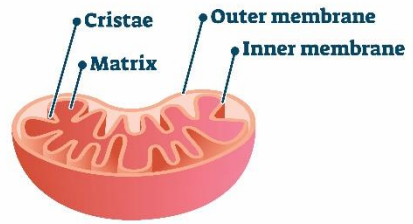
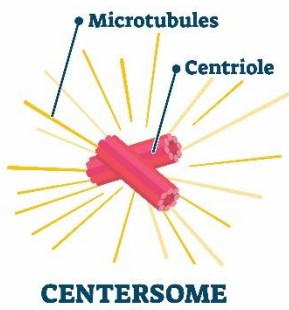
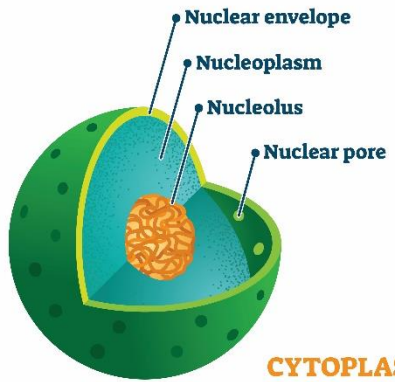
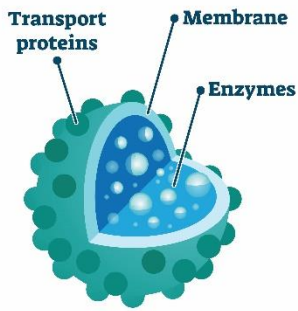
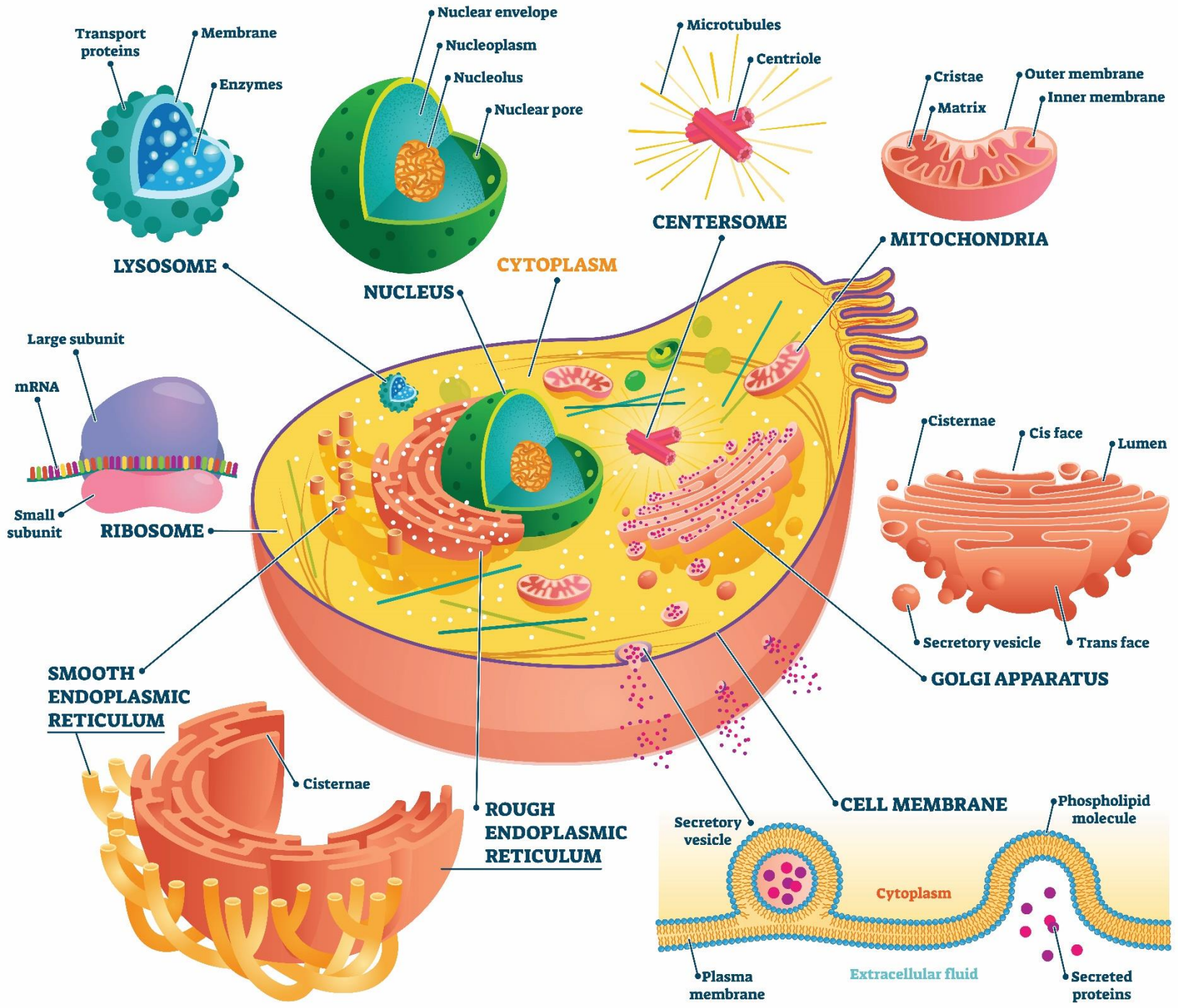


Cell structure and compartmentation of biochemical processes

1st week



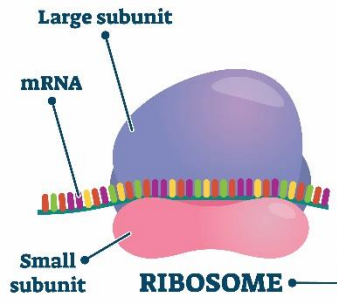
LYSOSOME

NUCLEUS

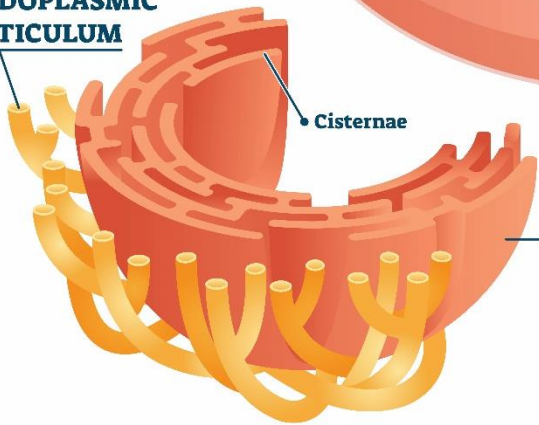
CYTOPLASM

CENTROSOME

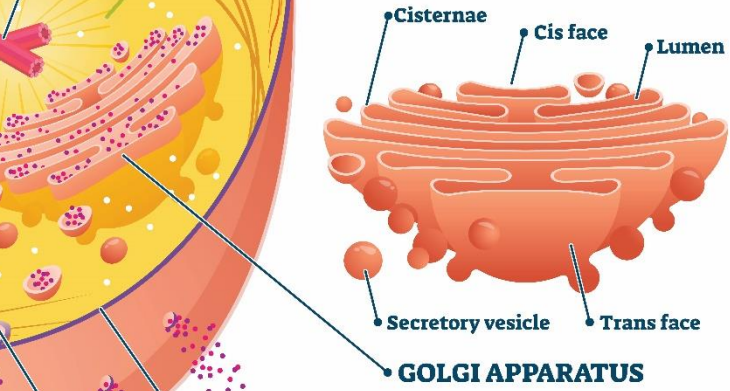
MITOCHONDRIA



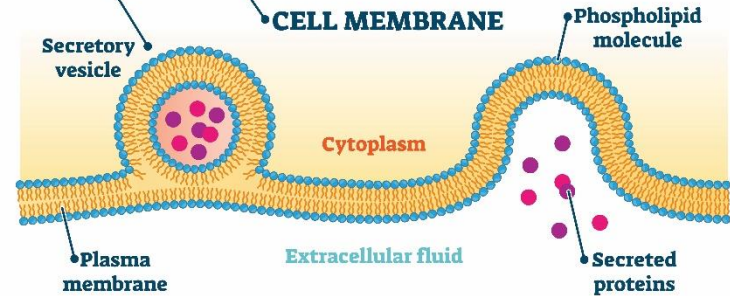
SMOOTH ENDOPLASMIC RETICULUM



ROUGH ENDOPLASMIC RETICULUM



CELL MEMBRANE



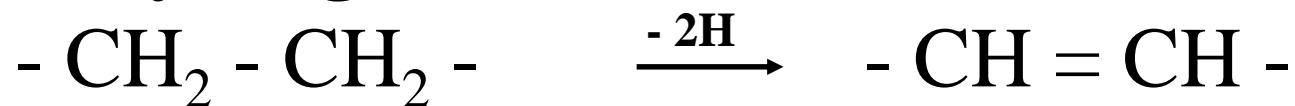
Biological oxidations

Types of oxidations

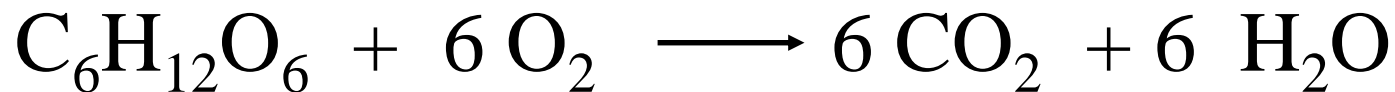
1. direct electron transfer :



2. dehydrogenation



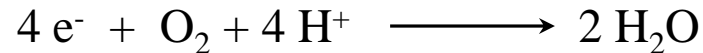
3. direct fusion with oxygen



Enzymes of redox reactions

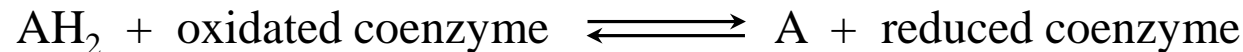
1. Oxidases

- take electrons from the substrate, which they transfer to an oxygen atom to form water



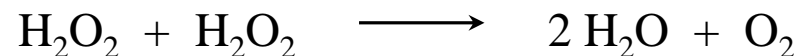
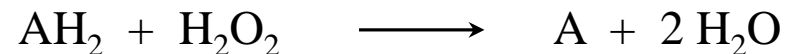
2. Dehydrogenases

- take 2 hydrogens from the substrate and transfer them to the dehydrogenase coenzyme, which is reduced in the process



3. Hydroperoxidases

- degradation of hydrogen peroxide in cells

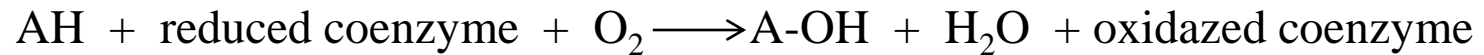


4. Oxygenases

- react with molecular oxygen, while 1 or 2 oxygen atoms are incorporated into the substrate molecule

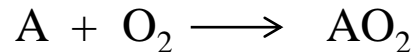
A. Monooxygenases

- catalyze the hydroxylation of the substrate, requiring a hydrogen donor (reduced coenzyme)



B. Dioxygenases

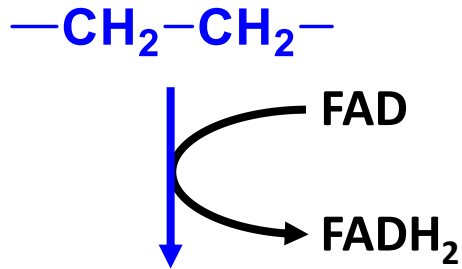
- catalyze the incorporation of the entire oxygen molecule into the substrate



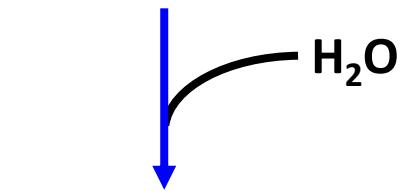
Coenzymes of redox processes

- nicotinamide: NAD^+ a NADP^+
- flavine: FAD a FMN
- oxidized coenzyme Q (ubiquinone)
- **by accepting 2 electrons, coenzymes change to their reduced forms :**
 - NADH a NADPH
 - FAD a FADH_2
 - reduced coenzyme Q (ubiquinol)

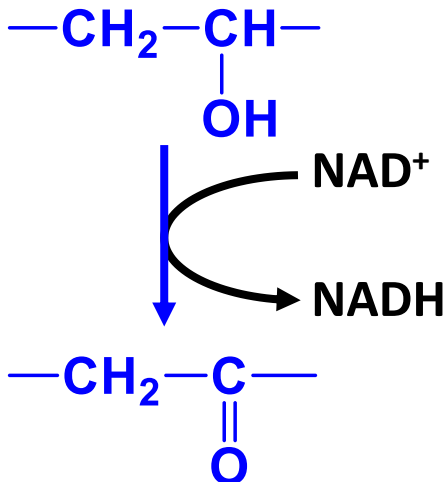
Oxidation of hydrocarbon chain *in vivo*



dehydrogenation
(double bond formation)



hydration
(hydroxy-group formation)



dehydrogenation
(oxy-group formation)

Diferences between oxidations

	<i>In vitro</i>	<i>In vivo</i>
Conditions	high temperature	„physiological“ conditions
Speed	rapid reaction	slow reaction
Release of energy	release of energy at one stroke, most as heat	transformation to chemical energy – creation of energy rich bonds
Course of reaction	always end products CO ₂ + H ₂ O	intermediates which can be used in other pathways
Mechanism of oxidation	C + O ₂ → CO ₂	dehydrogenation 2H + 1/2O ₂ → H ₂ O CO ₂ –decarboxylation

Macroergic compounds

Macroergic compounds

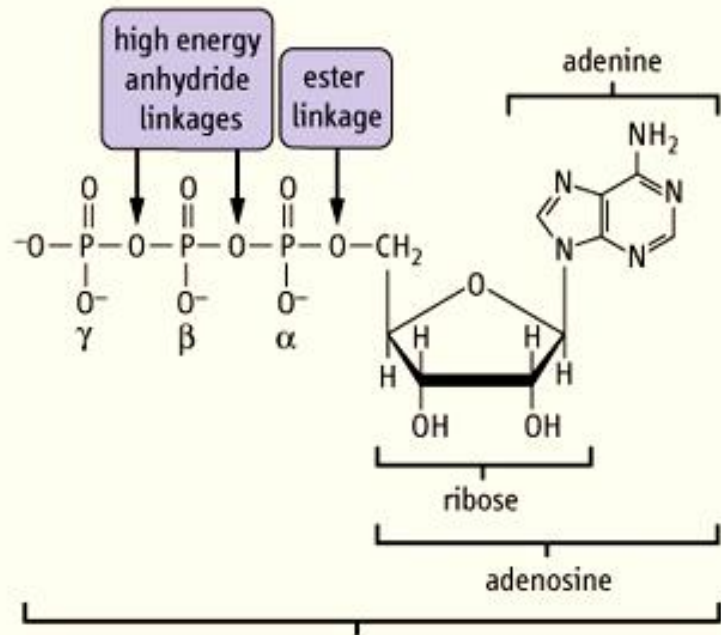
Energy rich compounds

1. **Endergonic metabolic processes** (synthesis of proteins, lipids, carbohydrates, nucleic acids)
2. **Transport processes** (active transport against the concentration gradient)
3. **Mechanical work** – muscles
4. **Nerve impulse transmission**

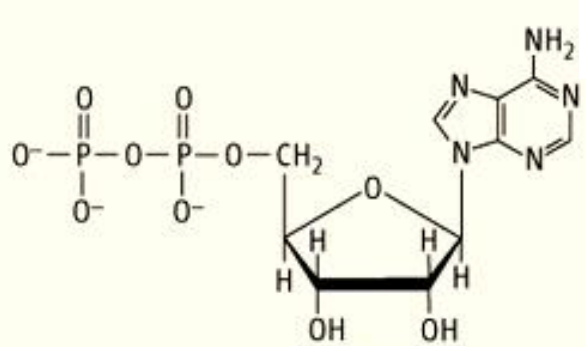
Types of macroergic bonds

1. phosphoanhydride (diphosphate)
2. enolphosphate
3. acylphosphate (carboxyphosphate)
4. guanidine phosphate
5. thioester

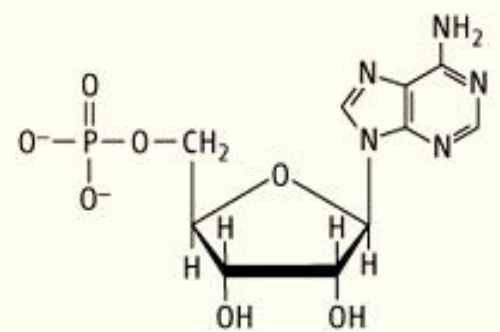
1. Phosphoanhydride bond (Diphosphate bond)



Other compounds:
GTP, GDP, UTP, UDP,
CTP, CDP, NAD^+ , FAD, CoA

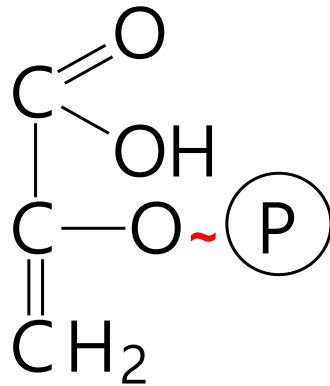


ADP



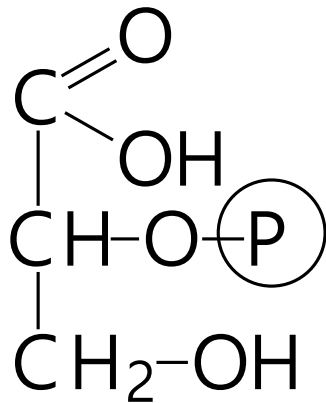
AMP

2. Enol phosphate bond

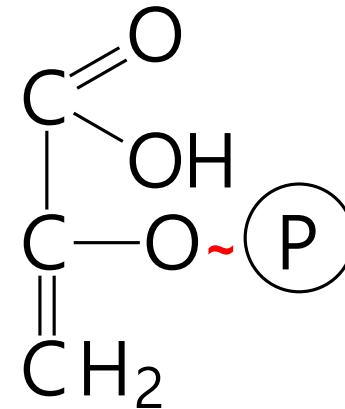
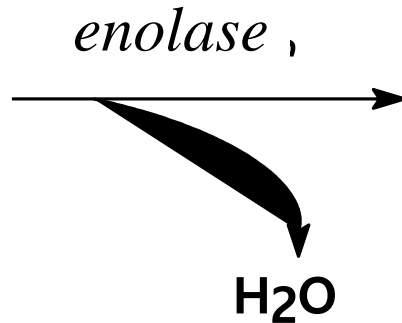


phosphoenolpyruvate (PEP)

Production: in glycolysis by *enolase*

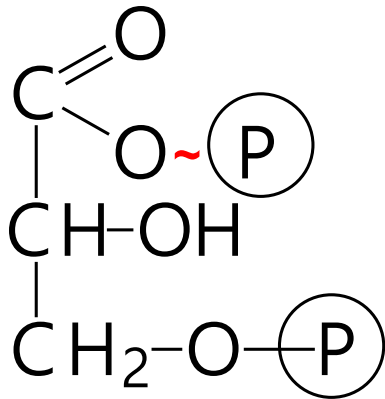


2-phosphoglycerate



phosphoenolpyruvate (PEP)

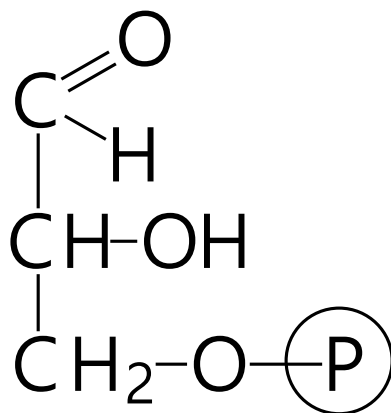
3. Acyl phosphate bond



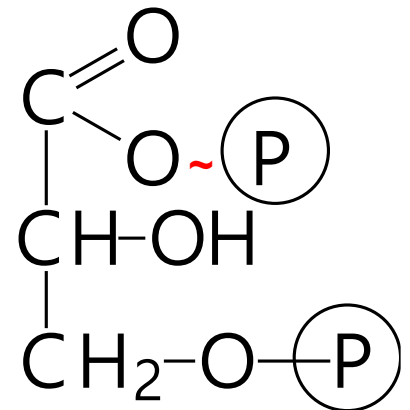
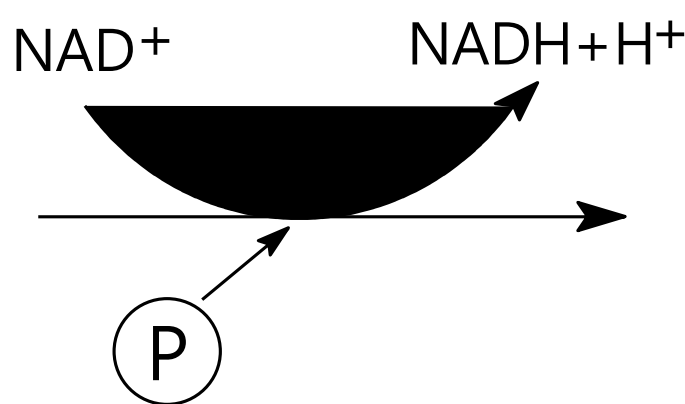
1,3-bisphosphoglycerate (1,3-bPG)

Other compounds: some intermediates in metabolism (aminoacyladenylate..), carbamoylphosphate

Production: in glycolysis by *glyceraldehyde dehydrogenase*

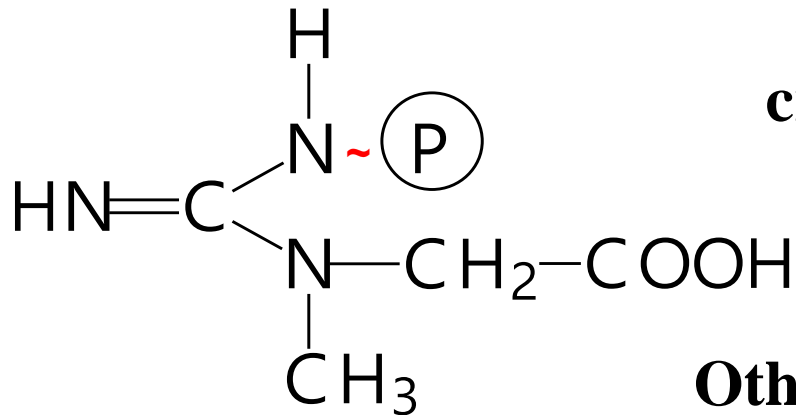


glyceraldehyde-3-P



1,3-bisphosphoglycerate (1,3-bPG)

4. Guanidine phosphate bond



creatine phosphate (Cr~P)

Other compound: arginine phosphate

Production: $\text{Cr} + \text{ATP} \rightarrow \text{Cr} \sim \text{P} + \text{ADP}$

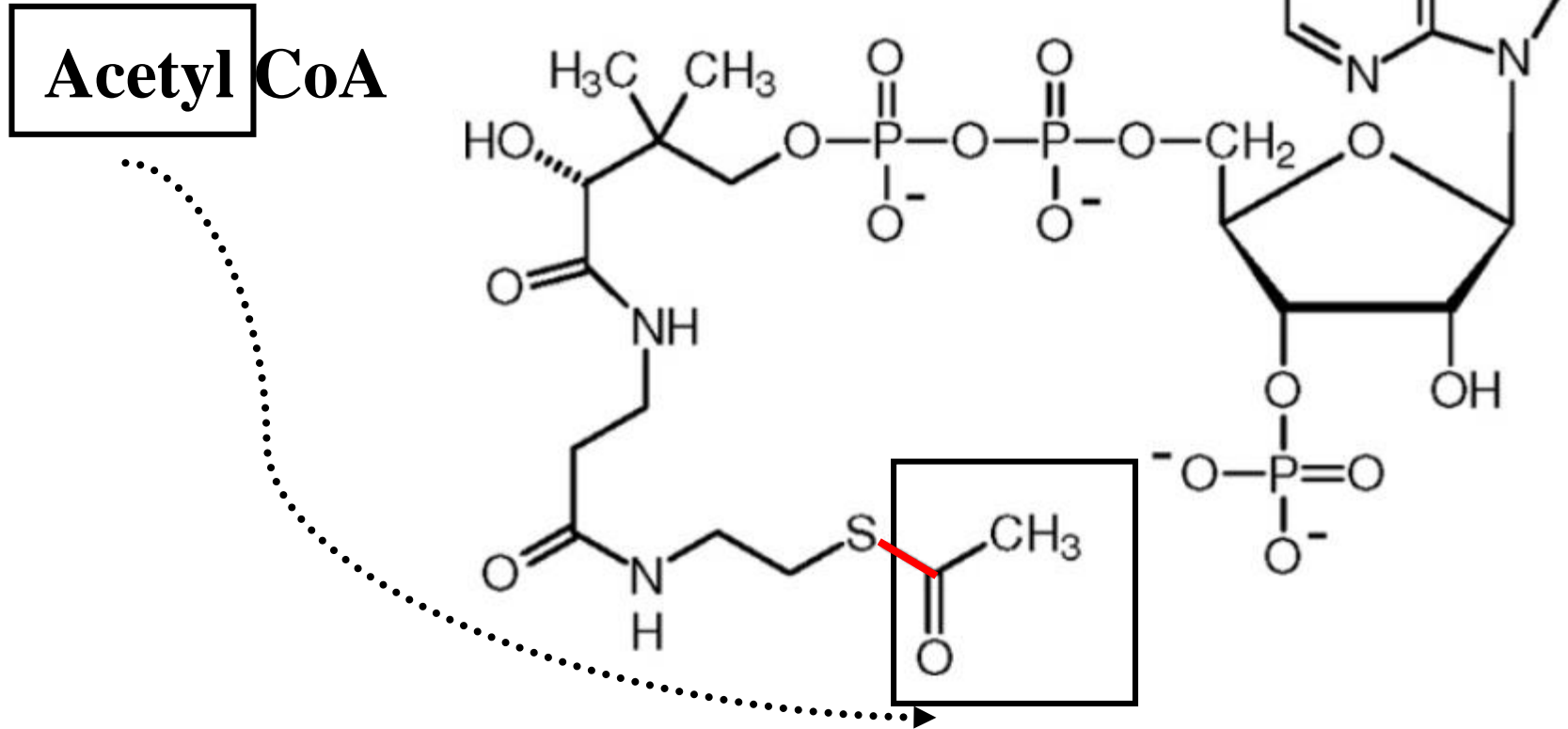
Utilization: storage form of energy, not a direct source!



direct source of energy

5. Thioester bond

(Activation of molecules)



Other compounds: acylCoA, some intermediates of metabolism – succinylCoA in Krebs cycle, intermediates of FA synthesis, S-acetyl lipoate in *pyruvate dehydrogenase* complex)

Aerobic / Anaerobic metabolism

