Take notes while watching the following video tutorials to prepare for the "Metabolism Part 1 Activity".

Carbohydrates Part 5:
Catabolism Stage 1 (Hydrolysis) & Stage 2 (Glycolysis)

An overview of catabolism

Stage 1 – Hydrolysis

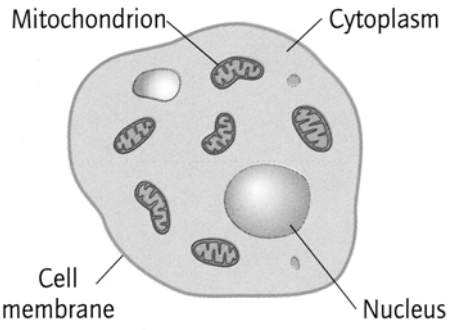
The 1st stage of carbohydrate catabolism is digestion – the breakdown into small molecules for energy production. Digestion begins in the mouth with the physical grinding, softening and mixing of food. Salivary α-amylase catalyzes the hydrolysis of the glycosidic bonds in carbohydrates. α-amylase is denatured in the stomach. Further digestion occurs in the small intestine where additional α-amylase is secreted by the pancreas along with maltase, sucrase and lactase from the mucous lining of the small intestine.

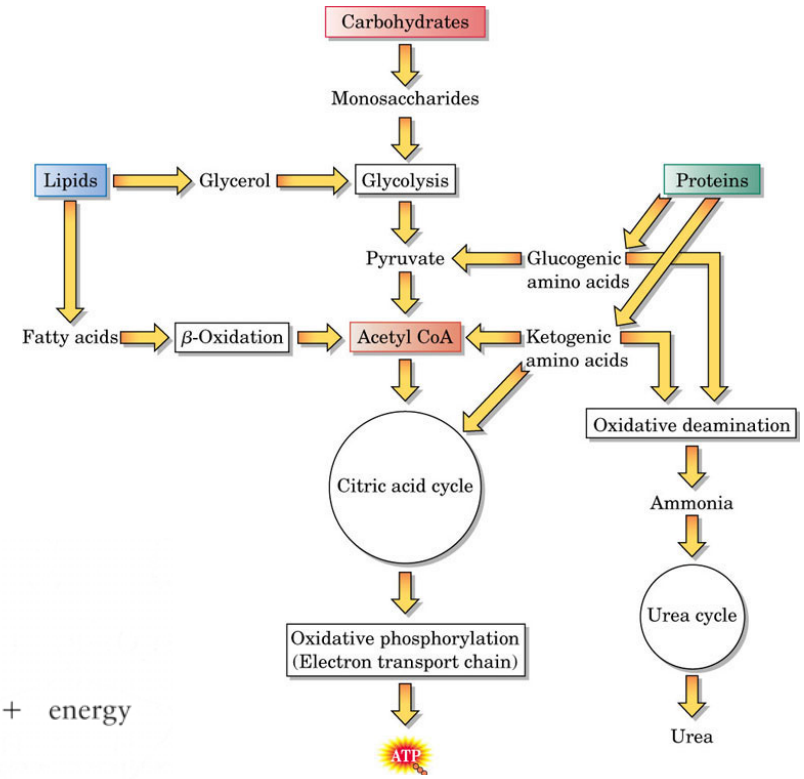
Stage 2 – Glycolysis

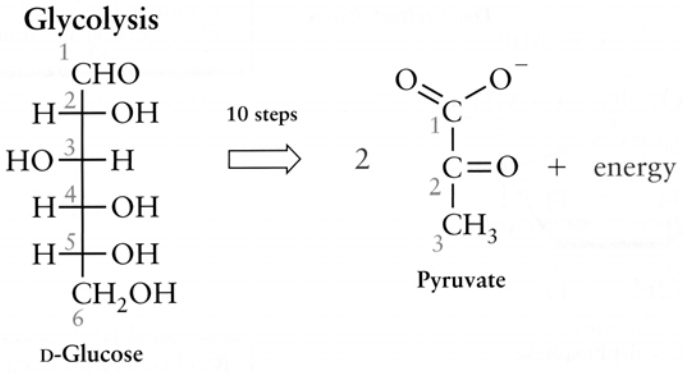
The conversion of glucose to pyruvate.

Stage 3 – Acetyl CoA enters Citric Acid Cycle to produce CO_2 & reduced coenzymes.

The reduced coenzymes transport the electrons to the electron transport chain (ETC) which ultimately reduces O_2 to H_2O and drives the formation of ATP from ADP & P1 in a process known as oxidative phosphorylation.



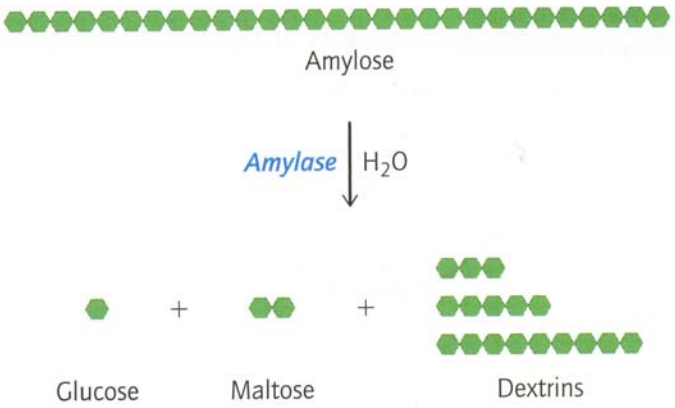




Stage 1

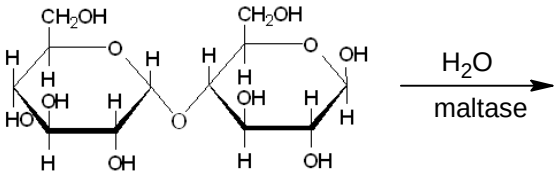
Hydrolysis of polysaccharides (starch, corn syrup & high fructose corn syrup)

breaking the glycosidic bond with water & enzymes



Hydrolysis of Disaccharides

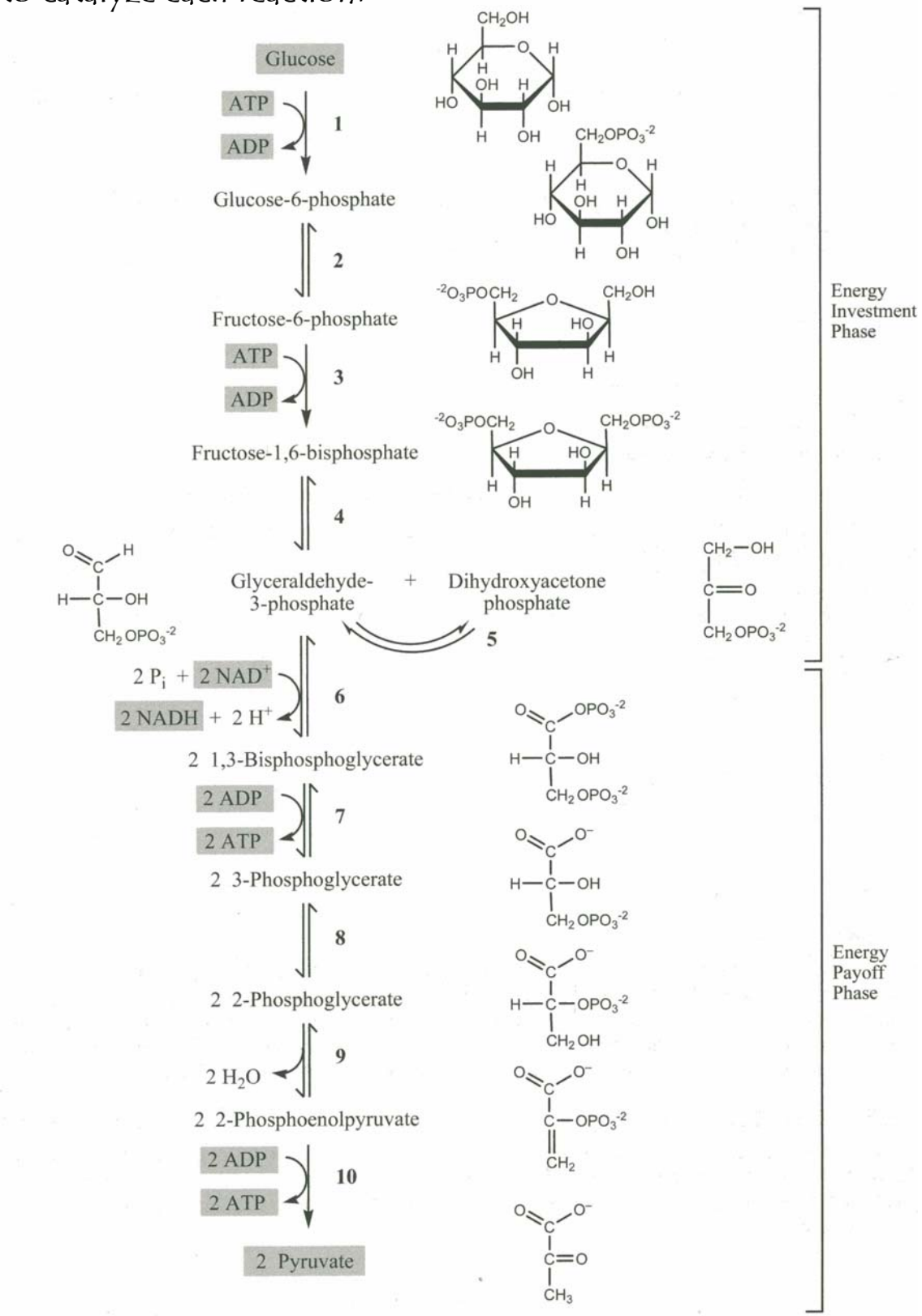
breaking the glycosidic bond with water & enzymes



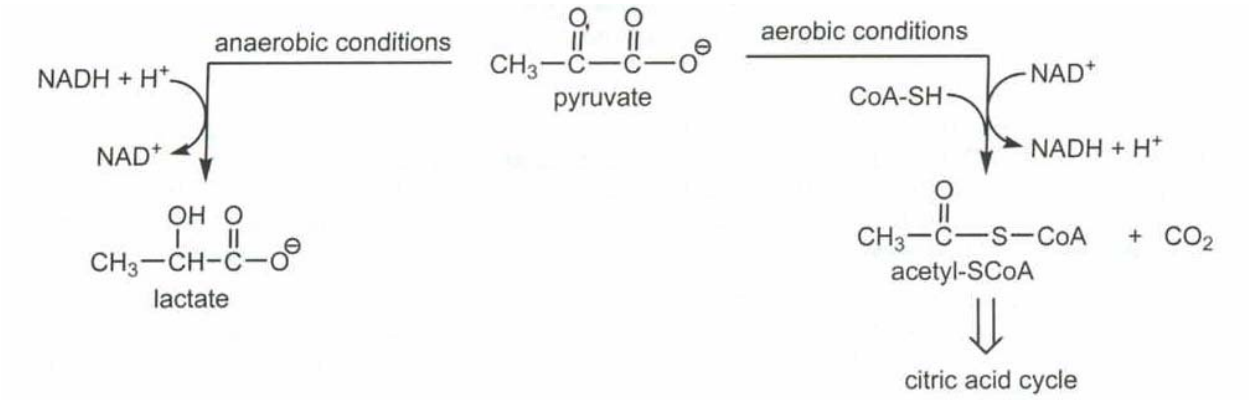
Draw the reaction for sucrose hydrolysis using Haworth projections. Name the products.

Stage 2 - Glycolysis

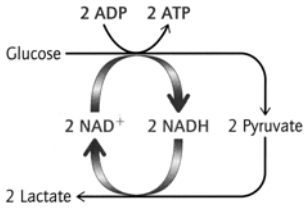
Look through the sequence of reactions in glycolysis and classify the enzymes (Hydrolase, Lyase, Ligase, Transferase, Isomerase, or Oxidoreductase) needed to catalyze each reaction.



Fate of Pyruvate



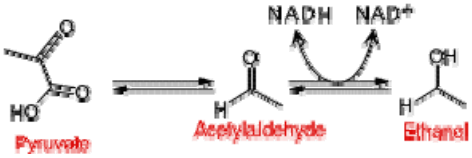
A lack of oxygen will slow down electron transport, causing a build-up of NADH and decreasing the amount of NAD^+ available for glycolysis. If there is no NAD^+, glycolysis cannot continue. The reduction of pyruvate to lactate will generate the NAD^+ needed for glycolysis (step 6).



Tissues with low oxygen content (such as skeletal muscle) rely on anerobic production of ATP by glycolysis.

Some bacteria can convert pyruvate to lactate under anaerobic conditions. The preparation of kimchee, sauerkraut, and yogurt involve these types of bacteria.

Yeast converts pyruvate to ethanol and CO_2 under anaerobic conditions.

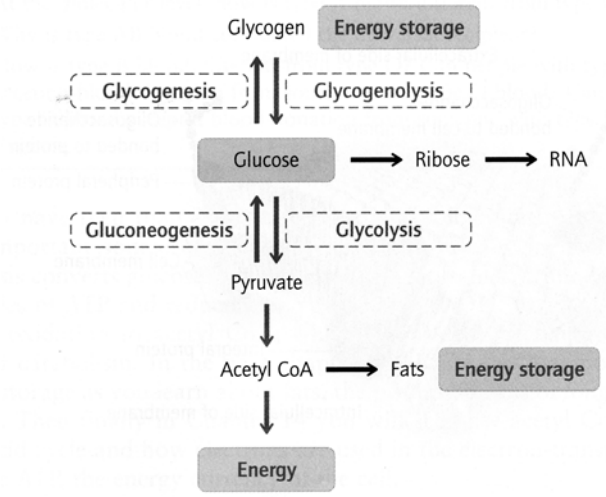


In the mitochondrial matrix, pyruvate is oxidized to form carbon dioxide and an acetyl group (acetyl-CoA).

Pyruvate must diffuse into the mitochondria from the cytosol. It is then transported by a membrane protein across the inner mitochondrial membrane into the matrix.

What happens to the acetyl-CoA?

Carbohydrates Part 6: Other Metabolic roles of Glucose & Ketone Bodies



Glycogenesis: synthesis of glycogen from glucose

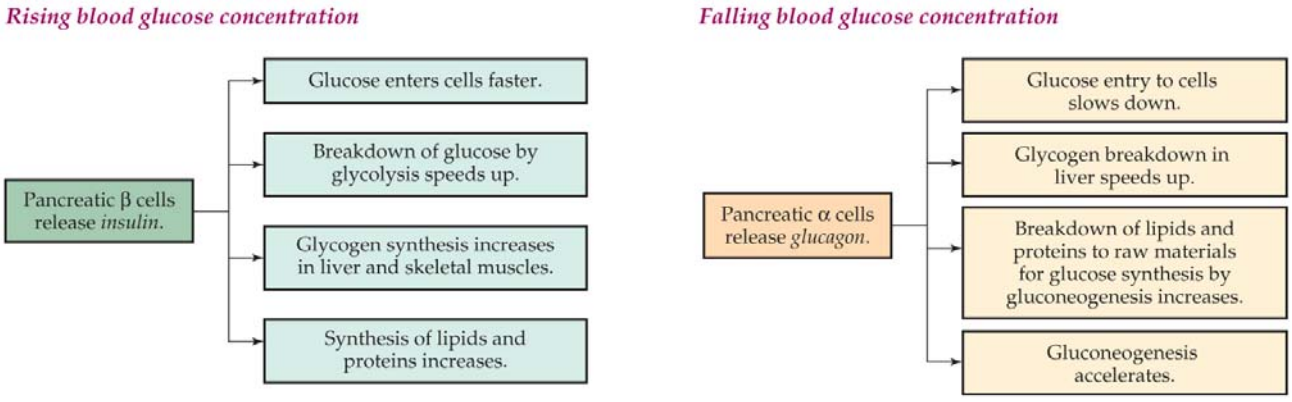
Glycogenolysis: Degradation of glycogen to produce glucose

Gluconeogenesis: Breakdown of proteins; in the liver, formation of glucose from amino acids

Glycolysis: breakdown of glucose to pyruvate

Regulation of Glucose Metabolism and Energy Production

Two hormones secreted by the pancreas play a major role in glucose metabolism.

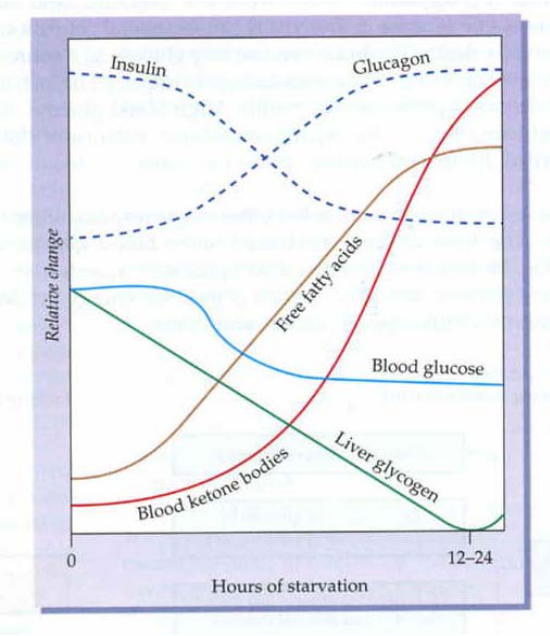


Metabolism during Fasting or Starving

Gradual decline in blood glucose triggers the catabolism of lipids causing a build-up of acetyl CoA.

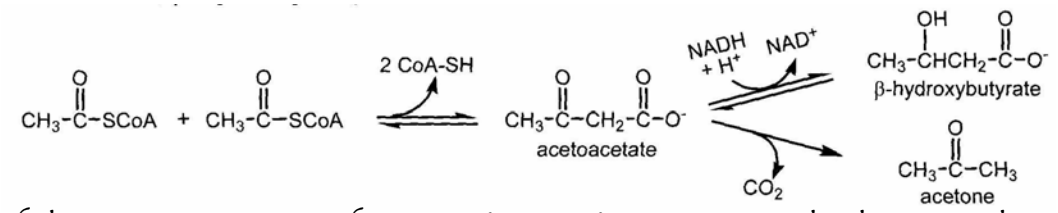
Ketone bodies form from the excess acetyl CoA. (see rxn below)

Brain and other tissues catabolize ketone bodies to produce ATP.



Diabetes Mellitus - glucose cannot be utilized or stored as glycogen because insulin is not secreted or does not function properly, therefore, insufficient amounts of glucose are present in tissues. Liver cells synthesize glucose from non-carbohydrate sources (gluconeogenesis) and break down fat. Acetyl-CoA builds up and ketone bodies accumulate (ketosis). Glucose appears in the urine because it is not undergoing glycolysis.

Formation of Ketone Bodies from Acetyl CoA



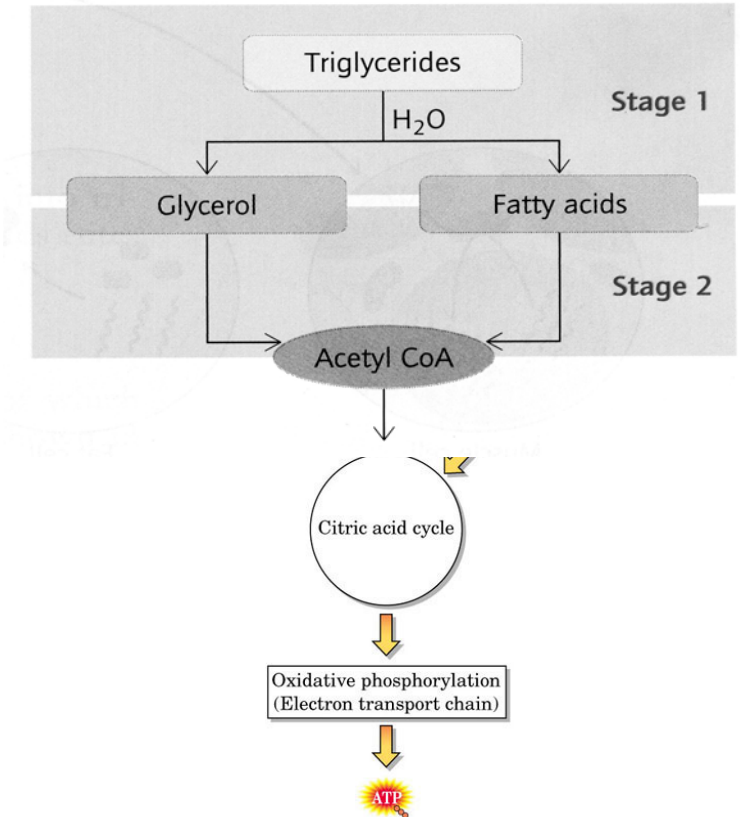
If the concentration of ketone bodies becomes too high, as can happen in diabetics or on a very low carbohydrate diet, a condition called ketoacidosis can result. Since the ketone bodies are not completely metabolized, acetone can diffuse out of the bloodstream into air in the lungs. The acetone can be smelled on the breath. Also, since ketone bodies acetoacetate and β-hydroxybutyrate are acidic, the pH of the blood decreases. This affects the ability of the hemoglobin to carry oxygen – breathing can become difficult. Untreated, ketoacidosis can lead to coma or death.

Lipids Part 7: Fatty Catabolism

Stage 1:

Stage 2:

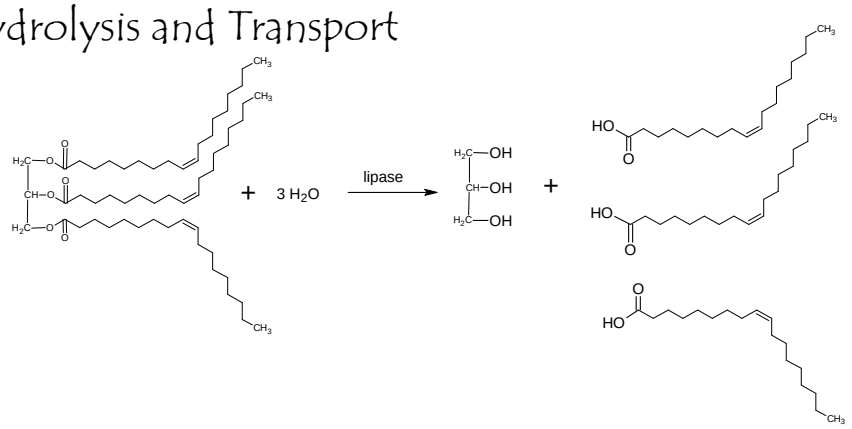
Stage 3:



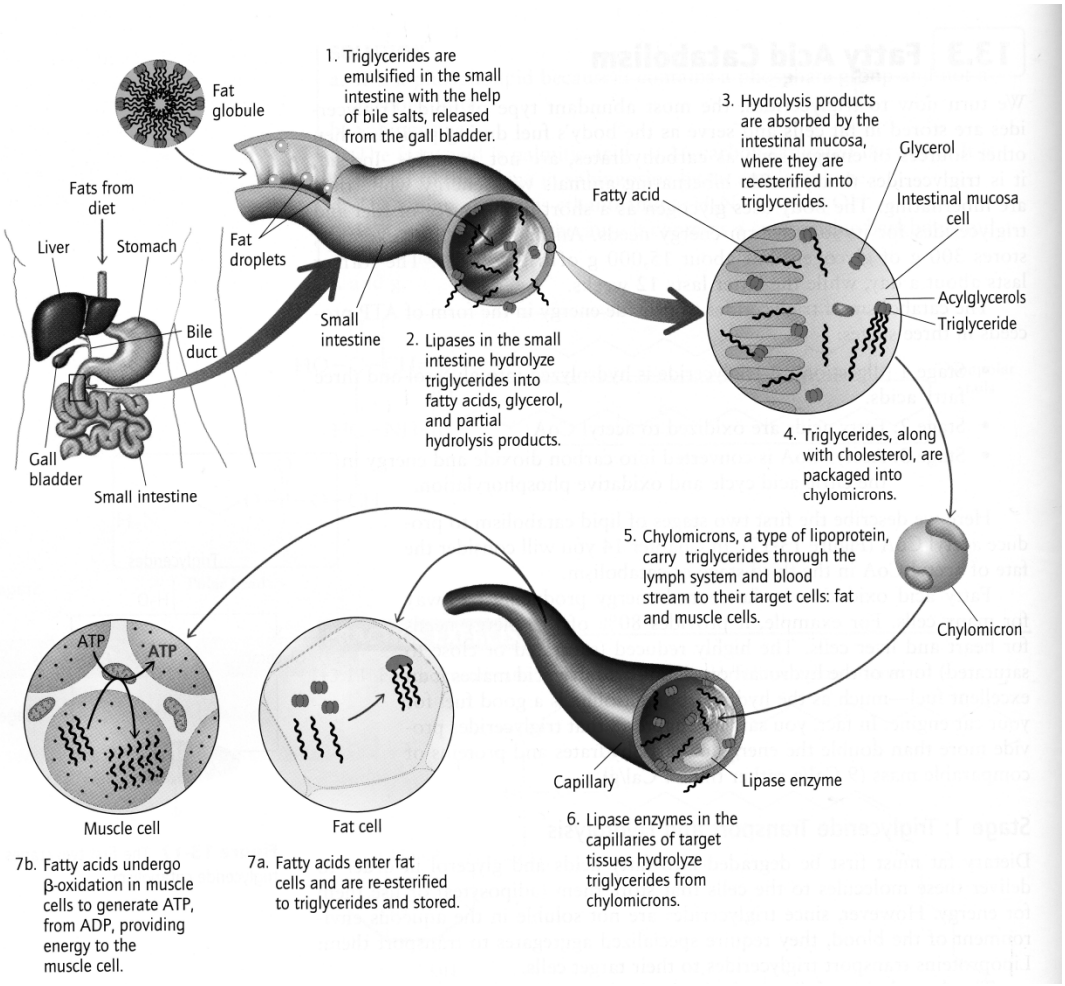
When eaten, fats pass through the mouth unchanged and enter the stomach.

How is this process different than carbohydrates?

Stage 1: Hydrolysis and Transport



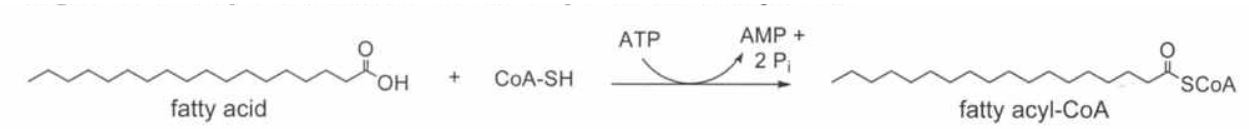
- The bile acids (salts) and phospholipids help emulsify the fatty acids so that the enzymes can breakdown the lipids.
- The hydrophobic lipids are transported by various lipoproteins.
- As the fats are hydrolyzed, the smaller fatty acids and glycerol are water soluble and are absorbed directly through the surface of the villi that line the small intestine.
- Chylomicrons surround the still-insoluble larger fatty acids within the intestine for further hydrolysis by lipases.



Stage 2: Fatty Acid Oxidation

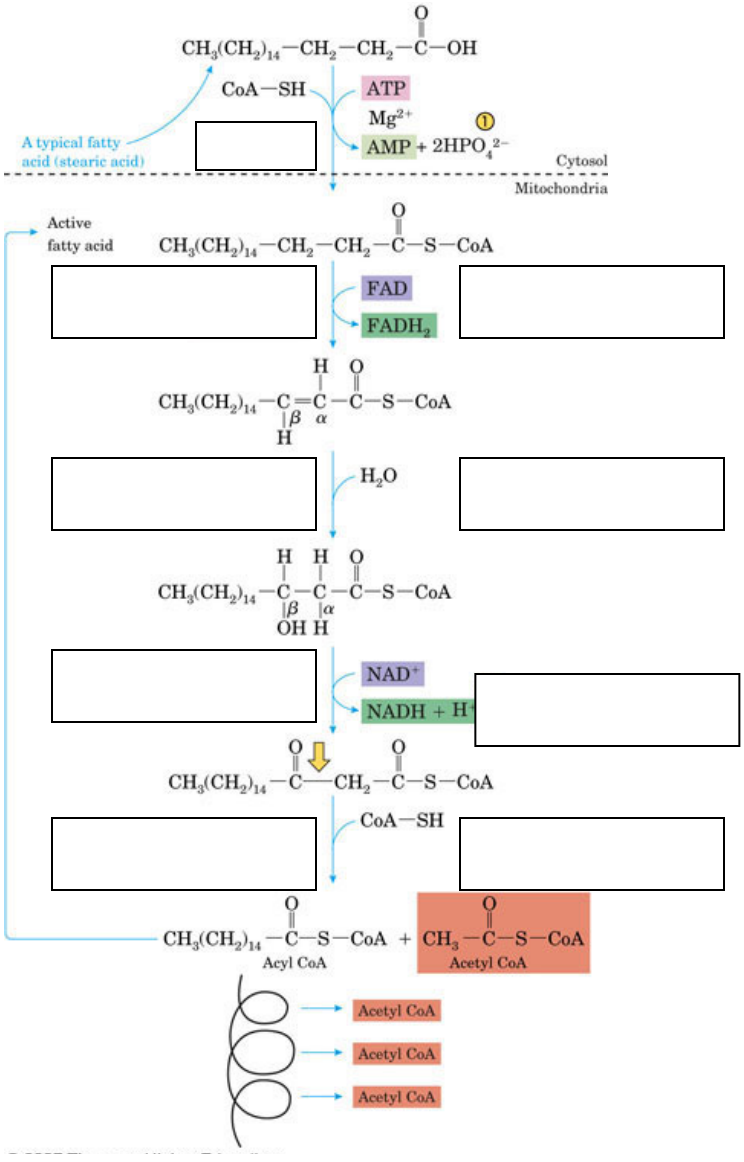
Once a fatty acid enters the cytosol of a cell that needs energy, three successive processes must occur:

1. Activation: The fatty acid is converted to Fatty Acyl-CoA.

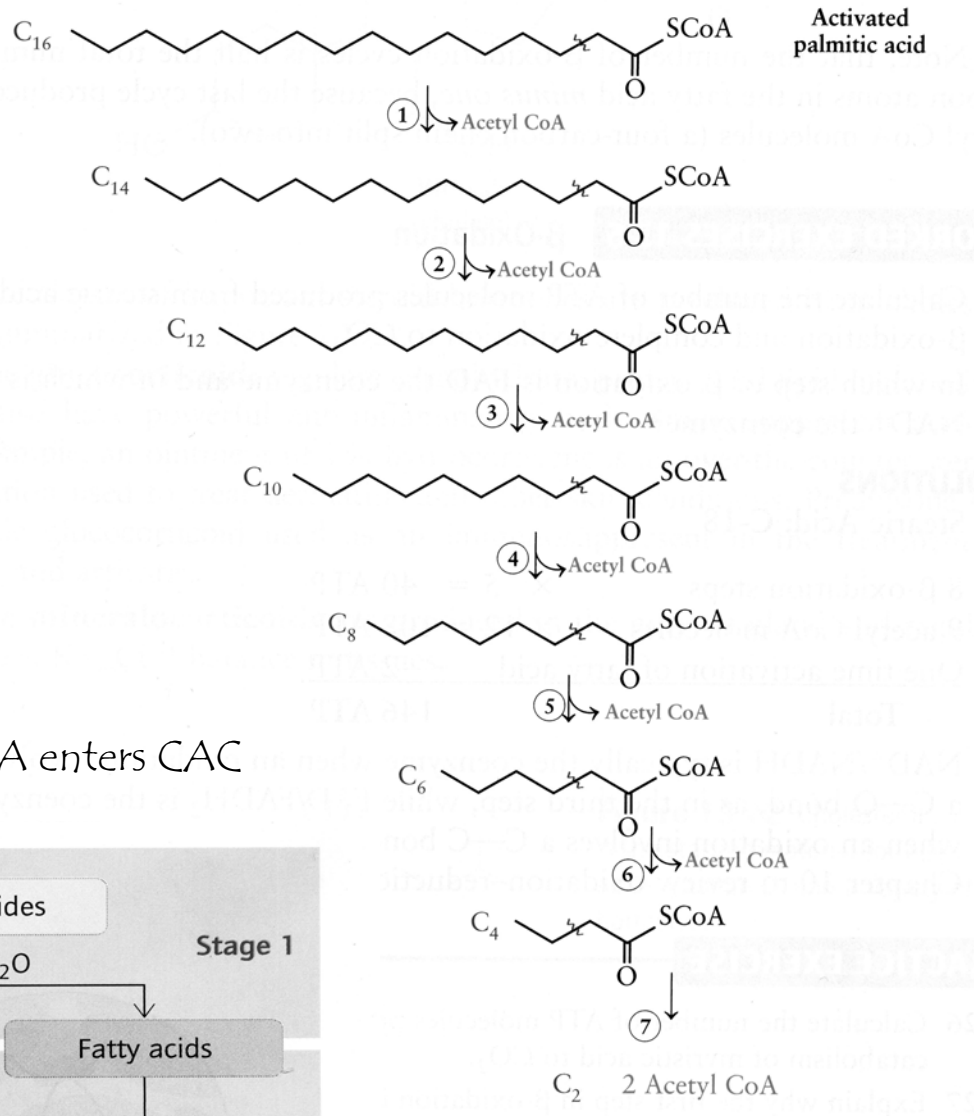


2. Transport: The Fatty Acyl-CoA is transported into the mitochondrial matrix where energy generation will occur.

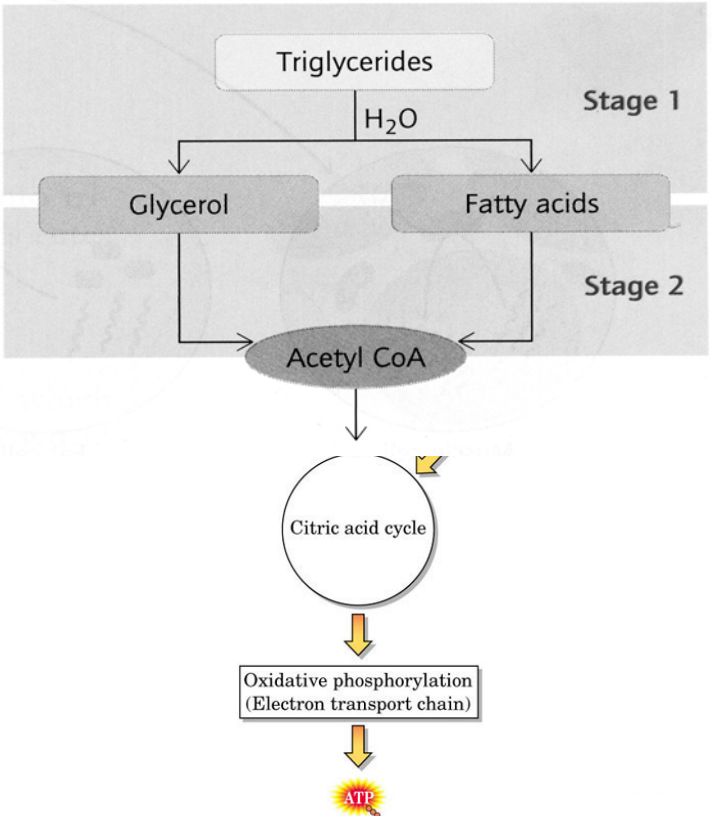
3. β-Oxidation: The Fatty Acyl-CoA is oxidized by enzymes to produce acetyl-CoA and reduced coenzymes. Each repetition of the oxidation cleaves a 2-carbon acetyl group.



Fatty acids undergo repeating cylcles of β-oxidation until the final four-carbon fatty acyl CoA molecule is converted into two acetyl CoA molecules.



Stage 3: Acetyl CoA enters CAC



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