Biosynthesis – Inspiration for Drug Discovery

Primary Metabolism & Enzyme Cofactor Chemistry

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Lessons in Synthesis - Azadirachtin

- **Azadirachtin** is a potent *insect anti-feedant* from the Indian *neem tree*:
  - exact biogenesis unknown but certainly *via* steroid modification:
    - Intense synthetic efforts by the groups of Nicolaou, Watanabe, Ley and others since structural elucidation in 1987.
    - 1st total synthesis achieved in 2007 by Ley following 22 yrs of effort
    - ~40 researchers and over 100 man-years of research! – 64-step synthesis

![Diagram](image_url)
Rational Agrochemical Development – Shikimate Pathway Intervention

- **The shikimate biosynthetic pathway is not found in animals/humans – only in plants**
  - selective intervention in these pathways allows development of agrochemicals with minimal toxicity

- **Glyphosate (‘Roundup’) – a Monsanto agrochemical is a potent inhibitor of the conversion of 3-phosphoshikimate (3-PS) → 5-enolpyruvylshikimate-3-phosphate (5-EPS-3P)**
  - a non-selective herbicide

\[
\begin{align*}
\text{P}_i & \quad \text{CO}_2 \quad \text{P}_i \\
\text{CO}_2 & \quad \text{P}_i \\
\text{CO}_2 & \quad \text{P}_i \\
\text{CO}_2 & \quad \text{P}_i \\
\end{align*}
\]

\[
\begin{align*}
\text{phosphoenol pyruvate (PEP)} & \quad \text{PO}^+ \quad \text{CO}_2 \\
\text{erythrose-4-phosphate (E-4-P)} & \quad \text{PO}^- \quad \text{HO}^- \\
\end{align*}
\]

\[
\begin{align*}
\text{3-phosphoshikimate (3-PS)} & \quad \text{OH} \\
\text{PEP} & \quad \text{OH} \\
\text{PEP} & \quad \text{OH} \\
\text{PEP} & \quad \text{OH} \\
\end{align*}
\]

glyphosate (Roundup®) inhibits this step

\[
\begin{align*}
\text{(S)-tyrosine} & \quad \text{NH}_3 \quad \text{CO}_2 \\
\text{(S)-phenylalanine} & \quad \text{NH}_3 \quad \text{CO}_2 \\
\end{align*}
\]
Inspiration for Med Chem - Statins

- **HMG CoA → MVA** is the *rate determining step* in the biosynthetic pathway to **cholesterol**

- ‘**Statins**’ inhibit HMG CoA reductase and are used clinically to treat **hypercholesterolemia** - a causative factor in **heart disease**
  - *e.g. Lipitor* (Atorvastatin calcium, Pfizer) is a competitive inhibitor of HMG-CoA reductase and the world's biggest selling drug [first drug to reach $10 billion sales (2004: $10.8 bn)]
Format & Scope of Lectures

• **What is biosynthesis?**
  – some definitions – phototrophs, chemotrophs; metabolism (catabolism/anabolism), 1° & 2° metabolites

• **Overview of primary metabolism → secondary metabolites**
  – photosynthesis & glycolysis → shikimate formation → shikimate metabolites
  – acetylCoA & the citric acid cycle → α-amino acids → penicillins, cephalosporins, alkaloids
  – acetylCoA → malonylCoA → fatty acids, prostaglandins, polyketides, macrolide antibiotics
  – acetylCoA → mevalonate → isoprenoids, terpenoids, steroids, carotenoids

• **Biological/biosynthetic reactions – enzyme & cofactor chemistry**
  – free energy source – ATP
  – C-C & C-O bond formation – CoASH, SAM, DMAPP, biotin
  – oxidation – NAD⁺, FAD/FMN, haem iron oxo monooxygenases
  – reduction – NADPH
  – C-N bond formation – pyridoxal
Metabolism & Natural Product Diversity

- lysergic acid
- camphor
- caffeine
- quinine
- clavulanic acid
- nicotine
- patulin
- androstenedione

Reactants: CO₂, H₂O, P_i, N₂

hv
Phototrophs & Chemotrophs

- **Living organisms** are not at equilibrium. They require a continuous influx of **free energy** to perform mechanical work & for cellular growth/repair:

  - **Phototrophs** (e.g. green plants, algae & photosynthetic bacteria): derive free energy from the sun via **photosynthesis** (‘CO₂ fixation’):
    - 10\(^{15}\) kg/year by green plants, which constitute 99% of Earth's biomass (i.e. 10\(^{12}\) tons of dry matter)
    - 1g of carbon processed = >6250 litres of air

\[
\text{CO}_2 + \text{H}_2\text{O} \xrightarrow{hv} (\text{CHO}) + \text{O}_2
\]

  - **Chemotrophs** (e.g. animals, fungi, most bacteria): derive free energy by **oxidising nutrients** (carbohydrates, lipids, proteins) obtained from other organisms, ultimately phototrophs
    - some bacteria & fungi require just D-glucose
    - mammals require sugars, essential amino acids (~half total used) & certain vitamins (enzyme co-factors or precursors)

- Degradation of the nutrients is coupled to the stoichiometric production of ‘high energy’ phosphate compounds, particularly **adenosine triphosphate** (**ATP**, see later). All metabolic function is underpinned by ATP energetic coupling.
  - By analogy with a money-based economy, the metabolic cost of production of a given metabolite from another can be **quantified in terms of ‘ATP equivalents’** defined as the # of moles of ATP consumed/produced per mole of substrate converted in the reaction or sequence
Metabolism

- **Metabolism** is the term used for *in vivo* processes by which compounds are degraded, interconverted and synthesised:
  - **Catabolic** or **degradative**: primarily to release energy and provide building blocks
    - generally **oxidative** processes/sequences (glycolysis, Krebs cycle)
  - **Anabolic** or **biosynthetic**: primarily to create new cellular materials (1° & 2° metabolites)
    - generally **reductive** processes/sequences
- These two types of process are coupled – one provides the driving force for the other:
Types of Metabolite & Biosynthesis

- **Biosynthesis** is the term for the in vivo synthesis of metabolites/natural products:
  - These are divided into two camps:
    - **Primary metabolites**: These are the universal and essential components for the survival of living organisms. *e.g.* sugars, amino acids, nucleotides, ‘common’ fats and polymers such as proteins, DNA, RNA, lipids and polysaccharides
    - **Secondary metabolites**: Compounds produced by organisms which are not required for survival, many of which have no apparent utility to the host organism. Frequently a given metabolite will only be produced in a single organism or in a set of closely related organisms. Provide a rich source of pharmacologically active compounds. *e.g.* shikimate derivatives, alkaloids, fatty acids, polyketides, isoprenoids
  - Although the boundary is imprecise the term **biosynthesis** is most commonly applied, by organic chemists, to the in vivo synthesis of secondary metabolites:

  "Now ever since Perkin, failing to make quinine, founded the dyestuffs industry, organic chemists have found the study of ‘natural products’ an inexhaustable source of exercises, which can be performed out of pure curiosity even when paid for in the hope of a more commercial reward. As a result the organic chemist’s view of nature is unbalanced, even lunatic but still in some ways more exciting than that of the biochemist. While the enzymologist’s garden is a dream of uniformity, a green meadow where the cycles of Calvin and Krebs tick round in disciplined order, the organic chemist walks in an untidy jungle of uncouthly named extractives, rainbow displays of pigments, where in every bush there lurks the mangled shapes of some alkaloid, the exotic perfume of some new terpene, or some shocking and explosive polyacetylene…”

  … Since these intriguing derivatives AND e.g. lysine or ATP are ALL in a sense ‘natural products’ we may prefer the term ‘secondary metabolite’ for the former

Primary Metabolism - Overview

**Primary metabolism**

\[ \text{CO}_2 + \text{H}_2\text{O} \]

**Photosynthesis**

1) 'light reactions': \( \text{hv} \rightarrow \text{ATP and NADPH} \)
2) 'dark reactions': \( \text{CO}_2 \rightarrow \text{sugars (Calvin cycle)} \)

**Glycolysis** & other 4, 5, 6 & 7 carbon sugars

- **glucose**
- **phosphoenol pyruvate**
- **erythrose-4-phosphate**

**Citric acid cycle (Krebs cycle)**

- **pyruvate**
- **acetyl coenzyme A**
- **mevalonate**
- **malonyl coenzyme A**

**Primary metabolites**

- Oligosaccharides
- Polysaccharides
- Nucleic acids (RNA, DNA)
- Peptides
- Proteins
- Tetrapyrroles (porphyrins)
- Saturated fatty acids
- Unsaturated fatty acids
- Lipids

**Secondary metabolites**

- Shikimate metabolites
- Cinnamic acid derivatives
- Aromatic compounds
- Lignans, flavonoids
- Alkaloids
- Penicillins
- Cephalosporins
- Cyclic peptides
- Fatty acids & polyketides
- Prostaglandins
- Polyacetylenes
- Aromatic compounds, polyphenols
- Macrolides
- Isoprenoids
- Terpenoids
- Steroids
- Carotenoids

For interesting animations of e.g. photosynthesis see: [http://www.johnkyrk.com/index.html](http://www.johnkyrk.com/index.html)
Biological/Biosynthetic Reactions – Enzyme Catalysis & Cofactors

• Most biosynthetic steps are catalysed by specific, individual enzymes. They generally perform familiar processes such as oxidation, reduction, alkylation, hydrolysis, acylation, hydroxylation, elimination etc.

• Different enzymes carrying out related reactions often employ common co-factors: small organic functional fragments and/or metal ions. e.g.

  – **FREE ENERGY RELEASING COUPLE**: Adenosine triphosphate (ATP)

  – **C-C & C-O BOND FORMATION**: Coenzyme A (CoASH); S-adenosyl methionine (SAM); dimethylallylpyrophosphate (DMAPP); biotin

  – **OXIDATION**: NAD(P)^+; FAD/FMN; Haem iron oxo species (e.g. P_{450})

  – **REDUCTION**: NAD(P)H; (FADH\_2/FMNH\_2)

  – **C-N BOND FORMATION**: Pyridoxal
Free Energy Releasing Couple - ATP

- **Adenosine triphosphate (ATP)**
  - phosphorylation of an alcohol by adenosine diphosphate (ADP) is highly **exothermic** (i.e. liberates energy):

  \[
  \Delta G^\circ = -31 \text{ kJ mol}^{-1}
  \]

  - The phosphorylated alcohol (ROP) is then activated towards nucleophilic displacement:

    \[
    \text{Nu}^\ominus + \text{ROP} \rightarrow \text{R-Nu} + ^\ominus \text{OP}
    \]

    - So, overall the **endothermic** process ROH + Y\(^-\) → RY + OH\(^-\) has been achieved by ‘coupling’ the process to the ‘hydrolysis of ATP’
    - The situation is analogous to the use of tosylate activation to achieve nucleophilic displacement of an alcohol
    - In general, the exothermicity associated with phosphorylation shifts the equilibria of ‘coupled’ process by a **factor of \(~10^8\)**
Acylation & C-C Bond Formation \( \alpha \) to C=O – CoASH

- **Coenzyme A (CoASH)**
  - Coenzyme A acts as an acyl transfer/\( \alpha \)-carbon activation reagent by forming reactive acyl thioesters:

    - Acyl CoA derivatives can act as nucleophiles or electrophiles depending on the circumstances
    - **These modes of reactivity reflect inherent properties of alkyl thioesters:**
      - The *good leaving group ability of RS\(^-\)* (cf. RO\(^-\)) reflects: \( pK_a \) (RSH) \( \sim \) 10 cf. \( pK_a \) (ROH) \( \sim \) 16
      - The *high electrophilic character of a thioester carbonyl carbon* (cf. normal esters) reflects the poor orbital overlap between the lone pairs on sulfur \( (n_S) \) [cf. \( n_O \)] and the carbonyl anti bonding molecular orbital \( \pi^*_{C=O} \)
      - The *enhanced acidity of protons \( \alpha \) to the carbonyl of thioesters* (cf. normal esters) reflects the same poor \( n_S \leftrightarrow \pi^*_{C=O} \) resonance:

\[
\begin{array}{c}
\text{H} \quad \text{R} \\
\text{R'} \\
\text{H} \\
\text{B:} \\
\text{H} \quad \text{X} \quad \text{R'} \\
\end{array}
\]
Methylation/Dimethylallylation – SAM & DMAPP

• **S-Adenosyl methionine (SAM)**
  - SAM acts as a versatile O-, C-, N- & S- methylating reagent *in vivo*
    - Equivalent to performing an $S_N2$ methylation using Mel in the laboratory

![Diagram of S-Adenosyl methionine (SAM)](image)

• **Dimethylallyl pyrophosphate (DMAPP)**
  - DMAPP acts a dimethylallylating reagent – the pyrophosphate (+ Mg$^{2+}$/Mn$^{2+}$) is an excellent leaving group
    - Equivalent to performing an $S_N2$ allylation using allyl bromide in the laboratory

![Diagram of Dimethylallyl pyrophosphate (DMAPP)](image)
Carboxylation – *Biotin*

- **Biotin**
  - Biotin in the presence of bicarbonate, ATP and Mg$^{2+}$ enables nucleophile carboxylation *in vivo*:
Oxidation – \( NAD^+ \)

- **Nicotinamide-adenine dinucleotide (NAD\(^+\))** [and its phosphorylated analogue (NADP\(^+\))] are mediators of biological oxidation (e.g. alcohol to ketone oxidation)
  - In general, the couple NAD\(^+\)/NADH is used by enzymes in **catabolic oxidation** (degradation)
  - The reagent is a stereospecific **hydride acceptor**:

\[
\text{nicotinamide-adenine dinucleotide (NAD\(^+\))} \quad R = H \\
\text{NB. NADP\(^+\)}: \quad R = \text{PO}_3\text{H}^-
\]

- Different enzymes show different absolute specificities but are generally specific for the pro-R or pro-S hydrogens both for removal and delivery
Oxidation – *Flavins (FAD & FMN)*

- Flavin adenine dinucleotide (FAD) and flavin mononucleotide (FMN) are also mediators of biological oxidations (e.g. dehydrogenations – alkane to alkene)
  - Unlike NAD\(^+\), which readily diffuses from enzyme to enzyme, FAD/FMN is usually tightly bound to a given enzyme, sometimes covalently

- Re-oxidation of the FADH\(_2\) back to FAD is generally by molecular oxygen (although NAD\(^+\) is also sometimes used). The intermediate peroxyflavin can also mediate hydroxylation, epoxidation & other oxygen transfer reactions (see next slide):
Oxidation Reactions Mediated by Flavins

- **Dehydrogenation by flavins** – e.g. dehydrogenation of succinate $\rightarrow$ fumarate:

  \[
  \text{succinate} \quad \xrightarrow{\text{gauche}} \quad \text{fumarate} \quad \text{(i.e. 100% trans)}
  \]

- **Baeyer-Villiger-type oxidation by peroxyflavins** – e.g. ketone monooxygenase:

  \[
  \text{peroxyflavin} \quad \xrightarrow{\text{NADPH} + H^+ + O_2} \quad \text{FAD} \quad \xrightarrow{\text{H}_2\text{O}} \quad \text{FADH}_2
  \]
Oxidation – Haem Iron oxo Species ($P_{450}$)

- **Haem iron oxo species** e.g. in cytochrome $P_{450}$ (a ubiquitous *heam monooxygenase*) are also mediators of biological oxidation (e.g. phenolic coupling, epoxidation, *hydroxylation*):

  - The porphyrin ring acts as a tetradentate ligand for the octahedral iron. The two axial positions are occupied by an enzyme amino acid ligand (typically a histidine nitrogen) and hydroxy/hydroperoxy residue respectively.
Reduction - \textit{NADPH}

- \textit{Dihydro-nicotinamide-adenine dinucleotide phosphate (NADPH)} [and its de-phosphorylated analogue (NADH)] are mediators of \textit{biological reduction} (e.g. ketone to alcohol reduction)
  - In general, the couple NAPH/NADP$^+$ is used by enzymes in \textit{anabolic reduction} (biosynthesis)
  - The reagent is a stereospecific \textit{hydride donor}:

\[
\begin{align*}
\text{enzyme} & \quad - H^\ominus & \quad + H^\ominus \\
\text{NADPH} & \quad \text{NADP}^+ \\
\text{NADPH} & \quad \text{NADP}^+ \\
\text{NADH:} & \quad \text{R} = H
\end{align*}
\]

- As for the reverse process, different enzymes show different absolute specificities but are generally specific for the pro-R or pro-S hydrogens both for removal and delivery
Transamination - PLP

- **Pyridoxine (vitamin B₆)** → **pyridoxal-5’-phosphate (PLP)**
  - PLP forms **amines** (Schiffs bases) with **primary amines**. This forms the basis of **in vivo transamination** of **α-ketoacids** to give **α-amino acids** (& also **racemisation/decarboxylation** processes, see ‘alkaloids’)

- The α-carbon protonation is stereospecific and gives the (S) configured chiral centre
Primary Metabolism - Overview

Primary metabolism

$$\text{CO}_2 + \text{H}_2\text{O} \rightarrow \text{glucose}$$

**PHOTOSYNTHESIS**
1) ‘light reactions’: $h\nu \rightarrow \text{ATP and NADPH}$
2) ‘dark reactions’: $\text{CO}_2 \rightarrow \text{sugars (Calvin cycle)}$

**glycolysis**
& other 4,5,6 & 7 carbon sugars

**phosphoenol pyruvate**

**erythrose-4-phosphate**

**pyruvate**

**Citric acid cycle**
(“Krebs cycle”)

**acetyl coenzyme A**

**malonyl coenzyme A**

**mevalonate**

**SHIKIMATE METABOLITES**
Cinnamic acid derivatives
Aromatic compounds
Lignans, flavonoids

**ALKALOIDS**
Penicillins
cephalosporins
Cyclic peptides

**FATTY ACIDS & POLYKETIDES**
Prostaglandins
Polyacetylenes
Aromatic compounds, polyphenols
Macrolides

**ISOPRENOIDS**
Terpenoids
Steroids
Carotenoids

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