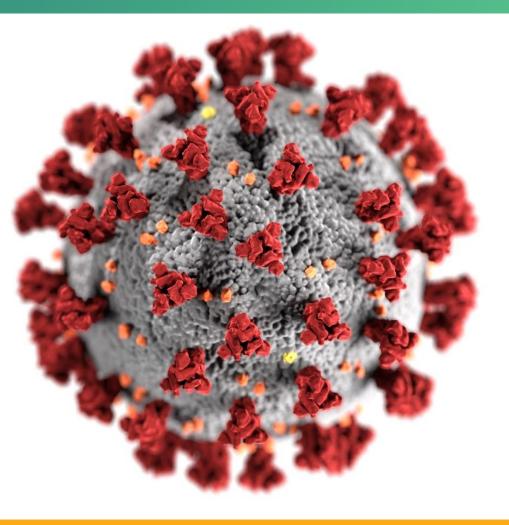
Epidemiology of COVID-19-Associated Hospitalizations, including in Pregnant Persons and Infants

Fiona Havers, MD, MHS, FIDSA

Team Lead, RESP-NET Hospitalization Surveillance Team Commander, US Public Health Service Coronavirus and Other Respiratory Viruses Division National Center for Immunization and Respiratory Diseases Centers for Disease Control and Prevention

Advisory Committee on Immunization Practices June 23, 2023





cdc.gov/coronavirus

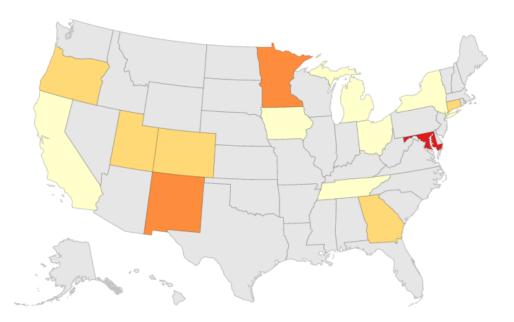
COVID-19-associated hospitalizations

COVID-NET: March 2020 – June 2023



COVID-NET Hospitalizations

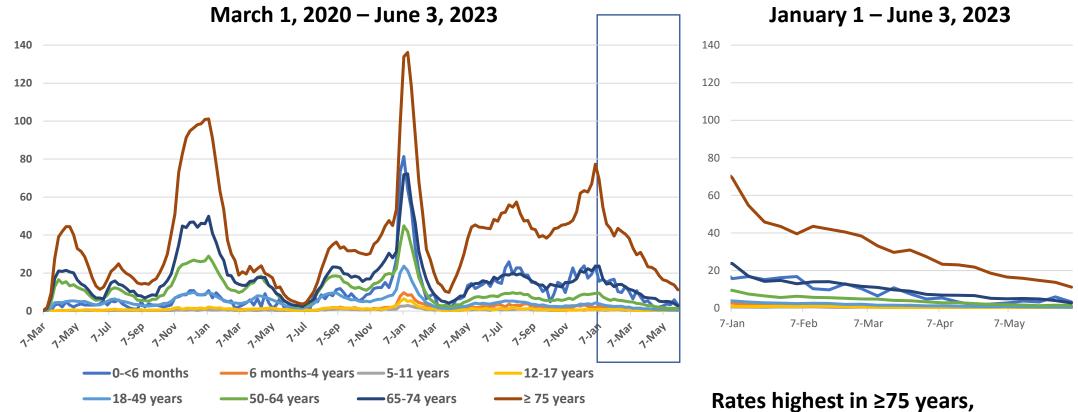
- >250 acute-care hospitals
- 98 counties in 13 states
- ~10% of U.S. population
- Positive SARS-CoV-2 within 14 days of or during hospitalization
- Screening or clinician-driven testing
- Clinical data is from representative sample of COVID-NET patients



Percentage of state population represented by participating COVID-NET counties • 4% - < 28% • 28% - < 52% • 52% - < 76% • 76% - 100%



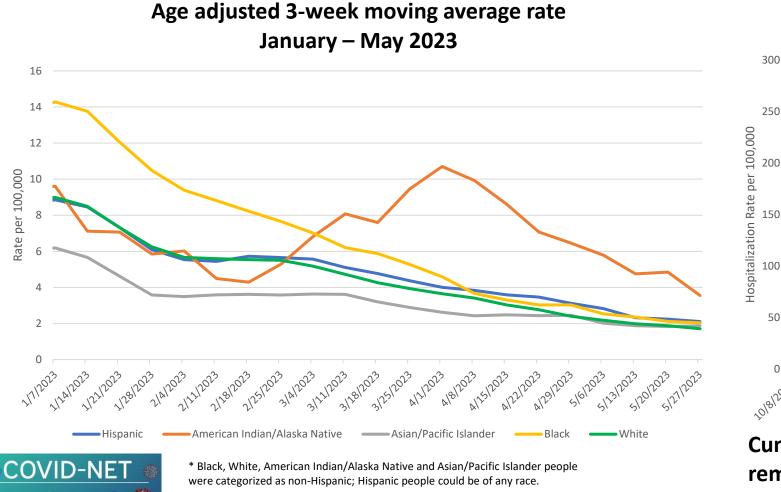
Weekly COVID-19-associated hospitalization rates — COVID-NET, 14 U.S. States



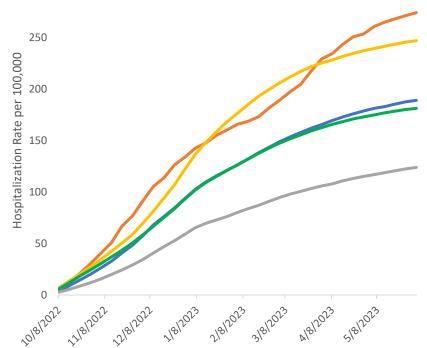


Rates highest in ≥75 years, followed by infants <6 months and those 65 – 74 years

COVID-19-associated hospitalization rates by race and ethnicity* — COVID-NET, 14 U.S. States, October 2022 – May 2023



Age adjusted cumulative rates October 2022 – May 2023



Cumulative hospitalization rates remain highest in American Indian/Alaska Native and Black persons

COVID-19-associated hospitalizations in pregnant persons

COVID-NET: January 2021 – April 2023



COVID-19-associated hospitalizations among pregnant persons aged 15–49 years — COVID-NET, 14 U.S. States, January 2021–April 2023

Most pregnant people hospitalized with a positive SARS-CoV-2 test had no respiratory symptoms

- 363 of 1,651 (21%) had respiratory symptoms recorded
- Proportion without respiratory symptoms increased from 73% to 82%

Among symptomatic patients, the proportion with underlying medical conditions increased



Table 1. Characteristics of COVID-19-associated hospitalized pregnant patients (15–49 years) with respiratory symptoms

	January – December 2021 –		July 2022 – April	
	November 2021	June 2022	2023	
	(pre-Omicron)	(early Omicron)	(later Omicron)	
	N (%)	N (%)	N (%)	
Total number	184	99	80	
Age group				
15-24 years	45 (16)	38 (31)	22 (21)	
25-34 years	4 years 98 (60)		45 (59)	
35-49 years	41 (25)	15 (15)	13 (20)	
Any underlying medical conditions	<mark>72 (33)</mark>	<mark>32 (33)</mark>	<mark>46 (56)</mark>	
Any pregnancy- associated complications*	37 (20)	25 (31)	23 (29)	

*Pregnancy-associated complications include hypertensive disorders of pregnancy, gestational diabetes, intrauterine growth restrictions, and pre-eclampsia. Note that percentages are weighted to account for sampling scheme.

Interventions and outcomes among pregnant patients with respiratory symptoms and a positive SARS-CoV-2 test

The proportion requiring ICU admission and vasopressor support decreased over time



COVID-19-Associated Hospitalization Surveillance Network: Respiratory Virus Hospitalization Surveillance letwork (RESP-NET) Platform

	January – November 2021 (pre-Omicron) N (%)*	December 2021 - June 2022 (early Omicron) N (%)*	July 2022 – April 2023 (later Omicron) N (%) *
Total Number	184	99	80
Interventions			
High flow nasal cannula	<mark>20 (12)</mark>	<mark>2 (2)</mark>	<mark>0 (0)</mark>
BIPAP/CPAP	5 (2)	1 (1)	1 (3)
Mechanical ventilation	15 (7)	3 (2)	1 (2)
Vasopressor	<mark>26 (15)</mark>	<mark>3 (2)</mark>	<mark>3 (6)</mark>
Dialysis or RRT	1 (0.5)	1 (1)	0 (0)
Severe outcomes			
ICU admission	<mark>31 (17)</mark>	<mark>7 (6)</mark>	<mark>2 (4)</mark>
In-hospital death	0 (0)	1 (1)	0 (0)

*Note that percentages are weighted to account for sampling scheme.

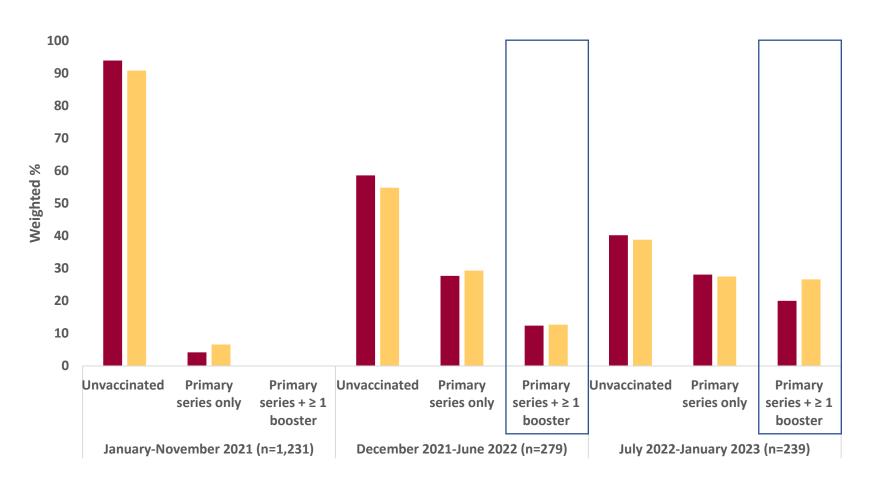
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Vaccination status among pregnant patients hospitalized with laboratory-confirmed SARS-CoV-2 infection by symptom status, COVID-NET, January 2021–April 2023

More asymptomatic pregnant patients received boosters compared with symptomatic patients

Most pregnant patients had not received booster doses

COVID-NET COVID-19-Associated Hospitalization Surveillance Network: A Respiratory Virus Hospitalization Surveillance Network (RESP-NET) Platform

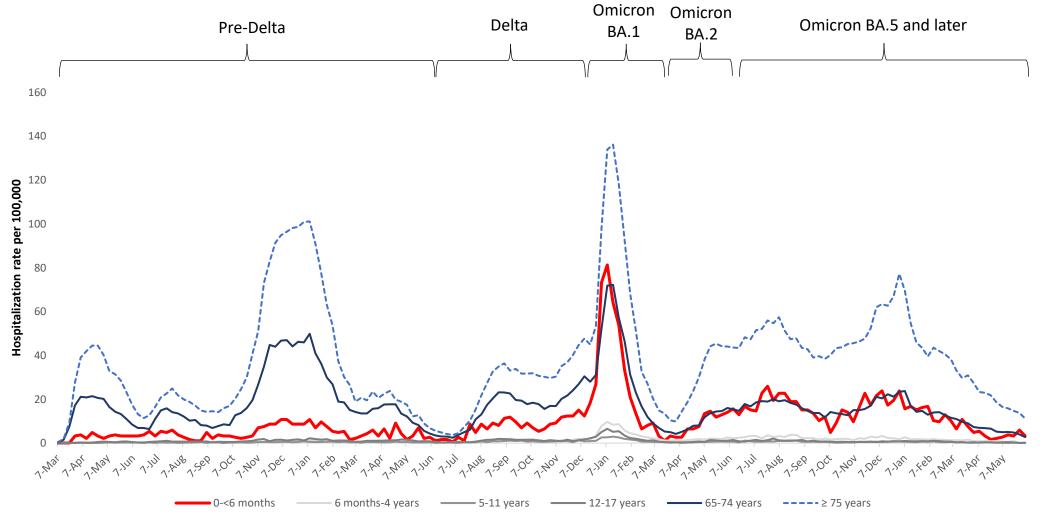


COVID-19-associated hospitalizations in infants <6 months

COVID-NET: March 2020 – May 2023

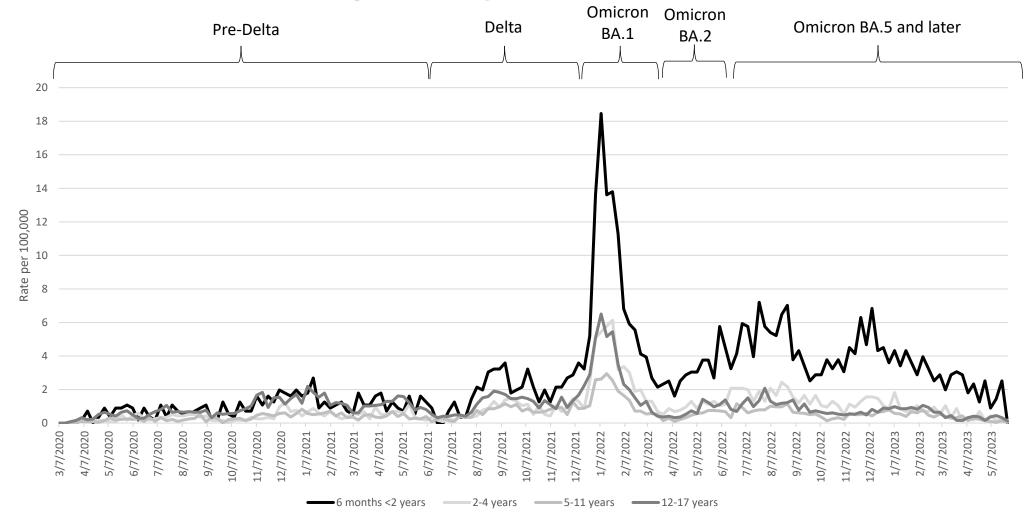


Infants <6 months old had similar COVID-19–associated hospitalization rates to adults aged 65–74 years old



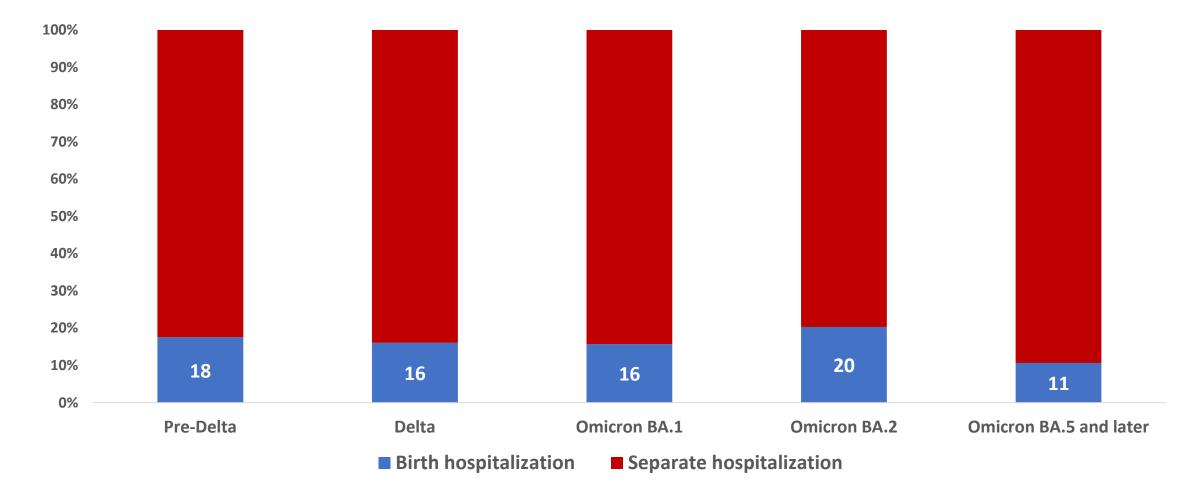
Source: COVID-NET: https://www.cdc.gov/coronavirus/2019-ncov/covid-data/covid-net/purpose-methods.html. Data March 1, 2020 through March 31, 2023. Pre-Delta: March 1, 2020 – 11 June 19, 2021; Delta: June 20–December 18, 2021; Omicron BA.1: December 19, 2021–March 19, 2022; Omicron BA.2: March 20–June 18, 2022; Omicron BA.5 (June 19, 2022–June 3, 2023)

Hospitalization rates in infants, children and adolescents aged 6 months through <18 years



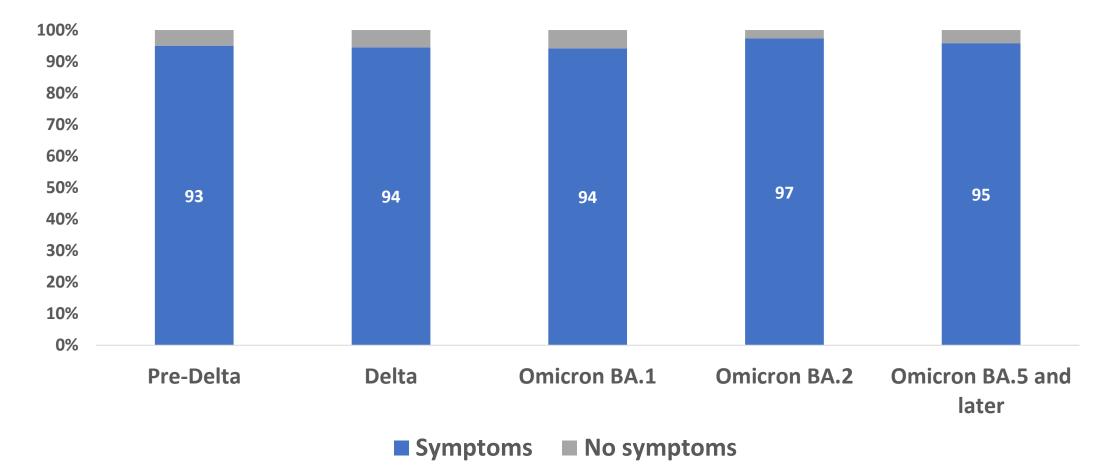
Source: COVID-NET: https://www.cdc.gov/coronavirus/2019-ncov/covid-data/covid-net/purpose-methods.html. Data March 1, 2020 through March 31, 2023. Pre-Delta: March 1, 2020 – June 19, 2021; Delta: June 20–December 18, 2021; Omicron BA.1: December 19, 2021–March 19, 2022; Omicron BA.2: March 20–June 18, 2022; Omicron BA.5 (June 19, 2022–May 27, 2023)

On average, 15% of hospitalized infants <6 months with COVID-19 were identified during their birth hospitalization*



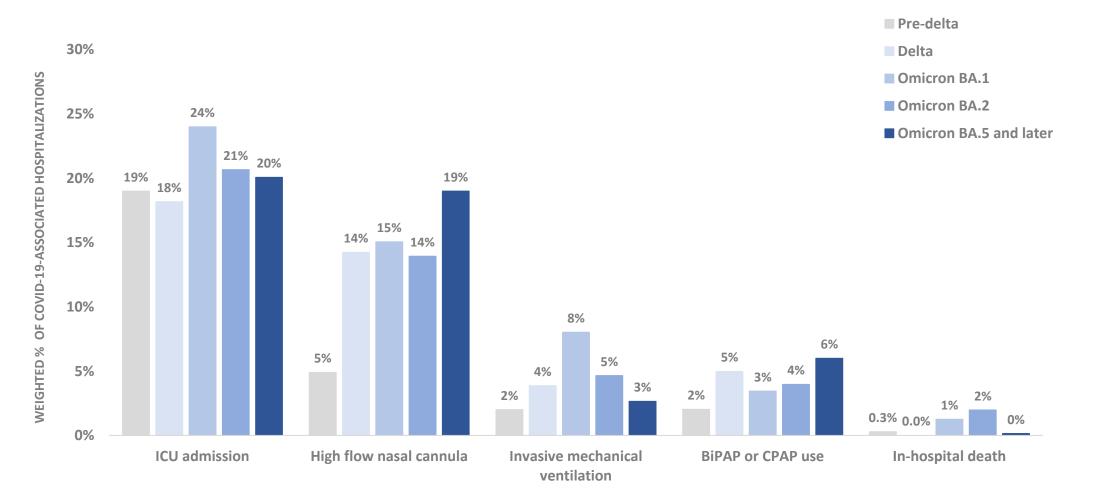
*Birth hospitalization was defined as admission date within 1 day of birth. Source: COVID-NET: https://www.cdc.gov/coronavirus/2019-ncov/covid-data/covid-net/purpose-methods.html. Data March 1, 2020 through March 31, 2023. Pre-Delta: March 1, 2020 – June 19, 2021; Delta: June 20–December 18, 2021; Omicron BA.1: December 19, 2021–March 19, 2022; Omicron BA.2: March 20–June 18, 2022; Omicron BA.5 (June 19, 2022–March 31, 2023).

95% of infants <6 months old with a separate hospitalization had COVID-19 symptoms during the Omicron BA.5 period



Source: COVID-NET: https://www.cdc.gov/coronavirus/2019-ncov/covid-data/covid-net/purpose-methods.html. Data March 1, 2020 through March 31, 2023. Pre-Delta: March 1, 2020 – June 19, 2021; Delta: June 20–December 18, 2021; Omicron BA.1: December 19, 2021–March 19, 2022; Omicron BA.2: March 20–June 18, 2022; Omicron BA.5 (June 19, 2022–March 31, 2023)

1 in 5 infants < 6 months old with COVID-19 were admitted to the ICU (excluding birth hospitalizations)



Source: COVID-NET: https://www.cdc.gov/coronavirus/2019-ncov/covid-data/covid-net/purpose-methods.html. Data March 1, 2020 through March 31, 2023. Pre-Delta: March 1, 2020 – June 19, 2021; Delta: June 20–December 18, 2021; Omicron BA.1: December 19, 2021–March 19, 2022; Omicron BA.2: March 20–June 18, 2022; Omicron BA.5 (June 19, 2022–March 31, 2023)

Trends in COVID-19-associated hospitalizations – COVID-NET, March 2020 – May 2023

- General: Hospitalization rates decreased in all age groups
 - The age distribution of persons hospitalized with COVID-19 has shifted such that the highest rates are in adults aged ≥75 years followed by those 65-74 years and infants ages <6 months

• Pregnant persons:

- Most pregnant persons hospitalized with a positive SARS-CoV-2 test had no symptoms recorded at admission and were likely identified through screening on admission
- Among those with respiratory symptoms, the proportion with underlying medical conditions has increased and the proportion with severe outcomes has decreased
- Most hospitalized pregnant persons with a positive SARS-CoV-2 test, regardless of symptoms or reason for testing, were not up to date with vaccinations
- Infants <6 months: Hospitalization rates increased in the Omicron period
 - Most hospitalized with COVID-19-like symptoms
 - Excluding birth hospitalizations, 20% admitted to the ICU since June 2022



Acknowledgments

Coronaviruses and Other Respiratory Viruses Division (CORVD): RESP-NET Team (COVID-NET/RSV-NET):

- Michael Whitaker
- Kadam Patel
- Christopher Taylor
- Huong Pham
- Onika Anglin
- Jenny Milucky
- Bhoomija Chatwani
- Michael Melgar
- Monica Patton

Many others in CORVD....



- State, Local, and Territorial health Department partners
- RESP-NET partners

Thank you.

Centers for Disease Control and Prevention

COVID-19 in pregnant people

Romeo Galang, MD MPH

Emergency Preparedness and Response Team

Field Support Branch

Division of Reproductive Health

National Center for Chronic Disease Prevention and Health Promotion



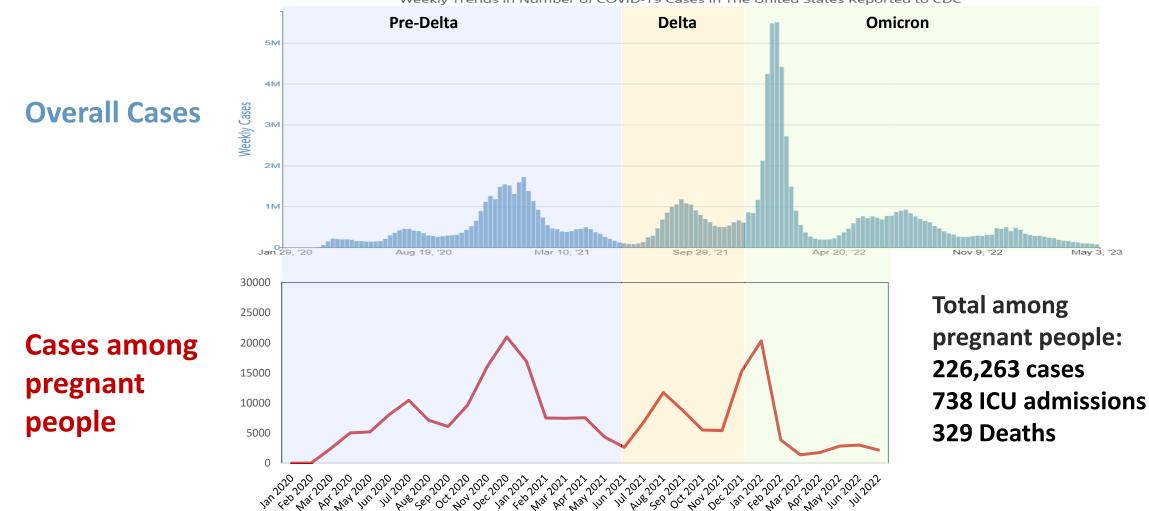


Disease burden and risks to maternal and infant health

COVID-19 in Pregnant People

- COVID-19 during pregnancy is associated with more severe maternal health outcomes
- COVID-19 during pregnancy is associated with adverse pregnancy outcomes (e.g., preterm birth, stillbirth)
- Adverse maternal, fetal, and infant outcomes differed according to the circulating variant

Reported COVID-19 cases overall and among pregnant people in the US (National COVID-19 Case Surveillance Data; Jan 22, 2020–May 3, 2023)



Weekly Trends in Number of COVID-19 Cases in The United States Reported to CDC

<u>Among people with COVID-19</u>, pregnancy increased the risk for ICU admission and invasive ventilation

(Living Systematic Review with data from 1 Dec 2019 - 27 Apr 2021)

		# with event/# i	Odds ratio (95% CI)		
Outcomes	# Studies	Pregnant with COVID -19 COVID-19			
All cause mortality	11	242/122 222 (0.2)	5252/2 138 726 (0.2)	1.48 (0.62 to 3.49)	
ICU admission	10	912/118 403 (0.8)	11 513/1 908 957 (0.6)	2.61 (1.84 to 3.71)	
Invasive ventilation 8		310/116 458 (0.3)	3607/1 772 716 (0.2)	2.41 (2.13 to 2.71)	
ECMO	5	19/30 694 (0.1)	122/432 623 (0.0)	3.71 (0.71 to 19.41)	
ARDS	4	22/197 (11.2) 45/418 (10.8)		1.19 (0.24 to 5.95)	
Major organ failure	4	5/197 (2.5)	28/418 (6.7)	0.39 (0.15 to 1.04)	

ECMO: extracorporeal membrane oxygenation; ARDS: Acute respiratory distress syndrome

Allotey, J et al. Clinical manifestations, risk factors, and maternal and perinatal outcomes of coronavirus disease 2019 in pregnancy: living systematic review and metaanalysis. *BMJ*. 2020. Final version 7-May 2022 <u>https://doi.org/10.1136/bmj.m3320</u>

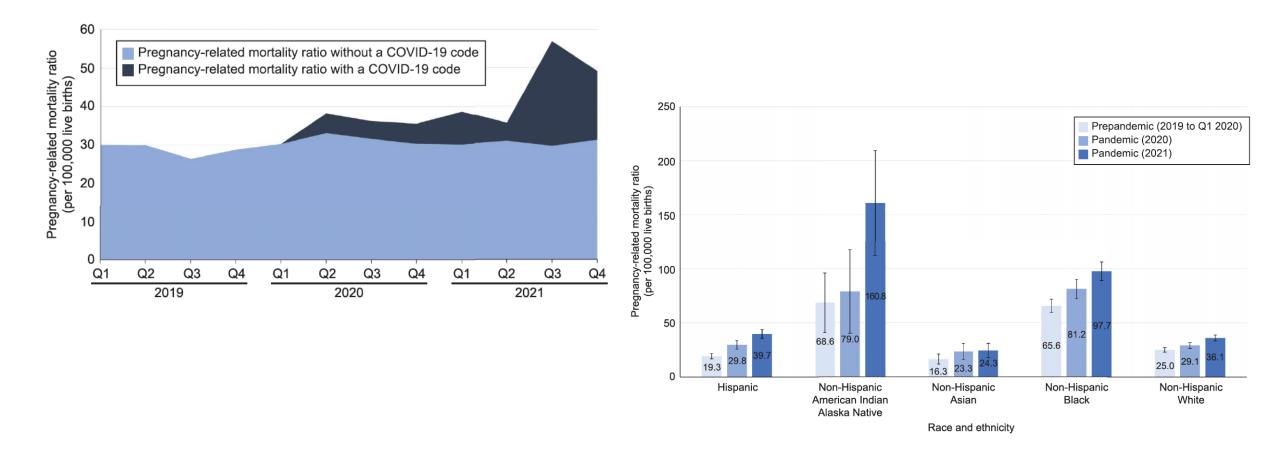
<u>Among pregnant people</u>, COVID-19 increased the risk for adverse maternal, fetal, and infant outcomes

(Living Systematic Review with data from 1 Dec 2019 - 27 Apr 2021)

Outcomes		# with event/# i			
	# Studies	Pregnant with COVID-19	Pregnant without COVID-19	Odds ratio (95% CI)	
Maternal outcomes:					
All cause mortality	21 47/11 362		37/411 126 (0.0)	6.09 (1.82 to 20.38)	
ICU admission	21	447/12 957 (3.4)	1962/459 359 (0.4)	5.41 (3.59 to 8.14)	
Preterm birth <37 weeks	48	1306/12 076 (10.8)	26 068/436 964 (6.0)	1.57 (1.36 to 1.81)	
Perinatal outcomes:					
Stillbirth	25	76/9338 (0.8)	1397/414 139 (0.3)	1.81 (1.38 to 2.37)	
Neonatal death	21	16/3153 (0.5)	28/9 263 (0.3)	2.35 (1.16 to 4.76)	
Admission to neonatal unit	29	687/4072 (16.9)	6968/193 124 (3.6)	2.18 (1.46 to 3.26)	
Fetal distress	6	131/1073 (12.2)	246/3933 (6.3)	2.22 (1.45 to 3.41)	

Allotey, J et al. Clinical manifestations, risk factors, and maternal and perinatal outcomes of coronavirus disease 2019 in pregnancy: living systematic review and metaanalysis. *BMJ*. 2020. Final version 7-May 2022 <u>https://doi.org/10.1136/bmj.m3320</u>

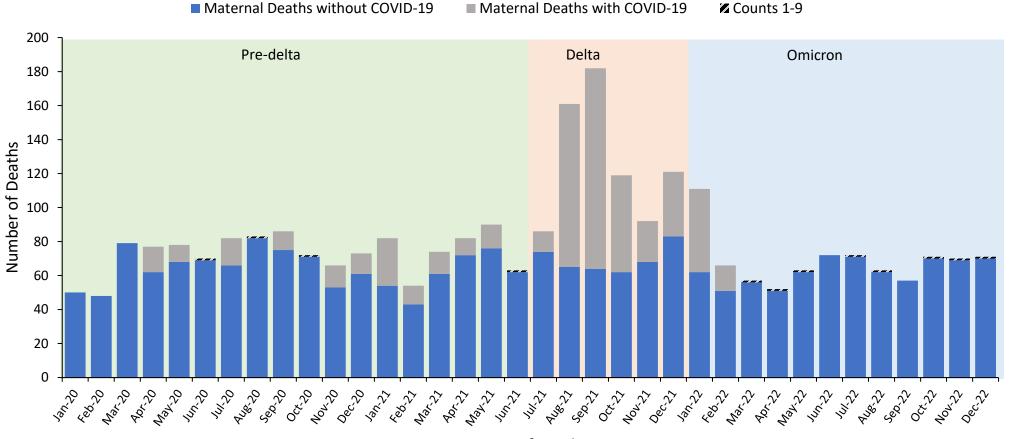
Pregnancy-related mortality increased rapidly in 2021 (pre-Omicron), consistent with rising rates of COVID-19 associated mortality



Thoma, M. E., & Declercq, E. R. (2023). Changes in Pregnancy-Related Mortality Associated With the Coronavirus Disease 2019 (COVID-19) Pandemic in the United States. *Obstetrics and gynecology*, 141(5), 911–917. <u>https://doi.org/10.1097/AOG.000000000005182</u>

Have risks evolved by COVID-19 variant?

Figure 1. Maternal deaths and maternal deaths mentioning COVID-19, by count, US 2020-2022

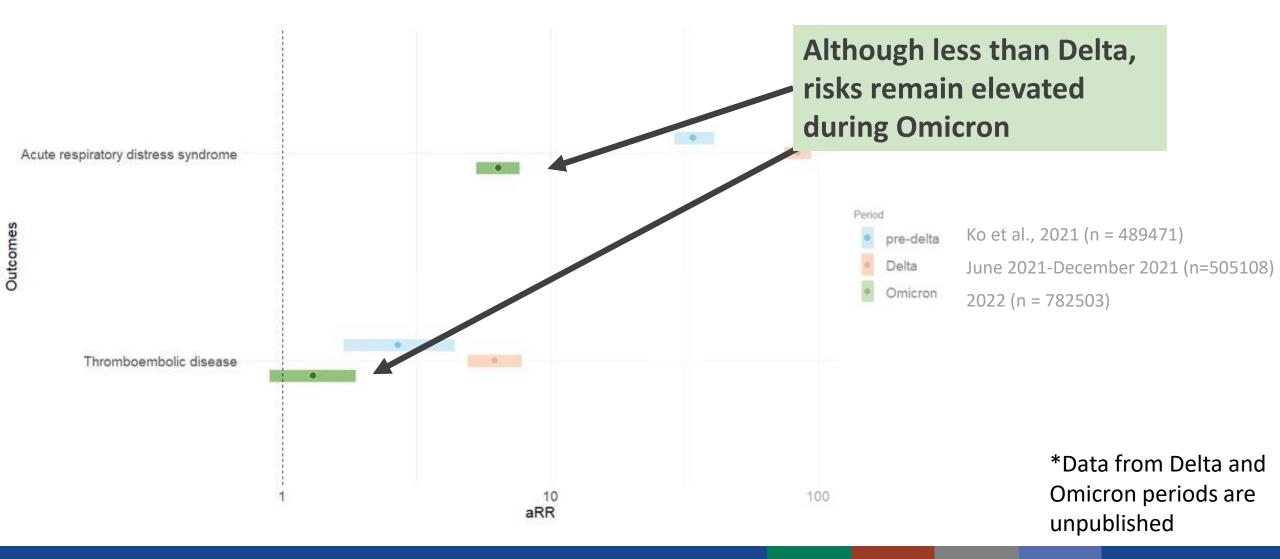


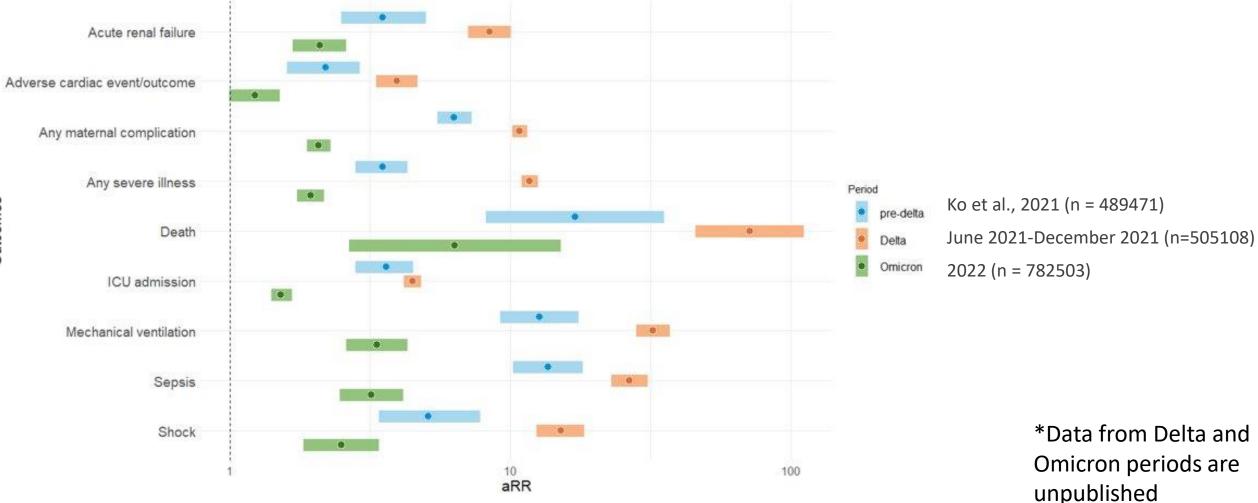
Date of Death

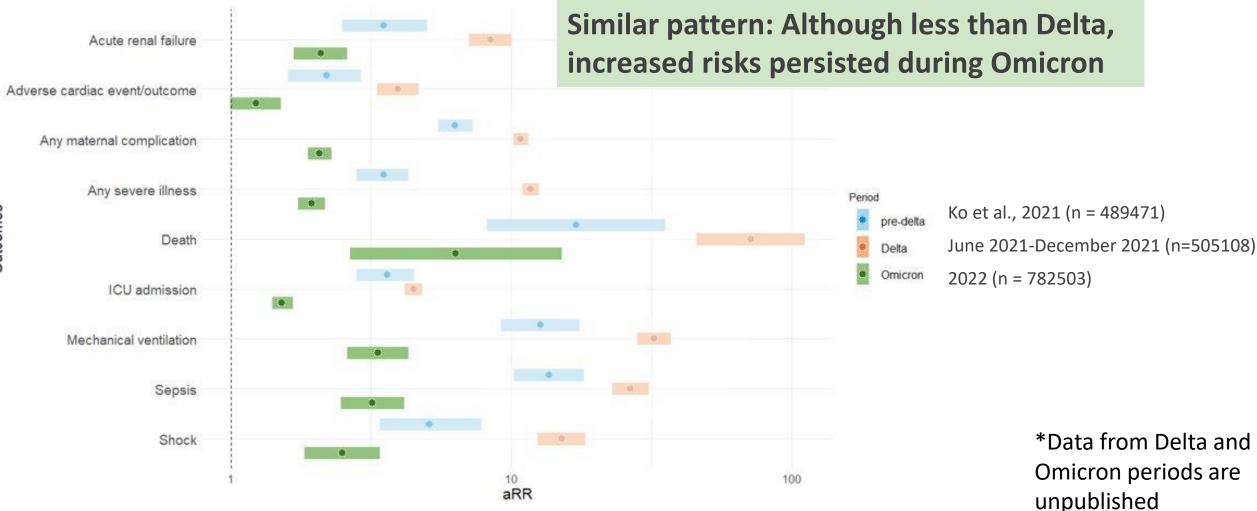
Note: Data for 2020-2021 are final and data for 2022-2023. Death counts between 1-9 are suppressed in accordance with NCHS confidentiality standards. Weeks with COVID-19 death counts 1-9 are noted on the figure with black and grey diagonal lines.

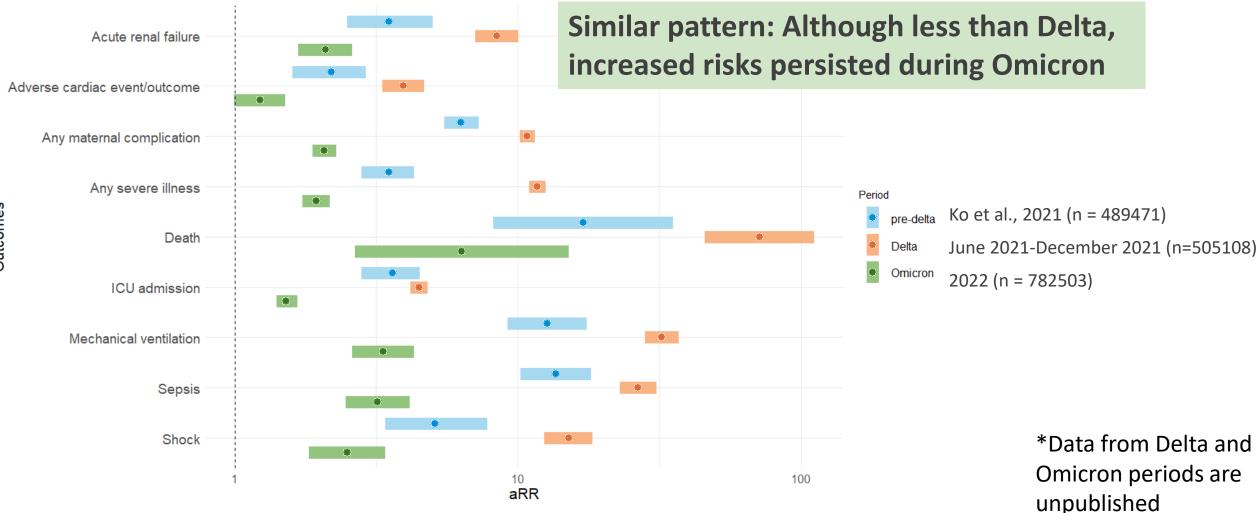
Source: CDC, National Center for Health Statistics. National Vital Statistics System, Provisional Mortality on CDC WONDER Online Database. Accessed at http://wonder.cdc.gov/mcd-icd10-provisional.html







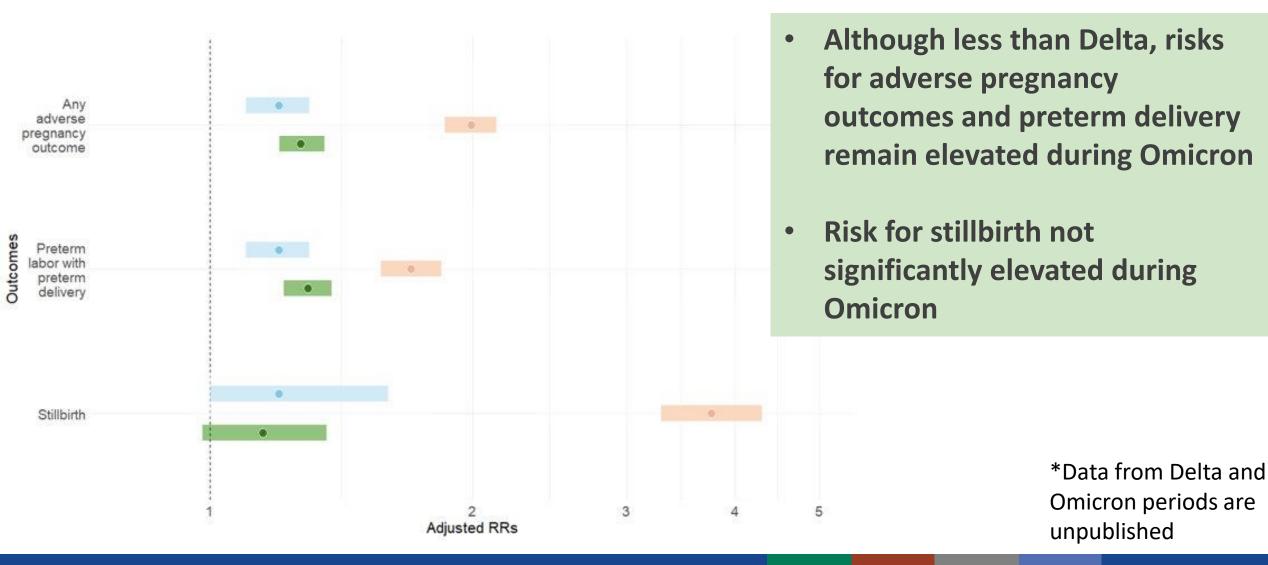




Adverse Pregnancy Outcomes Associated With a Documented COVID-19 Diagnosis at Delivery Hospitalization—United States, Pre-delta, Delta, and Omicron Periods, Premier



Adverse Pregnancy Outcomes Associated With a Documented COVID-19 Diagnosis at Delivery Hospitalization—United States, Pre-delta, Delta, and Omicron Periods, Premier



Trends (%) in pregnancy outcomes by maternal COVID-19 status: 14 states and the District of Columbia (July 2020-December 2022)

	pre-Delta July 2020-June 2021		Delta July 2021 - December 2021		Omicron January 2022 - December 2022	
	COVID-19 ¹	No COVID-19	COVID-19 ¹	No COVID-19	COVID-19 ¹	No COVID-19
ICU admission	0.6	0.1	1.2	0.2	0.2	0.2
NICU admission	10.3	8.7	10.9	8.5	9.0	8.7
Total preterm ²	11.9	9.9	13.0	10.1	10.3	10.1
Early preterm	3.2	2.6	3.8	2.7	2.4	2.7
Late preterm	8.8	7.2	9.2	7.4	7.9	7.4
Total low birthweight ³	8.8	7.9	9.8	8.1	7.9	8.2
Very low birthweight ⁴	1.5	1.3	1.6	1.3	1.1	1.3

¹ Confirmed or presumed COVID-19 during pregnancy. Confirmed cases only are included for California, Maryland, Ohio, North Dakota, and Tennessee.

² Gestational age in completed weeks; based on the obstetric estimate of gestation.

³ Less than 2,500 grams

⁴ Less than 1,500 grams

NOTES: Reporting area includes Alabama, Alaska, Arkansas, California, District of Columbia, Idaho, Maine, Maryland, New Hampshire, North Dakota, Ohio, Oklahoma, Oregon, Tennessee, and West Virginia. District of Columbia did not report for October-December 2022.

SOURCE: National Center for Health Statistics, National Vital Statistics System, Natality.

https://www.cdc.gov/nchs/data/health_policy/trends-in-outcomes.pdf

Summary

- Incidence of COVID-19 among pregnant people mirrors that of the general population
- Pregnancy remains a risk factor for severe maternal disease and adverse pregnancy outcomes, even with new variants
- Some maternal, fetal, and infant risks were lower with Omicron; cannot disentangle the impact of prior infection/vaccination

Conclusion

COVID-19 vaccination improves outcomes for pregnant people, their pregnancies and their infants; therefore vaccination should continue to be recommended for maternal and fetal benefit

Acknowledgements

National Center on Birth Defects and Developmental Disorders Jeffrey Carlson Amanda Cohn Dana Meaney-Delman Suzanne Gilboa Kara Polen Emily Reeves Van Tong Kate Woodworth

For more information, contact CDC 1-800-CDC-INFO (232-4636) TTY: 1-888-232-6348 www.cdc.gov National Center for Immunization and Respiratory Diseases Sascha Ellington Katherine Fleming-Dutra Sara Oliver Regina Simeone

National Center for Health Statistics Donna Hoyert Joyce Martin Michelle Osterman Claudia Valenzuela

The findings and conclusions in this report are those of the authors and do not necessarily represent the official position of the Centers for Disease Control and Prevention.



Centers for Disease Control and Prevention National Center for Immunization and Respiratory Diseases



COVID-19 vaccine effectiveness updates

23 June 2023

Ruth Link-Gelles, PhD, MPH LCDR, US Public Health Service COVID-19 Vaccine Effectiveness Program Lead Centers for Disease Control and Prevention

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U.S. COVID-19 Vaccination Coverage (%) of Total Population by Age Group — May 10, 2023

Coverage / Age (years)	<2	2-4	5-11	12-17	18-24	24-49	50-64	<u>></u> 65
At least one dose ⁺	8.9	10.9	40.0	72.2	82.3	85.5	95.0	95.0
At least one bivalent dose	0.6	0.6	4.8	7.8	7.4	12.1	21.7	43.3
Unvaccinated	91.1	89.1	60.0	27.8	17.7	14.5	†	+

Organization of vaccine effectiveness (VE) data

- Bivalent VE, by outcome and Omicron subvariant in adults
- VE in special populations:
 - *Monovalent* and *bivalent* VE in pregnant people
 - *Bivalent* people with immunocompromising conditions

Monovalent and bivalent VE, against hospitalization and critical illness by Omicron subvariant in adults ≥18 years, VISION Network

VISION Multi-State Network of Electronic Health Records



- Cases: COVID-like illness (CLI) with positive PCR for SARS-CoV-2 within 14 days before or 72 hours after the admission or encounter
- **Controls**: CLI with negative PCR for SARS-CoV-2

- Variant periods designated for analysis based on time when novel sublineage became predominant (>50%) at study site
- VE adjusted for age, sex, race and ethnicity, geographic region, and calendar time
- Vaccination documented by electronic health records and state and city registries

VISION: Absolute VE of monovalent and bivalent booster doses against hospitalization and critical illness among immunocompetent adults aged ≥18 years – September 2022 – May 2023

mRNA Dosage Pattern	Total tests	SARS-CoV-2- test-positive, N (%)	Median interval since last dose, days (IQR)	Adjusted VE (95% CI)	
Hospitalization					
Unvaccinated (ref)	16,219	1,835 (11)		Ref	
Monovalent doses only	38,843	4,086 (11)	381 (275-513)	21 (16-26)	⊢ ••
Bivalent booster, 7-59 days earlier	4,894	329 (7)	35 (21-47)	62 (57-67)	H0+
Bivalent booster, 60-119 days earlier	5,283	491 (9)	87 (73-103)	47 (41-53)	H H H
Bivalent booster, 120-179 days earlier	3,756	346 (9)	146 (132-161)	24 (12-33)	
Critical illness					
Unvaccinated (ref)	14,762	378 (3)		Ref	
Monovalent doses only	35,415	658 (2)	380 (275-514)	31 (21-40)	
Bivalent booster, 7-59 days earlier	4,614	49 (1)	34 (21-47)	69 (58-77)	⊢ ●-1
Bivalent booster, 60-119 days earlier	4,880	88 (2)	87 (73-103)	45 (29-58)	
Bivalent booster, 120-179 days earlier	3,445	35 (1)	146 (132-161)	52 (30-67)	—
					-20 0 20 40 60 80 100

Vaccine Effectiveness (%)

Critical illness defined as admission to intensive care unit or death; case-patients were persons admitted to ICU or who experienced death associated with COVID-19, and control patients were persons hospitalized without COVID-19. VE estimates adjusted for age, sex, race and ethnicity, geographic region, and calendar time. Updated from: Link-Gelles et al., MMWR, https://www.cdc.gov/mmwr/volumes/72/wr/mm7221a3.htm

VISION: Absolute VE of monovalent and bivalent booster doses against hospitalization and critical illness among immunocompetent adults aged ≥18 years, during BA.4/5 predominance – September 2022 – January 2023

mRNA Dosage Pattern	Total tests	SARS-CoV-2- test-positive, N (%)	Median interval since last dose, days (IQR)	Adjusted VE (95% CI)	
Hospitalization					
Unvaccinated (ref)	11,240	1,426 (13)		Ref	
Monovalent doses only	27,564	3,106 (11)	349 (238-460)	25 (19-30)	H - H
Bivalent booster, 7-89 days earlier	6,723	524 (8)	47 (28-67)	61 (56-65)	H e t
Bivalent booster, ≥90 days earlier	1,511	163 (11)	105 (96-115)	40 (28-50)	
Critical illness					
Unvaccinated (ref)	10,110	296 (3)		Ref	
Monovalent doses only	24,976	518 (2)	347 (236-460)	33 (21-42)	
Bivalent booster, 7-89 days earlier	6,199	91 (1)	47 (28-66)	61 (50-70)	⊢ ∎-1
Bivalent booster, ≥90 days earlier	1,348	25 (2)	105 (96-115)	49 (21-67)	

CDC unpublished data. VE estimates adjusted for age, sex, race and ethnicity, geographic region, and calendar time. Variant predominance based on regional circulation: <u>https://covid.cdc.gov/covid-data-tracker/#variant-proportions</u>

Vaccine Effectiveness (%)

40

60

20

-20

100

VISION: Absolute VE of monovalent and bivalent booster doses against hospitalization and critical illness among immunocompetent adults aged ≥18 years, during XBB predominance – January – May 2023

mRNA Dosage Pattern	Total tests	SARS-CoV-2- test-positive, N (%)	Median interval since last dose, days (IQR)	Adjusted VE (95% Cl)	
Hospitalization					
Unvaccinated (ref)	4,979	409 (8)		Ref	
Monovalent doses only	11,279	980 (9)	469 (375-605)	9 (-4 to 20)	
Bivalent booster, 7-89 days earlier	1,045	60 (6)	65 (43-79)	51 (35 to 63)	
Bivalent booster, 90-179 days earlier	4,654	419 (9)	139 (119-157)	20 (7 to 32)	
Critical illness					
Unvaccinated (ref)	4,652	82 (2)		Ref	
Monovalent doses only	10,439	140 (1)	469 (375-602)	28 (3 to 46)	
Bivalent booster, 7-89 days earlier	994	9 (1)	65 (43-78)	58 (15 to 79)*	
Bivalent booster, 90-179 days earlier	4282	47 (1)	139 (119-157)	48 (23 to 65)	

40 Vaccine Effectiveness (%)

60

20

-20

0

CDC unpublished data. VE estimates adjusted for age, sex, race and ethnicity, geographic region, and calendar time.

* These interim estimates are imprecise, which might be because of a relatively small number of persons in each level of vaccination or case status. This imprecision indicates the actual VE may be substantially different from the point estimate shown, and estimates should therefore be interpreted with caution. Additional data accrual should increase precision and allow appropriate interpretation.

Variant predominance based on regional circulation: https://covid.cdc.gov/covid-data-tracker/#variant-proportions

80

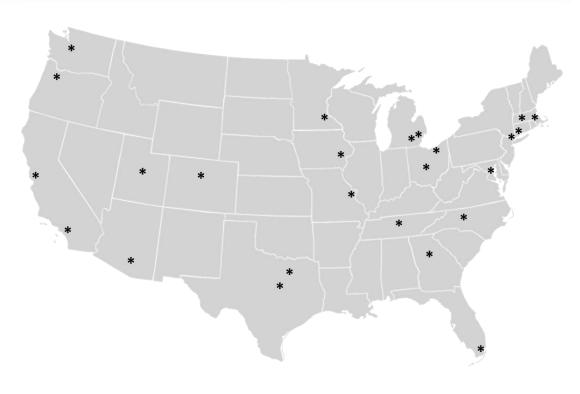
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Monovalent and bivalent VE against hospitalization among adults aged 218 years, IVY Network

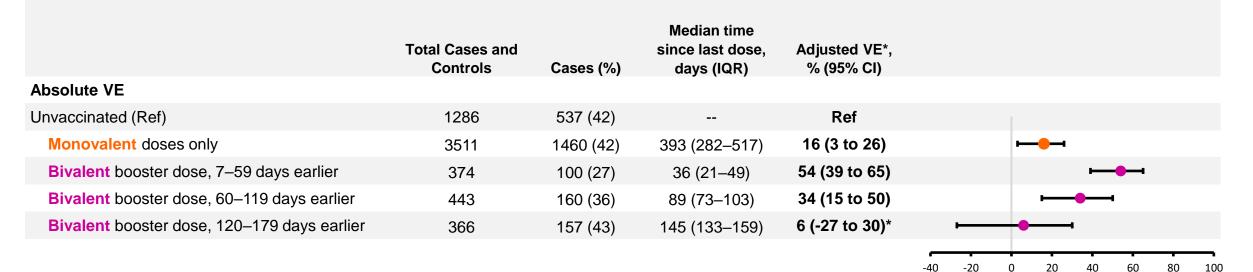
IVY Network — 25 hospitals, 20 U.S. States

- Design: Prospective, case-control
- Population: Adults aged ≥18 years hospitalized with Acute respiratory illness (ARI)*
 - Cases: ARI and test *positive* for SARS-CoV-2 by NAAT or antigen test within 10 days of illness
 - Controls: ARI and test *negative* for SARS-CoV-2 and influenza by NAAT within 10 days of illness
- Vaccination data: Electronic medical records (EMR), state and city registries, and self-report
- Specimens: Upper respiratory specimens obtained for central RT-qPCR testing and sequencing

I V Y INVESTIGATING RESPIRATORY VIRUSES IN THE ACUTELY ILL



IVY Network: *Absolute* VE against COVID-19 *hospitalization* among immuno*competent* adults *aged* ≥18 years —September 8, 2022 – May 29, 2023



Vaccine Effectiveness (%)

* These interim estimates are imprecise, which might be because of a relatively small number of persons in each level of vaccination or case status. This imprecision indicates the actual VE may be substantially different from the point estimate shown, and estimates should therefore be interpreted with caution. Additional data accrual should increase precision and allow appropriate interpretation. VE adjustments: Age, sex, race, ethnicity, admission date (biweekly), and HHS region

IVY Network: *Absolute* VE against COVID-19 *hospitalization* among immuno*competent* adults aged ≥18 years by lineage period — September 8, 2022 – May 24, 2023

	Total Cases and Controls	Cases (%)	Median time since last dose, days (IQR)	Adjusted VE*, % (95% Cl)		
BA.4/5 (September 8 – November 13, 2022)						
Unvaccinated (Ref)	313	138 (44)		Ref		
Monovalent doses only	1003	398 (40)	304 (188–386)	30 (8–47)		• •
Bivalent booster dose, 7–59 days earlier	83	26 (31)	25 (13–40)	59 (21–78)*		••
BQ.1 (November 14, 2022 – January 22, 2023)						
Unvaccinated (Ref)	458	190 (41)		Ref		
Monovalent doses only	1262	504 (40)	386 (297–518)	17 (-5 to 34)	-	
Bivalent booster dose, 7–59 days earlier	226	52 (23)	40 (25–52)	63 (44–75)		•
Bivalent booster dose, 60–119 days earlier	225	68 (30)	83 (69–95)	49 (24–66)		• • •••
XBB (January 23 – May 24, 2023)						
Unvaccinated (Ref)	514	209 (41)		Ref		
Monovalent doses only	1246	558 (45)	464 (378–590)	-8 (-34 to 13)		
Bivalent booster dose, 7-89 days earlier	155	56 (36)	64 (46–78)	29 (-8 to 53)*		
Bivalent booster dose, 90–179 days earlier	478	208 (44)	137 (118–154)	-8 (-44 to 19)*	·•	

* These interim estimates are imprecise, which might be because of a relatively small number of persons in each level of vaccination or case status. This imprecision indicates the actual VE may be substantially different from the point estimate shown, and estimates should therefore be interpreted with caution. Additional data accrual should increase precision and allow appropriate interpretation. VE adjustments: Age, sex, race, ethnicity, admission date (biweekly), and HHS region

-35 -15 5 25 45 65 85 Vaccine Effectiveness (%)

-55

VE in special populations: pregnant people

VISION Multi-State Network of Electronic Health Records



- Cases: COVID-like illness (CLI) with positive PCR for SARS-CoV-2 within 14 days before or 72 hours after the encounter
- **Controls**: CLI with negative PCR for SARS-CoV-2

- Among pregnant people 18-45 years at time of emergency department/urgent care encounter
- VE adjusted for age, ethnicity, race, underlying medical conditions, gestational age at encounter, site, Medicaid status, day of encounter, site facility urbanicity
- Vaccination documented by electronic health records and state and city registries
- Separate results for COVID-19 vaccine monovalent doses received prior to pregnancy and bivalent doses received during pregnancy due to timing of bivalent authorization/analysis (Sept 2022-May 2023)

VISION: *Absolute* VE of COVID-19 *monovalent* doses received *prior to* pregnancy against *ED/UC encounters* among immuno*competent* pregnant persons aged 18-45 years – June 2022 – May 2023*

Vaccine Dosage Pattern	Total tests	SARS-CoV-2- test-positive, N (%)	Median interval since last dose, days (IQR)	Adjusted VE (95% Cl)	
Absolute VE					
Unvaccinated (ref)	2238	317 (14)		Ref	
Monovalent received:					
<6 months before pregnancy	833	108 (13)	270 (216, 326)	27 (6, 44)	▶ →
≥6 months before pregnancy	1986	264 (13)	454 (375, 544)	5 (-15, 22)	0 20 40 60 80 100

Vaccine Effectiveness (%)

Adjusted for: Age, ethnicity, race, underlying medical conditions, gestational age at encounter, site, Medicaid status, day of encounter, site facility urbanicity *Unpublished CDC data.

VISION: *Absolute* VE of COVID-19 *bivalent* doses received *during* pregnancy against *ED/UC encounters* among immuno*competent* pregnant persons aged 18-45 years – September 2022 – May 2023*

Vaccine Dosage Pattern	Total tests	SARS-CoV-2- test-positive, N (%)	Median interval since last dose, days (IQR)	Adjusted VE (95% CI)	
Absolute VE					
Unvaccinated (ref)	1701	196 (12)		Ref	
Bivalent dose**	191	10 (5)	56 (29, 97)	61 (22, 81)***	0 20 40 60 80 100 Vaccine Effectiveness (%)

Adjusted for: Age, ethnicity, race, underlying medical conditions, gestational age at encounter, site, Medicaid status, day of encounter, site facility urbanicity

*Unpublished CDC data

Doses received **during pregnancy for bivalent group

***These interim estimates are imprecise, which might be because of a relatively small number of persons in each level of vaccination or case status. This imprecision indicates the actual VE may be substantially different from the point estimate shown, and estimates should therefore be interpreted with caution. Additional data accrual should increase precision and allow appropriate interpretation.

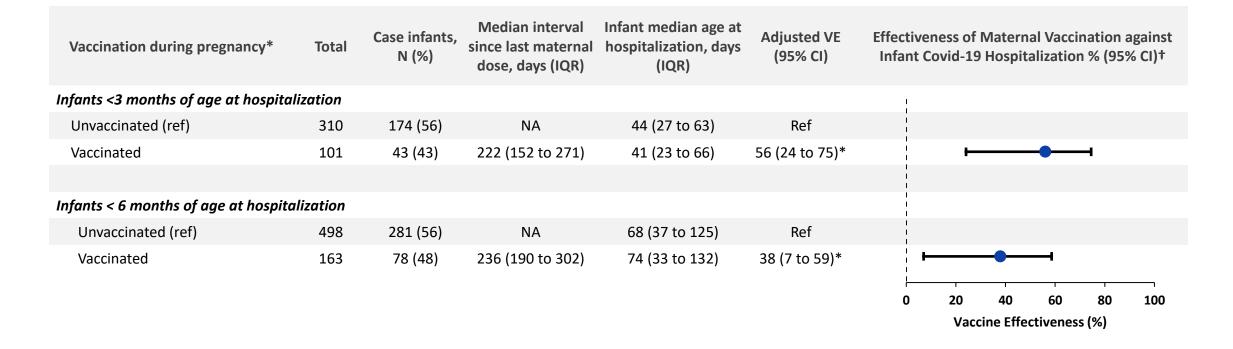
Overcoming COVID-19 network



- Cases infants: hospitalized with COVID-19 as the primary reason for admission and with a positive SARS-CoV-2 RT-PCR or antigen test result
- Control infants: hospitalized with or without COVID-19 symptoms and negative SARS-CoV-2 RT-PCR or antigen test result
 - Matched to case-infants by site; hospitalized within 4 weeks of case-infant admission

- Case-control study to assess effectiveness of maternal vaccination for COVID-19 in infants < 6 months of age
- 25 pediatric hospitals across 20 states
- Infants admitted between March 9, 2022, and May 9, 2023
- Baseline demographic and clinical characteristics obtained via parent interview
- Maternal vaccination status verified using state vaccination registries, electronic medical records, or other sources

Overcoming COVID-19: Effectiveness of maternal vaccination in prevention of hospitalization among infants – March 9, 2022 – May 9, 2023



*Last mRNA or viral vector vaccine dose received between the beginning of pregnancy and 14 days before delivery. 14 people received a bivalent mRNA vaccine. †These estimates are imprecise, which might be because of a relatively small number of persons in each level of vaccination or case status. This imprecision indicates the actual VE may be substantially different from the point estimate shown, and estimates should therefore be interpreted with caution. Additional data accrual should increase precision and allow appropriate interpretation.

CDC unpublished data. VE estimates adjusted for infant age, sex, race and ethnicity, census region, and month and year of hospitalization.

Bivalent VE in special populations: people with immunocompromising conditions

VISION: Absolute VE of monovalent and bivalent booster doses against hospitalization and critical illness among immunocompromised adults aged ≥18 years – September 2022 – May 2023

mRNA Dosage Pattern	Total tests	SARS-CoV-2- test-positive, N (%)	Median interval since last dose, days (IQR)	Adjusted VE (95% CI)	
Hospitalization					
Unvaccinated (ref)	3,240	322 (10)		Ref	
Monovalent doses only	11,623	1,169 (10)	359 (242-481)	3 (-12-16)	
Bivalent booster, 7-59 days earlier	1,627	144 (9)	33 (19-46)	27 (9-41)	
Bivalent booster, 60-119 days earlier	1,862	144 (8)	88 (74-104)	39 (24-51)	
Bivalent booster, 120-179 days earlier	1,448	118 (8)	146 (133-161)	11 (-13-31)	
Critical illness					
Unvaccinated (ref)	3,006	88 (3)		Ref	
Monovalent doses only	10,725	271 (3)	358 (241-481)	16 (-10-35)	
Bivalent booster, 7-59 days earlier	1,515	32 (2)	33 (19-46)	41 (8-62)*	· · · · · · · · · · · · · · · · · · ·
Bivalent booster, 60-119 days earlier	1,755	37 (2)	88 (74-104)	43 (13-62)	
Bivalent booster, 120-179 days earlier	1,348	18 (1)	146 (133-162)	51 (15-72)*	

* These interim estimates are imprecise, which might be because of a relatively small number of persons in each level of vaccination or case status. This imprecision indicates the actual VE may be substantially different from the point estimate shown, and estimates should therefore be interpreted with caution. Additional data accrual should increase precision and allow appropriate interpretation.

Critical illness defined as admission to intensive care unit or death; case-patients were persons admitted to an ICU or who experienced death associated with COVID-19, and control patients were persons hospitalized without COVID-19. VE estimates adjusted for age, sex, race and ethnicity, geographic region, and calendar time. Updated from: Link-Gelles et al., MMWR, https://www.cdc.gov/mmwr/volumes/72/wr/mm7221a3.htm

80

100

20 40 60

Vaccine Effectiveness (%)

-20

0

Summary and conclusions

Limitations of VE against severe disease

- For estimates of *absolute* vaccine effectiveness, if unvaccinated are meaningfully different from vaccinated individuals (e.g., by COVID-19 risk factors), estimates may be biased.
 - For estimates of *relative* vaccine effectiveness, residual protection from prior doses is an important consideration for interpretation.
- Information on prior infection is limited, although we know rates of prior infection in the U.S. population are high and vary by age.
- VE against COVID-19-associated hospitalization may underestimate protection against more severe COVID-19 disease.
- Lack of statistical power to estimate VE for maternal vaccination by timing of doses during pregnancy; could not separate monovalent and bivalent doses for protection against infant hospitalization

Conclusions: updates to VE of *bivalent* **COVID-19 boosters**

- Bivalent boosters are helping provide additional protection against hospitalization, though evidence of waning
- For most people who received *monovalent* doses and are eligible for a *bivalent* booster, more than a year has elapsed since their last monovalent dose. Because of waning, they may have limited remaining protection against hospitalization.
- Effectiveness against the most critical illness (ICU admission and death) more sustained compared to less severe illness
- VE during XBB predominance may wane more quickly against hospitalization compared to early variant predominant periods
- Vaccination during pregnancy provides protection against hospitalization for infants <6 months; protection may be highest in the first 3 months
- CDC will continue ongoing monitoring of VE, including for all outcomes of interest and for all authorized COVID-19 vaccines in the U.S. with a focus on assessing new policy recommendations and VE in populations at higher risk of severe COVID-19

Acknowledgements

CDC COVID-19 Vaccine Effectiveness and VE platforms teams, including: Policy Team

- Amadea Britton
- Allison Ciesla
- Monica Godfrey
- Eric Griggs
- Katherine Fleming-Dutra
- Dani Moulia
- Morgan Najdowski
- Erica Okwuazi

- Sara Oliver
- Josephine Mac
- Amanda Payne
- Lauren Roper
- Laura Steinhardt
- Evelyn Twentyman
- Megan Wallace
- Ryan Wiegand

- Sarah Ball
- Angela Campbell
- Jennifer DeCuir
- Monica Dickerson
- Margaret Dunne
- Kiara Everett
- Shikha Garg
- Victoria Lazariu
- Patrick Mitchell
- Palak Patel

- Caitlin Ray
- Sarah Reese
- Elizabeth Rowley
- Regina Simeone
- Zach Smith
- Diya Surie
- Mark Tenforde
- Zack Weber
- Laura Zambrano

And many more!!!

Questions?

For more information, contact CDC 1-800-CDC-INFO (232-4636) TTY: 1-888-232-6348 www.cdc.gov

The findings and conclusions in this report are those of the authors and do not necessarily represent the official position of the Centers for Disease Control and Prevention.

