



Chronic liver disease



Learning outcomes

- LO1: Definition of chronic liver disease
- LO2: Causes of chronic liver disease
- LO3: Clinical features of chronic liver disease
- LO4: Diagnosis of chronic liver disease
- LO5: Staging of chronic liver disease
- LO6: Treatment of chronic liver disease
- LO7: Complications of chronic liver disease

LO1: CLD



- A cirrhosis is consequence of CLD characterized by replacement of liver tissue by fibrosis and regenerative nodules leads to irreversible loss of liver function and its complications
- Micronodular - alcohol
- Macronodular - chronic viral hepatitis



L02: Causes

- Alcoholic liver disease
- Chronic viral hepatitis- C or B
- Autoimmune hepatitis
- PBC and PSC
- Hemochromatosis, Wilson's disease
- α 1-antitrypsin deficiency
- Cystic fibrosis



LO3: Stigmata of CLD

- Pallor, jaundice
- Scratch marks
- Muscle wasting
- Parotid enlargement
- Xanthelasma
- Clubbing
- Palmar erythema
- Dupuytren's contracture
- Spider nevi
- Petechiae, purpura
- Decreased body hair
- Gynecomastia
- Testicular atrophy
- Caput medusa
- Edema, ascites
- Splenomegaly
- Asterixis
- Fetor hepaticus

Chronic Liver Disease



Hepatomegaly and Ascites



Cirrhosis



Caput medusae
(dilated veins around the umbilicus due to portal htn)



Gynecomastia
(impaired breakdown of estrogens)



Icterus
(increased bilirubin due to dysfunction of bilirubin metabolism)



Palmar erythema
(impaired breakdown of sex hormones)



Spider nevi
(isolated telangiectasias)



Ecchymosis
(defective coagulation)



Leukonychia
(hypoalbuminemia)



Finger clubbing



Asterixis
(abnormal motor fct due to faculty metabolism)

FEOTOR HEPATICUS

(characteristic odor due to volatile aromatic compounds)



Itching sensation



Swelling in the lower legs



Jaundice



Getting bruised easily



Fluid build up



Development of blood vessel





L04: Investigation

- LFT
- \pm elevated SGPT, alkaline phosphatase, GGT
- Increased bilirubin
- Low albumin, increased globulins
- Increased PT/INR
- Thrombocytopenia
- Low sodium
- Ultrasound- shrunken liver, \pm portal HT/HCC
- OGD- varices
- Liver biopsy- but not always necessary

LO5: Staging of CLD

- Based on Child- Turcotte -Pugh scoring system

- Includes- each given score of 1-3
 1. Ascites
 2. Encephalopathy
 3. Bilirubin
 4. Albumin
 5. PT/INR
- Class- total score
 - A. A- 5-6
 - B. B- 7-9
 - C. C- 10-15



LO6: Treatment

General measure

To retard progression and reduce complications

- Abstinence from alcohol
- Vaccination- Hepatitis C and B
- Treat underlying cause



L06: Liver transplantation

- Option in ESLD
- Donor, cost, technical expertise
- GVHD, recurrence



L07: Variceal bleed

- Varices- dilated submucosal veins, in esophagus or stomach
- Cause- portal HT
- Causes ~80% of UGI bleed in CLD
- Risk factors for bleed-
 - Size of varices
 - Severity of liver disease
 - Continued alcohol intake
- Dx- OGD



L07: Complications

- Ascites
- Spontaneous bacterial peritonitis- SBP
- Variceal bleed
- Hepatic encephalopathy
- Hepatorenal syndrome
- Hepatocellular carcinoma- HCC



LO7: Ascites

- Diagnostic paracentesis- SAAG >1.1
- Cause-
 - Portal HT
 - Hypoalbuminemia
 - Raised renin-angiotensin-aldosterone levels causing Na retention by kidneys



L07: Ascites- treatment

- Salt \pm fluid restriction
- Diuretics- Spironolactone \pm Furosemide
- Large volume paracentesis-
 - With massive or refractory ascites
 - >5 lit. fluid removed
 - Albumin- ~ 8 gm/lit. fluid removed
 - Avoid hepatorenal syndrome
- TIPS- transjugular intrahepatic portosystemic shunt
 - For refractory ascites or refractory variceal bleed
 - Preferred for short duration, pending liver transplant
 - Increases risk of hepatic encephalopathy, shunt occlusion/infection



LO7: SBP

- symptoms
 - Abdominal pain, fever, worsened ascites and encephalopathy
- Dx- paracentesis
 - PMN >250/ microliter
 - Ascetic fluid culture- bedside, commonly Gram –ve bacteria
- Rx- Cefotaxime/Ciprofloxacin
- Prophylaxis- Ciprofloxacin/Co-trimoxazole
- Prognosis- 30% mortality during hospital stay & 70% within 1 year



LO7: Treatment of Variceal bleed

- Acute-
 - Resuscitation
 - FFP, platelets, Vit. K
 - Terlipressin/ octreotide
 - Lactulose
 - →OGD→
 - Banding/ sclerotherapy
 - Balloon tamponade
 - TIPS
 - Surgery
- Prevent rebleed-
 - Band ligation- over repeated sessions
 - Non-selective β - blockers- Propranolol/ Nadolol
 - TIPS- for recurrent bleed or bleed from gastric varices
 - Surgery- portosystemic shunts
 - Liver transplantation



L07: Hepatic encephalopathy

- Confusion → drowsiness → stupor → coma
- Ammonia is an identified/measurable toxin
- Precipitants-
 - GI bleed
 - Constipation
 - Alkalosis, hypokalemia
 - Sedatives
 - Paracentesis → hypovolemia
 - Infection
 - TIPS
- Dx- clinical- s/s of CLD
with asterixis and altered sensorium



L07: Hepatic encephalopathy

- Correct underlying precipitating factor
- Avoid sedatives
- Restrict dietary protein intake
- Lactulose- 2-3 loose stools a day
- Oral antibiotic- Metronidazole, Rifaximin, Neomycin



L07: Hepatorenal syndrome

- Occurs in patients with advanced CLD & ascites
- Marked by renal impairment in the absence of any renal parenchymal disease or shock
- Oliguria, Hyponatremia & low urinary Na accompany raised creatinine
- Albumin infusion, with vasoconstrictors (norepinephrine, terlipressin /ornipressin, octreotide) may help
- Liver transplantation is Rx of choice



L07: Hepatocellular carcinoma

- Associated with cirrhosis in ~80%
- Suspect if- worsening of CLD, enlarged liver, hemorrhagic ascites, weight loss
- Dx-
 - CT/MRI with contrast- vascular SOL in cirrhotic liver
 - Raised AFP- α -fetoprotein
 - Liver biopsy
- Rx-
 - Early-resection
 - Advanced- liver transplantation or local palliative treatment
- Screening- US and AFP q 6 months