

# Liver and biliary tract disorders

## **Synthesis of plasma proteins**

Albumin, prealbumin, transferin, coagulations factors....

## **Detoxication reactions of endogen toxic substances**

Ammonia-urea cycle, bilirubin-clearance, bilirubin-conjugation

## **Detoxication reactions of exogen toxic substances**

Toxic substances from environment; drugs ....

## **Barriere function**

Barriere between gut (environment) and systemic circulation

## **Digestive functions**

Bile production;

(bile is a complex mixture of bilirubin, bile acids, cholesterol, lecithin and electrolytes)

**Metabolic center for glycid, fats and proteins metabolism**

# **Bilirubin**

is metabolic product of heme catabolism.

It is conjugated with glucuronic acid in the liver and excreted into the bile.

Bilirubin mono- and di- glucuronid is more water soluble and is easily excreted .

# Hyperbilirubinemia

Icterus

Jaundice

**prehepatic**

**hepatic**

**posthepatic**



**causes of hyperbilirubinemia**

# **Prehepatic causes of hyperbilirubinemie**

adults  
newborns

## Adults

Increased production of unconjugated bilirubin

Intravascular hemolysis

Hemolytic anemia

### *Laboratory diferencial diagnosis.*

Increas of unconjugated bilirubin

Urine: bilirubin neg.; urobilinogen posit.

Signs of anemie

Decreas of haptoglobin i plasma

## **Newborns**

Neonatal jaundice; icterus neonatorum; physiologic jaundice-icterus

Massive destruction of erythrocytes with fetal hemoglobin

Low activity of bilirubin glucuronyltransferase

Pathological icterus associated with Rh incompatibility



# **Hepatic causes of hyperbilirubinemia**

Alcoholic liver disease

Acute and chronic alcoholic hepatitis

Viral hepatitis A, B, C

Toxins (fungi, mushrooms))

Drugs

Autoimmune diseases.

## **Inborn diseases**

Gilbert syndrom (unconjugated hyperbil.)

Crigler-Najjar syndrome (unconjugated hyperbil.)

Dubin-Jonson syndrom (conjugated hyperbil.)

Rotor syndrome (conjugated hyperbil.)

# **Posthepatic causes of hyperbilirubinemia**

Cholestasis is defined as impaired bile flow from the liver to the intestine.

## **Intrahepatic cholestasis**

Intrahepatic cholestasis of pregnancy,  
drug induced cholestasis-phenothiazines, androgens  
Cholestatic forms of hepatitis

## **Extrahepatic biliary obstruction**

Mechanical obstruction to the bile flow – choledocholithiasis (gallstones)

Carcinomas of the gallbladder, pancreatic head, Vaters papilla

Liver flukes (parasits)

Inborn anatomic deformations of the biliary tract

- Biliary atresia

## **Tests of hepatic synthetic capacity**

The liver synthesizes a number of plasma proteins.

Total plasma protein

Albumin

Prealbumin

Coagulation proteins (factors)

# **Albumin.**

is synthesized exclusively in the liver.

## **Causes of hypoalbuminemia**

decreased production by the liver

protein malnutrition

loss from the skin, kidney, gastrointestinal tract

## **Prealbumin**

is synthesized by the liver  
acts as a transport protein for thyroxin and vit. A

short half-life (1-2 days)  
small body pool

Good and reliable indicator of hepatic synthetic capacity

# Liver enzymes

enzymes indicate hepatocellular damage

ALT, AST

enzymes indicate cholestasis and obstruction

ALP, GGT





