

# Gluconeogenesis (Production of glucose from noncarbohydrate precursors)

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### **Glucose Synthesis is Required for Survival**

- Brain is dependent on glucose 120g/day
- Body glucose reserve is limited
  - $\approx$  20 g (extra cellular fluid)
  - $\approx 75~g\,$  ( liver glycogen); enough for 16 hours
  - ≈ 400 g (muscle glycogen); for muscle use only

Main source of energy for resting muscle in post-absorptive state

- 70 Kg man has ≈ 15 Kg fat
  - Fatty acids can not be converted to glucose
  - Utilization of FA is increased 4-5 X in prolonged fasting
  - In prolonged fasting;  $FA \rightarrow ketone bodies$  at high rate



### Where and when does gluconeogenesis occur?



- During an overnight fast, ~ 90% of gluconeogenesis occurs in the liver and10% by the kidneys
- During prolonged fasting kidneys become major glucoseproducing organs (40% of total glucose production)

# Entrance of substrates into gluconeogenesis Lactate — Pyruvate — Amino acids Amino acids ---> oxaloacetate Glycerol -----> Triosephosphates



### Reversing the irreversible steps

# 1. From pyruvate to phosphoenolpyruvate (PEP)

### Carboxylation of Pyruvate Produces Oxaloacetate (OAA)



# From OAA to PEP

#### Enzyme is found in both cytosol and mitochondria

- Acetyl CoA Pyruvate carboxvlase CO2 is activated and transferred to pyruvate The generated (with covalently by pyruvate carboxylase producing oxaloacetate. attached biotin) PEP in the mitochondria ATP ADP + P is transported Lysyl residue -0-0 of enzyme to the cytosol Pyruvate O-C-CH<sub>2</sub> Oxaloacetate by a specific Biotin Oxaloacetate NADH + H+ cannot cross transporter the mitochondrial membrane so it is NAD+ reduced to malate that can. Malate The PEP that MITOCHONDRION is generated CYTOSOL NADH + H+ NAD+ P-0-C-C-O GTP In the cytosol, malate in the cytosol is reoxidized to oxaloacetate, which is Oxaloacetate Malate Phosphoenolpyruvate requires the converted to phosphoenolpyruvate by PEP carboxykinase. transport of CO2
  - OAA from the mitochondria to the cytosol

### Reversing the irreversible steps

# 2. From fructose-1,6-bisphosphate to fructose-6-phosphate

### **Dephosphorylation of fructose 1,6-bisphosphate**



### Reversing the irreversible steps

### 3. From glucose-6-phosphate to glucose



### Dephosphorylation of glucose 6phosphate

- Bypasses the irreversible hexokinase reaction
- Only in liver and kidney
- Glucose 6-phosphate translocase is needed to transport G-6-P across the ER membrane

### Glucose 6-phosphatase in Endoplasmic Reticulum (ER)

Hint: Muscle lacks glucose 6-phosphatase, and therefore muscle glycogen can not be used to maintain blood glucose levels.

## Energy requirements of gluconeogenesis



# Regulation of gluconeogenesis

- Mainly by:
- 1. The circulating level of glucagon
- Glucagon lowers the level of fructose 2,6-bisphosphate, resulting in activation of fructose 1,6bisphosphatase and inhibition of PFK-1
- Inhibition of pyruvate kinase
- Glucagon increases the transcription of the gene for PEP-carboxykinase
- 2. The availability of gluconeogenic substrates



3.Slow adaptive changes in enzyme activity due to an alteration in the rate of enzyme synthesis or degradation, or both

# Regulation of gluconeogenesis

