Georgia Board of Public Health

November 8, 2022

GEORGIA DEPARTMENT OF PUBLIC HEALTH

Agenda

- Call to order
- Roll Call
- Approval/Adoption of Minutes
- New Business
 - Commissioner's Report
 - Epidemiology Updates
 - Influenza
 - o Early Hearing Detection and Intervention; Cytomegalovirus
 - Viral Hepatitis Prenatal Testing Requirements
- Board Comments
- Adjournment

Influenza Update

Board of Public Health / Cherie L. Drenzek DVM, MS, State Epidemiologist / Nov. 8, 2022

GEORGIA DEPARTMENT OF PUBLIC HEALTH

Introduction

- Flu is ever-changing, complex, and difficult to predict
- It remains a significant cause of morbidity and mortality globally
- Unlike COVID-19, individual influenza infections are not reportable to Public Health (not feasible--most are not lab confirmed)
- But flu surveillance is critical to inform prevention and control
- Flu "season" traditionally is October through May (but we can see flu all year round)

Goals of Influenza Surveillance

NOT to document every case of influenza, but to:

- Determine when and where influenza activity is occurring
- Determine what influenza viruses are circulating
- Determine the <u>severity</u> of influenza virus infections and who is most at risk
- Detect novel viruses or changes in influenza viruses

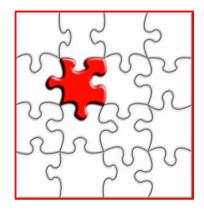
TO INFORM CONTROL AND PREVENTION EFFORTS!

What Does Flu Surveillance Entail?

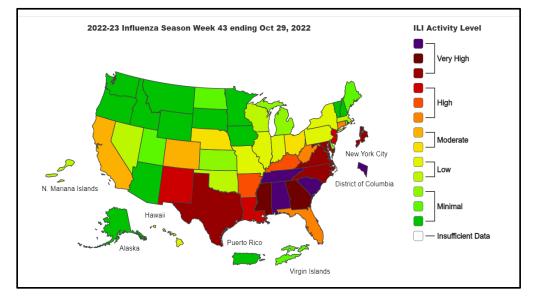
Pieces of the Puzzle:

- Virologic surveillance (lab characterization of what flu viruses are circulating)
- Proportion of outpatient visits for ILI (influenza-like illness-fever, cough, sore throat)
- Proportion of outpatient visits for ILI in Emergency Departments
- Comparing the level of ILI to baseline levels (non-flu season)
- Percent of flu tests that are positive
- Influenza hospitalizations (metro Atlanta only)
- Influenza-associated deaths (all ages)
- Influenza outbreaks

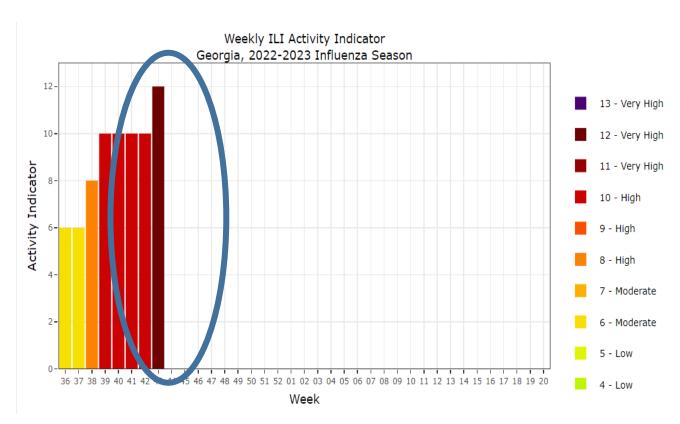
"Picture" of Flu Activity



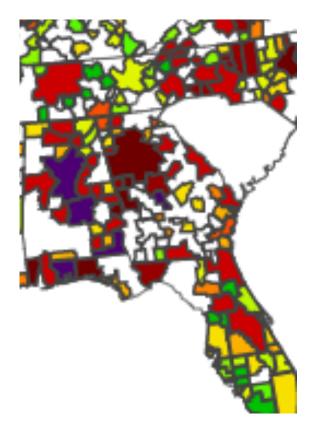
National Flu Snapshot



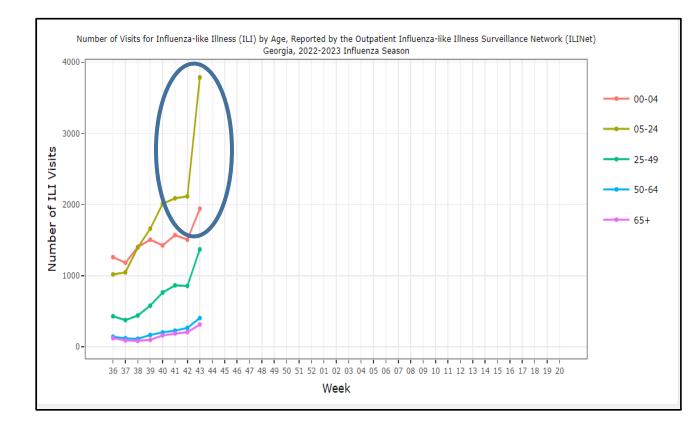
- The South and mid-Atlantic areas have the highest flu activity in the nation (this compares proportion of visits for ILI to non-flu season)
- This is very early and has been rising for two months
- About 75% of flu viruses have been Influenza A H3N2
- 4.3% of outpatient visits were for ILI (national baseline is 2.5%)
- About 9% of all tests for flu were positive (the week before was 6%)
- Hospitalization rate (3/100K) hasn't been this high this early since 2010



- Flu activity has been rising in Georgia since mid-August, was HIGH for 5 weeks, and now is VERY HIGH.
- 100% of viruses are Influenza A, and most are A (H3N2)
- The percent of positive flu tests is 18% (9% nationally)
- The proportion of outpatient visits for ILI is 10% (baseline is 3%, the earliest it has been this high since the 2009 pandemic)



 We see flu activity all around Georgia, but have VERY HIGH activity in large areas of metro Atlanta, Macon, and Columbus (see maroon, purple areas)



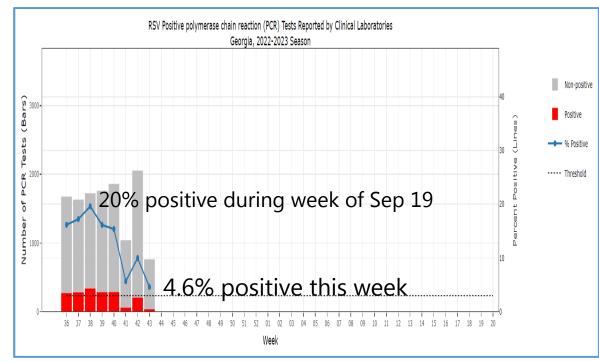
- The vast majority of outpatient visits for ILI have been among the "young" (25% among 0-4 years, 50% among 5-24 years old)
- Underscores importance of flu vaccination in this population
- "H3N2" seasons tend to disproportionately affect children and the elderly

- Flu-related hospitalizations (metro Atlanta) nearly doubled over the last two weeks; 30% were in those older than 65; our overall hospitalization rate is 10/100K.
- We have confirmed **2** flu-related deaths so far this season and **31** last season (both this season occurred in those >50 years).
- We documented 58 flu outbreaks in institutional settings in the last week—about half in K-12 schools and half in LTCFs. This was up from 5 outbreaks each in the previous two weeks.
- Again, underscores particular importance of flu vaccination in old and young (but certainly for everyone!!)

Influenza: Prevention

- Annual vaccination remains the <u>best method</u> for preventing seasonal flu (and protecting against serious outcomes like hospitalization and death) and is recommended for **all persons aged 6 months and older.**
- The flu vaccine is quadrivalent—it protects against **four** flu strains: Influenza A H3N2, Influenza A H1N1, Influenza B/Victoria, Influenza B/Yamagata
- For this flu season, individuals over 65 are recommended to get one of the 3 high-dose or adjuvanted vaccines
- With this early flu activity, important to get vaccinated now (takes a few weeks for max immunity)
- In addition, flu prevention also relies on the pillars of handwashing, staying home if you're sick, and respiratory etiquette.

Respiratory Syncytial Virus (RSV) in Georgia



- RSV infection can be very serious or even deadly in young children, those under 12 months most at risk for hospitalization.
- No vaccine for RSV
- The RSV season has also been severe and early this year (usually peaks late winter)
- Similar to flu in that we don't count individual cases, but use percent of lab specimens that are positive for RSV to look at trends
- RSV activity was very high in Georgia during September and October, with a peak of 20% positive on Sep 19, but appears to be declining

Closing Comments

- 1. Infectious diseases are ever-changing and unpredictable!
- **2.** Surveillance and epidemiologic investigation are the cornerstones of prevention and control recommendations.

Questions

For more information, please contact:

Cherie Drenzek, DVM, MS

State Epidemiologist & Chief Science Officer Georgia Department of Public Health (404) 657-2609 <u>cherie.drenzek@dph.ga.gov</u>

Early Hearing Detection and Intervention

Board of Public Health Meeting / Tina Turner, MPA / Nov. 8, 2022

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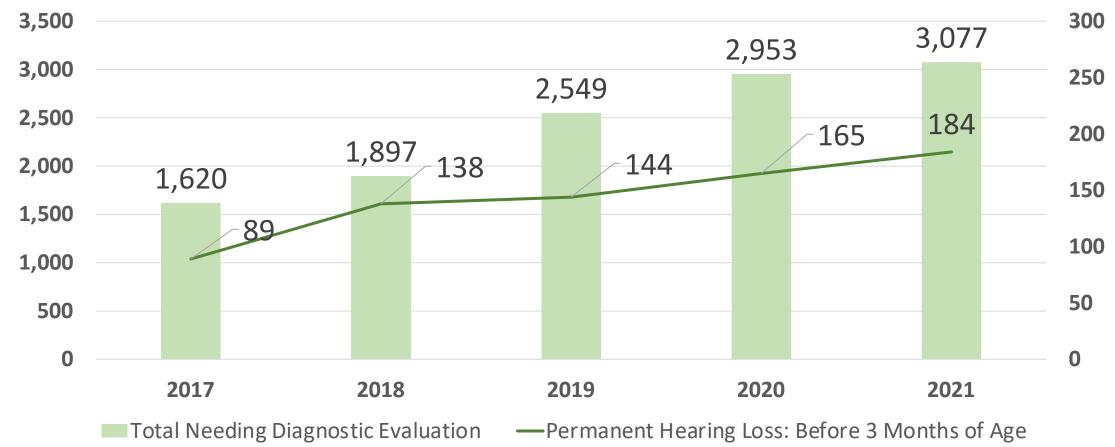
Georgia EHDI Program

- Public health program since 2001
- 2014 hearing screening added to Newborn Screening Rule 511-5-5
 Mandated hospital screening and reporting for every infant
- Purpose of the program:
 - $\ensuremath{\circ}$ Statewide screening and referral system
 - Linkages to appropriate intervention
 - Technical assistance and training for hospitals, physicians, audiologists, early intervention and public health

General EHDI Protocols in Georgia

- Initial hearing screening prior to discharge
- ONE follow up screen as an outpatient (well baby nurseries)
- Diagnostic Evaluation
- Early Hearing Orientation Specialist (EHOS) visit
- Enrollment in Early Intervention and Support Services
- Transition to educational services / support

Number of Babies Identified with Hearing Loss Before 3 Months of Age is Increasing



Source: 2018-2021 HSFS Data for PHIP Request as of 8-17-2022

Achieving Goals

- Babies are screened at birthing hospitals prior to discharge
 - Automated screen result: pass/fail/refer
 - Technicians, nurses or audiologists perform screen
 - Only two screens allowed before discharge
- Final screen/hearing screening status at discharge is reported to public health through SendSS

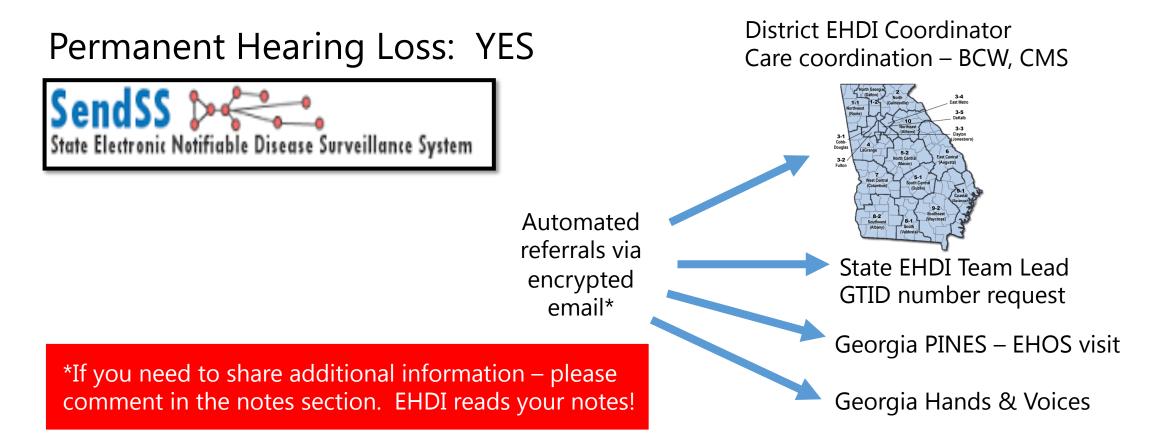
District Follow Up Coordinators

- A designated coordinator in every health district as of 2009
 Work with birthing facilities, families, primary care physicians, and audiologists
- Enhance service delivery to meet the overall goals of the program in a timely manner
- Provide technical assistance, when needed

Diagnostic Evaluations

- Diagnostics are completed by a licensed Audiologist, either in a natural sleep "nap study" or with sedation
- Providers are mandated to report results to DPH within 7 days of testing
 - $\circ~$ Web-based reporting directly into the SendSS via the
 - Audiology Portal
 - Paper surveillance form or report

Early Intervention Referrals



Congenital Cytomegalovirus

GEORGIA DEPARTMENT OF PUBLIC HEALTH

Cytomegalovirus

- Cytomegalovirus (CMV) is a virus that infects people of all ages
- If you are pregnant, CMV is cause for concern because the infection can be passed on to the baby
- If a baby is born with a CMV infection, it is called congenital CMV (cCMV)
- 1 in 5 children with congenital CMV will have a permanent disability, such as hearing loss or developmental delay

Goals for cCMV Training and Education

Goal 1: Train all applicable hospital providers and screening personnel and district staff on how to educate families and care givers of infants who fail/refer their newborn hearing screening for one or both ears prior to hospital discharge regarding cCMV as a potential cause for not passing. December 2022 – August 2023.

Goal 2: Provide education for families and caregivers of infants who fail/refer their newborn hearing screening for one or both ears prior to hospital discharge regarding cCMV as a potential cause for not passing. April 2023 – August 2023

Questions

For more information, please contact:

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Brandt Culpepper, Ph.D., CCC-A

Early Hearing Detection and Intervention Team Lead Maternal and Child Health Section 404-651-5482 Brandt.Culpepper@dph.ga.gov

Viral Hepatitis

Updated Prenatal Testing Requirements

Board of Public Health / Tracy Kavanaugh, MS, MCHES; Megan Andrews, J.D. / Nov. 8, 2022

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Reportable Conditions – Perinatal Hepatitis B & C

Hepatitis B

- HBsAg(+) pregnant women
- Perinatal HBV exposure •

Hepatitis C

- anti-HCV(+) or HCV RNA • detected pregnant women
- anti-HCV(+) or HCV RNA ٠ detected children ages <3 years

Must be reported within **7** days

		DISEA	SE	AIDS#	-anti-HCV(+) or HCV RNA
CONDITION REPORTING			ed by law to report	acute flaccid myelitis anaplasmosis aseptic meningitis babesiosis	detected children ages <3 years hepatitis D (Delta virus present with HBsAg);
REPORT IMMEDIATELY REPORT WITHIN 7 DAYS					
To Report Immediately Call: District Health Office		ADS*	- anti-HCV(+) or HCV RNA	blood lead level (all)	acute and chronic hepatitis E
or 3-866-PUB-HLTH (3-866-782-4584)		anaplasmosis <3	detected children ages <3 years	campylobacteriosis	(acute) influenza-associated
any cluster of illnesses	novel influenza A virus	aseptic meningitis babesiusis	hepatitis D (Delta virus present with HBsAg)	Carbapenem-resistant	death
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- bruceflosis	polomyelitis	Chlamydia trachomatis (penital infection)	(Mycobacterium leprae) Lyme disease	chancroid	leprosy or Hansen's disease
cholera	+ O fever	Creutzfekitt Jakob Disease	lymphogranuloma venereum malaria		(Mycobacterium leprae)
diphtheria E-mil-0157	rabies (human & animal)	(CJD), suspected cases, under age 55	maternal deaths	Chlamydia trachomatis	Lyme disease
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hantawinus pulimonary syndrome	MIC ≥ 4µg/mi	gonorrhea HIV infection*	children) mumps	(CJD), suspected cases,	malaria
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(HUS)	syphilis (adult)	(permanent under age 5)** hepatitis 8	rubella (including congenital) salmonellosis	cryptosporidiosis	(during pregnancy or within
hepatitis A (acute)	syphilis during pregnancy	- acute hepatitis B	higeliesis treptococcal disease.		1 year of end of pregnancy)##
meanim (rubecka)	fuberculosis	 dwonic HEsAg(*) or HEV DNA detected infections 	Group A or B (invasive)** Streptopopous preumoniae		MIS-C (multi-system
melioidosis	latent TB infection in children-5 years old	 HBsAg(+)pregnant women Perinatal HBV exposure 	(invasion)**	ehrlichiosis	inflammatory syndrome in
meningitis (specify agent)	+ b/avnia	hepatitis C (past or present) - anti-HCV(+)	Stitunce information	giardiasis	children)
meningococcal disease (invasive)	 viral hemorrhagic fevers 	- HCV RNA detected - HCV genotype detected	oxic shock syndrome typicid aricella (Chickerpox) Vibio	gonorrhea	
		 anti-HCV(*) or HCV 854A detected prepriet average 	infections	HIV infection#	mumps
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or more information				- anti-HCV(+)	tetanus
ww.dph.ga.gov/disea	se-reporting			 HCV RNA detected 	toxic shock syndrome typhoid
			Streep severage of the ground	 HCV genotype detected 	Varicella (Chickenpox) Vibrio
and Jacomban 2011 Secure Separtment of Auto-Hauto-				 anti-HCV(+) or HCV RNA 	infections
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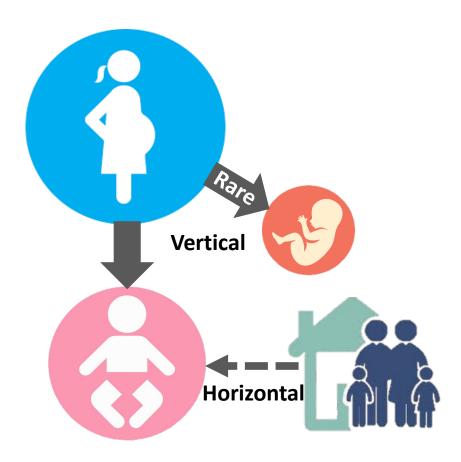
Perinatal Hepatitis Transmission

Vertical

- Contact with infected blood or body fluids
- In utero transmission is rare

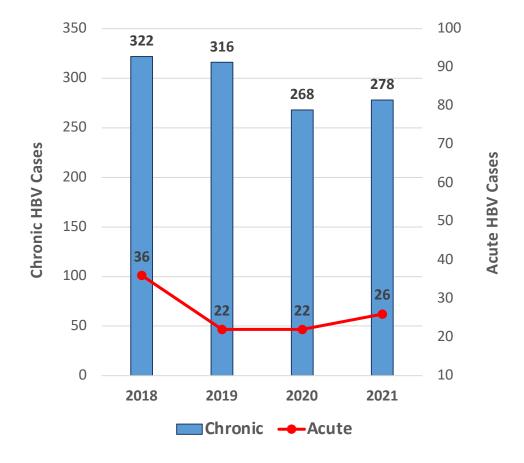
Horizontal

• Interpersonal contact

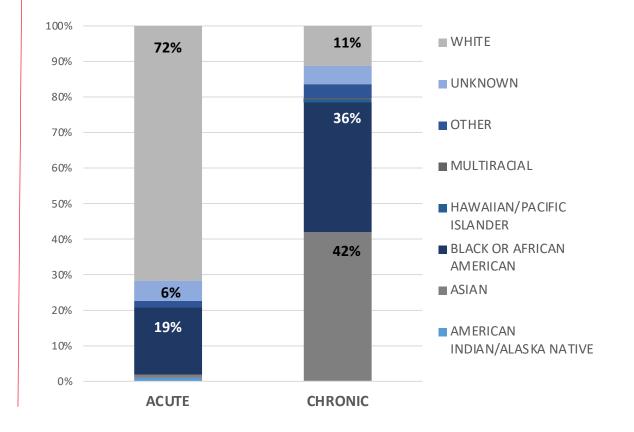


HBV Cases in Females Ages 13-47 Years

Hepatitis B Infections in Females 13-47 Years of Age by Diagnosis, Georgia, 2018-2021



Hepatitis B Infections in Females 13-47 Years of Age by Race and Diagnosis, Georgia, 2018-2021



Perinatal Hepatitis B Infection

• Perinatal Transmission Risk:

○ 70%-90% HBsAg(+) and HBeAg(+)

○ 10% HBsAg(+) only

Perinatal HBV Outcomes:

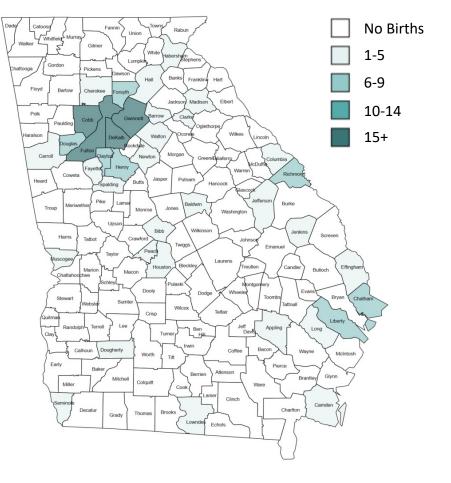
- o 90% of infected infants become chronically infected
- o 25% may die prematurely

Post-exposure Prophylaxis:

o 94% effective in reducing perinatal transmission

Perinatal Hepatitis B Exposures in Georgia

HBV-Exposed Births by Infant's County of Residence, Georgia, 2020





75% of HBV-Exposed Infants were born to a Mother Born Outside of the USA in 2020

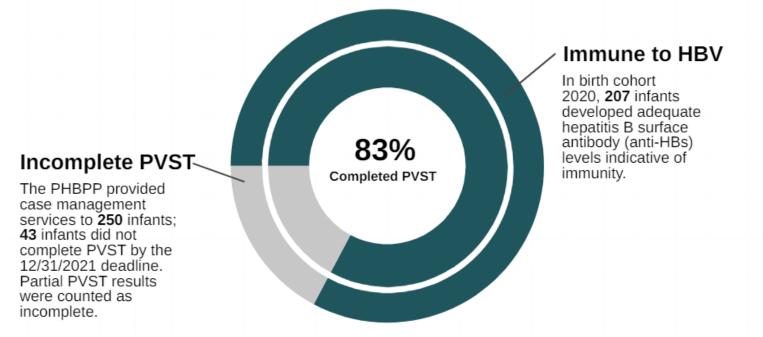
Georgia Perinatal Hepatitis B Prevention Program

- Identify hepatitis B-infected pregnant women
- Ensure hepatitis B immune globulin (HBIG) and hepatitis B vaccine are administered within 12 hours of birth
- Ensure hepatitis B vaccines are administered at the recommended intervals
- Coordinate post-vaccination serologic testing (PVST) at 9-12 months of age

Birth Cohort 2020 Outcomes

Postvaccination Serologic Testing (PVST)

Postvaccination serologic testing (PVST) is recommended for infants and children born to hepatitis B-infected mothers. The PVST must include the hepatitis B surface antigen (HBsAg) and hepatitis B surface antibody (anti-HBs). Postvaccination serologic testing confirms whether the child has developed immunity, needs additional vaccine or has been infected with the hepatitis B virus (HBV).



No HBV-infected infants were identified in birth cohort 2020.

9 Infected Infants 2011-2020

Perinatal Hepatitis C Infection

• Perinatal Transmission Risk¹:

- $\circ~$ 7.2%% with no HIV infection
- 12.1% with HIV co-infection

Perinatal HCV Outcomes:

- 20%-40% of infected infants spontaneously clear the infection by age 5
- o 50% have chronic asymptomatic infection
- 30% have chronic active infection

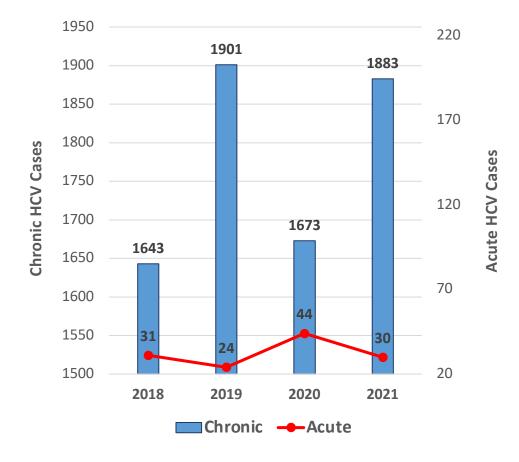
Post-Exposure Prophylaxis

 \circ None

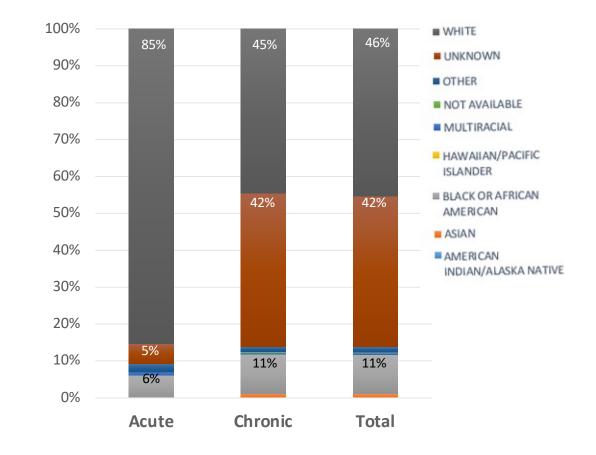
¹Ades, A.E., et al., Overall vertical transmission of HCV, transmission net of clearance, and timing of transmission. Clin Infect Dis, 2022.

HCV Cases in Females Ages 13-47 Years

Hepatitis C Infections in Females 13-47 Years of Age by Diagnosis, Georgia, 2018-2021

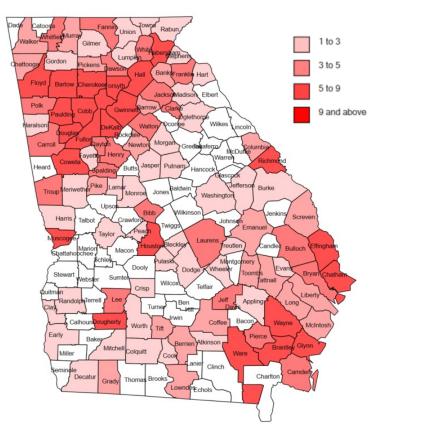


Hepatitis C Infections in Females 13-47 Years of Age by Race and Diagnosis, Georgia, 2018-2021



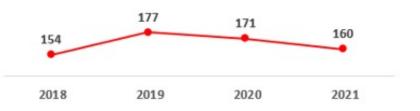
HCV RNA-Exposed Births 2018-2021

Maternal HCV RNA(+) Result <365 Days of Birth



Perinatal Hepatitis C Exposures, Georgia, 2018-2021

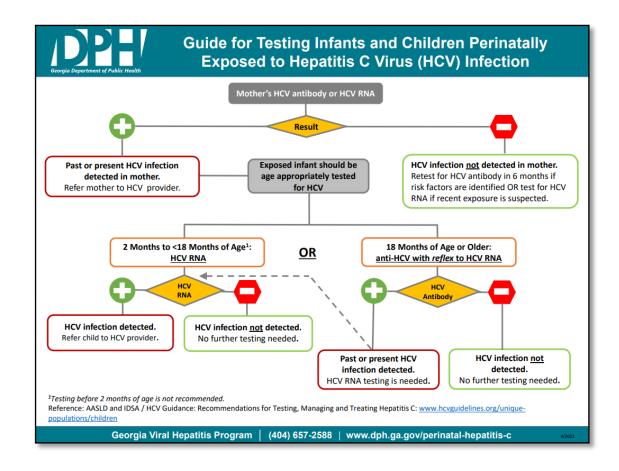
- 2,172 births to persons with an HCV(+) history
- 662 births to persons with HCV RNA(+) lab test <365 days of birth



Source: Perinatal Hepatitis C SendSS Data *Preliminary data and subject to change

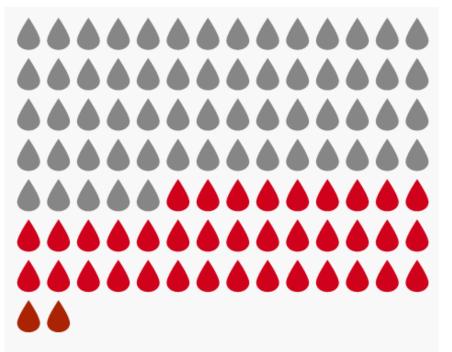
Perinatal Hepatitis C Program

- Identify the birth
- Notify Pediatric Provider
- Coordinate testing
- Verify laboratory results



Case Management Outcomes, 2018-2021

Case Managed Infants/Children Testing Outcomes, Georgia, 2018-2021



Infants Not Tested (60.96%)
 Infants Not Infected (37.24%)
 Infected Infants (1.8%)

Birth Cohort	Case Managed	Not Case Managed
2018	4	5
2019	3	4
2020	3	1
2021	1	0
2022	0	2
Total	11	13
Reported by Birth Cohort 24 Reported by Year of Onset		

Includes cases that may be in case management database but did not have a maternal HCV RNA+ result within 365 days of infant's birth and were not actively being case managed

Source: Perinatal Hepatitis C SendSS Data *Preliminary data and subject to change

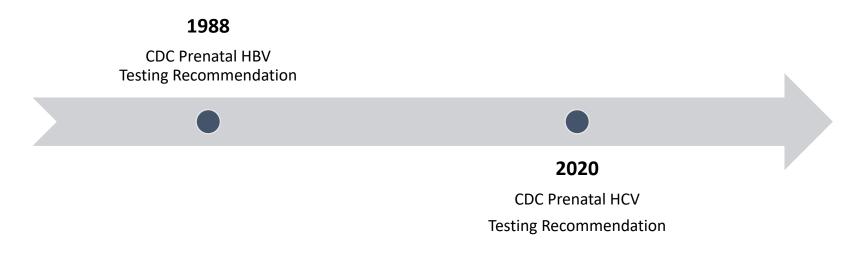
National Testing Recommendations

Prenatal Hepatitis B Testing

- Centers for Disease Control and Prevention (CDC)
- The American College of Obstetricians and Gynecologists (ACOG)

Prenatal Hepatitis C Testing

- Centers for Disease Control and Prevention (CDC)
- The American College of Obstetricians and Gynecologists (ACOG)
- American Association for the Study of Liver Disease (AASLD)



Serologic Tests for Pregnant Women

Chapter 511-5-4 / Authority: O.C.G.A. §§ 31-2A-6, 31-12-3 and 31-17-4.

Effective: August 31, 2022

First Prenatal Visit:

• HBV and HCV testing

Third Trimester:

 HCV testing for women at continued risk or with exposure to hepatitis C

At Delivery:

- HBV testing for women not tested prenatally, with signs or symptoms of hepatitis, or at high risk for hepatitis B
- HCV testing for delivery for women not tested prenatally



Serologic Testing Requirements

FIRST PRENATAL VISIT

TEST ALL PREGNANT WOMEN:

Hepatitis B (HBV): HBsAg
Hepatitis C (HCV): anti-HCV with reflex to HCV RNA
HIV: HIV EIA or Rapid Assay (fingerstick preferred)
Syphilis: Non-treponemal (RPR) with reflex to treponemal test



Serologic Testing Requirements

THIRD TRIMESTER

TEST ALL PREGNANT WOMEN:

HIV: HIV EIA or Rapid Assay (fingerstick preferred) before **36 weeks Syphilis:** Non-treponemal (RPR) with reflex to treponemal test ideally at **28 to 32 weeks** of gestation

TEST SELECT PREGNANT WOMEN AT CONTINUED RISK OR WITH KNOWN EXPOSURE:

Hepatitis C: anti-HCV with reflex to HCV RNA



Serologic Testing Requirements

AT DELIVERY

ASSESS ALL PREGNANT WOMEN:

Hepatitis B, Hepatitis C, HIV, and Syphilis testing

TEST SELECT PREGNANT WOMEN:

Hepatitis B (HBV): HBsAg

- No evidence of screening during pregnancy Persons at high risk
- Signs or symptoms of hepatitis
- Hepatitis C (HCV): anti-HCV with reflex to HCV RNA
- No evidence of screening during pregnancy
- **HIV:** HIV EIA or Rapid Assay (fingerstick preferred)
- No evidence of screening during pregnancy
 Persons at high risk
- Persons not tested in the third trimester
- Syphilis: Non-treponemal (RPR) with reflex to treponemal test
- No evidence of screening during pregnancy
 Persons who deliver a stillborn infant(s)

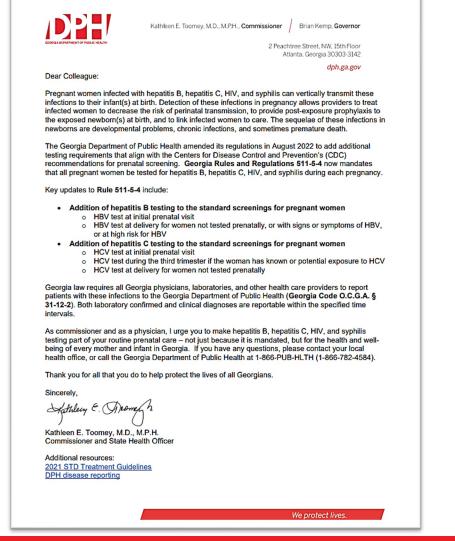
• Persons at high risk

Persons not tested in the third trimester



Outreach

- Dear Colleague Letter
- Partner organization collaborations
 - Georgia Hospital Association
 - o Georgia OB/Gyn Society
 - Georgia Academy of Family Physicians
 - Georgia Chapter of the American Academy of Pediatrics
 - Medical Association of Georgia
 - Department of Community Health
- Webinars
 - Viral Hepatitis Project ECHO sessions in October and November



Questions

For more information, please contact:

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Perinatal Hepatitis B Program Coordinator Phone: (404) 651-5196 <u>Tracy.Kavanaugh@dph.ga.gov</u>

Megan Andrews, J.D.

Assistant Commissioner for Policy Georgia Department of Public Health Phone: (404) 657-27278 <u>Megan.Andrews@dph.ga.gov</u>

Perinatal Hepatitis B Web Page: <u>www.dph.ga.gov/perinatal-hepatitis-b</u> Perinatal Hepatitis C Web Page: <u>www.dph.ga.gov/perinatal-hepatitis-c</u>

Next Meeting

The next Board of Public Health Meeting will be held Jan. 10, 2023