BACKGROUND

PERINATAL HEPATITIS B

 Pregnant women should be universally screened for hepatitis B surface antigen (HBsAg) during each pregnancy to prevent perinatal transmission of hepatitis B virus (HBV) to their infant

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- Infants born to HBV-infected women are at risk for perinatal HBV transmission and subsequent chronic liver disease
- Appropriate and timely immunoprophylaxis of HBV-exposed infants is up to 95% effective in preventing infection
- Immunoprophylaxis includes: the hepatitis B immunoglobulin (HBIG) and HBV vaccine birth dose within 12 hours of birth, followed by completion of HBV vaccine series (Figure 1)

Fig. 1: HBV vaccine	Birth	1 mo	2 mos	4 mos	6 mos	9 mos	12 mos	15 mos	18 mos
schedule	≪ -1st dose-≯	<> ^{2nd} dose>			<3 rd dose3				>

NJDOH PERINATAL HEPATITIS B PREVENTION PROGRAM (PHBPP)

- Identifies HBsAg positive (+) pregnant women and their infants to ensure appropriate clinical management (i.e. immunoprophylaxis) (Fig. 2)
- According to CDC, only half of the 800-1000/year expected infants born to HBsAg + women in NJ are identified
- In 2009, PHBPP began using NJDOH's electronic web-based Communicable Disease Reporting and Surveillance System (CDRSS) for surveillance and case management; CDRSS is also used for other reportable communicable diseases

OBJECTIVES

- . Describe NJ's comprehensive perinatal HBV surveillance and case management system for HBV-infected mothers and their infants
- 2. Evaluate the sensitivity of case reporting HBV-infected mothers
- 3. Describe the vaccination and post-vaccine serology rates of HBVexposed infants

METHODS

DATA SOURCES

A. Surveillance data from CDRSS for:

- 1. HBV-infected women, who were reported during prenatal period and delivered a liveborn infant (LBI) from Jan. 1, 2010 through Dec. 31, 2011
- 2. Infants born Jan. 1, 2010 through Dec. 31, 2011 to HBV- infected women
- 3. All HBV cases from hepatitis B surveillance reported through Dec. 31, 2011

B. Hospital discharge data from the NJ Discharge Data Collection System for admissions during Jan. 1, 2010 through Dec. 31, 2011

• Using ICD-9 diagnosis codes, women with LBI deliveries and a HBV diagnosis

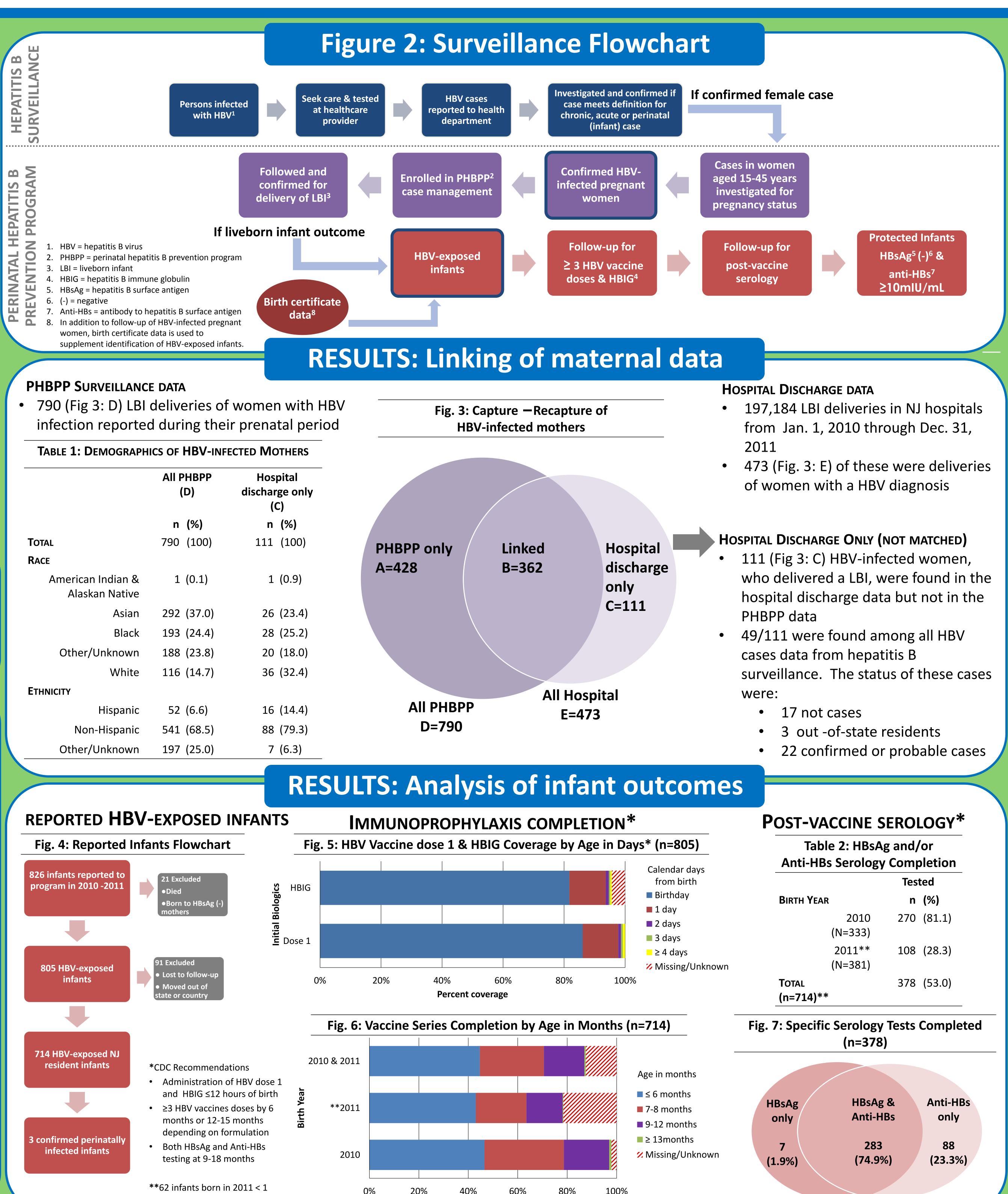
DATA ANALYSIS

A. LINKING OF MATERNAL DATA

- Using capture-recapture method to identify further cases of HBVinfected mothers, hospital discharge data was matched to surveillance data using Link Plus 2.0 and manually reviewed
- Matching variables: Mother's first name, last name, date of birth and date of delivery or admission
- **B.** ANALYSIS OF INFANT CLINICAL OUTCOMES
- Infant case management completion, including receipt of HBIG, birth dose, vaccination series and serology testing analyzed using SAS 9.2

Baby on Board? Evaluating Perinatal Hepatitis B Surveillance, New Jersey 2010-2011

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Percent coverage

year of age when data analyzed



DISCUSSION

LINKING OF MATERNAL DATA

- In the PHBPP program data (Fig. 3: D), 37% of the women were Asian and 14% White compared to 23% Asian and 32% White in hospital discharge only data (Fig. 3: C). 24% of the women in PHBPP were categorized as Other/Unknown race.
- Differences in proportions of Asian and White race may reflect PHBPP efforts to identify pregnant women from Asian HBV endemic countries
- 111 women in hospital discharge only (Fig. 3: C) and not PHBPP data
- 49/111 women found only in hospital data were found among all HBV cases data from hepatitis B surveillance
- This suggests confirmation of pregnancy status for each pregnancy in a HBV-infected woman is a limiting step for PHBPP enrollment; however, 17/49 were categorized as "not cases" among all HBV cases data from hepatitis B surveillance

ANALYSIS OF INFANT CLINICAL OUTCOMES

- Receipt of HBIG (95.7%) & vaccine birth dose (97.8%) within 1 calendar day of birth was high and close to national rate of 96.8% in 2008
- 23% (Fig. 7) of infants who received post-vaccine serology completed only the Anti-HBs test; CDC recommends both HBsAg and Anti-HBs testing at 9-18 months after completion of vaccine series

LIMITATIONS

- Hospital discharge data is limited by ICD-9 diagnosis coding and not a gold standard to identify all cases
- Exact time for administration of HBV vaccine doses and HBIG not captured and could not be analyzed for receipt within 12 hours
- Immnunoprophylaxis data may not be captured by PHBPP program due to reporting lags and follow-up delays

RECOMMENDATIONS

- Hospital discharge data can be an additional reporting source to identify HBV-infected women, who delivered a LBI
- Training for public health staff regarding the PHBPP surveillance to ensure data quality and completion
- Education and outreach for physicians and public health staff to ensure timely completion of vaccination & post-vaccination serology, particularly HBsAg testing

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