HEPATIC JAUNDICE - A REVIEW

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ARTICLE INFO

Aim : To do review on hepatic jaundice
Objective : To create an awareness and causes of hepatic jaundice
Background : Hepatocellular (hepatic) jaundice can be caused by acute or chronic hepatitis, hepatotoxicity, cirrhosis. Hepatic jaundice if not treated properly may cause permanent damage to liver. Thus an awareness on causes and prevention of hepatic jaundice is very essential.

INTRODUCTION

Jaundice, also known as icterus, is a yellowish or greenish pigmentation of the skin and whites of the eyes due to high bilirubin levels.[1][2] It is commonly associated with itchiness.[3] The feces may be pale and the urine dark.[4] Jaundice in babies occurs in over half in the first week following birth and in most is not a problem.[1][2] If bilirubin levels in babies are very high for too long a type of brain damage, known as kernicterus, may occur.[5] Causes of jaundice vary from non-serious to potentially fatal.[6] Levels of bilirubin in blood are normally below 1.0 mg/dL (17 μmol/L) and levels over 2-3 mg/dL (34-51 μmol/L) typically results in jaundice.[7][4] High bilirubin is divided into two types: unconjugated (indirect) and conjugated (direct).[8] Conjugated bilirubin can be confirmed by finding bilirubin in the urine.[9] Other conditions that can cause yellowish skin but are not jaundice include carotenemia from eating large amounts of certain foods and medications like rifampin.[4] High unconjugated bilirubin may be due to excess red blood cell breakdown, large bruises, genetic conditions such as Gilbert's syndrome, no eating for a prolonged period of time, newborn jaundice, or thyroid problems.[4][6] High conjugated bilirubin may be due to liver diseases such as cirrhosis or hepatitis, infections, medications, or blockage of the biliary tract.[4] In the developed world it is more often infections such as viral hepatitis, leptospirosis, schistosomiasis, or malaria.[4] Blockage of the bile duct may occur due to gallstones, cancer, or pancreatitis.[4] Medical imaging such as ultrasound is useful for detecting bile duct blockage.[9] Treatment of jaundice is typically determined by the underlying cause.[10] If a bile duct blockage is present surgery is typically required, otherwise management is medical.[10] Medical management may involve treating infectious causes and stopping medication that could be contributing.[10] Among newborns, depending on age and prematurity, a bilirubin greater than 4-21 mg/dL (68-360 μmol/L) may be treated with phototherapy or exchanged transfusion.[7] The itchiness may be helped by draining the gallbladder or ursodeoxycholic acid.[3] The word jaundice is from the French jaunisse, meaning "yellow disease".[11]

Types

On the basis of causes Jaundice can be classified into three types.4

- Pre-hepatic Jaundice
- Hepatic Jaundice
- Post hepatic Jaundice
- Pre-hepatic Jaundice

Pre hepatic jaundice is such type of jaundice which is caused due to hemolysis therefore it is also known as hemolytic jaundice. The major cause of enhanced hemolysis is defective plasma membrane of red blood cells. This vulnerable cell membrane cannot bear the shear stress and hence ruptures resulting in hemolysis thus causing the increased serum bilirubin level.16,17
Etiology

The pre hepatic jaundice is mainly caused due to hemolysis. The causes of pre-hepatic/hemolytic jaundice are classified into two groups:

**Congenital causes**

*Congenital causes of hepatic jaundice involve following:*19,20

- Spherocytosis
- Elliptocytosis
- Congenital LCAT deficiency
- Sickle cell anemia
- Stomatocytosis
- Acanthocytosis
- Echinocytes
- GSH synthase deficiency
- Pyruvate kinase deficiency
- G6PD deficiency
- Erythroblastosis fetalis

**Acquired causes**

*Acquired causes of pre-hepatic jaundice involve following:*19,20

- Resorption of extensive hematomas
- Auto immune hemolysis
- Transfusion reactions
- Trauma
- Microangiopathy
- Hemolytic uremic syndrome
- Long distance runners
- Disseminated intravascular clot
- Infections e.g. malaria, etc.
- Toxins e.g. snake venoms, etc.
- Chemicals e.g. nitrates, aniline dyes, etc.
- Paroxysmal nocturnal hemoglobinuria
- Thrombotic thrombocytopenic purpura
- Hypophosphatemia
- Vitamin B12 deficiency
- Folic acid deficiency

Clinical presentations

Patients with hemolytic jaundice are presented with Anemia, Yellowing of sclera, dark yellow-brown colored urine, yellowish skin and high bilirubin levels.21

Hepatic jaundice

Hepatic jaundice is a type of jaundice in which the basic defect lies within the liver mainly in the hepatocytes. The liver captures bilirubin from plasma proteins mainly albumin, then after conjugation excretes in the bile via biliary system. Any pathology of the liver leading to defect in capture, conjugation and excretion can cause hepatic jaundice. Main enzyme of conjugation is UDP- Glucuronyltransferase. This is commonly immature at birth and its under-activity can cause so called Neonatal Physiological Jaundice. Further this enzyme can be defective due to the genetic mutation of the UTG1A gene. on chromosome 2. This gene encodes for UDP- Glucuronyltransferase and thus the defective conjugating enzyme leads to the hepatic jaundice.22-24 Any defect in the hepatic excretory mechanism of bilirubin can also cause hepatic jaundice. The excretory mechanisms involve hepatic bile acid-independent secretion, hepatocytic bile acid-dependent secretion and bile ductular secretion. Any defect in the above mentioned excretory mechanisms can lead to the accumulation of bilirubin in blood causing hepatic jaundice.25-34.

Etiology

Hepatic jaundice is caused due to the defect in capture, conjugation and excretion of bilirubin by liver.35-38 Hepatic causes of the jaundice can be classified in to two types:

**Congenital causes**

*Congenital causes of hepatic jaundice are following:*38,39

- Wilson’s Disease
- Rotor’s Syndrome
- Haemochromatosis
- CriglerNajar syndrome
- Gilbert’s syndrome
- Dubin-Johnson’s syndrome

**Acquired causes**

*Acquired causes of hepatic jaundice are following:*38,39

- Viral Hepatitis
- Alcoholic Hepatitis
- Auto immune Hepatitis
- Drug related Hepatitis (e.g. NSAIDs)
- Sepsis
- Pregnancy
- Systemic Diseases (e.g. celiac disease)
- Malnutrition
- Physical Trauma
- Hepatic Adenoma

Clinical presentations

The clinical presentations of hepatic jaundice include abdominal pain, fever, vomiting and nausea along with the complications involving satiety, gastrointestinal bleeding, diarrhea, anemia, edema, weight-loss and associated weakness, if unchecked leading to mental disturbances like kernicterus, coma or even death.40

Post hepatic jaundice

Post hepatic jaundice is such type of a jaundice in which the cause lies in the biliary portion of hepatobiliary system. The major cause of post hepatic jaundice is extra- hepatic biliary obstruction. Therefore it is also Known as obstructive jaundice.41

Etiology

The major cause of post hepatic jaundice is extra-hepatic biliary obstruction.41 The causes of obstruction may be classified into two types:

**Congenital causes**

The congenital obstruction involves following:41,42

- Biliary Atresia
- Cystic Fibrosis
- Idiopathic dilation of common bile duct
- Pancreatic biliary malfunction
- Choledochal Cyst
Acquired Causes

The acquired obstruction involves following:
- Portal bilipathy
- Cholecystitis
- Trauma
- Pancreatitis
- Strictures
- Cholelithiasis
- AIDS
- Intra-Abdominal Tuberculosis • Tumors
- Common Abdominal Obstruction

Clinical presentation

The clinical manifestations of obstructive jaundice are dark urine, pale stools and generalized pruritus. History of fever, biliary colic, weight loss, abdominal pain and abdominal mass are also the representatives of obstructive jaundice. Obstructive Jaundice may lead to various complications including cholangitis, pancreatitis, renal and hepatic failure.

DISCUSSION

Differential diagnosis

The pre-hepatic jaundice can be differentiated from hepatic and post hepatic jaundice exclusively on the basis of elevated serum levels of unconjugated bilirubin and urobilinogen, which are raised in case of pre-hepatic jaundice. The serum levels on conjugated bilirubin, alkaline phosphatase, Alanine transferase and Aspartate transferase are seen normal in the case of pre-hepatic jaundice. The urinary excretion of conjugated bilirubin is also not present in pre-hepatic jaundice.

Hepatic jaundice can be differentially diagnosed from post hepatic jaundice on the basis of abdominal ultrasoundography and other radiological technique. However the hepatic jaundice can be differentiated from pre-hepatic jaundice on the basis of diagnostic markers, like alpha-1 Antitrypsin, Ceruloplasmin, Immunoglobulins, etc. Elevated serum bilirubin level along with the conjugation is a key diagnosis of post hepatic jaundice. Serum bilirubin is usually less than 20 mg/dL. In pancreatic cancer the serum bilirubin may rise up to 40 mg/dL. Serum gamaglutamyltranspeptidase (Serum GGT), alkaline phosphatase and transaminases may be elevated. Tumour markers like CA-125, CA19-9 and CEA are usually elevated in cancerous obstruction. The diagnosis of obstructive jaundice can further be confirmed by ultrasonography, plain abdominal x-ray, computed tomography, contrast-enhanced multi sliced computed tomography, endoscopic retrograde cholangiopancreatography (ERCP), Percutaneous trans-hepatic cholangiography (PTC), Endoscopic Ultrasound, Magnetic Resonance cholangiopancreatography (MRCP), Cholescintigraphy, Radionuclide scanning angiography and Staging Laparoscopy.

Therapeutic approaches and management

Pre-hepatic jaundice

Infusion of immunoglobulins is used as primary treatment for pre-hepatic jaundice. Phototherapy is considered as an effective treatment of high levels of bilirubin in pre-hepatic jaundice. Bilirubin rapidly decreases within two hours of onset of phototherapy. However the duration of therapy and the strength of light treatment depend upon the severity of hyperbilirubinemia. Metaloporphyrins are also considered as a treatment possibility of pre-hepatic jaundice, because these metaloporphyrins target the hemeoxygenase enzyme to limit the production of bilirubin.

Hepatic jaundice

Treatment and Management of hepatic jaundice involves

- Phototherapy - for neonatal jaundice.
- Phenobarbital can be used for treatment of neonatal physiological jaundice however it is not frequently used due to certain drawbacks involving somnolence and febrile seizures.
- Supportive therapy - fluids, rest, pain relief - for Hepatitis A.
- Abstinence from alcohol and cessation of medications contributing to liver dysfunction.
- Steroids - for autoimmune hepatitis.
- Immunosuppressant - for autoimmune hepatitis.
- Interferon - for chronic hepatitis B and C.
- Liver transplantation for fulminant hepatitis and end stage liver failure.

Post hepatic jaundice

Low fat diet should be given to patient suffering from post-hepatic jaundice to minimize the discomfort due to fat ingestion and diarrhea. The treatment of the post hepatic obstructive jaundice is mechanical decompression however the complications and other symptoms are also necessarily treated. Decompression can be done by surgical bypass, percutaneous insertion of stents, removal of lesions and endoscopic insertion of stents. Dexchlorophenramine, Hydroxyzine, Cholestyramine, Ursodeoxycholic acid and Naltrexone are used as a therapeutic approach in treatment and management of post hepatic jaundice.

CONCLUSION

Jaundice is very common disease. Yellowing of skin, sclera and mucous membranes are common manifestations of jaundice due to defect in production, metabolism and excretion of bilirubin. The causes of jaundice are either congenital or acquired. Serum bilirubin level and ultrasonography are used for differential diagnosis. High water intake and low fat diet are best proper managements of jaundice. The treatment of jaundice varies with the type of jaundice.

Reference

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