

The the liver converts
a portion of the



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The liver carries many important physiological functions which include: processing of absorbed substances; synthesis and secretion of bile acids; bilirubin production and excretion; participation in metabolism of key nutrients, including carbohydrates, proteins, and lipids; and detoxification and excretion of waste products. LIVER METABOLIC FUNCTIONS The liver participates in carbohydrates, proteins, and lipids metabolism.

During carbohydrate metabolism, the liver performs gluconeogenesis, stores glucose as glycogen, and releases stored glucose into the bloodstream, when glucose level is low. During protein metabolism, the liver synthesizes non-essential amino acids and modifies amino acids so that they may enter biosynthetic pathways for carbohydrates. In addition, the liver also synthesizes almost all plasma proteins, including albumin and the clotting factors. Patients with liver failure develop hypoalbuminemia which may lead to edema due to loss of plasma protein oncotic pressure and clotting disorders. The liver also converts toxic ammonia which is byproduct of protein catabolism, to urea, which is later excreted in the urine.

When it comes to lipid metabolism, the liver participates in fatty acid oxidation and synthesizes lipoproteins, cholesterol and phospholipids. The liver converts a portion of the cholesterol to bile acids, which participate in lipid digestion and absorption. BILIRUBIN PRODUCTION AND EXCRETION The reticuloendothelial system (RES) processes senescent red blood cells. When hemoglobin is degraded by the RES, one of the byproducts is biliverdin which is later converted to yellow-colored bilirubin.

Later Bilirubin is bound to albumin in the circulation and carried to the liver, where it is taken up by the hepatocytes. In hepatic microsomes, bilirubin is conjugated with glucuronic acid via the enzyme UDP glucuronyl transferase. (Because UDP glucuronyl transferase is synthesized slowly after birth, some newborn babies develop "newborn jaundice".

“) Conjugated bilirubin is water-soluble and a portion of it easily is excreted in the urine. The remainder of the conjugated bilirubin is secreted into bile and then, via bile, into the small intestine. The conjugated bilirubin travels down to the terminal ileum and colon, where it is deconjugated by bacterial enzymes and metabolized to urobilinogen, some of which is absorbed via the enterohepatic circulation and delivered back to the liver; the remainder is converted to urobilin and stercobilin, which are excreted in the feces. BILE SECRETION Bile is necessary for the digestion and absorption of lipids in the small intestine. In contrast with carbohydrates and proteins, lipids pose special problems for digestion and absorption because they are insoluble in water. Bile, a mixture of bile salts, bile pigments, and cholesterol, solves this problem of insolubility. Bile is produced and secreted by the liver, stored in the gallbladder, and ejected into the lumen of the small intestine when stimulated to contract.

In the lumen of the intestine, bile salts emulsify lipids to prepare them for digestion and then solubilize the products of lipid digestion in packets called micelles. The hepatocytes of the liver continuously synthesize and secrete the constituents of bile. The components of bile include the bile salts, cholesterol, phospholipids, bile pigments, ions, and water. Bile flows out of the liver through the bile ducts and fills the gallbladder, where it is stored .

The gallbladder then concentrates the bile salts by absorption of water and ions. When chyme reaches the small intestine, CCK is secreted.

. In the small intestine, the bile salts emulsify and solubilize dietary lipids.

When lipid absorption is complete, the bile salts are recirculated to the

liver via the enterohepatic circulation . The steps involved in the

enterohepatic circulation include absorption of bile salts from the ileum into

the portal circulation, delivery back to the liver, and extraction of bile salts

from the portal blood by the hepatocytes . DETOXIFICATION OF

SUBSTANCES The liver protects the body from potentially toxic substances

that are absorbed from the GI tract. These substances are presented to the

liver via the portal circulation, and the liver modifies them in so-called “ first

pass metabolism,” ensuring that little or none of the substances make it into

the systemic circulation. For example, bacteria absorbed from the colon are

phagocytized by hepatic Kupffer cells and thus never enter the systemic

circulation.

In another example, liver enzymes modify both endogenous and exogenous

toxins to render them water soluble and thus capable of being excreted in

either bile or urine. Phase I reactions, which are catalyzed by cytochrome P-

450 enzymes, are followed by phase II reactions that conjugate the

substances with glucuronide, sulfate, amino acids, or glutathione