

Cirrhosis of liver

- It is a end stage liver disease characterized by fibrosis and regenerative nodule formation.
- Major Cause of Premature Death.

Causes

A - Alcohol, Non-Alcoholic Fatty liver Disease.

B - Primary Cholangitis.

Biliary cirrhosis, lytic fibrosis.

C - Cryptogenic, cholangitic venous outflow

obstruction, chronic liver Disease.

- Chronic Viral Hepatitis (B or C)

G1 - Genetic - Haemochromatosis

Wilson's Disease α_1 antitrypsin deficiency.

I - Immune \rightarrow Autoimmune liver disease.

Sclerosing cholangitis.

Pathophysiology

During recurrent injury

↓
Activation of stellate cells (usually stores Vitamin A) by cytokines by Kupffer cells & Hepatocytes.

↓
Transformation into myofibroblast.

↓
Collagen synthesis

↓
Fibrosis (Cirrhosis)

Portal

Hypertension

↓
Colones of living liver cells (Nodules)
surrounded by



Histologically classified into two

- i) Micronodular cirrhosis \Rightarrow $< 1 \text{ mm}$ in size
- ii) Macro nodular cirrhosis \Rightarrow Large nodules of various shapes

① Complication

Fibrosis \rightarrow ↑ Pressure \rightarrow ↓ Liver Function

① ↓ Detoxification \rightarrow ↑ Ammonia \Rightarrow BB

Hepatic Encephalopathy

② ↓ Estrogen Metabolism

\hookrightarrow Not excreted \rightarrow ↑ Estrogen in blood

i) Cystoconstrictio, Testicular atrophy, Impotence,

Unusual menses, amenorrhoea

ii) Spider angioma or spider naevus

\hookrightarrow Dilatation of arteriole mostly in

sun exposed areas, 1-2mm in diameter around nipple

iii) Palmar erythema

③ ↓ Bilirubin Conjugation

↑ Unconjugated Bilirubin

\downarrow Jaundice

④ ↓ Albumin production \rightarrow Hypoalbuminaemia

⑤ ↓ Clotting factors \rightarrow Haemorrhagic

⑥ Blood Backed Up in Spleen

\downarrow Splenomegaly

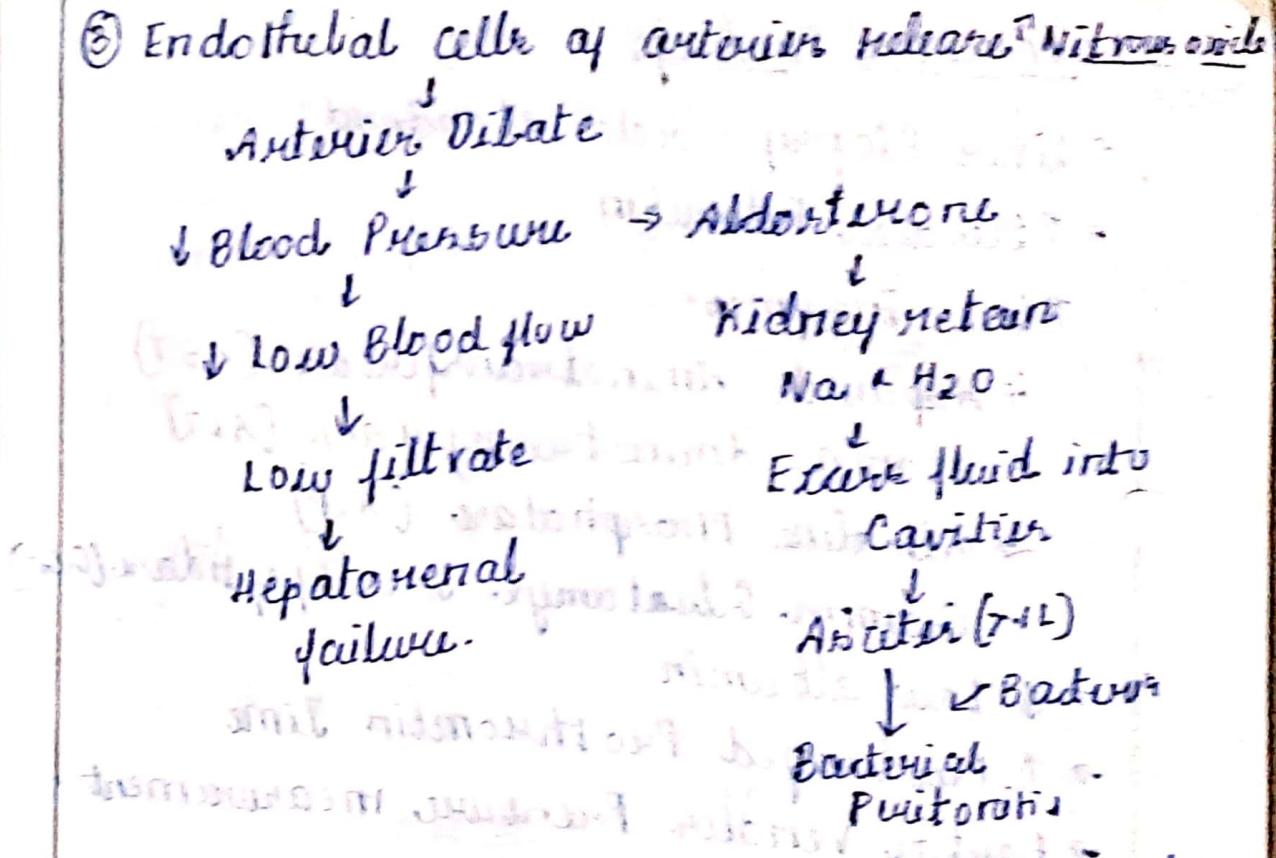
\downarrow Hypersplenism \leftarrow Anemia

Leukopenia

Thrombocytopenia



(Anemia) also leads to ↓ platelets \rightarrow ↓ clotting factors



④ Oesophageal varices, Haemorrhoids, Caput Medusae

Clinical features

- 1) Early → still do the work (Compensated)
 - ↳ Non specific (Weight loss, weakness, fatigue)
 - Asymptomatic

2) Later → Decompensated

- 1) Jaundice → Pruritis, Yellow skin.
- 2) ~~As~~ Hepatic Encephalopathy → Asterixis, Coma.
Altered Consciousness, Confusion, lethargy.
- 3) Men - Gynecomastia, Impotence, Testicular atrophy
Women - Irregular menses, Amenorrhoea
- 4) Haemorrhagic tendency → Purpura, Early bruising, Epistaxis
- 5) Spider Naevi, Palmar Erythema,
- 6) Anemia, Infection, Bleeding
- 7) Ascites, 8) others → Pigmentation, Dupuytren's Contracture

→ Liver Biopsy → Gold standard

→ ↑ Elevated Bilirubin

↑ Liver Enzymes.

→ Aspartate Aminotransferase (AST)

→ Alanine Aminotransferase (ALT)

→ Alkaline Phosphatase (ALP)

→ Gamma-Glutamyl Transpeptidase (GGT)

↓ Total albumin

→ ↑ Prolonged Prothrombin Time

→ Portal Venous Pressure measurement

Prognosis

5 to 10 years

adult family

adult disease (childhood) & non-hereditary

hereditary (adult)

child walls & adults & children

adults & children & progressive fibrosis

adults (adolescence) & children (adolescence)

adults (adolescence) & children (adolescence)