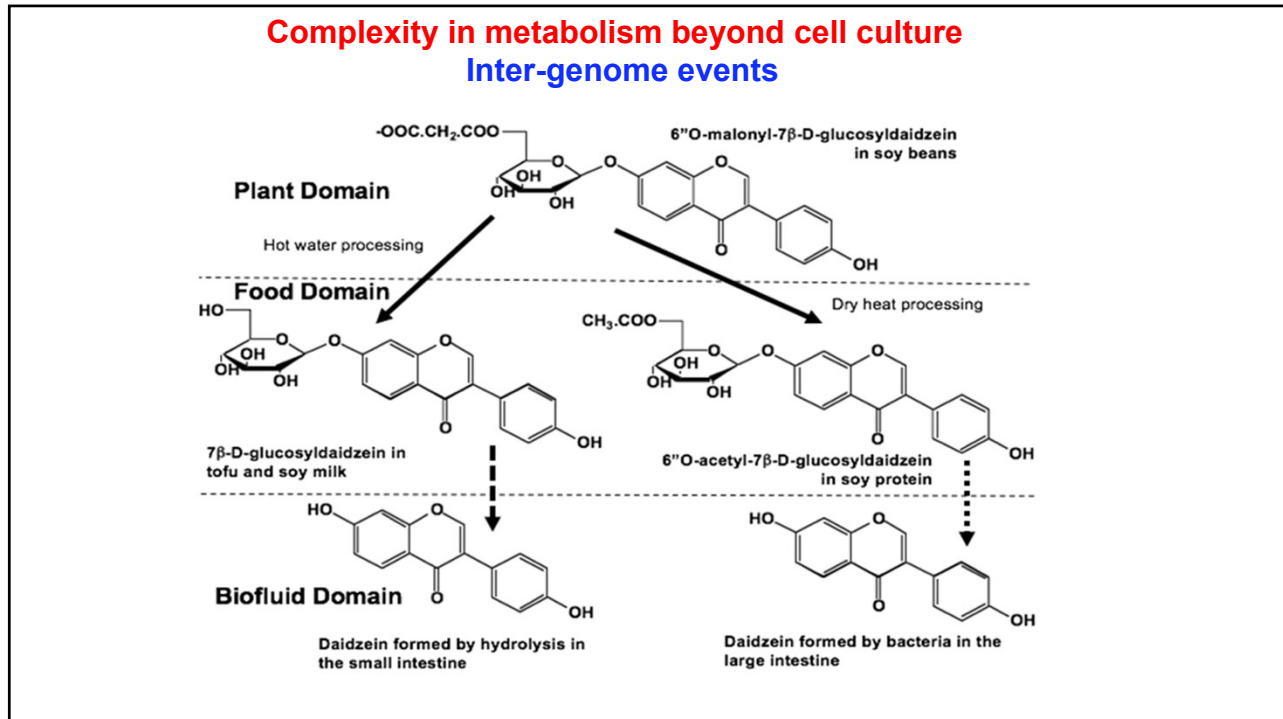


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## Complexity in metabolism beyond cell culture Inter-genome events

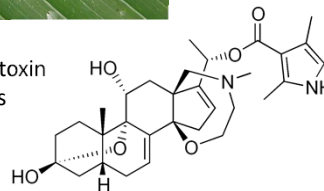


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## The Amazonian poison dart frog



Their skin contains molecules like batrachotoxin which irreversibly poisons the  $\text{Na}^+$ -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

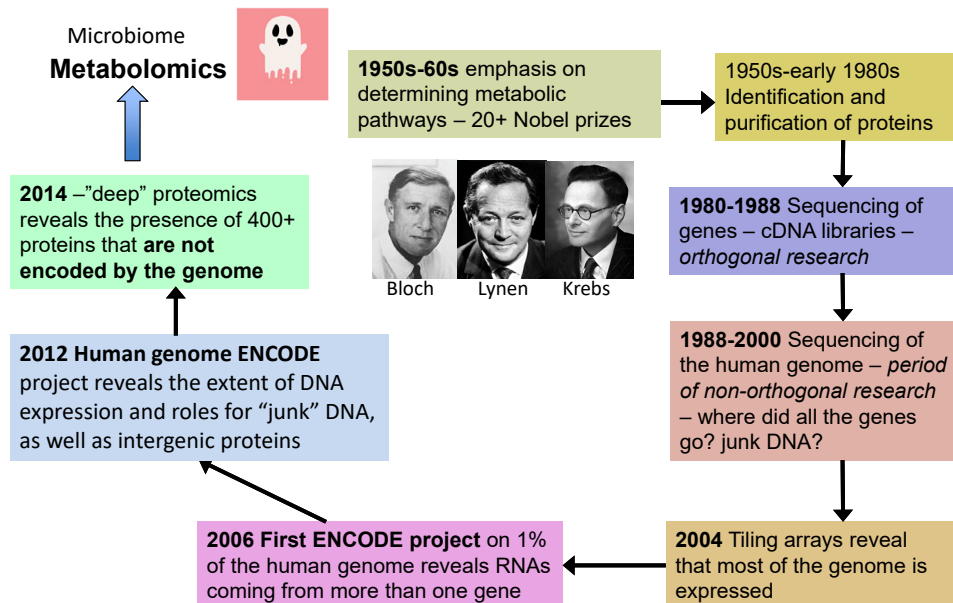


David Rittenberg

- 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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## Structure of NIH Research

Individual Research institutes (27)  
NIDDK, NHLBI, NIAID, etc.

National Centers  
CIT, CSR, FIC, NCCIH, NCRR, NIHCC

Office of the Director

2005 Review by Congress

Division of Program Coordination, Planning, and Strategic Initiatives, DPCPSI

NIH Common Fund

NIH Council of Councils

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NIH National Institutes of Health  
Office of Nutrition Research

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- Get the latest public health information from CDC »
- Get the latest research information from NIH »
- NIH staff guidance on coronavirus (NIH Only) »

**ONR**  
Office of  
Nutrition  
Research

DPCPSI - ONR

## Office of Nutrition Research (ONR)

*Advancing nutrition science to promote health and reduce the burden of diet-related diseases*

The Office of Nutrition Research:

- Advises the National Institutes of Health (NIH) Director, Division of Program Coordination, Planning, and Strategic Initiatives (DPCPSI) Director, and other key officials on matters relating to research on nutrition
- Coordinates implementation of the Strategic Plan for NIH Nutrition Research
- Coordinates research projects in nutrition science conducted or supported by the NIH Institutes and Centers (IC)
- Identifies research projects that deserve expanded effort and support by the ICs
- Develops, leads, and manages trans-NIH nutrition research projects in cooperation with the ICs
- Represents the NIH on intradepartmental or interagency committees on nutrition research and related policy issues

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## Technologies in Metabolomics

- **Gas-chromatography-mass spectrometry (GC-MS)**
- **Liquid chromatography-mass spectrometry (LC-MS)**
- **Capillary electrophoresis-mass spectrometry (CE-MS)**
- **Nuclear magnetic resonance (NMR)**

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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

Single quadrupole LC-MS analysis

LC-time-of-flight (TOF)-MS

FT-ICR MS

Orbi-trap

Triple quadrupole LC-MS analysis

Multiple reaction monitoring (MRM)

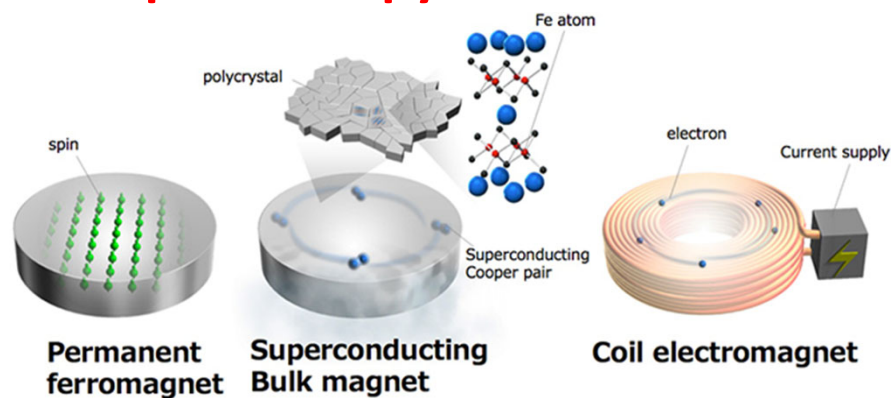
Q-TOF

TripleTOF

Ion Mobility

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## NMR spectroscopy and metabolomics



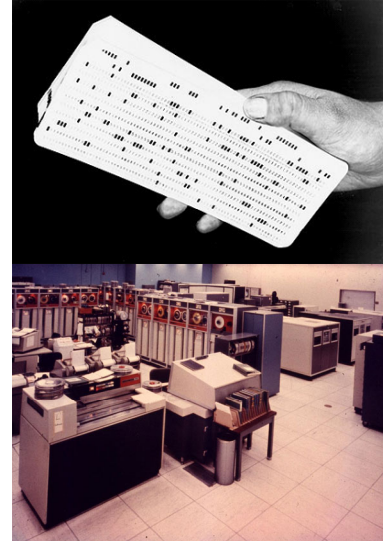
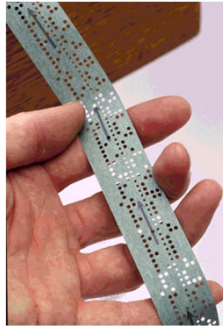
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NMR has had several critical development steps – Fourier Transform analysis of collected data, increase in field strength with superconducting magnets, micro-coil, cryogenic analysis, and hyperpolarization.

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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 2021

- 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
  - 100 Gbs Thunderbolt I/O
- Also cost ~\$2,000



### Cheaha high-performance computing

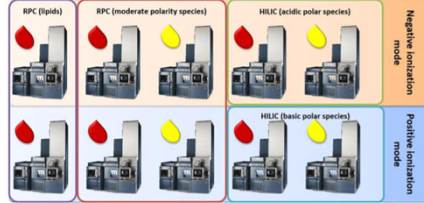
- Initially IBM Blue-Gene operating at 4.733 Tflop/s
- Replaced by Cheaha, in its current configuration it has 2800 conventional CPU cores and 6.6 PB raw storage
- It operates at 468 Tflop/s (max)

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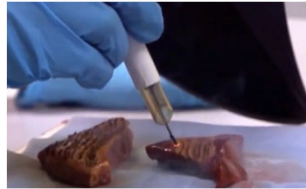
## MRC-NIHR National Phenome Centre



600 MHz NMR instruments in surgical suite



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



Iknife - revolutionizing surgery

This is Next-GEN precise medicine

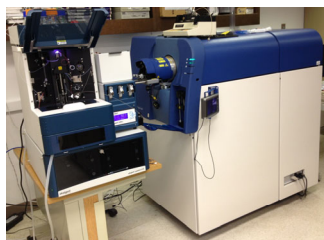
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## The UK National Phenome Center, LC-MS labs



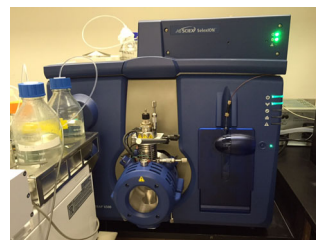
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## UAB capabilities in metabolomics

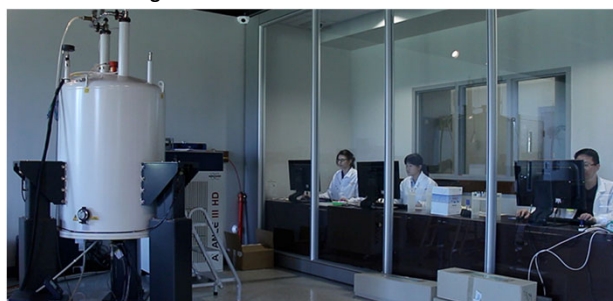


SCIEX 5600 TripleTOF  
with Eksigent nanoLC

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](#)