

Prenatal Syphilis in the US: Characterizing Screening and Treatment During Pregnancy in Publicly and Commercially Insured Individuals

Elizabeth A. Suarez¹, Christian Hague², June O'Neill², Katherine E. Round², Sarah K. Dutcher³, David Moeny³, Lucia Menegussi³, Terrence Lee³, Jamal T. Jones³, Nahida Chakhtoura⁴, Juanita J. Chinn⁴, Alexander C. Ewing⁵, Elizabeth B. Gray⁵, Phoebe Thorpe⁵, Catherine J. Vladutiu⁶, Rachel L. Abbey⁷, Judith C. Maro² 1. Department of Biostatistics and Epidemiology, Rutgers School of Public Health, New Brunswick, NJ; 2. Department of Population Medicine, Harvard Pilgrim Health Care Institute, Boston, MA; 3. Office of Surveillance and Epidemiology, Center for Drug Evaluation and Research, Food and Drug Administration, Silver Spring, MD; 4. Eunice Kennedy Shriver National Institute of Child Health and Human Development, National Institutes of Health, Bethesda, MD; 5. Centers for Disease Control and Prevention, Atlanta, GA; 6. Maternal and Child Health Bureau, Health Resources and Services Administration, Rockville, MD; 7. Office of the National Coordinator for Health Information Technology, Washington, D.C.

OBJECTIVES

To assess syphilis screening and treatment during pregnancy among publicly and commercially insured pregnant individuals in the US

BACKGROUND

- Cases of congenital (CS) syphilis have risen 10-fold in the US between 2012-2022
- A 2018 national study suggested that 28% of CS cases were due to a lack of timely prenatal care and syphilis testing, and 31% were due to inadequate maternal treatment in pregnancy
- Current CDC recommendations: 1st trimester screening and again at 28-weeks and delivery if high risk
- Between 2017-2021 in southern US states, first trimester testing rates were 41-64%; third trimester testing was performed in less than 50% of the pregnancies
- Data on syphilis screening rates in pregnancy are lacking from more recent time periods and from across the US and trends in the use and timing of treatment during pregnancy have not been examined in a large national study

METHODS

Cohort identified in Sentinel Distributed Database:

- US claims data: Medicaid (public) and commercial insurers
- Pregnancies resulting in live birth in individuals aged 10-54 years
- Continuous insurance coverage throughout pregnancy

Syphilis screening: 1+ procedure codes

Syphilis case: 2+ dates with diagnosis codes and no screening codes on the same date

Treatment:

- Dispensing records and administration billing codes
- Recommended treatment: Benzathine penicillin G
- Not recommended treatment: other antibiotics used for syphilis
- Included if treatment occurred up to 30 days following syphilis diagnosis and no recommended treatment received in pregnancy

RESULTS

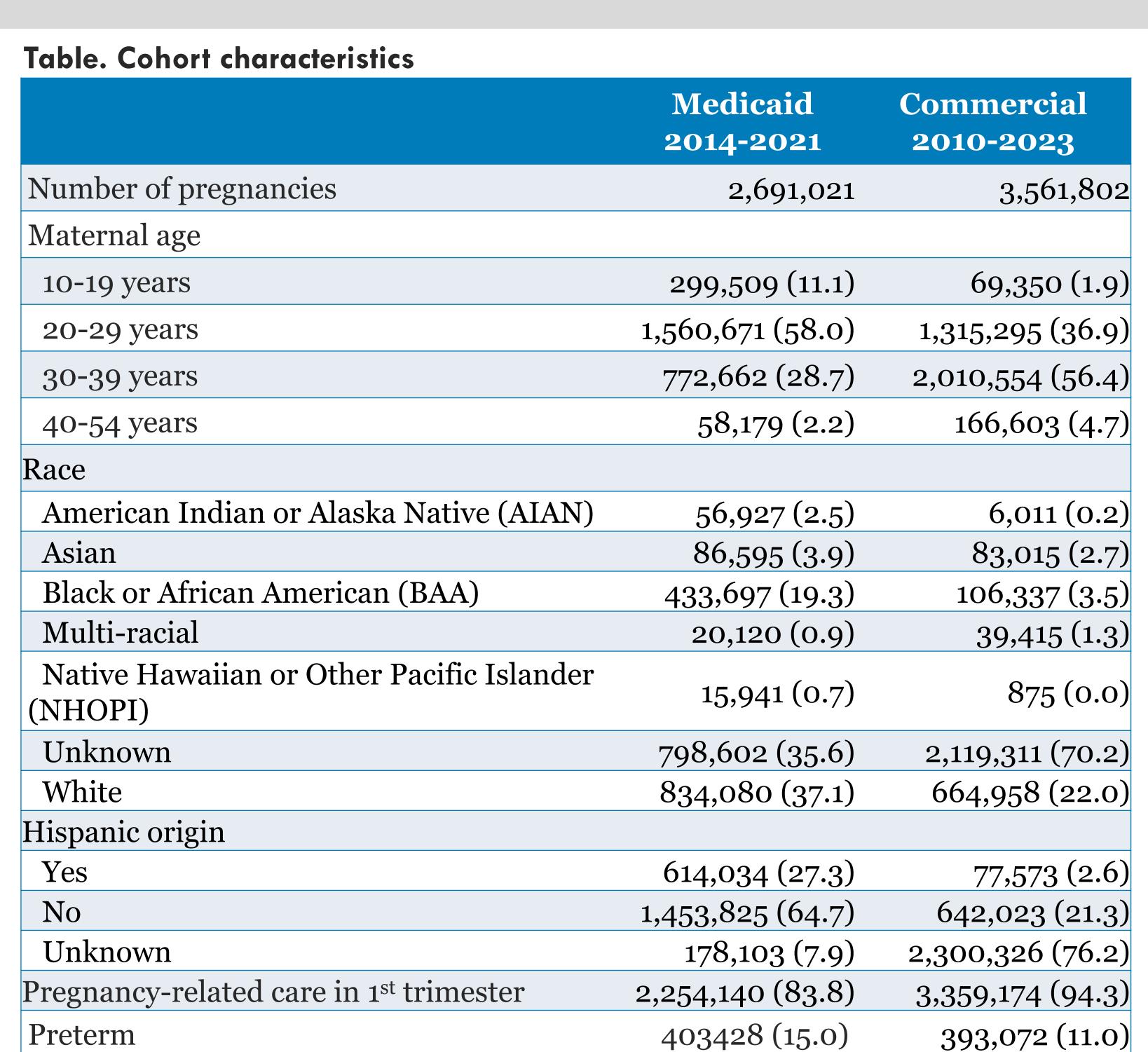


Figure 1. Screening prevalence in pregnancy by select covariates

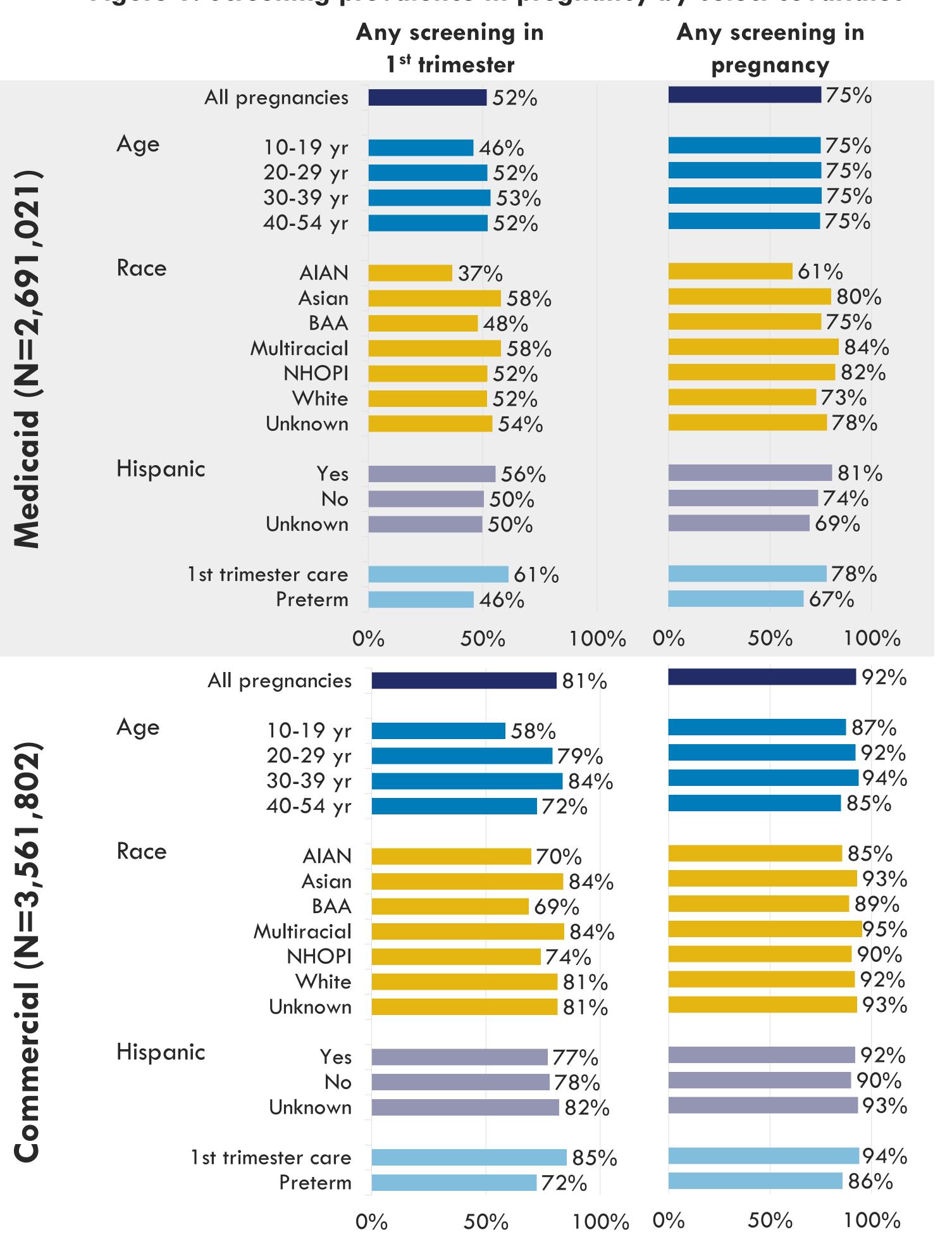
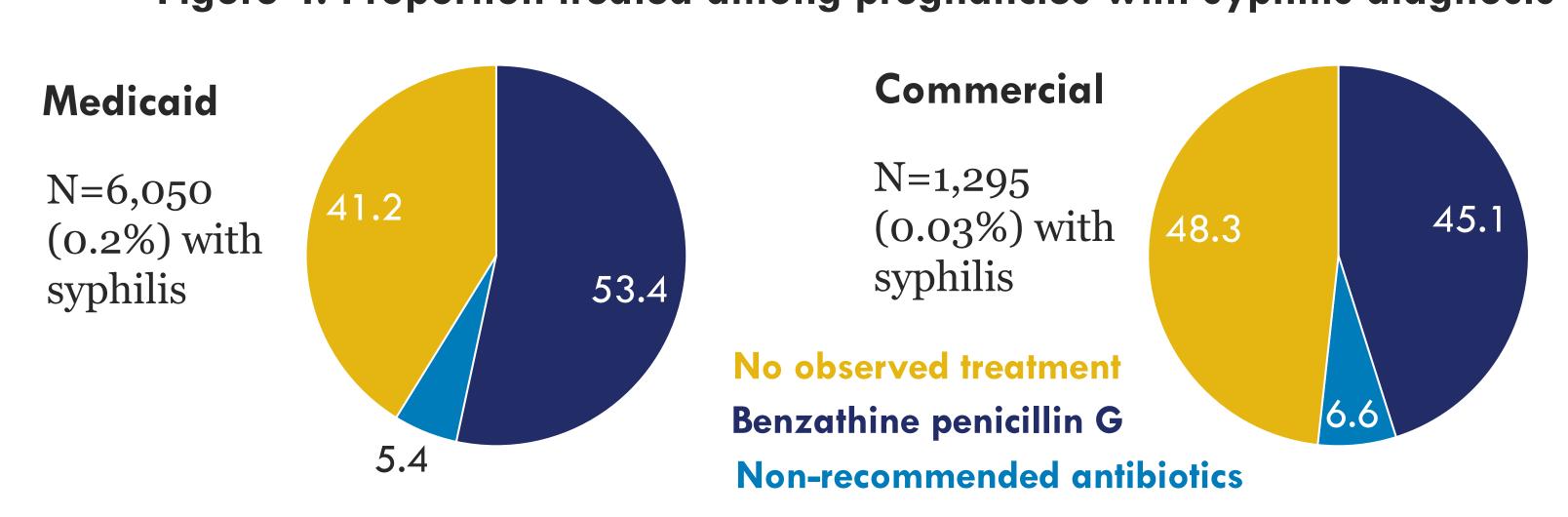


Figure 2. Timing of first screening Figure 3. Screening prevalence over time in pregnancy Any screening in 1st trimester 100% 90% 81% 90% 80% 80% 70% 70% 60% 60% 52% 50% 50% 40% 40% ййййййййййййййй 30% 25% Any screening in pregnancy 17% 20% 100% 10% 90% 6% 10% 2% 0% 80% 0% 70% Medicaid Commercial 60% 50% 1st trimester 3rd trimester 40%

Figure 4. Proportion treated among pregnancies with syphilis diagnosis

Medicaid Commercial



CONCLUSION

2nd trimester

- Medicaid-insured pregnant individuals were less likely be screened in first trimester (52%) vs 81%) and less likely to have any screening in pregnancy (75% vs 93%) than commercially-insured pregnant individuals
- Screening prevalence did not vary notably over time or by covariates.

At delivery

No screening

• 53% of Medicaid-insured and 45% of commercially-insured syphilis-diagnosed pregnant individuals had billed treatment with benzathine penicillin G; this is likely an undercount due to limitations in capturing treatment in these data.

LIMITATIONS

- Results of syphilis testing are not available in claims data; therefore, positive cases were approximated using diagnosis codes.
- Treatment may be under-captured due to receipt of treatment outside of traditional health care (i.e., at local health department) and inpatient setting due to bundled payments.

ACKNOWLEDGEMENTS/DISCLOSURES

endorsement, by HSA, FDA, CDC, NIH, HRSA or the U.S. Gov't.

- This work was funded by the Office of the Secretary Patient-Centered Outcomes Research Trust Fund (OS-PCORTF). The OS-PCORTF funding was made available to US Food and Drug Administration (FDA) by the Office of the Assistant Secretary for Planning and Evaluation (ASPE) through Interagency Agreement 750121PE080007. This work was the US Food and Drug Administration (FDA).
- supported by Task Order 75F40119F19001 under Master Agreement 75F40119D10037 from • The contents are of the authors and do not necessarily represent the official views of, nor an
- Some co-authors on this abstract are employed at organizations which conduct work for government and private organizations, including pharmaceutical companies. The authors have no conflicts of interest to disclose.
- Many thanks are due to Sentinel Data Partners who provided data used in the analysis.