

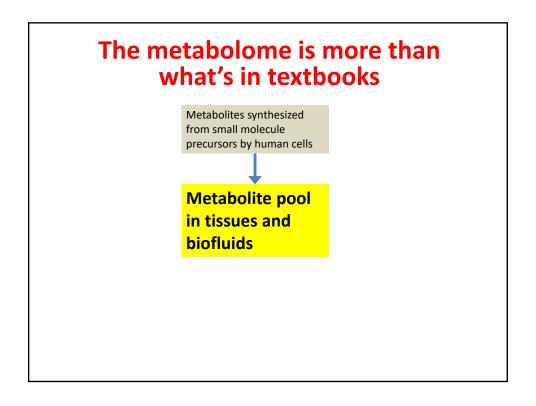
Knowledge that will change your world

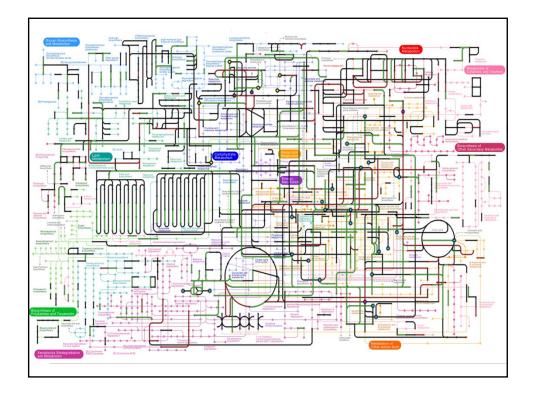
The Chemistry of the metabolome

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What is a component of the metabolome?

- In the context of metabolomics, it is compound of any origin that has a molecular weight <1,500 Da that can be detected in the biological system being studied
- · This is an arbitrary definition

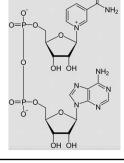


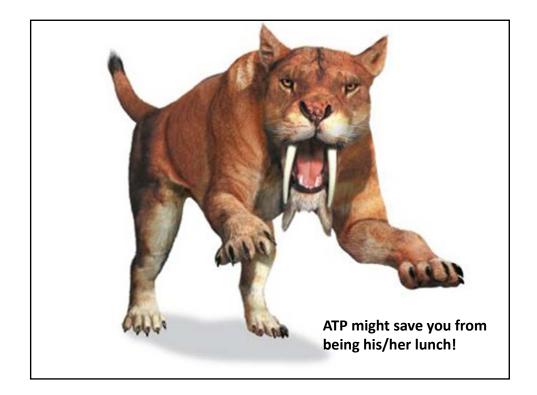


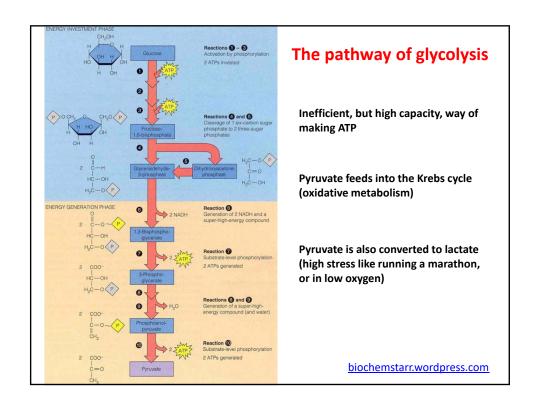
Critical metabolites

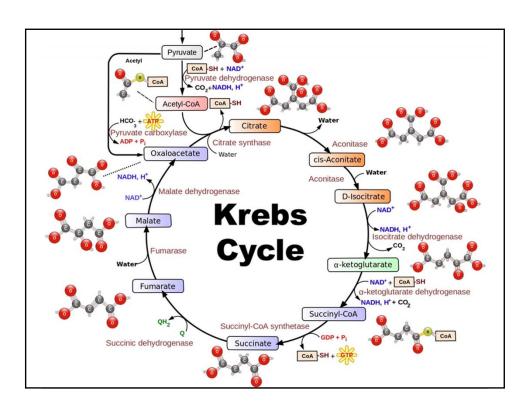
ATP: adenosine-5-triphosphate

NAD+/NADH: nicotinamide adenine dinucleotide









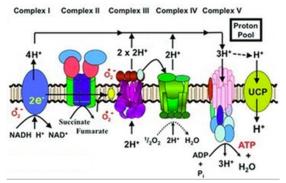
Sir Hans Krebs





Had the pleasure as a graduate student of introducing him at a seminar

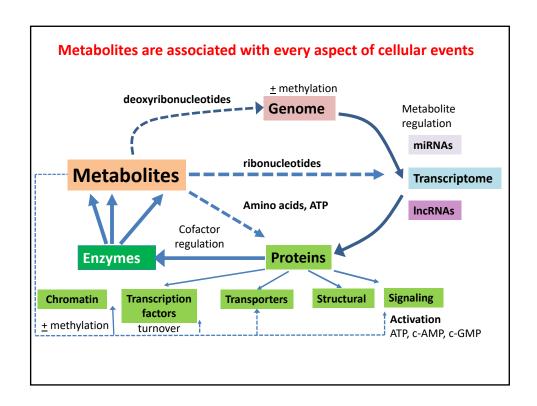
Mitochondrial oxidative phosphorylation



NADH from the Krebs cycle, as well as succinate, generate a proton (H⁺) gradient (upper region) that drives rotation of one of the subunits of ATP synthase. This exposes the catalytic domain of this enzyme and makes ATP.

Understanding metabolites

- Metabolites represent the action items that come from gene expression and protein activity
 - They are found in the same range of concentrations as drugs
 - $-\,$ Metabolites (µM or mM) (acetyl CoA, ATP, SAdMet, α KG) are regulators of epigenetics
 - Bile acids (μ M) are the natural ligands of FXR and LXR
 - Other metabolites (pM or nM) may be exquisite physiological regulators of kidney function (prostaglandins, F₂-isoprostanes)
- Studying the metabolome requires multiple levels of science from the analytical to the physiologic to the computational

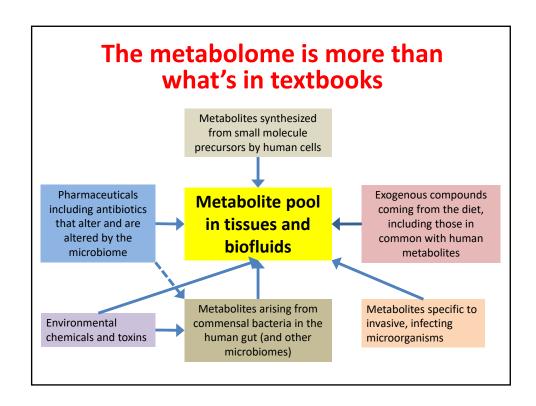


Metabolism and time

- Not only should metabolites appear in the right place, there is also the question of the importance of the timescale
- Metabolism defects in the heart may be only seconds away from death – rogue waves in metabolism??



- Irreversible damage to the brain may occur in minutes
- Go/No-Go decisions for a cell to divide or apoptose may occur in tens of mins



Overview of metabolome chemistry

Metabolites encompass an enormous range of chemistries

- Gaseous
 - H₂, H₂S
- Volatile
 - Butyric acid, acetone, skatole
- Hydrophilic (water-loving)
 - Glucose
- Charged-positive/negative
 - Amino acids, nucleotides, organic acids, amines
- Hydrophobic (fat-loving)
 - Lipids, steroids, hydrocarbons

Gases and volatiles

- In breath
 - H₂ from reductive anerobic bacteria
 - Lactose-intolerant
 - Measure of gut transit (typically 4-6 hours)
 - $-CO_2$
 - From all carbon-containing substrates
 - From specific ¹³C-labelled substrates
 - Acetone (in diabetics)
 - Trimethylamine
 - From fish, or flavin monooxygenase (FMO3)-deficient subjects

Gases and volatiles

- Sweat gland
 - Sweaty socks syndrome
 - Isovaleric acid (leucine metabolism)
 - Caused by bacteria or enzyme defect

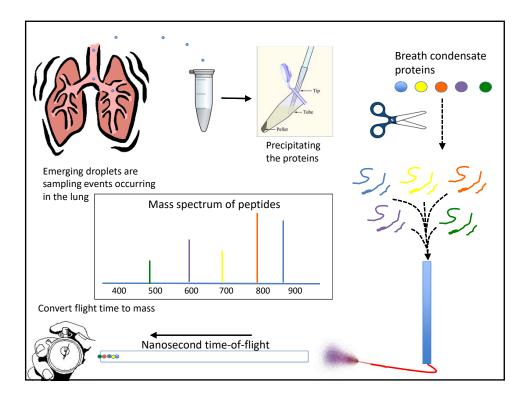
- Flatulance
 - Mostly gases (H₂, CO₂ and H₂S), but with volatiles produced by colonic bacteria (skatole, from the amino acid tryptophan)

Other volatiles

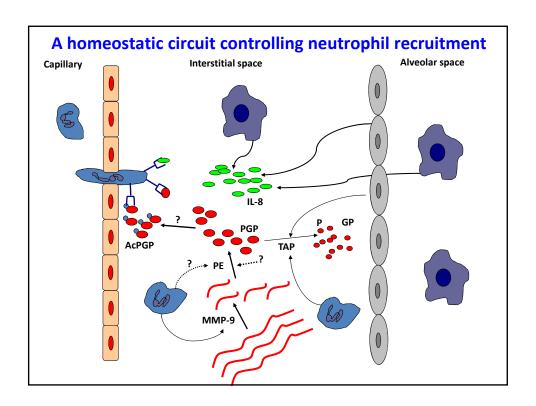
- Short chain, unsubstituted fatty acids
 - Formic, acetic, propionic, butyric acids
- Will evaporate in the acidic form
 - Formic acid, b.p. 101°C
 - Acetic acid, b.p. 118°C
 - Propionic acid, b.p. 141°C
 - Butyric acid, b.p. 163.8°C
 - Isobutyric acid, b.p. 155°C
- · Convert to ammonium salts before evaporating

Breath condensates

- Not strictly consisting of volatiles
- A mist or spray created by the frothing of the fluids inside the lung
 - Condensable using a dry-ice cooled trap
 - Several ml of condensate can be easily collected in 5-8 min

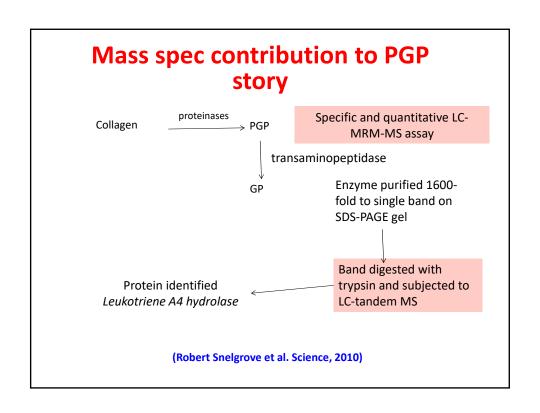


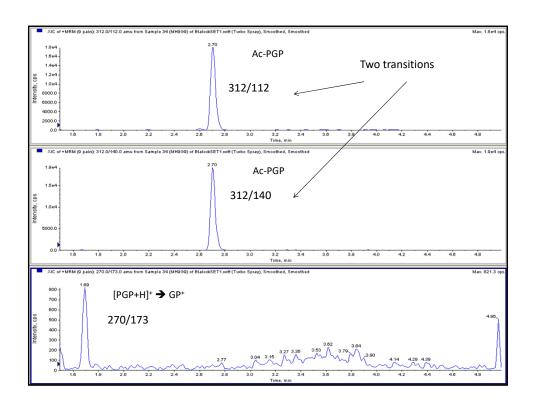
Certain metabolites are peptides



PGP is a common peptide in human collagen

MFSFVDLRLLLLLAATALLTHGOEEGOVEGODEDIPPITCVONGLRYHDRDVWKPEPCRI ${\tt CVCDNGKVLCDDVICDETKNCPGAEVPEGECCPVCPDGSESPTDQETTGVEGPKGDTGPR}$ $\texttt{MGPSGPRGL} \textcolor{red}{\textbf{PGP}} \texttt{PGAPGPQGFQGPPGEPGEPGASGPMGPRGP} \textcolor{red}{\textbf{PGP}} \texttt{PGKNGDDGEAGKPGR}$ MGPRGLPGERGRPGAPGPAGARGNDGATGAAGPPGPTGPAGPPGFPGAVGAKGEAGPQGP ${\tt RGSEGPQGVRGE} {\tt PGPPGP} {\tt AGAAGPAGNPGADGQPGAKGANGAPGIAGAPGFPGARGPSGP}$ $\tt QGPGGP{\color{red}PGP}KGNSGEPGAPGSKGDTGAKGE{\color{red}PGP}VGVQGP{\color{red}PGP}AGEEGKRGARGE{\color{red}PGP}TGL$ PGP PGERGG PGS RGF PGADGVAG PKG PAGERGS PGPAG PKG S PGEAG R PGEAG L PGAKG L TGSPGSPGPDGKTGPPGPAGQDGRPGPPGPPGARGQAGVMGFPGPKGAAGEPGKAGERGV ${\tt PGP} {\tt PGAVGPAGKDGEAGAQGP} {\tt PGP} {\tt AGPAGERGEQGPAGSPGFQGL} {\tt PGP} {\tt AGPPGEAGKPGE}$ ${\tt QGVPGDLGA} {\tt PGP} {\tt SGARGERGFPGERGVQGP} {\tt PGP} {\tt AGPRGANGAPGNDGAKGDAGAPGAPGS}$ $\tt QGAPGLQGMPGERGAAGL{\color{red}PGP}KGDRGDAGPKGADGSPGKDGVRGLTGPIGPPGPAGAPGD$ KGESGPSGPAGPTGARGAPGDRGE**PGPPGP**AGFAGPPGADGOPGAKGEPGDAGAKGDAGP **PGP**AGPAGP**PGP**IGNVGAPGAKGARGSAGPPGATGFPGAAGRVGP**PGP**SGNAGP**PGPPGP** ${\tt AGKEGGKGPRGETGPAGRPGEVGP{\tt PGPPGP}AGEKGSPGADGPAGAPGT{\tt PGP}QGIAGQRGV}$ VGLPGORGERGFPGLPGPSGEPGKOGPSGASGERGPPGPMGPPGLAGPPGESGREGAPGA $VGARGPAGPQGPRGDKGETGEQGDRGIKGHRGFSGLQGP \\ \textcolor{red}{\textbf{PGP}} PGSPGEQGPSGASGPAGP$ ${\tt RGPPGSAGAPGKDGLNGL} {\tt PGP} {\tt IGPPGP} {\tt RGRTGDAGPVGPPGPPGPPGPPGPPSAGFDFSF}$ LPQPPQEKAHDGGRYYRADDANVVRDRDLEVDTTLKSLSQQIENIRSPEGSRKNPARTCR DLKMCHSDWKSGEYWIDPNQGCNLDAIKVFCNMETGETCVYPTQPSVAQKNWYISKNPKD KRHVWFGESMTDGFQFEYGGQGSDPADVAIQLTFLRLMSTEASQNITYHCKNSVAYMDQQ TGNLKKALLLQGSNEIEIRAEGNSRFTYSVTVDGCTSHTGAWGKTVIEYKTTKTSRLPII DVAPLDVGAPDQEFGFDVGPVCFL





Metabolopeptidomics or peptidometabolomics

- Are peptides metabolites?
- Are the tripeptides real?
 Or is their mass simply coincident with the empirical formula of another metabolite?

Considering the case for tripeptides

- Examine the basic physiology and pharmacology
- Are there examples of bioactive tri-peptides?
- What about other oligopeptides?
- Where would they come from?
- Why does METLIN seem to always have triand not other oligopeptides?

Tripeptides could come from foods, but are hydrolyzed by peptidases in the enterocyte to amino acids

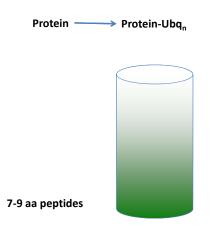
Deficiencies in the peptidases could lead to food and bacterial peptides entering the systemic circulation

Can tripeptides have biological activity?

- For toxicologists, there is one very familiar tripeptide without whom, I would not be giving this talk, or you to listen to it.
- Glutathione (GSH) glutamyl-cysteinyl-glycine
 - GSH reacts with free radicals to generate GSH conjugates and therefore protects many organs
- It is synthesized from small molecule precursors
 - However, it is a true metabolite, i.e., it is made from smaller precursors without the direct aid of ribosomes

Are there other sources of short peptides?

- Proteins undergo degradation in the proteasome caused by targeted ubiquitination
 - The digested products are peptides (escapees?)
- Lysosomes
- Autophagosome
- Neutrophil attack
- Other proteases (in renal tubules?
- Foreign antigens hydrolyzed and presented on surface of cells



Proteasome

Hydrophilic metabolites

- The most extreme hydrophilic metabolites without charged groups are the polyols:
 - Monosaccharides
 - Glucose
 - Fructose
 - Disaccharides
 - Lactose
 - Maltose
 - Oligosaccharides

Organic acids

- Besides the short chain fatty acids mentioned earlier, there are many organic acids representing important cellular pathways
 - Glycolytic intermediates
 - Glucose-1-P, Glucose-6-P, Fructose-6-P, Fructose-1,6-DP, Glyceraldehyde-3-P, Dihydroxyacetone-P, Glycerate-3-P, Phosphoenol-P, Pyruvate, Lactate
 - Krebs cycle
 - Citrate, cis-Aconitate, Iso-Citrate, α-ketoglutarate, Succinate, Fumarate, Malate, Oxaloactate and those resulting from pathway defects
 - Nucleotides
 - ATP, ADP, AMP, GTP, etc.

How could we isolate organic acids?

- Organic acids at neutral pH are negatively charged
- They will bind to anion exchange resins in say the formate form

AG-1

 Can be eluted with ammonium formate or ammonium acetate (mass spec compatible)

Amino acids

- All the α -amino acids found in proteins and their precursors and metabolites
 - Mostly L-isomers, but there are D-isomers in nature
- Tryptophan is oxidized to kynurenine and is a precursor to NAD(H) and NADP(H), serotonin melatonin and niacin
- β-Alanine is formed from uracil
 - 5-fluorouracil (anticancer drug) is converted to 2-fluoro- β -alanine which is in turn converted to bile acid conjugates

How could we isolate amino acids?

- Amino acids at neutral pH are positively charged
- They will bind to cation exchange resins in the H⁺ form

AG-50

 Can be eluted with ammonium hydroxide (mass spec compatible)

Hydrophobic metabolites

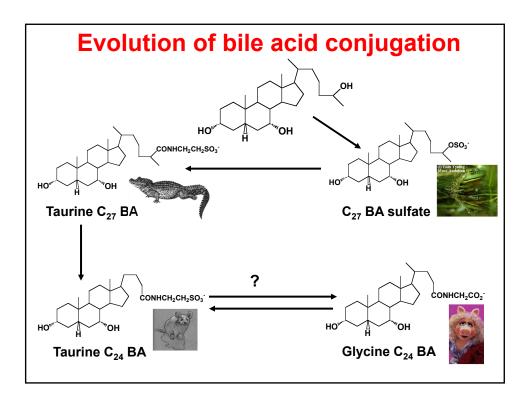
 These include sterols, steroid hormones, terpenoids, bile acids, vitamins A, D, E and K, and a vast array of lipids

Importance of sterols and other compounds in lens cataracts

E F OH O OH

Structures A, B and C (all sterols) have recently been shown to have the property of "dissolving" lens cataracts. Cholesterol, on the other hand, has no effect. Other sterols observed in *cerebrotedinous* xanthamatosis promote cataracts.

D, E and F all promote lens cataracts. D is prednisone (an anti-inflammatory steroid), E is ciprofloxacin (an antibiotic) and F is hypericin from the botanical, St. John's wort.



The vitamins

Lack of these leads to serious illness, but not death

Vitamin B

• They are all water-soluble

Vit B₁ – thiamine

- Vit B₂ - riboflavin

- Vit B₃ - niacin

- Vit B₅ - pantothenic acid

Vit B₆ – pyridoxine

Vit B₇ – biotin

Vit B₉ – folic acid

- Vit B₁₂ - cobalamins

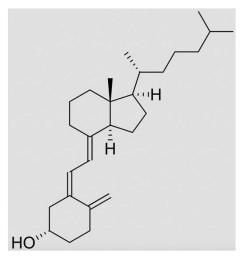
They are not made by human enzymes and if deficient in the diet cause disease

Vitamin C

Ascorbic acid

dehydroascorbic acid

Vitamin D



In fish, supplemented in milk, made in skin by UV light

Vitamin E

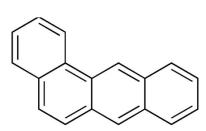
Found in oils from plants

Vitamin K

Is an anticoagulant – needed to stop bleeding

Phospholipids

Hydrocarbons



Benz[a]anthracene
In smoke from barbecued meat



Cetyl palmitate In hair shampoo

Solubilities of the metabolites

- Those in biological fluids are "in solution", but may not be soluble in water or methanol alone
 - Are glucose or amino acids soluble in methanol?
 - Are cholesterol esters in plasma soluble in methanol or water?
 - If a metabolite binding protein is precipitated by methanol, does the metabolite still bind to it?
 - Does pH have an effect on solubility?