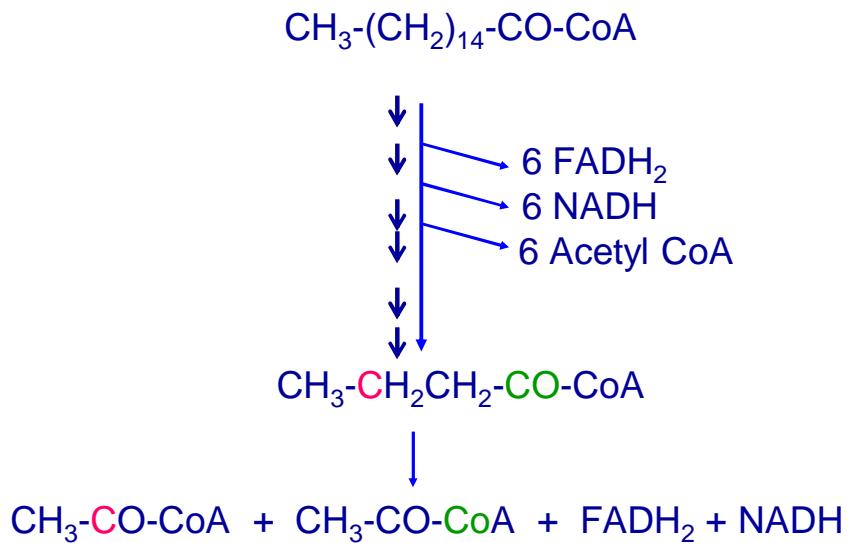


## Energy Yield from FA Oxidation



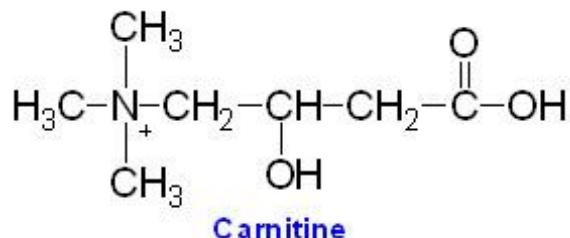
## Energy Yield from FA Oxidation (cont.)

- Oxidation of C 16 FATTY ACID

$$\begin{array}{lcl} - 7 \text{ FADH}_2 & \rightarrow & 14 \text{ ATP} \\ - 7 \text{ NADH} & \rightarrow & 21 \text{ ATP} \\ - 8 \text{ Acetyl CoA} & \rightarrow & 96 \text{ ATP} \end{array}$$

- Activation of the Acid consumes 2 ATP
- Net 129 ATP mole per mole of C16 Fatty Acid

# Carnitine



- \* Other functions:

- Export of branched chain acyl groups from mitochondria
- Excretion of acyl groups that cannot be metabolized in the body

## Carnitine Deficiencies

- Secondary deficiencies:  
Liver disease, malnutrition, ↑ requirements
- Congenital Deficiencies:  
↓ Enzyme, ↓ uptake, ↓ tubular reabsorption
- ↓ Ability to use FA as a fuel
- Accumulation of F.A and branched Acyl groups in cells



## Oxidation of unsaturated F.A: Oleic Acid

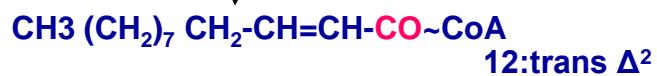


3 rounds of  $\beta$  oxidation

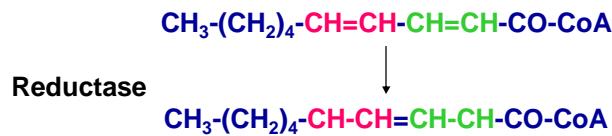
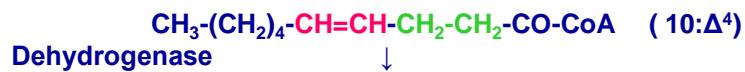
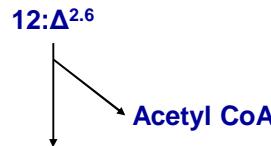
3 Acetyl CoA



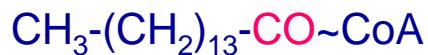
isomerase



## Oxidation of Unsaturated F.A: Linoleic Acid



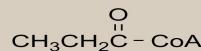
## Oxidation of FA with odd number of carbons



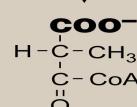
Six Cycles of  $\beta$  oxidation  $\downarrow$



Propionyl CoA



**Propionyl CoA**



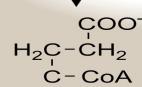
**D- Methylmalonyl CoA**

*Methylmalonyl CoA racemase*

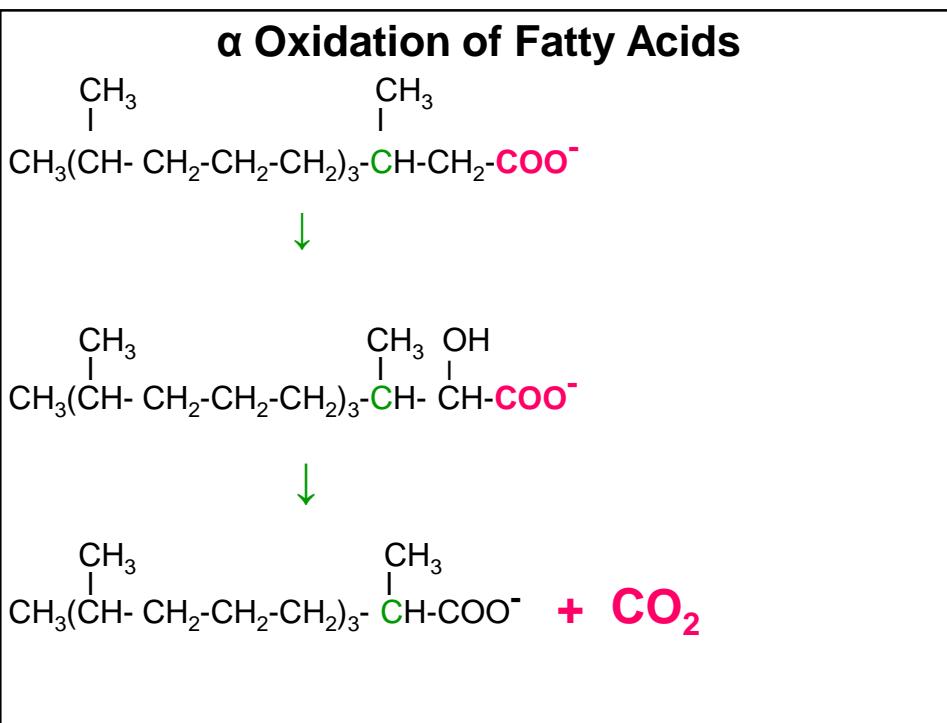
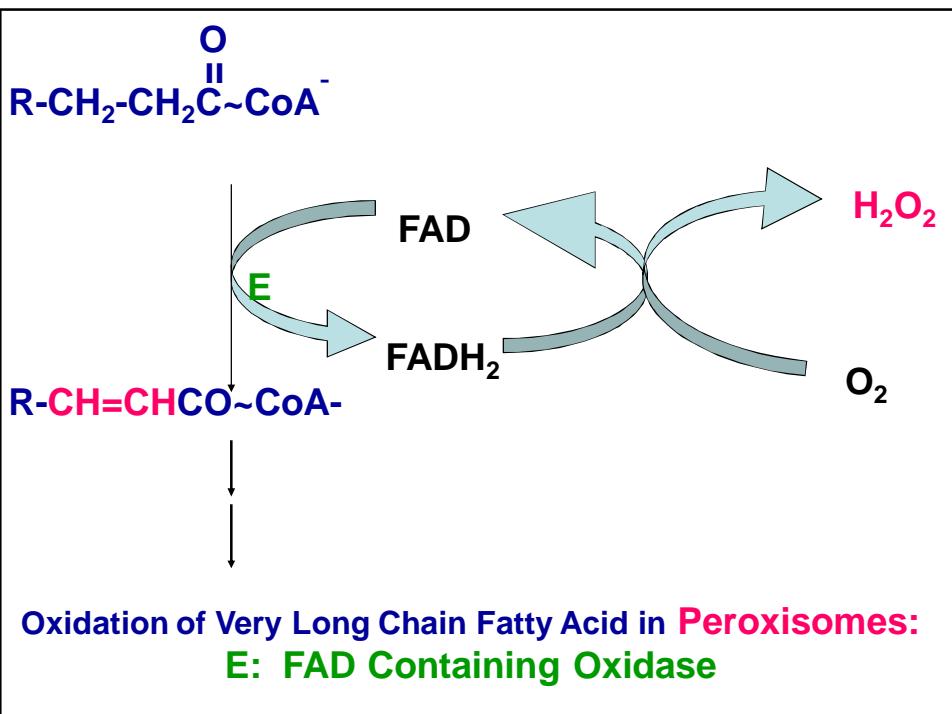
**L-Methylmalonyl CoA**

*Methylmalonyl CoA mutase*

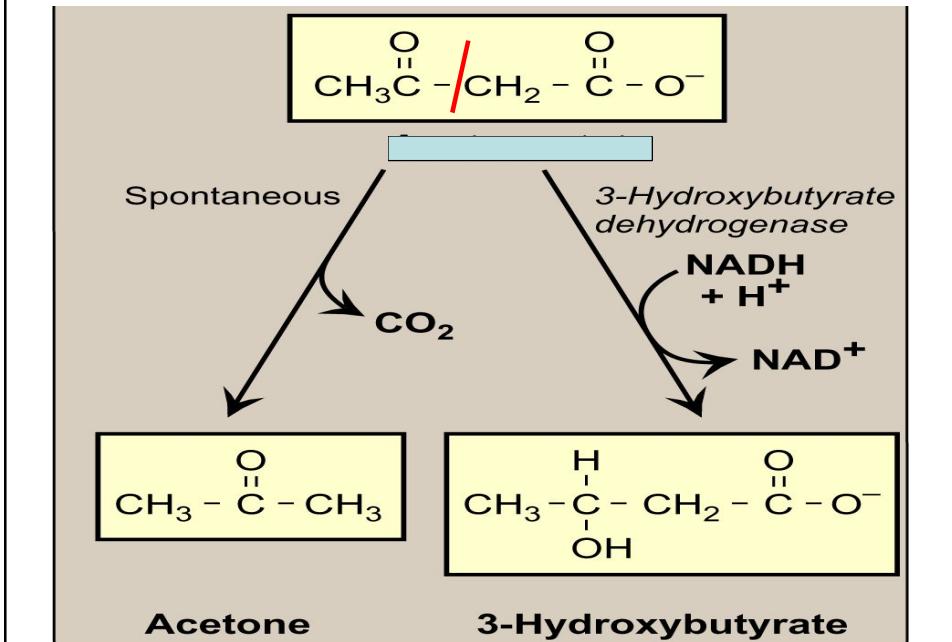
**Coenzyme form of vitamin B<sub>12</sub>**  
(Deoxyadenosyl cobalamin)



???



## Ketone Bodies



## Ketone Bodies

- Synthesis:  
In Liver
- Precursor:  
Acetyl CoA
- At high rate during:
  - Fasting
  - Uncontrolled Diabetes Mellitus

