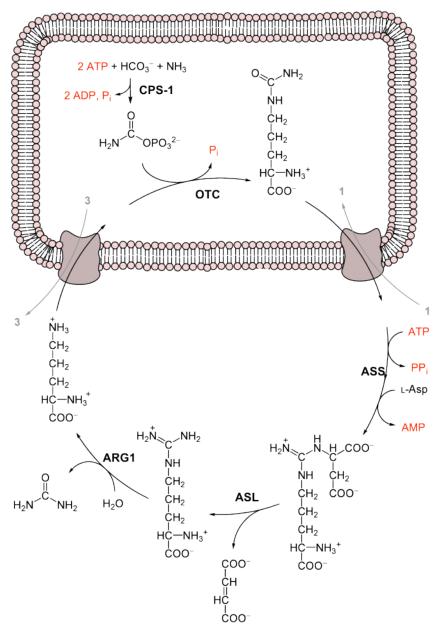
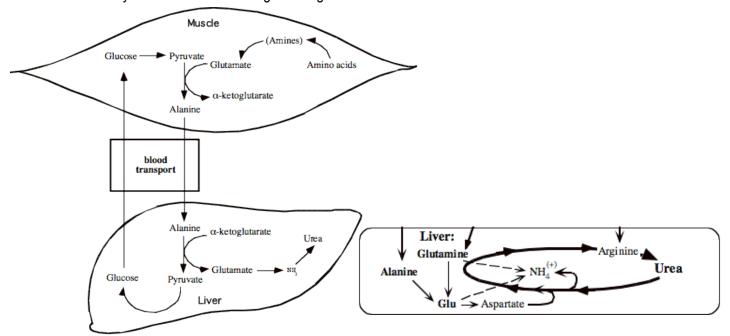
CONCEPT: AMINO ACID OXIDATION

- Urea cycle occurs in liver, removes amino groups from amino acids so they may enter the citric acid cycle
 - □ 2 nitrogen enter the cycle to ultimately leave the body as urea, and it costs 3 ATP
- 1. Carbamoyl phosphate is formed from HCO₃- + NH₄+, 2ATP is consumed in the process
- 2. Ornithine enters the mitochondria and combines with carbamoyl phosphate, releasing P_i
- 3. Citrulline moves back to cytosol and combines with Asp, ATP → AMP + PP_i (pyrophosphatase hydrolyzes into 2 P_i)
- 4. Arginosuccinate is cleaved into fumarate and arginine
- 5. Arginine is cleaved into urea and ornithine

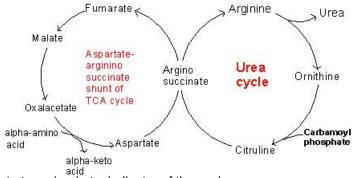


CONCEPT: AMINO ACID OXIDATION

- Glutamine synthetase makes glutamine to send to the liver (glutamate + ATP + NH₄⁺ → glutamine + ADP + Pi)
- Glutamine enters the mitochondria, and is broken down into glutamate and NH₄⁺ by glutaminase
- Glutamate dehydrogenase converts glutamate to α-ketoglutarate, releases NH₄⁺ and reduces NAD(P)⁺ → NAD(P)H
- Some glutamate is used to add NH₄+ to oxaloacetate, forming aspartate
- Glucose-alanine cycle occurs in muscles only, and can send alanine to liver
 - □ Convert pyruvate to alanine via a transaminase that transfers an amino group from glutamate
 - □ In liver, alanine is converted to pyruvate by transferring the amino group to α-ketoglutarate, forming glutamate
 - Pyruvate can be used for gluconeogenesis in the liver



• Fumarate can enter the citric acid cycle, and oxaloacetate can be converted to aspartate to enter the urea cycle



- Transaminases amino-keto to amino-keto, indicator of tissue damage
 - □ (S)GPT and (S)GOT indicate liver damage (S)CK indicate heart attack or infection
- N-acetylglutamate stimulates carbamoyl phosphate synthetase
- \square N-acetylglutamate synthase acetyl-CoA + glutamate \rightarrow N-acetylglutamate (stimulated by R)