SCHOOL OF MEDICINE

Background

- Mother-to-child transmission (MTCT) of hepatitis B virus (HBV) is 100% preventable.
- Guidelines from the American Association for the Study of Liver Diseases (AASLD) recommend the following:
 - Timely (within 12 hours) birthdose HBV vaccination
 - Timely hepatitis B immunoglobulin Nu (HBIG)
 - Anti-viral prophylaxis in women with *high-risk* HBV (defined as viral load ≥200,000 IU/mL and/or HBV e antigen [HBeAg] positivity)

Objectives

 To assess UNC Hospital's care of HBV-infected mothers and their exposed neonates in comparison to AASLD guidelines for the prevention of MTCT.

Methods

- Retrospective chart review of all HBV-positive mothers who gave birth at UNC hospitals from April 1, 2014 through December 31, 2019.
- Data metrics included:
 - Maternal HBV and HIV status
 - Maternal HBV viral load and HBV e antigen (HBeAg) testing
 - Time to neonatal HBIG and birth dose vaccination
 - Receipt of tenofovir by *high-risk* women
- Risk status and receipt of tenofovir were assessed for women who delivered on or after January 1st, 2016 (release of AASLD guidelines).

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Prevention of Mother-to-Child Transmission of Hepatitis B at UNC Hospitals

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haracteristic	No (%) or Mean ± SD
laternal Age	33.7 ± 5.14
1other HIV-Positive	1 (1.0)
ype of Delivery	
Spontaneous Vaginal	68 (68.7)
Caesarean Section	25 (25.3)
VBAC	3 (3.03)
Vaginal Assisted (forceps and/or vacuum)	3 (3.03)
Gestational Age	39w6d ± 15d
lumber of Premature [*] Infants	7 (7.07)
-minute APGAR	7.7 ± 1.7
-minute APGAR	8.8 ± 0.9
Sirth Weight (grams)	3149 ± 587

Table 1. Baseline Delivery Data for 99 HBsAg-positive Pregnant Women and Their Exposed Infants

Abbreviations: SD, standard deviation; VBAC, vaginal birth after C-section; APGAR, Appearance, Pulse, Grimace, Activity and Respiration; w, weeks; d, days. * Premature defined as born at <37w0d.

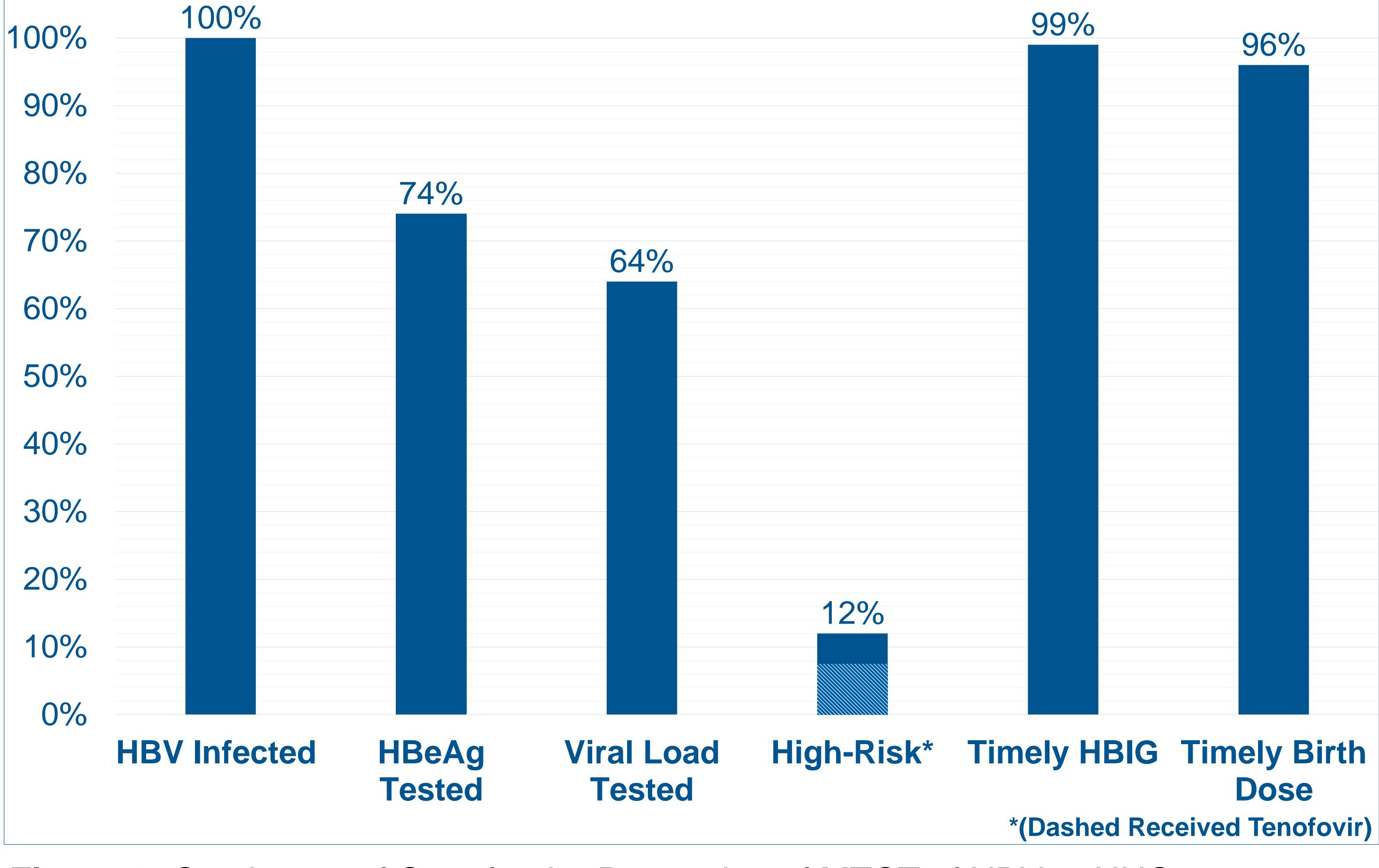


Figure 1: Continuum of Care for the Prevention of MTCT of HBV at UNC



Discussion

- UNC Hospitals were generally compliant with AASLD guidelines regarding:
 - Timely HBIG administration (99% of all neonates)
 - Timely neonatal vaccination (97%)

 Anti-viral use in women with high-risk HBV (83% of those identified as *high-risk*)

- Assessment of HBV risk status is an area for improvement, as HBV viral load and HBeAg were not routinely tested in this population. Without first identifying women with *high-risk* HBV, prophylactic tenofovir cannot be provided.
- Several infants did not receive timely birth dose vaccine because of prematurity. All HBV-exposed infants, regardless of birth weight, should receive birth dose vaccine within 12 hours.

Future Directions

- Future directions for improving care for this patient population include:
 - Increasing risk status assessments of HBV-positive mothers (viral load, HBeAg)
 - Assurance of follow-up care for HBV-positive mothers (GI, ID)
 - Follow-up testing for HBV-exposed infants (HBsAg, anti-HBs at 9-12 months)

References

Terrault NA, Bzowej NH, Chang K, et al. AASLD guidelines for treatment of chronic hepatitis B. *Hepatology* 2016;63(1):261-283.