



CLDF

Chronic Liver Disease Foundation

3RD ANNUAL
LIVER C^ONNECT
CONFERENCE

Accredited by:



Presented by:





CLDF

Chronic Liver Disease Foundation

3RD ANNUAL

LIVER C**NNECT**

CONFERENCE

WHISE



Viral Hepatitis B and C: Gender Issues, Care Opportunities and Pregnancies

Tatyana Kushner, MD, MSCE

Associate Professor

Division of Liver Diseases

Department of Obstetrics, Gynecology and Reproductive Sciences

Icahn School of Medicine at Mount Sinai

Objectives

- Delineate global burden of HBV and HCV in women of childbearing potential (WOCP) and in pregnancy
- Outline recommendations for screening for HBV and HCV in WOCP and pregnant people
- Outline recommendations for treatment in WOCP and pregnant people
- Delineate gaps in care

World Health Organization: Global Health Strategy Interventions 2022-2023

1. Primary prevention (i.e. vaccination)
2. Harm reduction for PWIDs
3. Prevention of vertical transmission
4. Prevention and treatment and care for children and adolescents
5. Infection prevention and control
6. Integrated testing
7. Partner notification/ services
8. Stigma and discrimination in healthcare settings

GLOBAL VISION

A world where viral hepatitis transmission is halted and everyone living with viral hepatitis has access to safe, affordable and effective prevention, care and treatment services.

GOAL

Eliminate viral hepatitis as a major public health threat by 2030.⁸

Priority populations:

1. Pregnant breastfeeding women
2. Women and girls, including adolescent girls and young women

Burden of HBV and HCV Among Women of Childbearing Potential (WOCP)

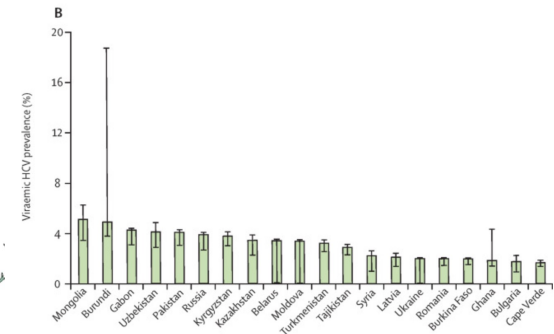
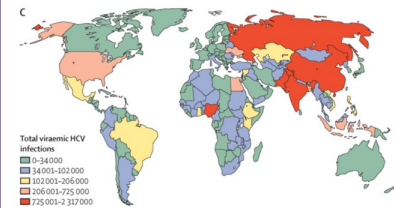
Women of childbearing age account for a quarter of the world's population

Hepatitis B

- 65 million women of childbearing potential globally

Hepatitis C

- Polaris Estimated 14.9 million women age 15-49 globally → 1/5 global HCV infection

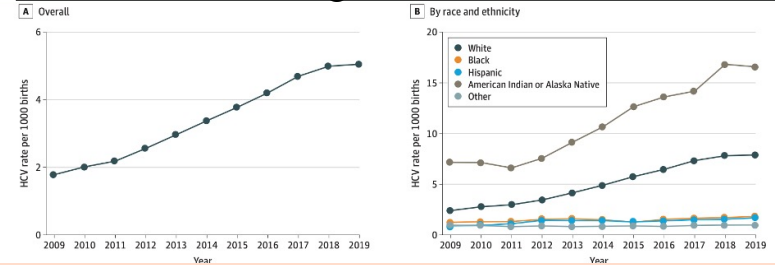


Global hepatitis report. WHO; 2017. Available from: <https://www.who.int/hepatitis/publications/global-hepatitis-report2017/en/>

Burden of HBV and HCV in Pregnant People

- **4.5 million** patients with HBV give birth annually
- HCV: **0.1% to 4.1%** prevalence of HCV in pregnant people
- Burden of HBV/ HCV in pregnant people varies by geographical region

HCV Rate among all US Births 2009-2019



HCV in pregnancy rate up from 1.8 to 5.1/ 1000

HBV among WOCP, 2011-2017



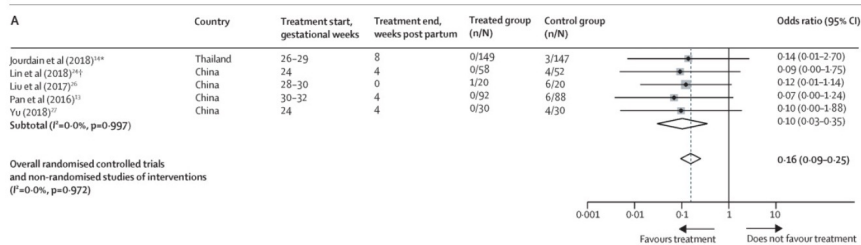
Chronic HBV rate down from 0.83% to 0.19%, but up in MS, KY, WV

Pregnancy and Mother-to-Child Transmission (MTCT)

MTCT is a major contributor to global burden of disease

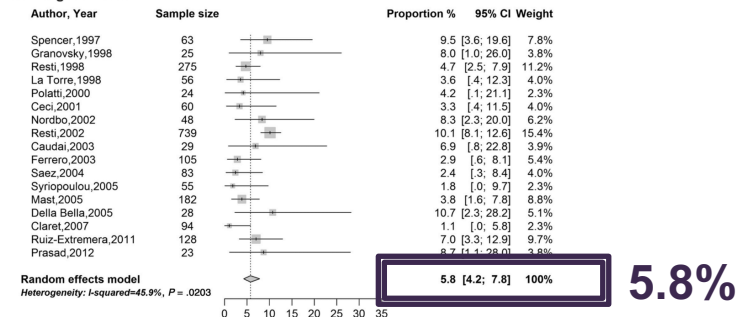
Hepatitis B

- Without immunoprophylaxis, HBV transmission is up to 90%
- Neonatal HBIG and HBV Vaccine decrease risk; Maternal antiviral therapy further decreases risk

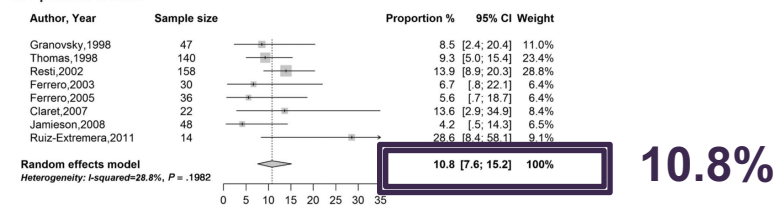


Hepatitis C

HIV-negative women



HIV-positive women



Risk Factors for MTCT

Risk Factors for MTCT	Increases transmission?
Biological	
High viral load (HBeAg+) / HCV (HCV RNA > 10 ⁶ log)	Yes
HIV Coinfection	Yes for HCV
Mode of delivery	Inconclusive
Breastfeeding	No
Invasive Fetal Testing	Potentially for patients with high viral load
PROM	No (with proper prophylaxis)
Social/environmental	
Lack of maternal knowledge	Yes
Lack of formal hospital policies	Yes
High cost/other barriers to access	Yes
Perceived stigma (particularly in developing countries)	Yes

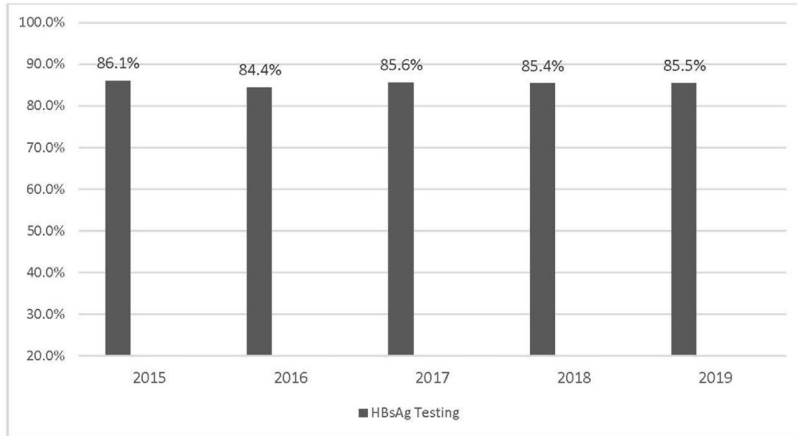
Recommendations for HBV/ HCV Testing in Women

	AASLD	ACOG	EASL	APASL	CDC
HCV					
Women	Screen all age 18+	N/a	Risk-based	Risk-based	Screen all age 18+
During Pregnancy	✓	✓ (2022)	N/A	N/A	✓ (2020)
HBV					
Women	Risk-Based	Risk-Based	Risk-Based	Risk-Based	✓ (2023)
During Pregnancy	✓	✓	✓	✓	✓

Is screening actually being done?

HBV

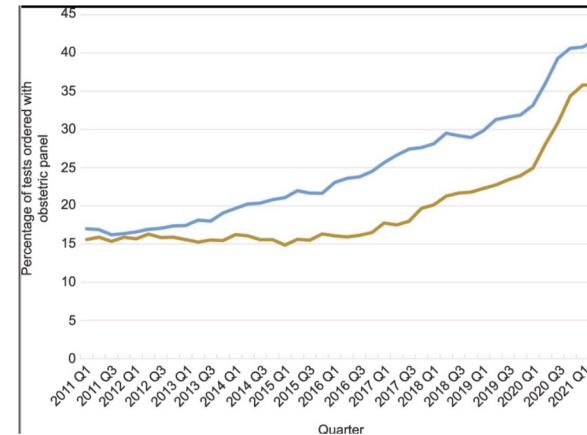
Optum Clinformatics Database; 500,000+ pregnancies 2015-2020



0.5 million (»14%) pregnant persons who gave birth annually were **not** tested for HBsAg to prevent perinatal transmission

HCV

Retrospective study using Quest laboratory data, 2011-2021

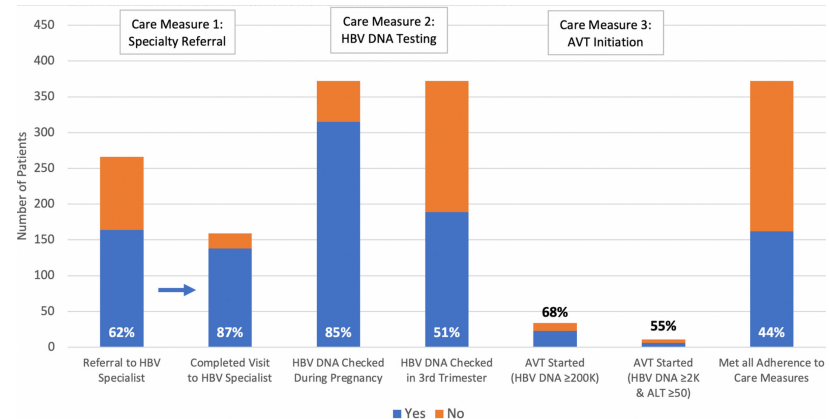


Pregnant persons HCV Ab screens (percentage), by commercial (*blue line*) and Medicaid insurance (*brown line*).

“Despite progression in pregnant persons screened for HCV, current testing rates fall short of universal recommendations”

Treatment of HBV in Pregnant People

- The infants of all HBsAg-positive women should receive immunoprophylaxis
- Antiviral therapy should be started at 28-32 weeks of gestation in most of the studies if HBV DNA >200,000 IU/mL (1 million copies/mL) is a conservative recommendation to start treatment
- For pregnant women with immune-active hepatitis B, treatment should be based on recommendations for nonpregnant women.
- Breastfeeding is not contraindicated. C-section is not indicated owing to insufficient data to support benefit.



Treatment of HCV in Pregnant People

Treatment with DAAs during pregnancy



1. Maternal cure while engaged in pregnancy care
2. Possible decrease in MTCT
3. Maternal treatment while under insurance coverage
4. Decrease in community transmission
5. Potential decrease in HCV-associated adverse pregnancy outcomes?

1. Human safety in pregnancy not established
2. Safety during breastfeeding not established
3. More established data for treatment prior to pregnancy or children starting at age 3
4. Difficulty in accessing DAA therapy in time (prior to delivery)
5. Cost-effectiveness not established

Barriers to Care in WOCA and Pregnant Women

- Stigma
- Difficulty in engagement of women postpartum
- “Fears” of offering treatment to pregnant women
- Fragmented health care system – no communication between PCP, Ob/Gyn, and liver specialists → co-location of care may help
- Disconnect between care of mothers and infants

Where Do We Go From Here to Optimize Care of Women With HBV and HCV?

Need to develop systems to improve screening for HBV and HCV, particularly in pregnancy setting

Efforts to combat stigma in WOCB and in pregnant women living with HBV/HCV to improve care

Need more robust safety data to inform HCV treatment in pregnant people

Increased interdisciplinary involvement to engage Ob/ GYN and primary care women's health providers in HBV and HCV screening and treatment



Thank You!

- Tatyana.Kushner@mssm.edu