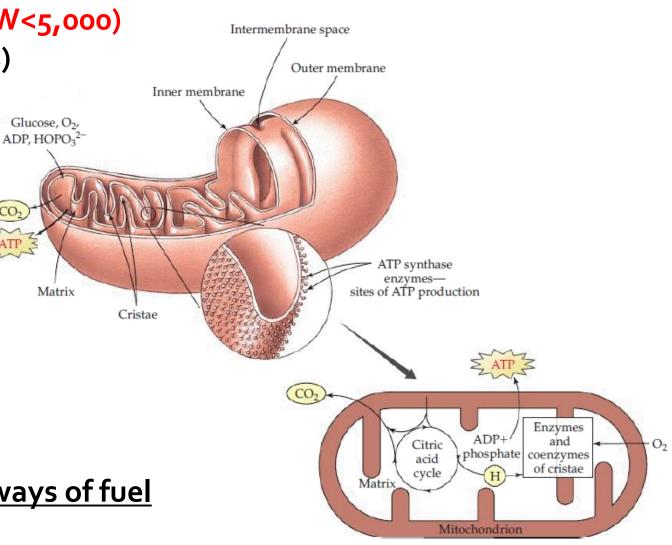
Nafith Abu Tarboush DDS, MSc, PhD <u>natarboush@ju.edu.jo</u> <u>www.facebook.com/natarboush</u>

Oxidative Phosphorylation

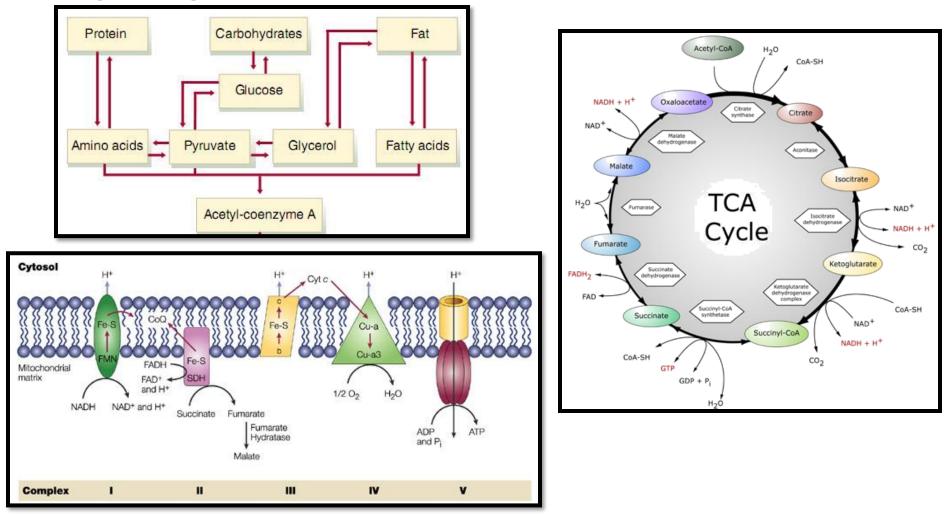
The Mitochondria

- OMM: <u>permeable</u> to small molecules (MW<5,000) & ions, <u>porins</u> (transmembrane channels)
- IMM: <u>impermeable</u> even to H+; specific transporters
- IMM bears the components of the respiratory chain and the ATP synthase
- Matrix: contains pyruvate dehydrogenase complex & TCA cycle enzymes, fatty acid β-oxidation pathway, and the pathways of amino acid oxidation
- In other words: <u>matrix contains all pathways of fuel</u> <u>oxidation except glycolysis (cytosol)</u>



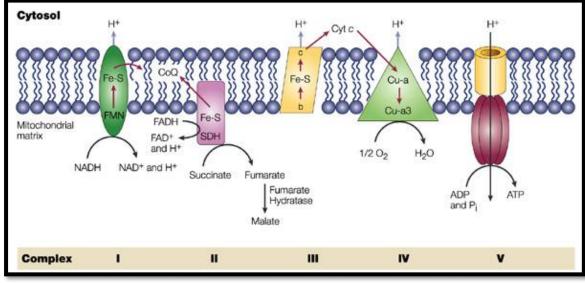
The oxidative phosphorylation, Where are we?

Stages: Digestion; Acetyl-CoA, TCA, OxPhos

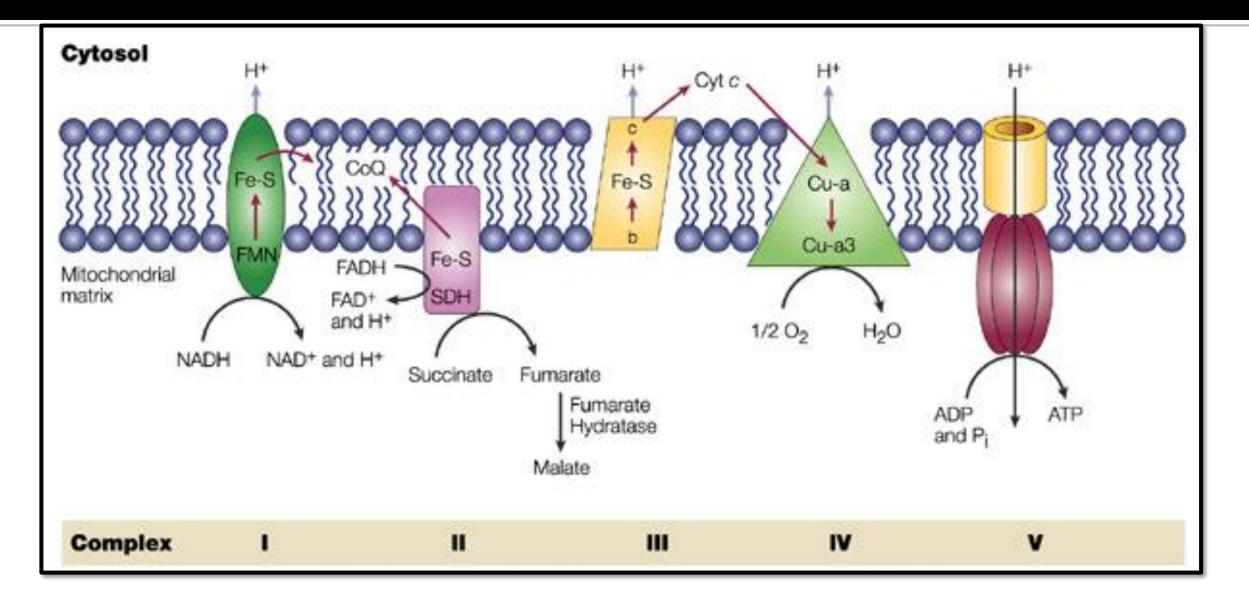


Oxidative phosphorylation (OxPhos)

- \succ Generation of ATP aided by the reduction of O_2
- Peter Mitchell (1961): the chemiosmotic theory
- Oxidative phosphorylation have 3 major aspects:
 - (1) It involves <u>flow of electrons</u> through a chain of membrane-bound carriers (<u>prosthetic groups</u>)
 - (2) The free energy available (exergonic) is <u>coupled to transport protons across</u> a proton-impermeable membrane
 - (3) The transmembrane <u>flow of protons</u> down their concentration gradient provides the free energy for synthesis of ATP (ATP synthase)

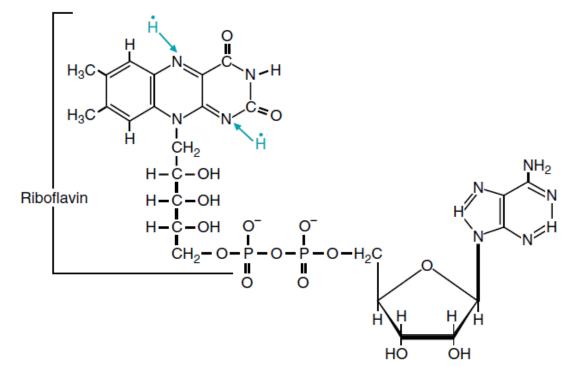


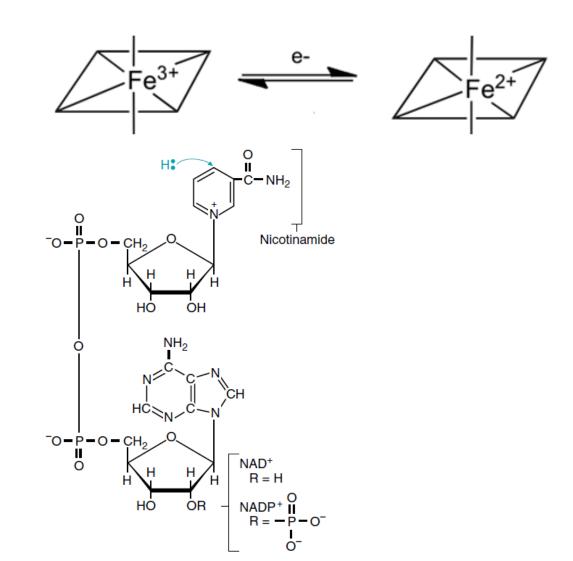
Oxidative phosphorylation (OxPhos)



Types of electron transfer (ET) through the electron transport chain (ETC)

- > 3 types of ET occur in OxPhos:
 - ✓ Direct ET, as in the reduction of Fe⁺³ to Fe⁺²
 - ✓ Transfer as a hydrogen atom {(H⁺) + (e⁻)}
 - ✓ Transfer as a hydride ion (:H⁻)





Electrons are funneled to a universal electron acceptors

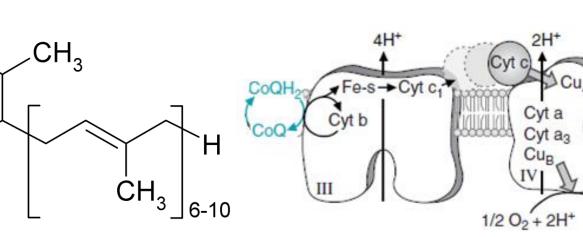
COENZYME	AS OXIDIZING AGENT	AS REDUCING AGENT
Nicotinamide adenine dinucleotide	NAD+	NADH/H+
Nicotinamide adenine dinucleotide phosphate	NADP+	NADPH/H+
Flavin adenine dinucleotide	FAD	FADH ₂
Flavin mononucleotide	FMN	FMNH ₂

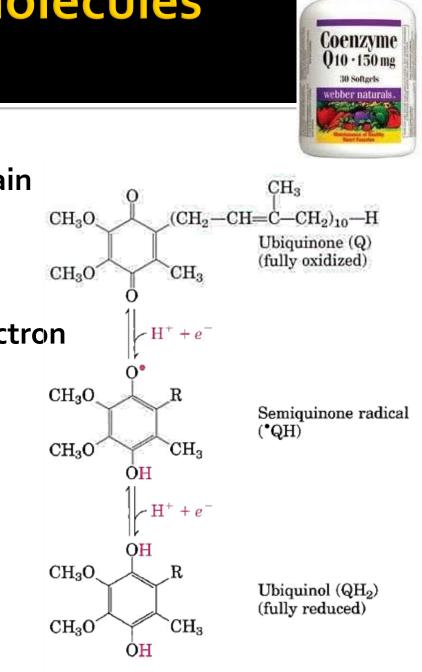
Other electron-carrying molecules "Ubiquinone"

- > Also called coenzyme Q, or Q
- Lipid-soluble benzoquinone with a long isoprenoid side chain
- Small & hydrophobic (freely diffusible)
- Carries electrons through the IMM
- Can accept either 1 e- or 2 e-

H₃C

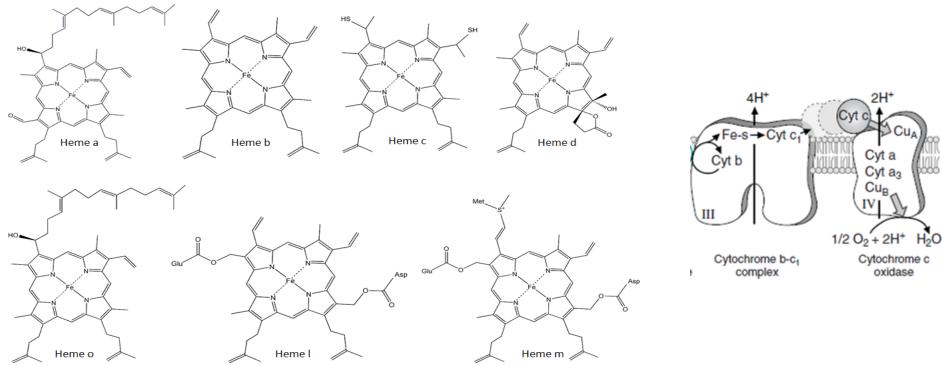
- Act at the junction between a 2-electron donor and a 1-electron acceptor
- Sometimes prescribed for recovering MI patients





Other electron-carrying molecules "Cytochromes"

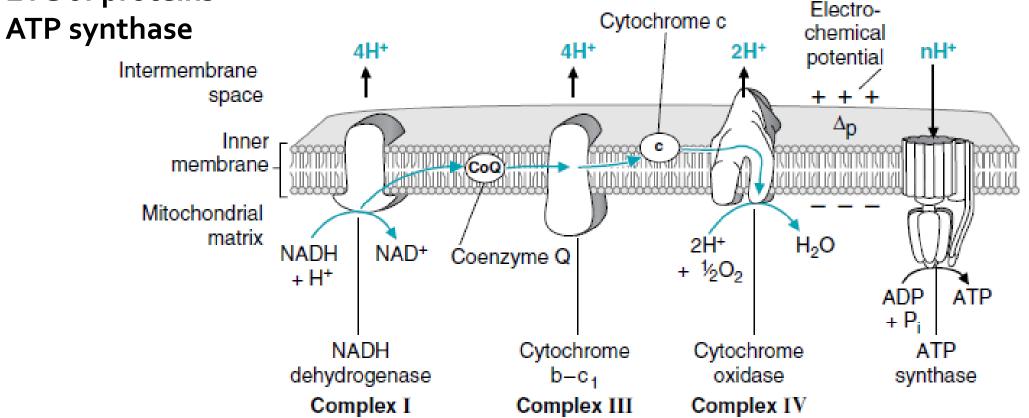
- Proteins with characteristic strong absorption of visible light (Fecontaining heme prosthetic groups)
- Classification based on light absorption
- Mode of binding (a, b, c)
- Mitochondria contain three classes o f cytochromes (a , b, & c)



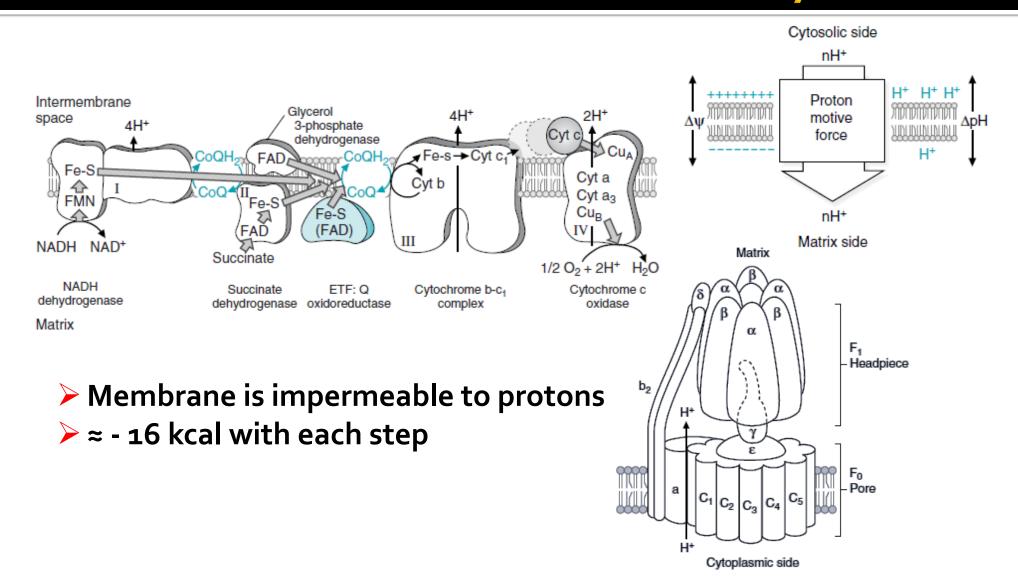
Requirements of OxPhos

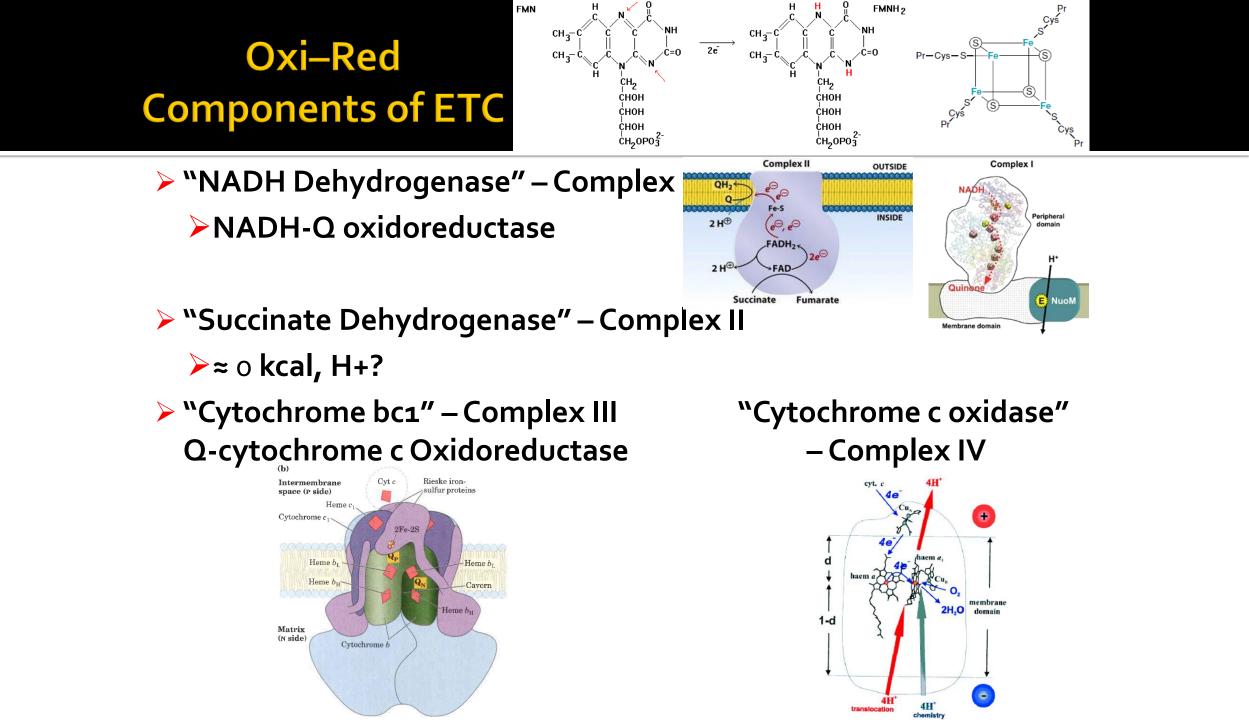
- Redox reaction: electron donor (NADH or FADH2) & electron acceptor (O2)
- An intact IMM
- ETC of proteins

>

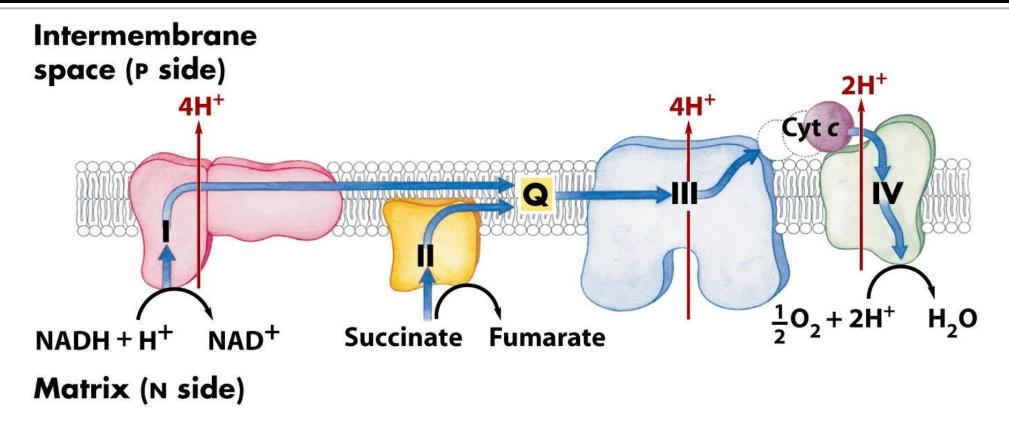


ET to O₂, how does the process occurs? "The chemi-osmotic theory"





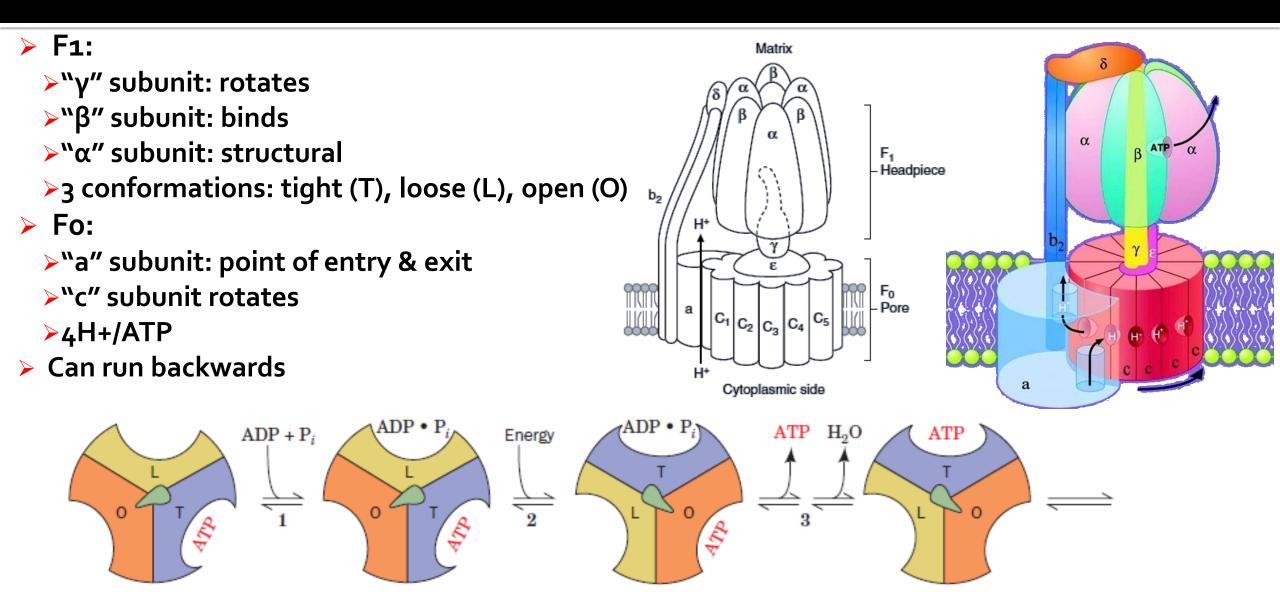
Pumping of Protons



For every 2 electrons passing:

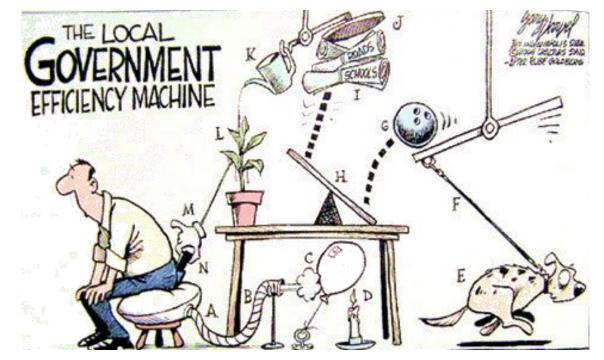
4H⁺ (complex I); 0H⁺ (complex II); 4H⁺ (complex III), 2H⁺ (complex IV)

ATP Synthase



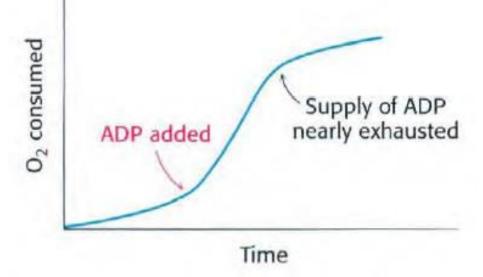
Energy yield from the ETC

- NADH, -53 kcal, ATP?
- FADH2, -41 kcal, ATP?
- $> \Delta G^{\circ}$ is so negative, never reversible
- > ATP machine efficiency, (anions, Ca⁺², heat, phosphate, substrates)
- Electron transport chain is our major source of heat



Regulation – the need for ATP

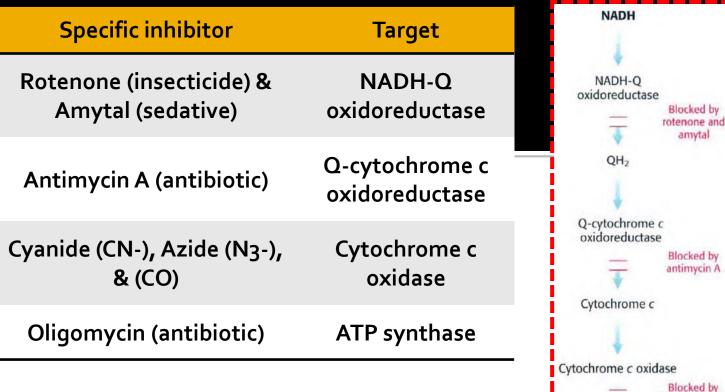
- What OxPhos needs? (NADH, O2, ADP, and Pi)
- In skeletal muscles, 20% drop in ATP concentration
- In the heart, Ca⁺² activates TCA enzymes for extra push (NADH, ATP), no drop
- ET is tightly coupled to phosphorylation (simultaneously)
- ADP is the most important factor in determining the rate
- The regulation of the rate of oxidative phosphorylation by the ADP level is called <u>respiratory control</u>



Regulation – inhibition (coupling)

- Can occur at any stage
- Specific inhibitors:
- Cyanoglycosides such as amygdalin are present in edible plant pits
- Oligomycin prevents the influx of H+





أشهر جرائم القتل العائلية في المملكة

جراسا نيوز -

ية الرئيسية > محليات

CNT, N3T, and CO

جراسا -نعرض فيما يلي قائمة بأشهر جرائم القتل العائلية التِّي حدثتُ في الاردن خلال السنواتُ المَّاضيَة ، والتي كان لكل منها وقع الصدمة حين وقوعها لما تمثله من فعل غريب على المجتمع وأعرافه ، فضلًا عن مخالفتها الشرائع السماوية والقوانين النافذة والطبيعة الإنسانية بعامة.

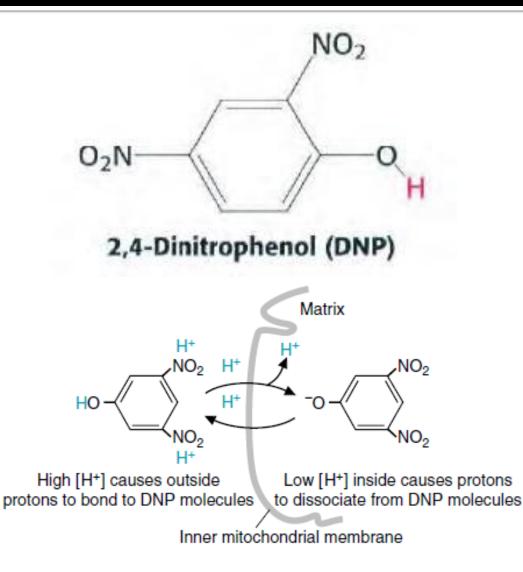
قضبة السيانيد

أول جريمة من نوعها يرتكبها أب ضد ولديه ، اذ قام الاب بوضع مادة السيانيد في كأس الحليب وطلب من طفليه ان يشربا منه ، حيث فارقا الحياة بعد 10 دقائق من مغادرة الام المنزل لتعود وتجدهما جثتين هامدتين.

وقد ادين الاب يعقونة الاعدام شنقا الا ان والده اسقط الحق الشخصي كونه وليا عن الطفلين وحكم عليه بالاشغال المؤيدة.

Regulation – Uncoupling Unregulated – chemical uncouplers

- What is uncoupling?
- How does it occur? Dissipation of PMF
- What is the result?
- Is it physiological or not?
- 2,4-dinitrophenol (DNP) & other acidic aromatic compounds
- What changes happen? ↑ O2 consumption,
 ↑NADH oxidation
- Soviet soldiers were given DNP, FDA banned DNP (1938)



Regulation – Uncoupling Regulated - Uncoupling proteins (UCPs)

- Short-circuiting ATP synthase
- > UCP1 (thermogenin):
 - Brown adipose tissue, non-shivering thermogenesis
 - Infants: neck, breast, around kidneys
 - Fatty acids directly activates UCP1
- UCP2 (most cells); UCP3 (skeletal muscle); {UCP4, UCP5} (brain)

ATP

vnthase

ANT

ATP

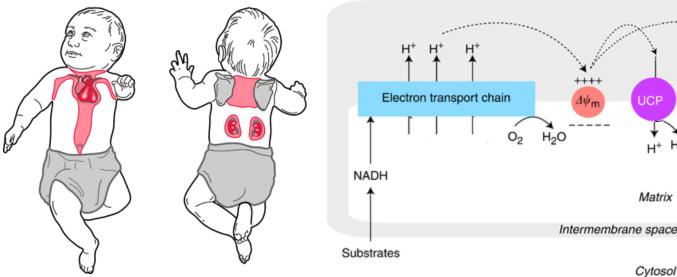
ADP

ADP

H+ Heat

¥ ¥ _H+ ATP

> Obesity tendency in some populations



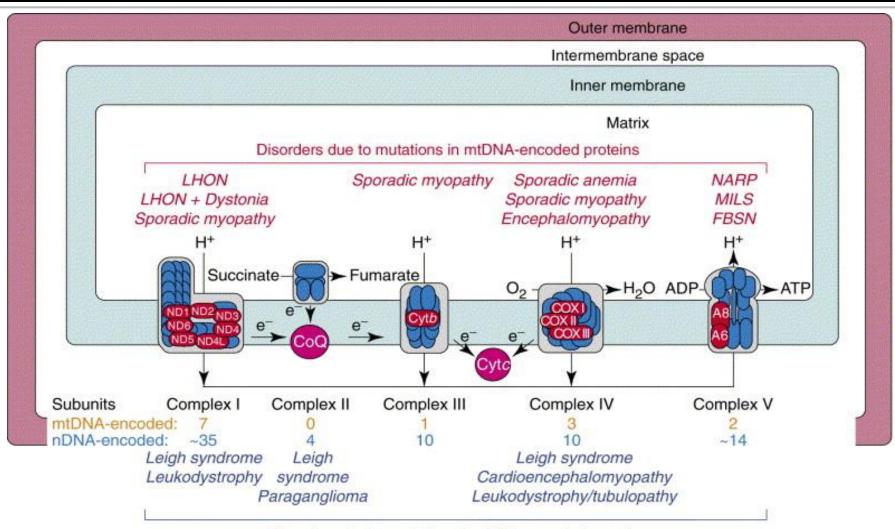
OxPhos Diseases (Genetic)

- A. Mitochondrial DNA and OXPHOS Diseases
 - Small (16,569) base pair, double-stranded, circular DNA
 - Encodes 13 subunits: 7 (I), 1 (III), 3 (IV), 2 (Fo)
 - Also encodes necessary components for translation of its own mRNA: a large and small rRNA and tRNAs
 - Maternal inheritance, heteroplasmy
 - Accumulation of somatic mutations with age
 - Highest ATP demands: CNS, heart, skeletal muscle, and kidney, liver

OxPhos Diseases (Genetic)

- B. Nuclear Genetic Disorders of Oxidative Phosphorylation
 1,000 proteins (50% of the mitochondria is protein)
 Usually autosomal recessive
 - Expressed in all tissues
 - Phenotypic expression with high ATP demand

OxPhos Diseases (Genetic)

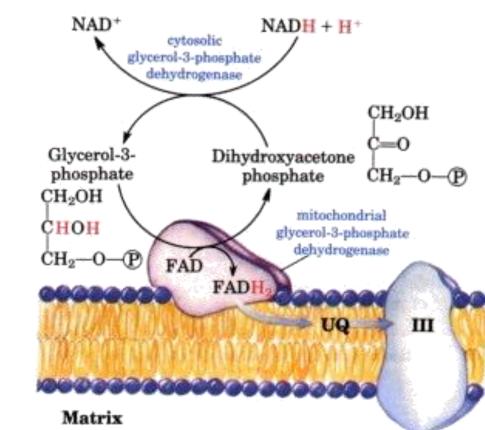


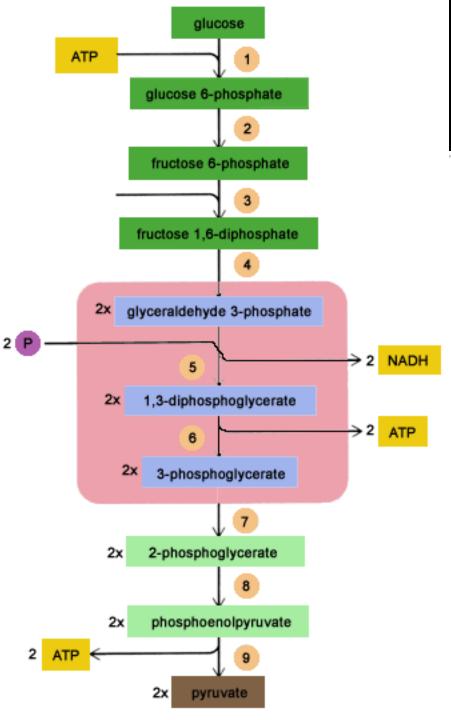
Disorders due to mutations in nDNA-encoded proteins

Mitochondrial shuttling systems "Cytosolic NADH"

- Glycerol 3-phosphate shuttle (Sk. muscles & brain)
- Glycolytic pathway as an example
- How NADH passes?

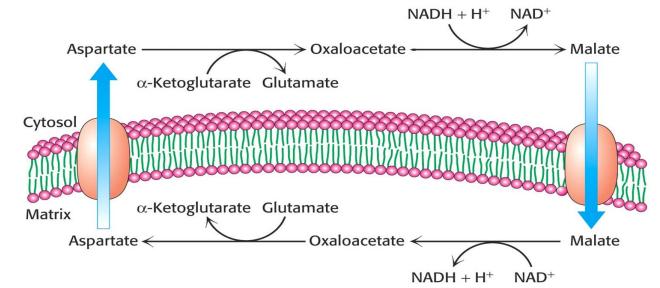
> ATP yield?





Mitochondrial shuttling systems "Cytosolic NADH"

- Malate-Aspartate shuttle
- Heart & liver
- > 2 membrane carriers & 4 enzymes
- Readily reversible (vs. Glycerol 3-phosphate shuttle)
- NADH can be transferred only if the NADH/NAD+ ratio is higher in the cytosol than in the mitochondrial matrix
- Exchange of key intermediates between mitochondria & cytosol



Mitochondrial shuttling systems "ATP/ADP"

- ATP-ADP Translocase (also called adenine nucleotide translocase or ANT)
- The flows of ATP and ADP are coupled (ADP enters only if ATP exits, and vice versa)
- Highly abundant (14% of IMM proteins)
- Contains a single nucleotide-binding site (alternates)
- Similar affinity to ATP and ADP
- Endergonic (25% of ETC)
- Inhibition leads to subsequent inhibition of cellular respiration

