





BASICS

Epidemiology and complications

- \sim 0.6% of the population of England have hepatitis B (1.5% in London) \rightarrow 270,000 people.
- <50% of these are aware of their diagnosis; work is needed to meet WHO target of 90% aware of their diagnosis and 80% being treated.
- Complications:
 - Chronic infection develops in 90% of those infected vertically, up to 50% of those infected before the age of five and about 5% of previously healthy adults (more if immunocompromised).
 - Up to 20% with chronic hep B will develop cirrhosis in five years.
 - Those with cirrhosis who are untreated have an up to 5% risk of hepatocellular carcinoma.
 - Chronic hep B also a risk factor for other malignancies e.g. cervical, gastric and lymphoma.

Transmission

- Infected blood (sharing needles, contaminated medical/dental equipment, tattoos, piercings, acupuncture, blood transfusions if blood not screened).
- Vertical transmission (reducing over time due to vaccination of mothers).
- Unprotected sex (vaginal, anal, oral).
- Household contacts of an infected person.
- Needlestick injuries.
- In the UK, most transmission is sexual or by sharing injecting drug use equipment; in countries where hep B is endemic, most transmission is vertical, or by close contact as a child.

DIAGNOSIS

Symptoms

- Acute symptoms:
 - Most children/up to 50% of adults asymptomatic and will clear infection spontaneously.
 - o Prodrome (fever, arthralgia, rash, malaise).
 - o RUQ pain and jaundice ~ 2/52 after prodrome.
 - <1% fulminant hepatitis, rapid progression to lifethreatening liver failure.
 - Transaminases may be in hundreds/thousands, prolonged clotting time.
- Chronic infection:
 - Often asymptomatic for years.
 - Signs of chronic liver failure (spider naevi, clubbing, jaundice, palmar erythema, hepatosplenomegaly, easy bruising, ascites).
 - Transaminases may be normal or mildly raised.

- Who to test

 Current/past injectable drug use.
- Born/brought up in ↑ prevalence country, and those who adopt/foster children from these countries.
- Multiple sexual partners, sex workers, anyone who has been sexually assaulted.
- Household contacts of those with hep B.
- Patient with chronic liver disease or on dialysis.
- Those in supported living or residential care for learning disabilities.
- Those who have had a needlestick injury or bite.
- All those with hepatitis C, HIV or ↑ transaminases.
- See green book for information on who to vaccinate.

Management

MANAGEMENT

(anti-HBs)

Investigations

| Test | Interpretation if positive |
|-------------------|-------------------------------------|
| Surface antigen | Acute infection. HBV DNA used |
| (HBsAg) | to quantify viral load. |
| e antigen | High levels of viral |
| (HBeAg) | replication/infectivity. |
| e antibody (anti- | e antigen has cleared; viral |
| HBe) | replication controlled. |
| Core antibody | Current/past infection – antibody |
| (anti-HBc) | persists for life; doesn't indicate |
| | immunity. IgM = infection in the |
| | last 6/12. |
| Surface antibody | Immunity due to past infection or |

vaccination.

Primary care:

- Refer to hepatology and request FBC, U&E, clotting, LFTs, GGT, hep C/HIV, serum alphafetoprotein and liver USS at time of referral.
- Advise on ↓ progression (e.g. ↓ alcohol) and preventing transmission - test household/sexual contacts. Consider STI screen.
- Signpost to support e.g. <u>Hepatitis B trust</u> or <u>British</u> Liver Trust.
- o Notify public health if diagnosed in acute phase.
- Vaccinate against hep A if ongoing risk.
- Secondary care:
 - Antivirals will be offered, with ongoing monitoring for progression from acute to chronic infection and screening for complications.