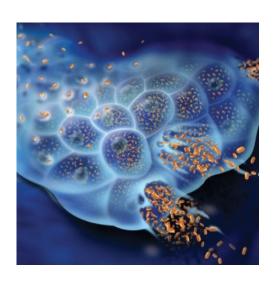
Viral Hepatitis

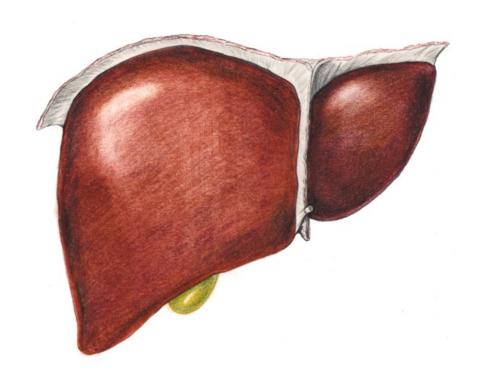


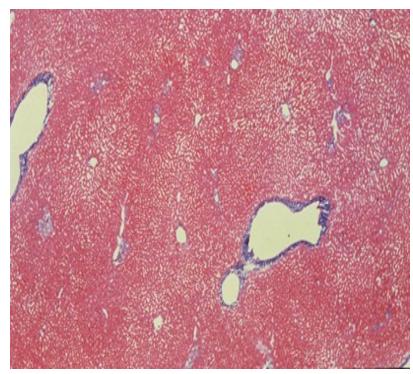
prof. MUDr. Petr Husa, CSc. Klinika infekčních chorob, FN Brno

Viral Hepatitis

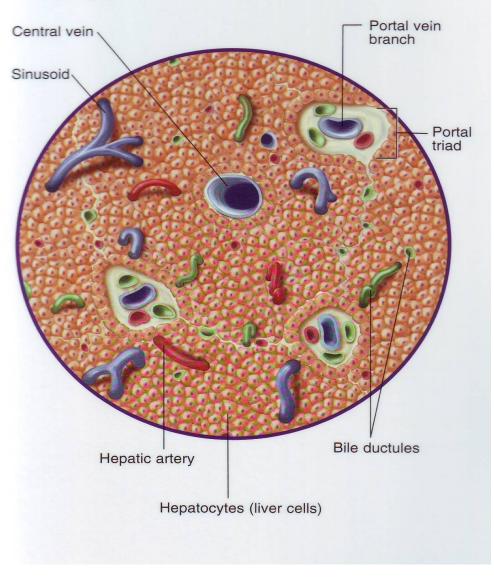
- Diffuse necrotic and inflammatory liver process
- On the opposite bacterial infections lead to formation of liver abscesses
- Division of viral hepatitis
- 1. Enterically transmissible
- HEP A only acute
- HEP E chronic in immunosuppressed pts. with G-3 or 4
- 2. Parenterally transmissible possible chronic stage
- HEP B
- HEP C
- HEP D

Healthy liver

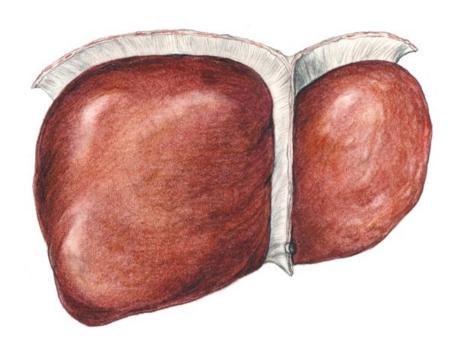


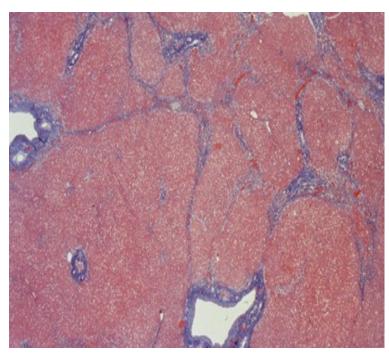


Normal Biopsy

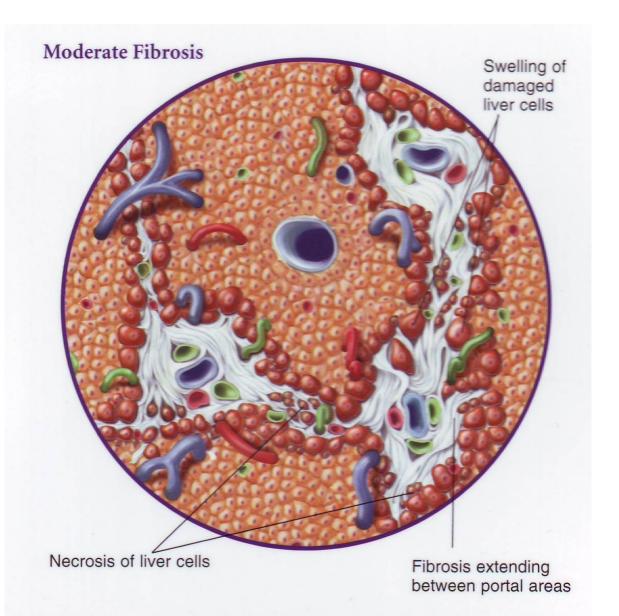


Liver fibrosis

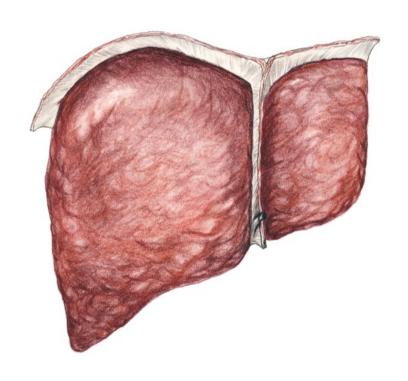


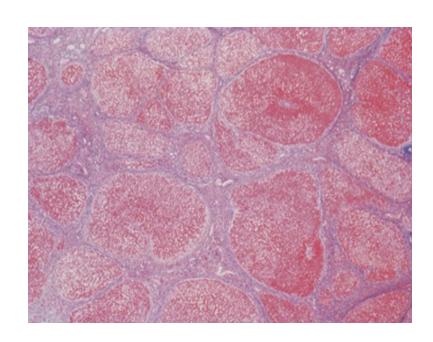


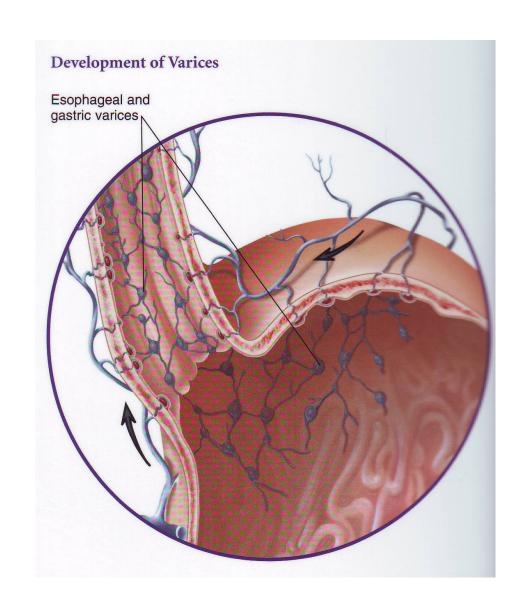
Mild Fibrosis Mild swelling and inflammation of Development of damaged liver cells around portal areas scar tissue (fibrosis) Normal hepatocytes (liver cells)



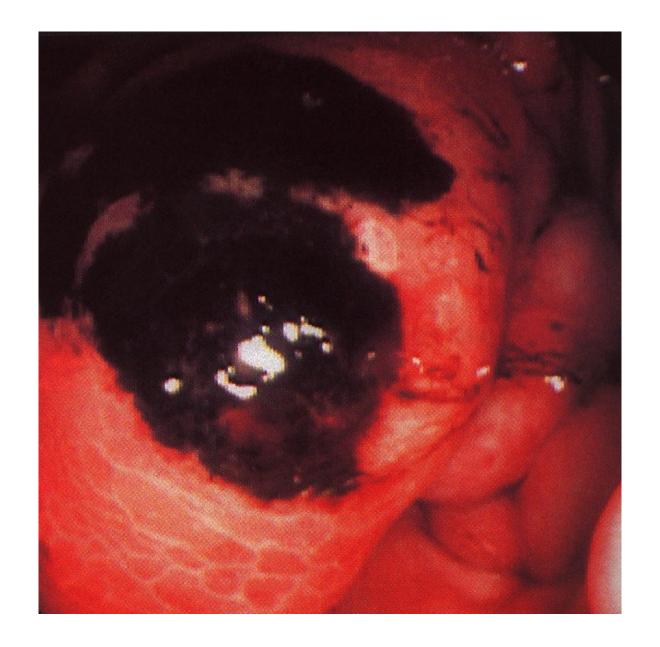
Liver cirrhosis

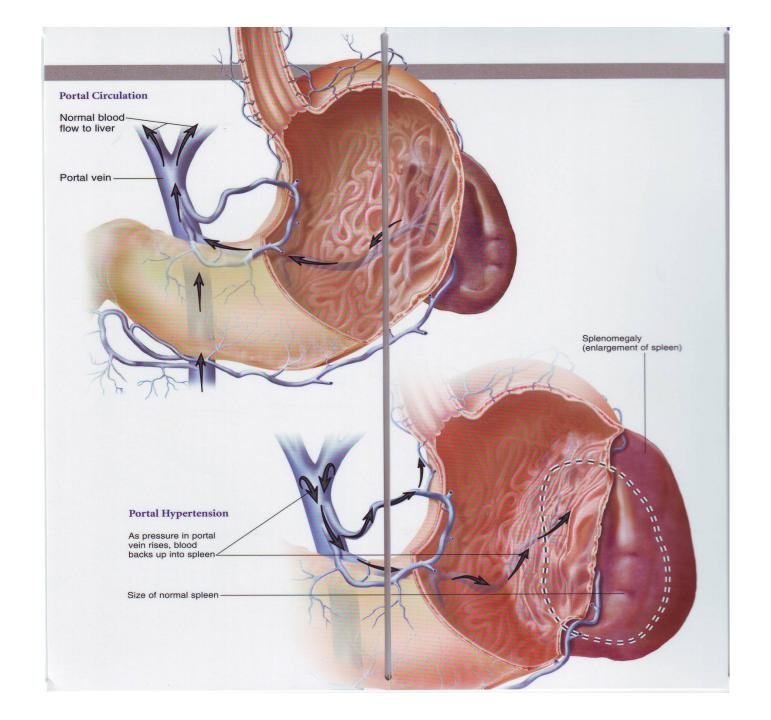










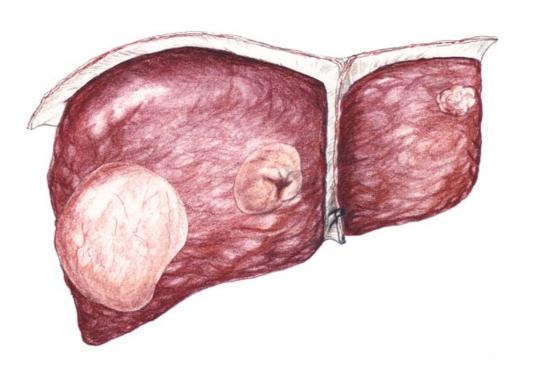


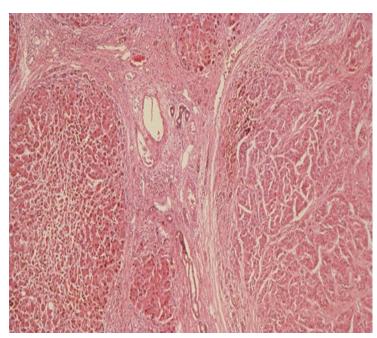






Hepatocellular carcinoma







Viral hepatitis in CR 2014-2024

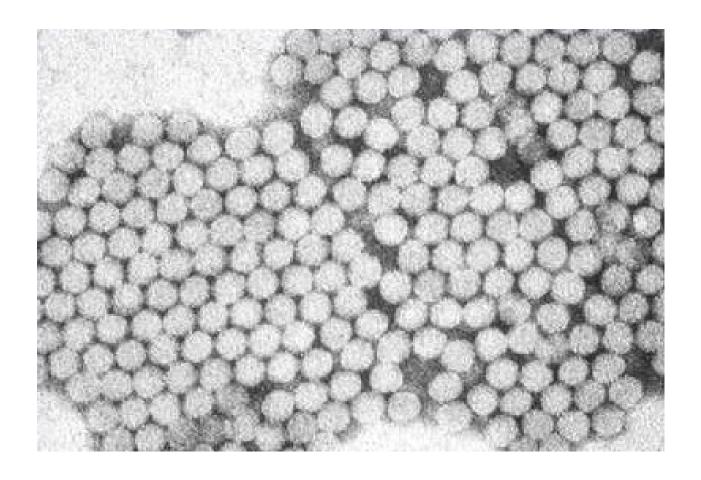
	2014	2015	2016	2017	2018	2019	2020	2021	2022	2023	2024*
VH A	673	723	930	772	211	240	183	210	70	66	239
akutní VH B	105	90	73	85	54	41	27	17	48	37	30
chronická VH B	193	193	208	248	269	276	142	127	244	378	305
VH C	867	945	1103	992	1050	1138	771	665	921	1301	1088
VH E	299	409	339	344	272	268	223	201	319	684	466
VH D					1	2	2	3	8	12	10

^{*} Do 8/2024 Zdroj: ISIN



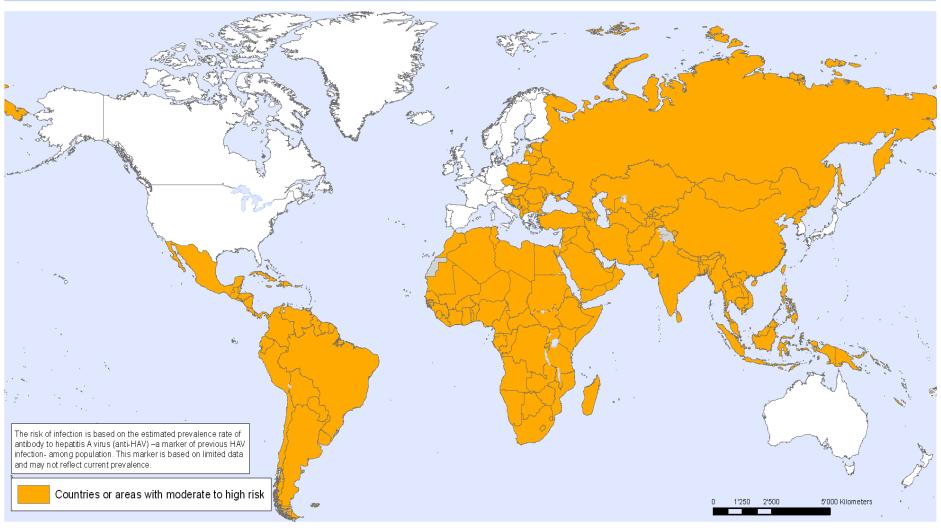


Hepatitis A virus (HAV)



Family Picornaviridae, genus *Hepatovirus* – non-enveloped RNA, 27 nm 3 human genotypes (I-III), worldwide G-I dominates, subtypes A a B, 3 exclusively simian genotypes (IV-VI), 1 serotype

Hepatitis A, countries or areas at risk



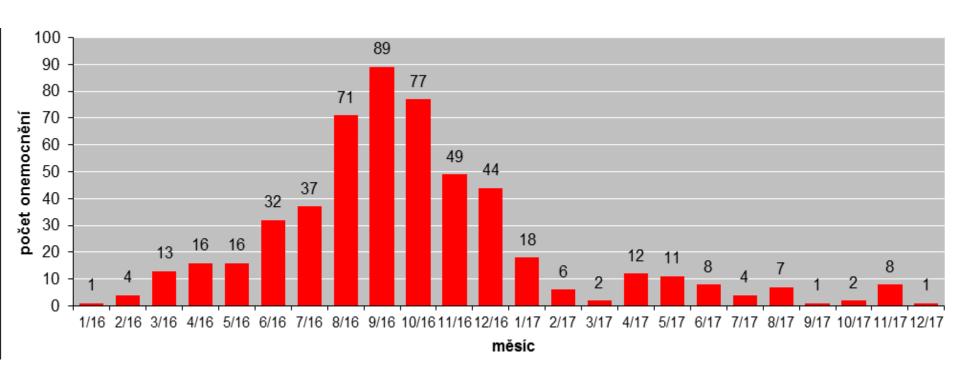
The boundaries and names shown and the designations used on this map do not imply the expression of any opinion whatsoever on the part of the World Health Organization concerning the legal status of any country, territory, city or area or of its authorities, or concerning the delimitation of its frontiers or boundaries. Dotted and dashed lines on maps represent approximate border lines for which there may not yet be full agreement.

Data Source: World Health Organization.
Jacobsen KH, Wiersma ST. Hepatitis A virus seroprevalence by age and world region,
1990 and 2005. Vaccine 2010 Sep;28(41):6653-7
Map Production: Public Health Information and Geographic Information Systems (GIS)
World Health Organization



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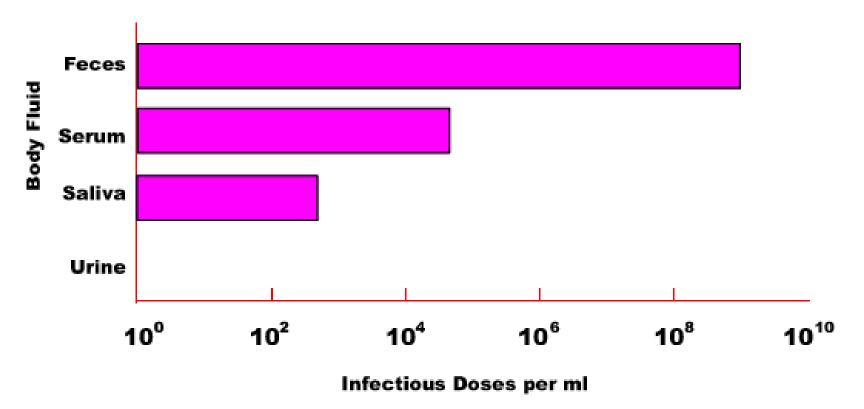
HAV epidemic in the South Moravia 2016-2017



Epidemiology

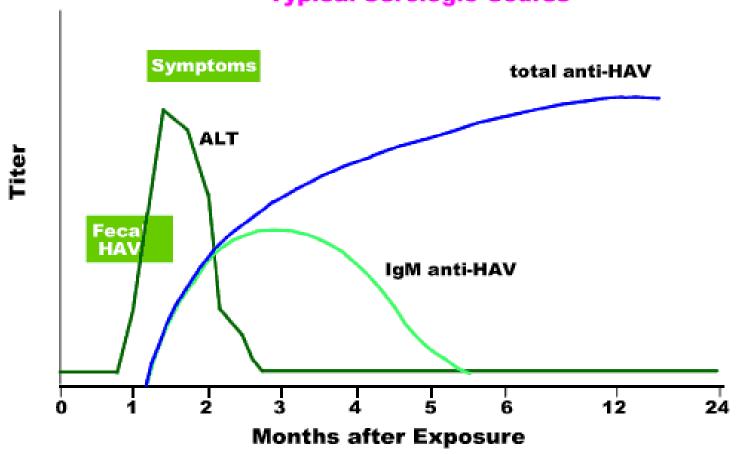
- Fecal –oral route of transmission
- ✓ Contaminated hands or daily used instruments
- ✓ Contaminated drinking water
- ✓ Contaminated food
- Vaccination available, recommended especially fore travelers to countries with lower standard of hygiene

Concentration of Hepatitis A Virus in Various Body Fluids

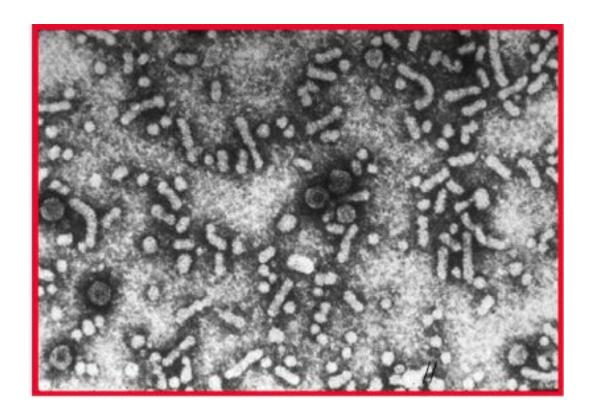


Source: Viral Hepatitis and Liver Disease 1984;9-2 J Infect Dis 1989; 160:887-890

Hepatitis A Virus Infection Typical Serologic Course



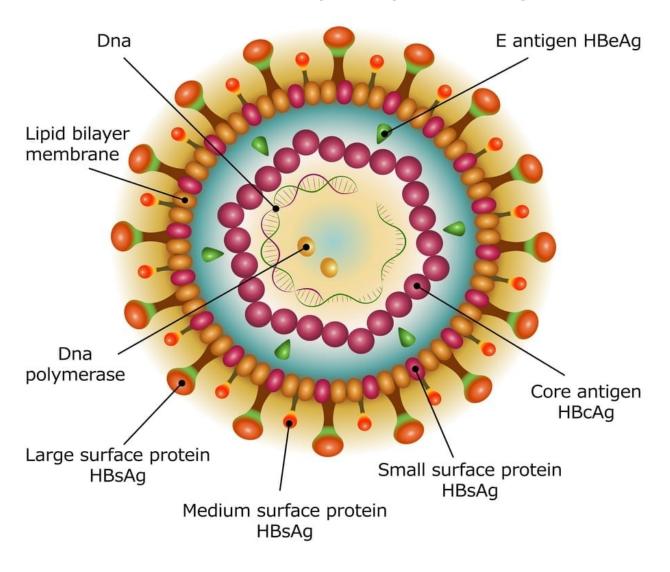
Hepatitis B Virus (HBV)



Family Hepadnaviridae, genus *Orthohepadnavirus*, enveloped DNA, 42 nm, 9 genotypes (A-I), Europe A,D, Asie B,C, several subtypes

Hepatitis B Virus

Baltimore Group VII (dsDNA-RT)



Estimated chronic HBV infection prevalence

• Worldwide: 240-350 million^[1] USA: 1.5-2.2 million^[2]

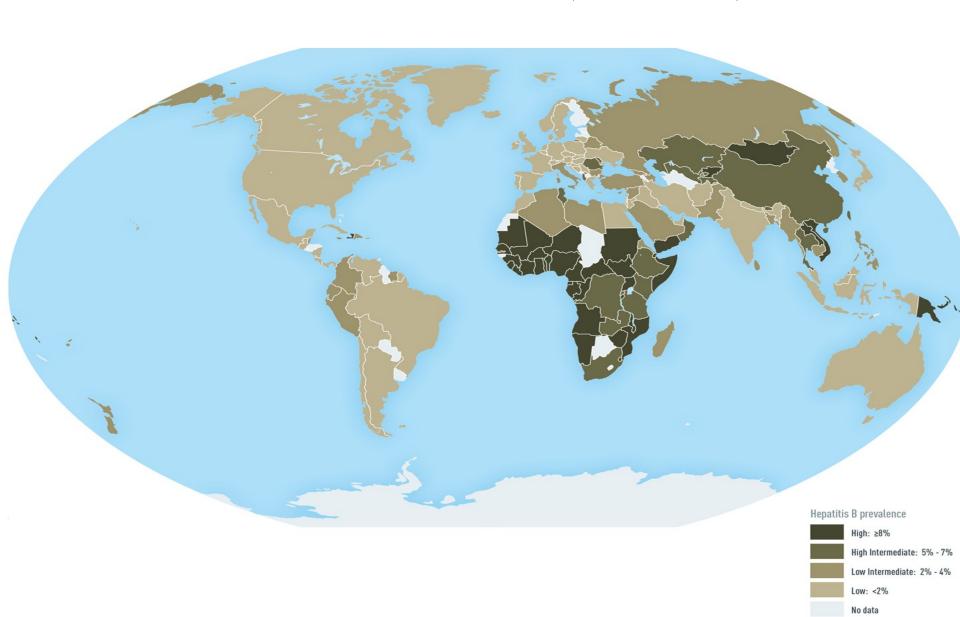


^{1.} MacLachlan. Cold Spring Harb Perspect Med. 2015;5:a021410. 2. Kim. Hepatology. 2009;49:S28. 3. Nguyen. Clin Microbiol Rev. 2020;33:e00046.

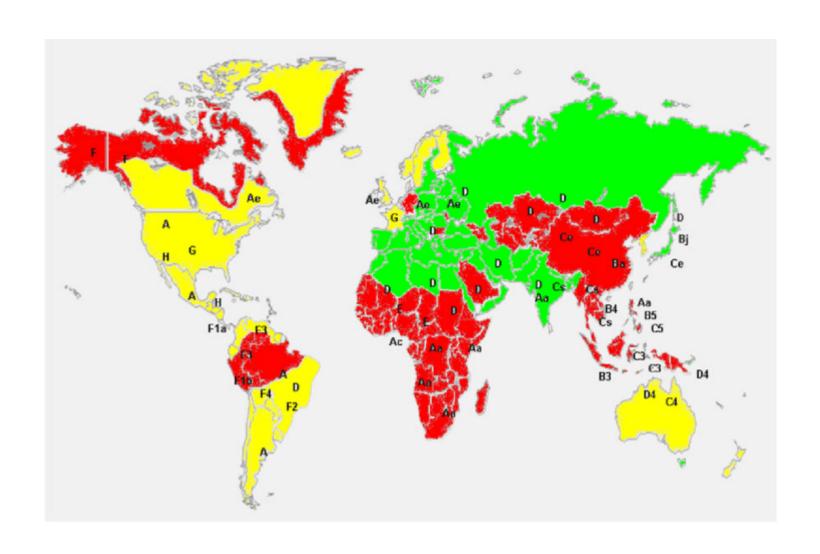
Global significance of HEP B

- One of the biggest global health problems
- ✓ More than 2 billions of infections during the life
- ✓ 240-350 million chronic HBsAg carriers
- ✓ Indication for 5-10 % liver transplantations globally
- ✓ Global vaccination in almost all countries

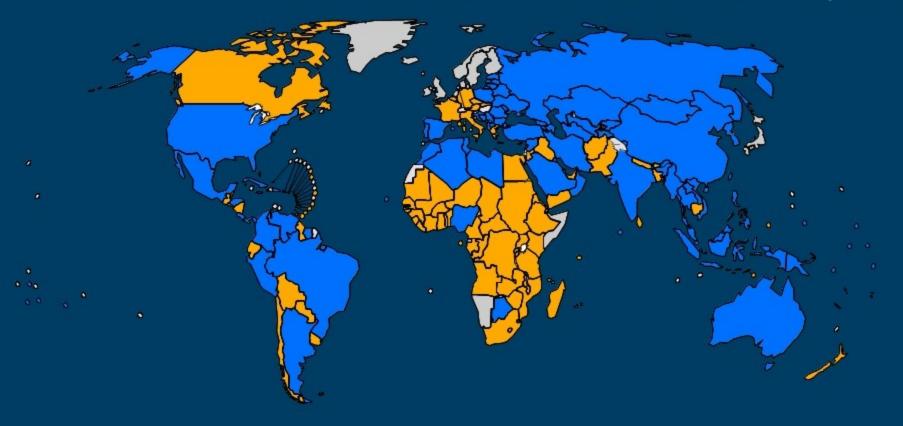
Chronic HBV infection (CDC 2020)



HBV genotypes (A-I)



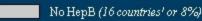
Countries using HepB in national immunization schedule, 2008



Source: WHO/IVB database, 193 WHO Member States. Data as of August 2009

Date of slide: 24 November 2009

The boundaries and names shown and bedesignous as used as day may do as ringly the expression of any apiene whatever as the part of the World Health Organization executing the legal status of any cases y, containly, cay or a color of as authorises, a color and the delimination of the financian in boundaries. Calculations as among representation among baseds io ca foi which there every our year be full agreement. D WHO 2009 Allinghance and



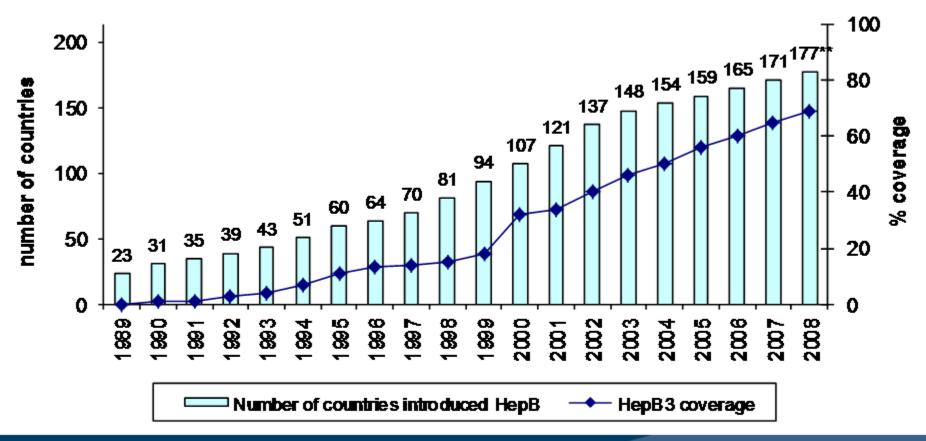
HepB no Birth Dose (92 countries? or 48%)

HepB with Birth Dose (85 countries or 44%)

finally dies three countries with adolescent investment on Analudia Audan with partial introduction includes has with pomal introduction



Number of countries having introduced HepB vaccine* and global infant coverage, 1989-2008



^{*} Year of introduction can be the year of partial introduction



^{**} Includes India and Sudan with partial introduction excluding 3 countries where HepB administered for adolescence

Hepatitis B in Czech Republic

- Still important infection but incidence and prevalence are gradually decreasing
- ✓ Prevalence of chronic carriers was 0.56 % (2001) ...0.064 % (2013)
- ✓ Decrease of prevalence and incidence due to vaccination of high-risk persons (health care workers, newborns of HBsAg-positive mothers, before hemodialysis)
- ✓ Global vaccination of all newborns and 12-years old children 2001-2013, now only newborns (hexavaccine)

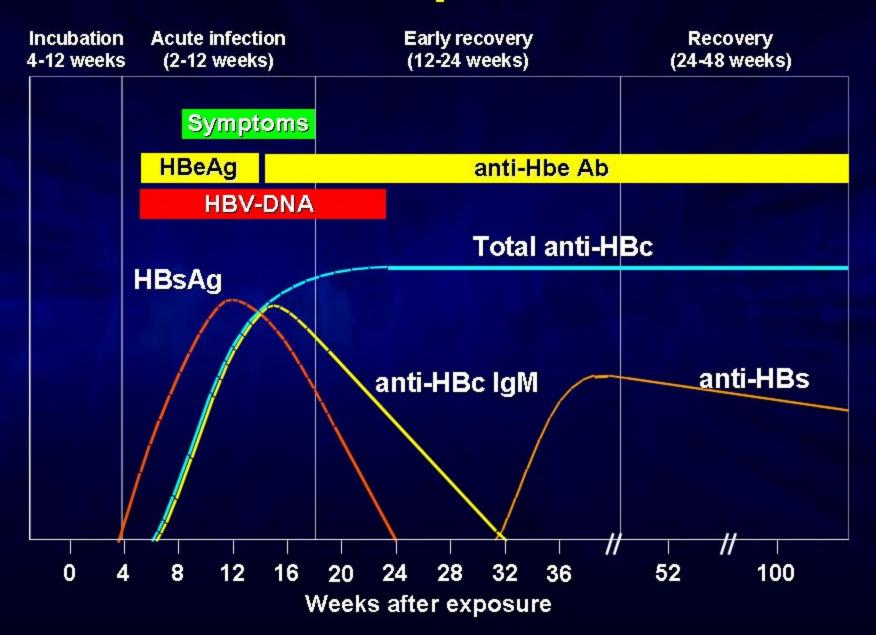
Epidemiology of HBV

- HBV transmission
- ✓ sexual intercourse
- ✓ vertically from mother to newborn during delivery or in the last trimester
- ✓ sharing of instruments among IUDs
- ✓ blood and blood products
- ✓ organ and tissue transplant recipients

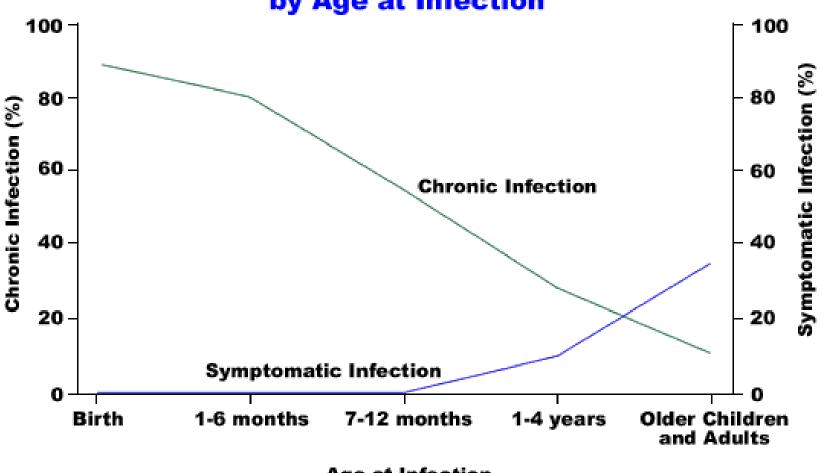
Clinical findings in acute HEP B

- IP: 30–180 days (mostly 2–3 months)
- Prodromal stage flu-like syndrome
- Fulminant hepatitis: < 1 %
- Chronic HBV infection mortality: 15 25 %

Acute Hepatitis B

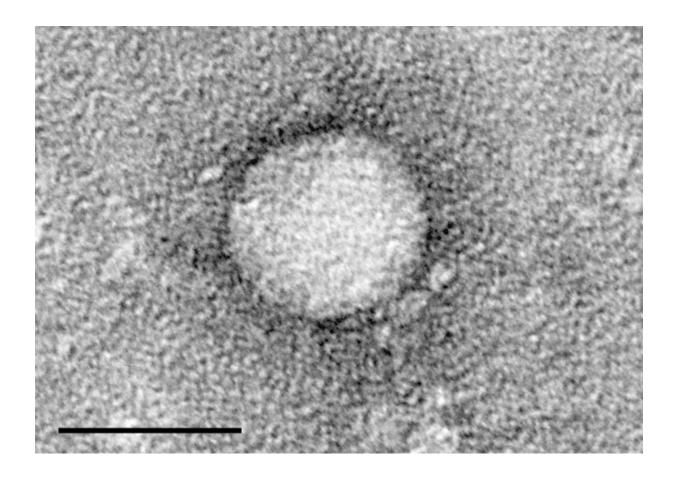


Outcome of Hepatitis B Virus Infection by Age at Infection



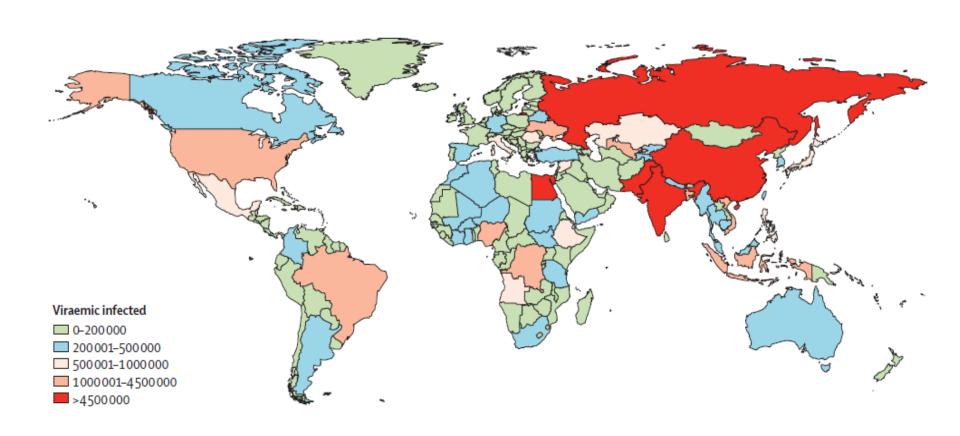
Age at Infection

Hepatitis C Virus (HCV)



Family Flaviviridae, genus *Hepacivirus*, enveloped RNA virus 60 nm, 8 genotypes (1-8), many subtypes (a...)

Global estimations: 56,8 million persons with chronic HCV infection



Approximately 700 000 persons died anualy due to complications related to HCV infection: liver cirrhosis, HCC, and liver failure

Risk groups of inhabitants for HCV infection



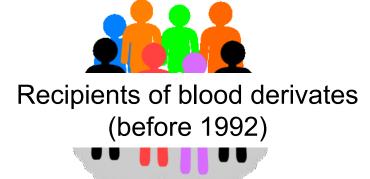




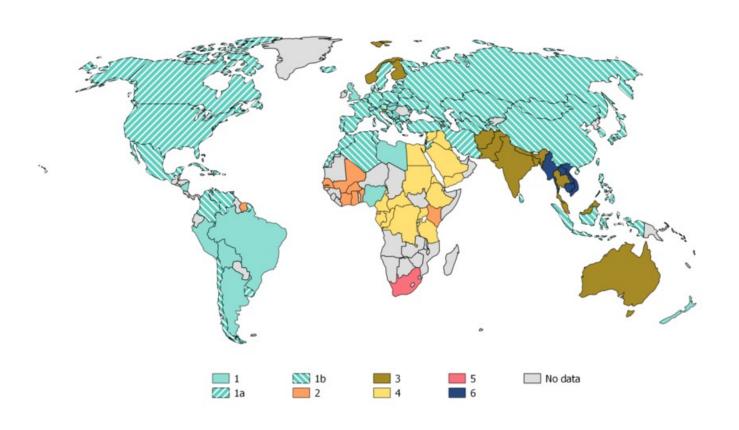






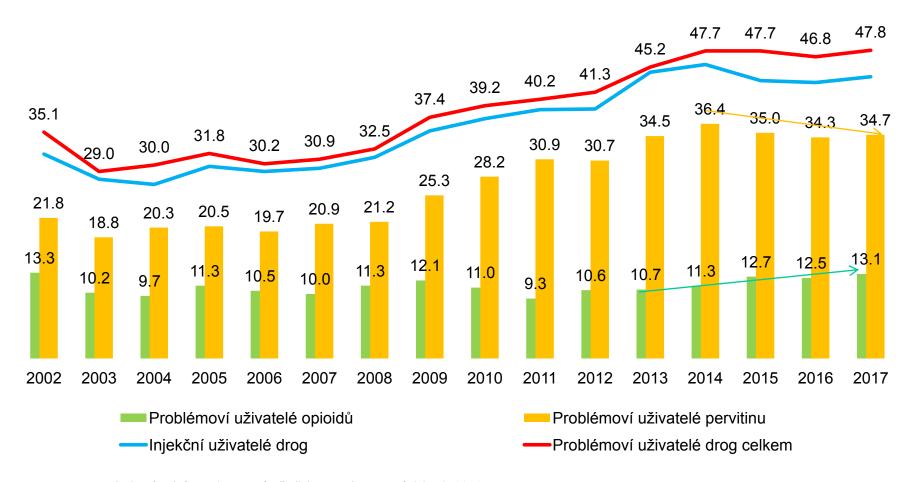


HCV genotypes distribution



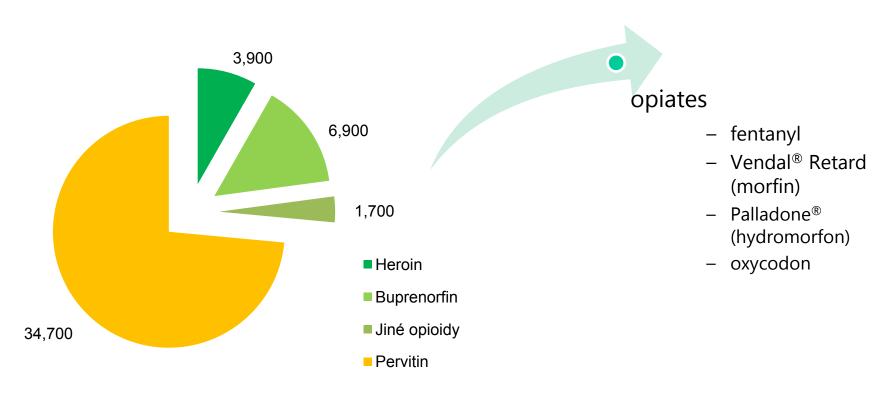
Problematic drug abusers in the CR (2017)

About 44 000 intravenous drug abusers (cca 90 % of all)



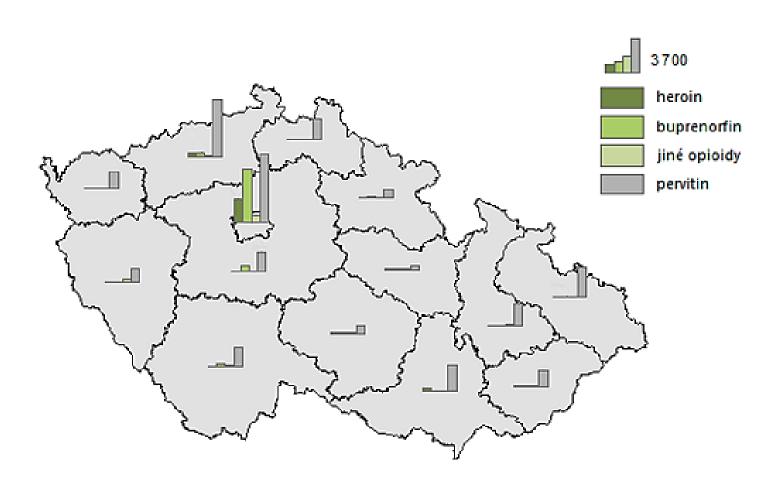
Problematic drug abusers in the CR (2017)

2017 - Estimation in the CR - 47 800 problematic drug abusers, about 90 % intravenous

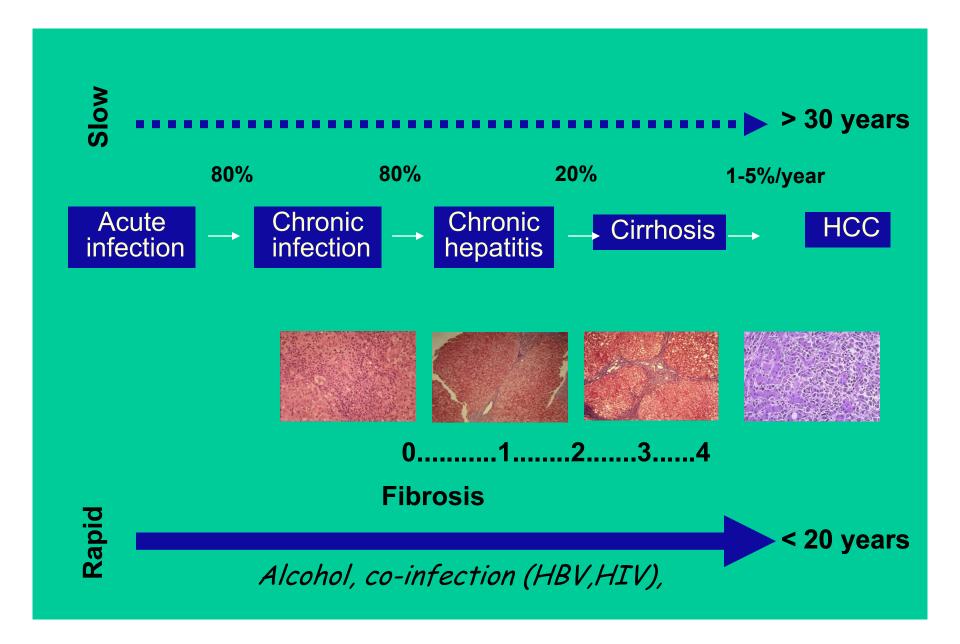


Zdroj: Národní monitorovací středisko pro drogy a závislosti (2018)

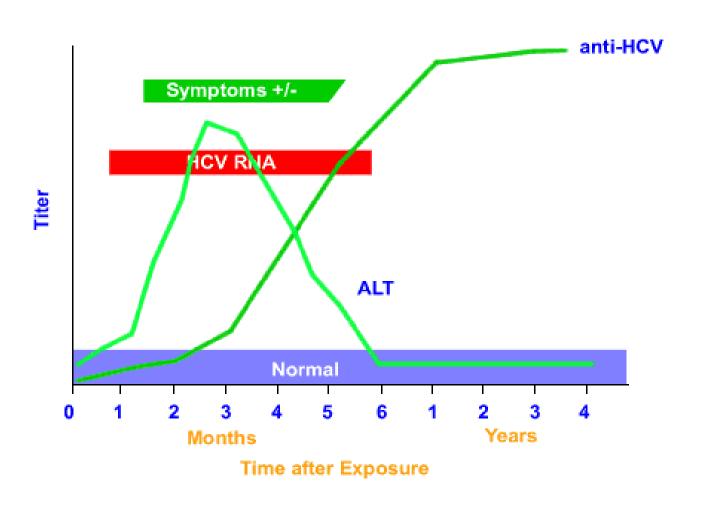
Opiates and pervitin abusus



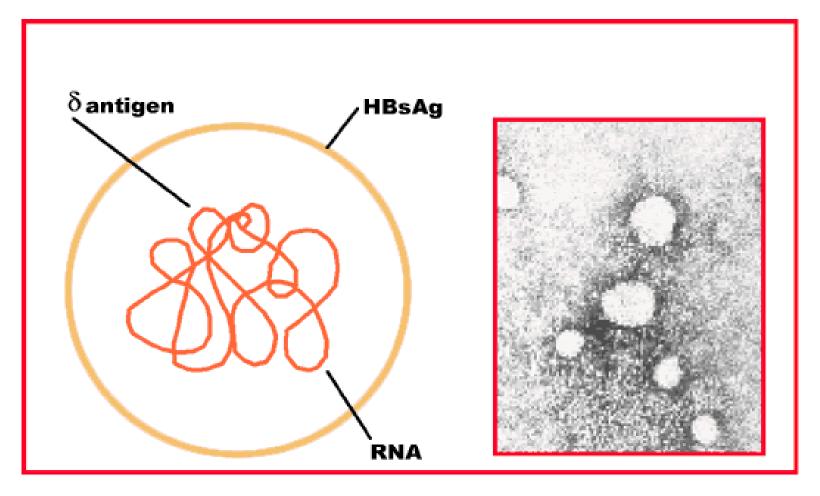
Clinical course of HCV infection



Diagnosis of HCV infection



Hepatitis D (Delta) Virus (HDV)



Satelite virus, family Deltaviridae, genus *Deltavirus*, enveloped RNA, 36 nm, 8 genotypes (I-VIII), genotype I the most common worldvide

Hepatitis D: fast facts



9-60 million people infected with HDV globally

Defective RNA virus, requiring HBV for infection

4.5-13% of HBV carriers co-infected with HDV



Most severe form of viral hepatitis

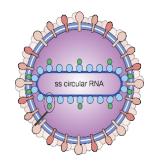
Increased risk of cirrhosis/HCC and higher mortality vs HBV

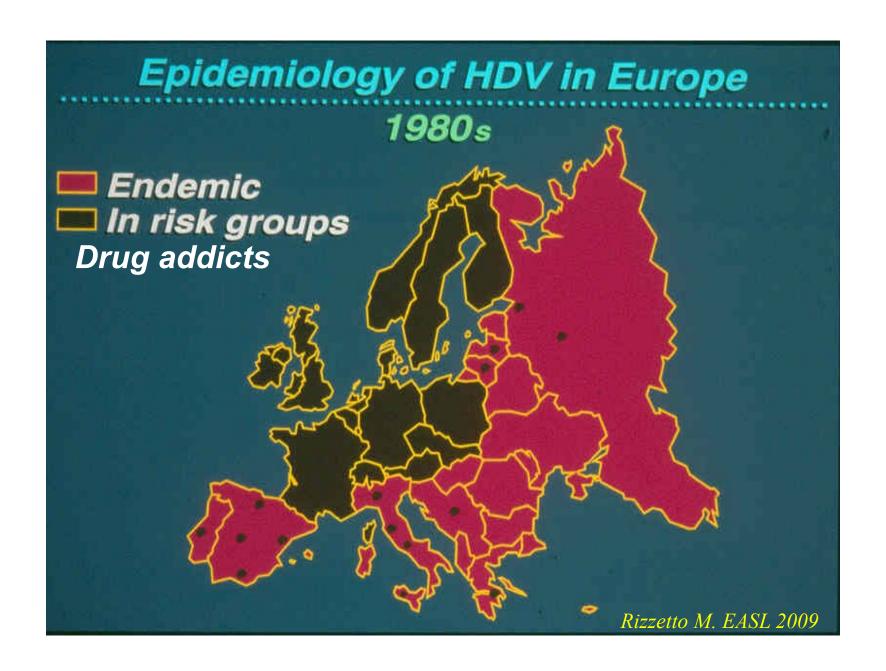
Progression to cirrhosis within 5 years and to HCC within 10 years



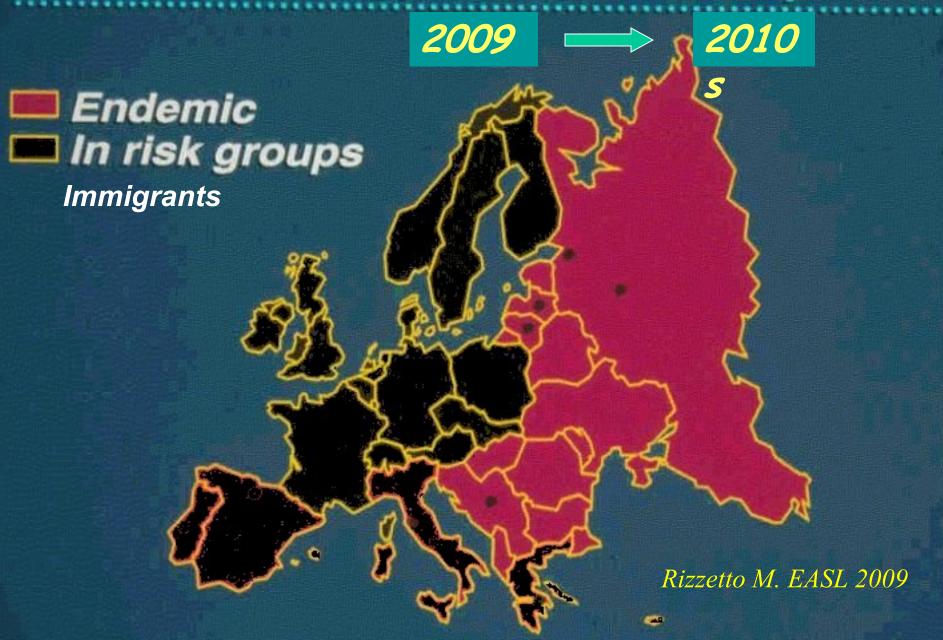
Eight HDV genotypes

Bulevirtid approved for therapy



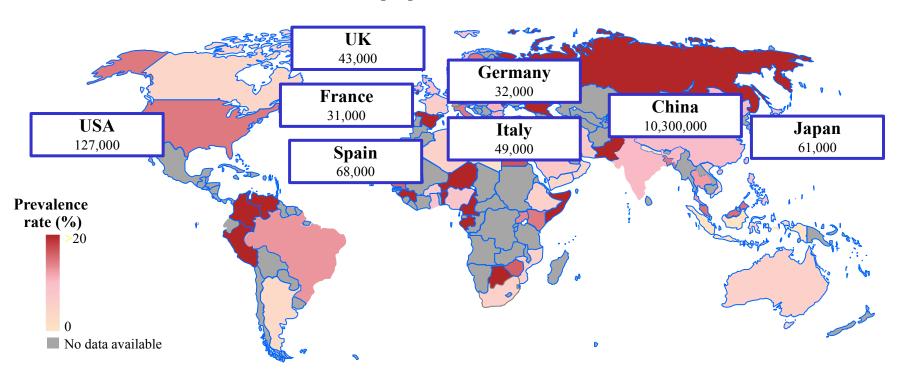


Epidemiology of HDV in Europe



Estimated number of individuals with HDV in selected countries

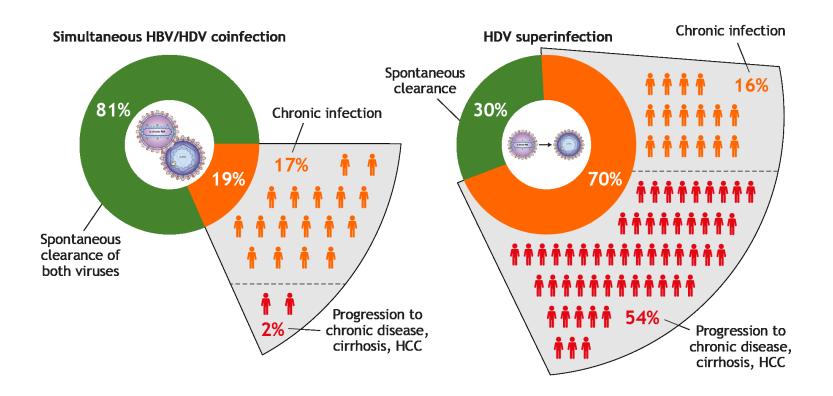
An estimated 48-60 million people are infected with HDV worldwide



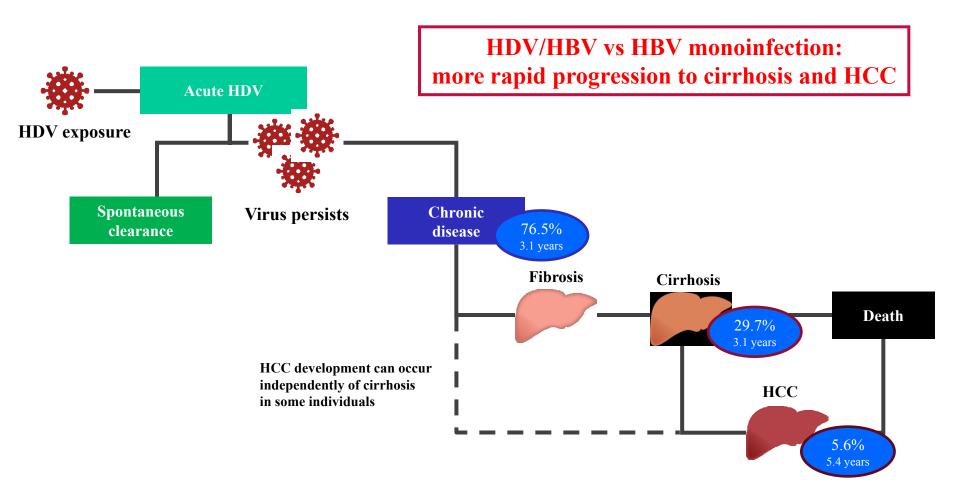
Numbers shown are patient numbers, ie prevalence of HDV in HBsAgpositive patients.

HBsAg: hepatitis B surface antigen; HDV: hepatitis delta virus.

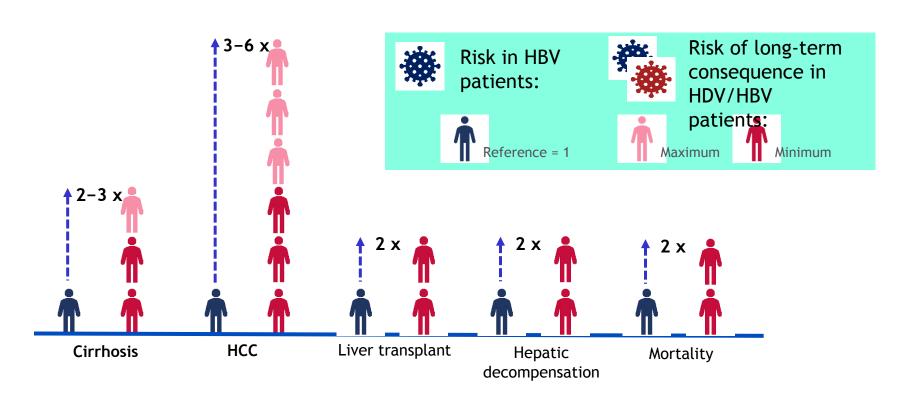
Clinical course of HBV/HDV infection depends on timing of HDV infection



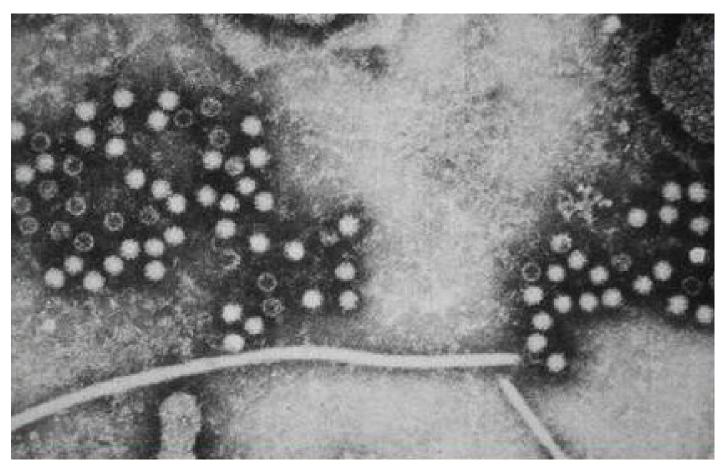
Clinical course of HDV superinfection



Increased risk of long-term consequences of viral hepatitis in HBV/HDV patients versus HBV monoinfection

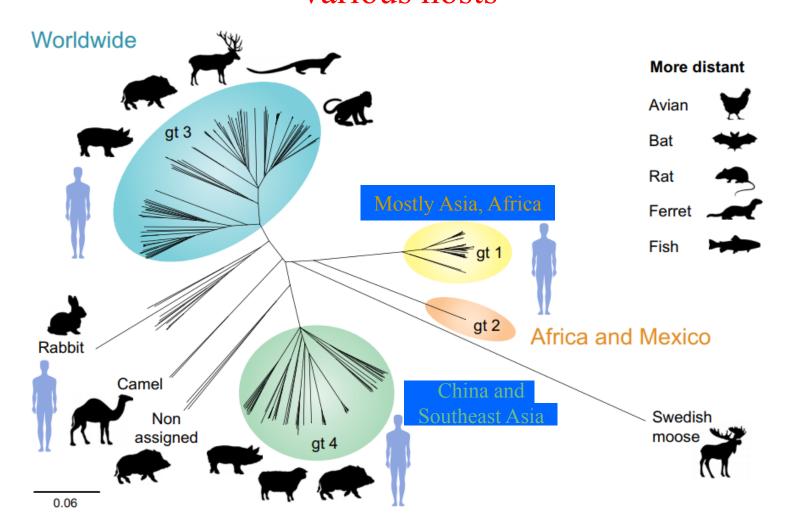


Hepatitis E virus

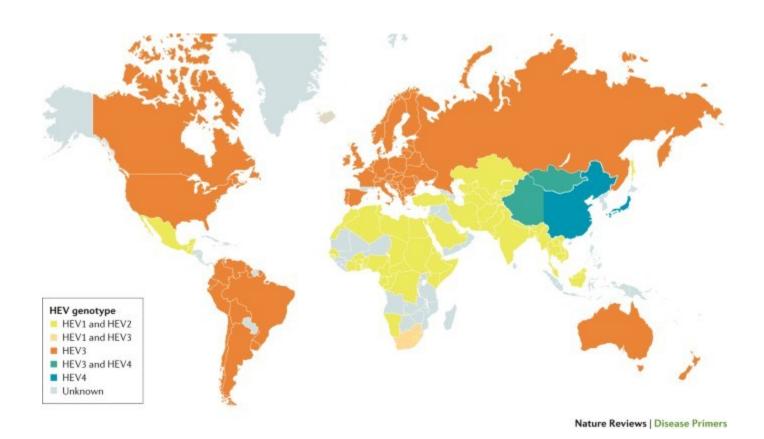


Non-enveloped RNA virus, family Hepeviridae, *genus Orthohepevirus*, 27-34 nm, 8 genotypes (1-8), human infections by G1-4

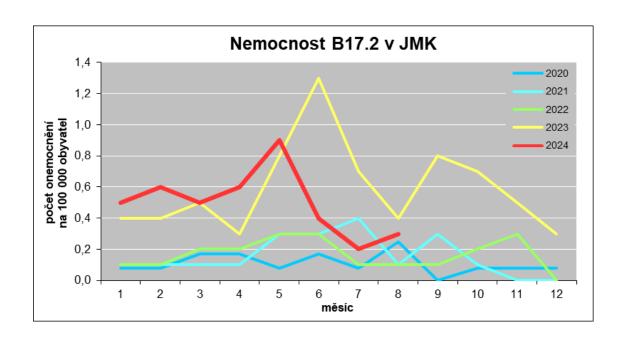
Phylogenetic relationship of hepeviruses identified in various hosts



HEV genotypes (1-4)



Hepatitis E in South Moravia 2019-2024



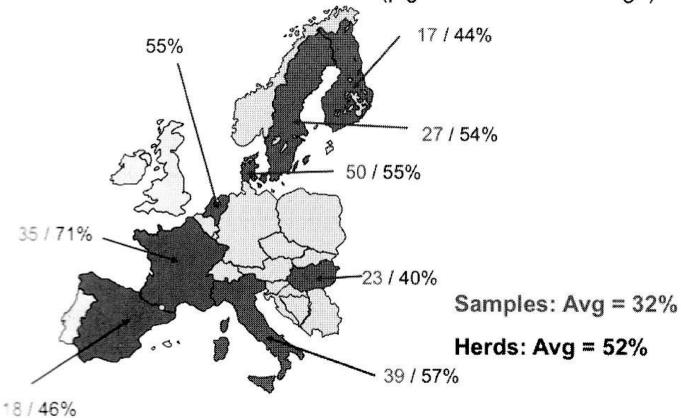
Infection with G-1,2 HEV

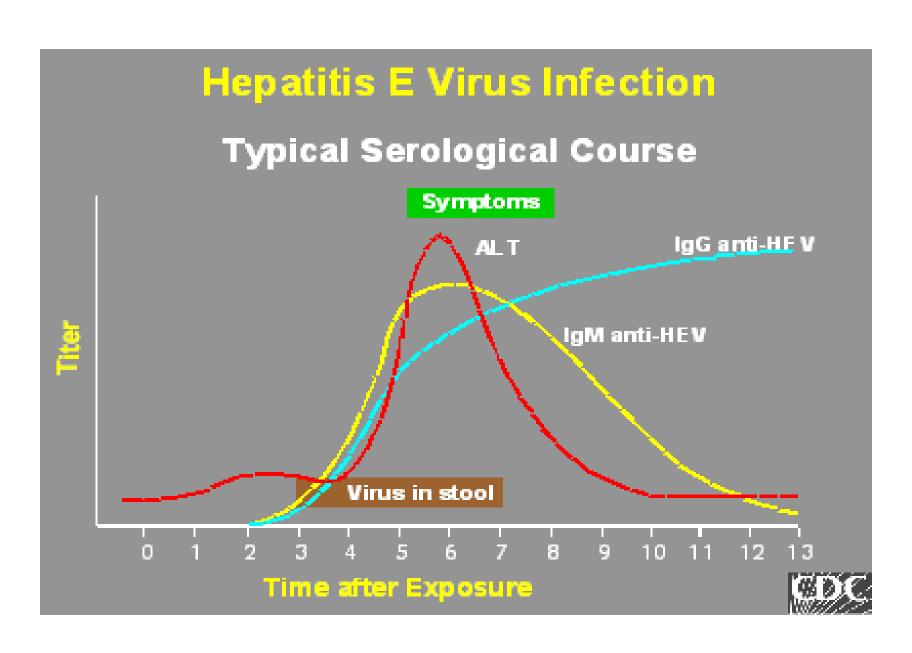
- Only human infection
- Mostly Asia, Africa
- Extremely serious clinical course in late pregnancy (mortality about 25 %)
- No chronicity
- Possibility of acute-on-chronic liver failure

Infection with G-3,4 HEV

- Both human and zoonotic infection
- Pigs are the main reservoir (venison as well)
- G-3 worldwide distribution, G-4 China a southeast Asia
- ≥ 2 million locally acquired HEV infections/year in Europe (G-3), mostly asymptomatic (minimally 95 %), tend to affect older males
- Possibility od chronic infection in persons with immunosuppression (after solid organ transplantation 50-66% probability of chronicity, patients with haematological disorders, individuals living with HIV, patients with rheumatic disorders receiving heavy immunosuppression)
- High mortality in patients with liver cirrhosis (60-70 %) acute-on-chronic liver failure

Prevalence of HEV in swine herds (pigs 1 to 5 months of age)





Rapid progression of chronic hepatitis E

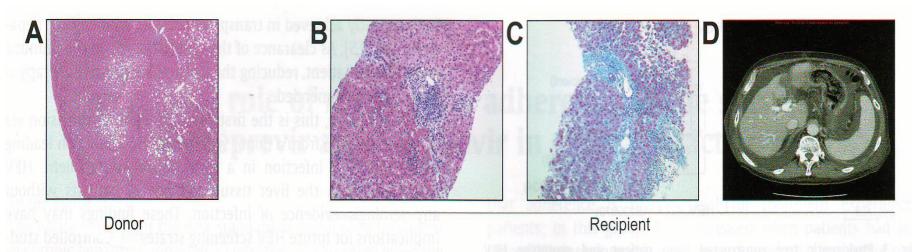


Fig. 1. Histologic assessment of the liver tissue before and after OLT and CT scan after OLT. (A) The liver tissue of the donor revealed absence of significant signs of chronic hepatitis but vesicular fatty liver disease was diagnosed. (B) Second biopsy. One hundred and fifty days after OLT, chronic inflammation with portal and interface hepatitis was described which was interpreted as an acute rejection. (C) Third biopsy. Three hundred and forty seven days after OLT, persistence of chronic hepatitis was associated with portal and septal bridging signs of fibrosis. (D) CT scan performed 1 year after liver transplantation revealed signs of portal hypertension including ascites, splenomegaly and gastric varices compatible with decompensated liver cirrhosis.

Treatment of acute hepatitis (all types)

- Symptomatic for all hepatitis types
- ✓ physical and mental rest
- ✓ no alcohol, no hepatoxic drugs
- ✓ diet (?)
- ✓ supportive treatment (silymarin, essential phosholipids) (?)
- ✓ antivirals for complicated acute HEP B and E

Therapy of acute HEP B

- Antiviral therapy is indicated only in serious (INR > 1,5) or prolongated
 (pronounced icterus > 4 weeks) clinical course od acute hepatitis B
- Therapy only with oral virostatics (NA)
- ✓ tenofovir
- ✓ entecavir

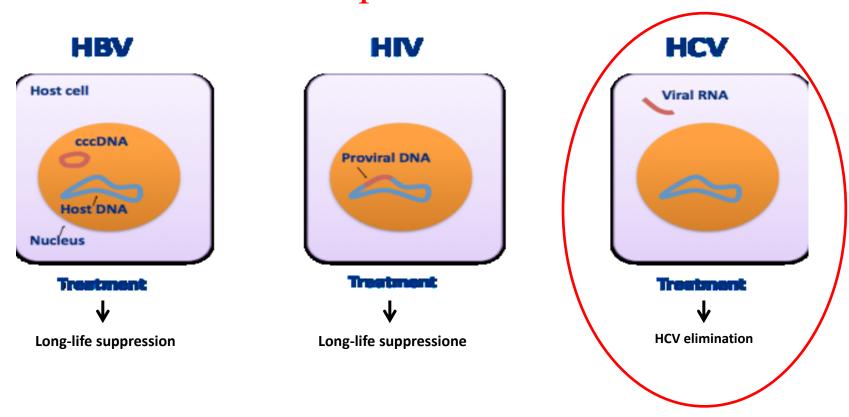
Current possibilities of treatment of chronic HEP B

- tenofovir
- entecavir

IFN-free regimens for HCV infection

- Current standard of HCV therapy
- Combination of oral drugs DAA direct-acting antivirals
- High efficacy minimally 99 %
- Almost no adverse events
- Short duration of therapy -8 or 12 weeks

HCV infection is curable in majority of patients



SVR – sustained virological response = the definite eradication of HCV infection

Hepatitis D treatment

- PEG-IFN $-1 \times$ weekly s.c.
- ✓ duration of therapy minimally 1 year
- ✓ In most cases only temporary effect frequent relapses after treatment discontinuation
- Bulevirtid (entry inhibitor) s.c. 1× daily, duration of therapy was not definitely established

Hepatitis E therapy

- Acute hepatitis E
- ✓ Spontaneous infection elimination without therapy
- ✓ fulminant course ribavirin mortality lowering
- Chronic hepatitis E
- ✓ Reduction of immunosupression infection elimination in about 30 % patients
- ✓ ribavirin for 3-6 months
- ✓ PEG-IFN for 3 months only after liver transplantation



Thank you for your attention!

