



Sensitivity of Diagnostic Codes in Identifying Laboratory Confirmed Congenital **Cytomegalovirus Infections in Electronic Health Record Database** Alexandra Campione^{1,2}, Tatiana M Lanzieri³, Emily Ricotta², Scott D. Grosse⁴, Scott Quinlan¹, Sameer Kadri⁵, Veronique Nussenblatt⁶, D Rebecca Prevots²

¹ Milken Institute School of Public Health, The George Washington University, ² Epidemiology Unit, Division of Intramural Research, National Institutes of Allergy and Infectious Disease (NIAID), National Institutes of Health (NIH), ³ National Center for Immunization and Respiratory Diseases, Centers for Disease Control and Prevention (CDC)⁴ National Center on Birth Defects and Developmental Disabilities, CDC, ⁵ Critical Care Medicine Department, NIH Clinical Center, ⁶ Infectious Disease, NIAID, NIH

Introduction

- About 1 in 200 babies are born with congenital Cytomegalovirus (cCMV) in the US and 90% of infections are asymptomatic¹
- Infections can result in permanent sequelae such as sensorineural hearing loss, vision impairment, and developmental disabilities
- More than 50% of symptomatic infections result in permanent sequelae²



- Studies using administrative databases have used CMV diagnostic codes assigned within the first 90 days of life to identify cCMV infections and found a prevalence of 1-4 per 10,000 births³
- ICD-9-CM and ICD-10-CM diagnostic codes are used primarily for billing or administrative purposes, and their sensitivity in identifying cCMV infections has not been evaluated

Objectives

- 1. To evaluate the sensitivity of diagnostic codes in identifying infants with laboratoryconfirmed cCMV infections at healthcare facilities that provide CMV lab data
- 2. To determine the prevalence of laboratoryconfirmed cCMV
- 3. To investigate conditions associated with cCMV using diagnostic codes

Methods

Patient Population: Infants with at least one enco life at a facility that provided CMV data to Cerner 2017

Laboratory-confirmed cCMV infected infants: Infants with a positive CMV laboratory test meeting inclusion criteria in the first 21 or 90 days of life

Inclusion Criteria: PCR, DFA, or culture from saliva, urine, respiratory secretion, CSF, or blood samples or IgM serology

Diagnostic Codes (ICD-9-CM and ICD-10-CM) corresponding to CMV and CMVrelated conditions were investigated among the laboratory-confirmed cCMV infected infants. A restriction of 21 or 90 days was applied for CMV diagnostic codes to account for delay in assignment and 90 days for all other diagnostic codes.

Results

7,908,711 infants with encounters in first 90 day life (349 hospitals)



668 infants were within 21 days at time of te

CNAV Tocting Critoria	CMV Diagnostic	Tota
Civity resting Criteria	Code Assignment	of In
CMV test in first 21 days	≤21 days	668
CMV test in first 21 days	≤90 days	668
CMV test in first 90 days	≤90 days	838
The sensitivity of CMV dia infants using 21- and 90-d	gnostic codes amo lay cut-offs for CM	ong la V tes

ounter in the first 21 or 90 days of
r Health Facts Database from 2010-

Sorv	Prevalen confirme Using 90 tests	<u>ce of Laboratory-</u> <u>ed cCMV infected</u> <u>infants:</u>)-day cut-offs for /encounters:	 Wa Ma <
est	Using 21 tests 0.9 cases	L-day cut-offs for /encounters: per 10,000 infants	1. Go Key
l No. fants	No. with CMV codes	Sensitivity % (95% CI)	- Inf 2. Do pre mo
	69 74	10.3 (8.0 <i>,</i> 12.6) 11.1 (8.7 <i>,</i> 13.5)	inf 3. Gro CN

12.8 (10.5, 15.0) 107 aboratory-confirmed cCMV infected sts and diagnostic code assignment

Conclusions

• About 1 in 10 infants with laboratory-confirmed cCMV also had CMV diagnostic codes in this population of infants at 349 US hospitals The sensitivity of CMV diagnostic codes did not substantially change with different age restrictions

• The administrative prevalence of cCMV was like other studies, but laboratory results were used instead of diagnostic codes to identify infants • Low birth weight (31%), jaundice (26%), and thrombocytopenia (15%) were the most common cCMV associated conditions in those less than 21 days at time of test. Microcephaly as present in about 4% and hearing loss in 6%. lost laboratory-confirmed infants (21 days) ere identified through culture alone (77%) nd only 10% had a positive PCR test fants in the Northeast represented 20% of the tal patient population and 55% of the boratory-confirmed cCMV infected population hich indicates missing laboratory data.

References

oderis J, De Leenheer E, Smets K, Van Hoecke H, eymeulen A, Dhooge I. Hearing loss and congenital CMV fection: a systematic review. Pediatrics 2014;134:972-82. Ilard SC, Grosse SD, Ross DS. New estimates of the evalence of neurological and sensory sequelae and ortality associated with congenital cytomegalovirus fection. Rev Med Virol 2007;17:355-63

rosse SD, Leung J, Lanzieri TM. Identification of congenital CMV cases in administrative databases and implications for monitoring prevalence, healthcare utilization, and costs. Curr Med Res Opin 2021;:1-23.

Corresponding author: Alexandra Campione email: acampione@gwu.edu