



Indiana
Department
of
Health

CLINICIAN UPDATES

GUY CROWDER, M.D., MPHTM
CHIEF MEDICAL OFFICER

9/27/2024

OUR MISSION:

To promote, protect, and improve the health and safety of all Hoosiers.

OUR VISION:

Every Hoosier reaches optimal health regardless of where they live, learn, work, or play.



Conflict of interest

I have no conflicts of interest to disclose

CMEs



CME credits are available for physicians participating in this webinar.

Once you complete the REDCap survey (link will be added to the chat during the Clinician Update), the IDOH enters your name into the Accreditation Council for Continuing Medical Education (ACCME) Program and Activity Reporting System (PARS). PARS is your entry point into the digitized world of CME.

To access the CME credit from this webinar, please go to [PARS - ACCME](#) (This will allow you to monitor CMEs awarded and entered into ACCME's PARS) and/or [Homepage \(cmepassport.org\)](http://cmepassport.org) (This will allow you to monitor CME credits and find other available opportunities to gain CMEs.)



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BOTULISM RESPONSE FOR HEALTHCARE PROVIDERS

NICOLE STONE, MPH
INFECTIOUS DISEASE EPIDEMIOLOGY
& PREVENTION DIVISION

9/27/2024

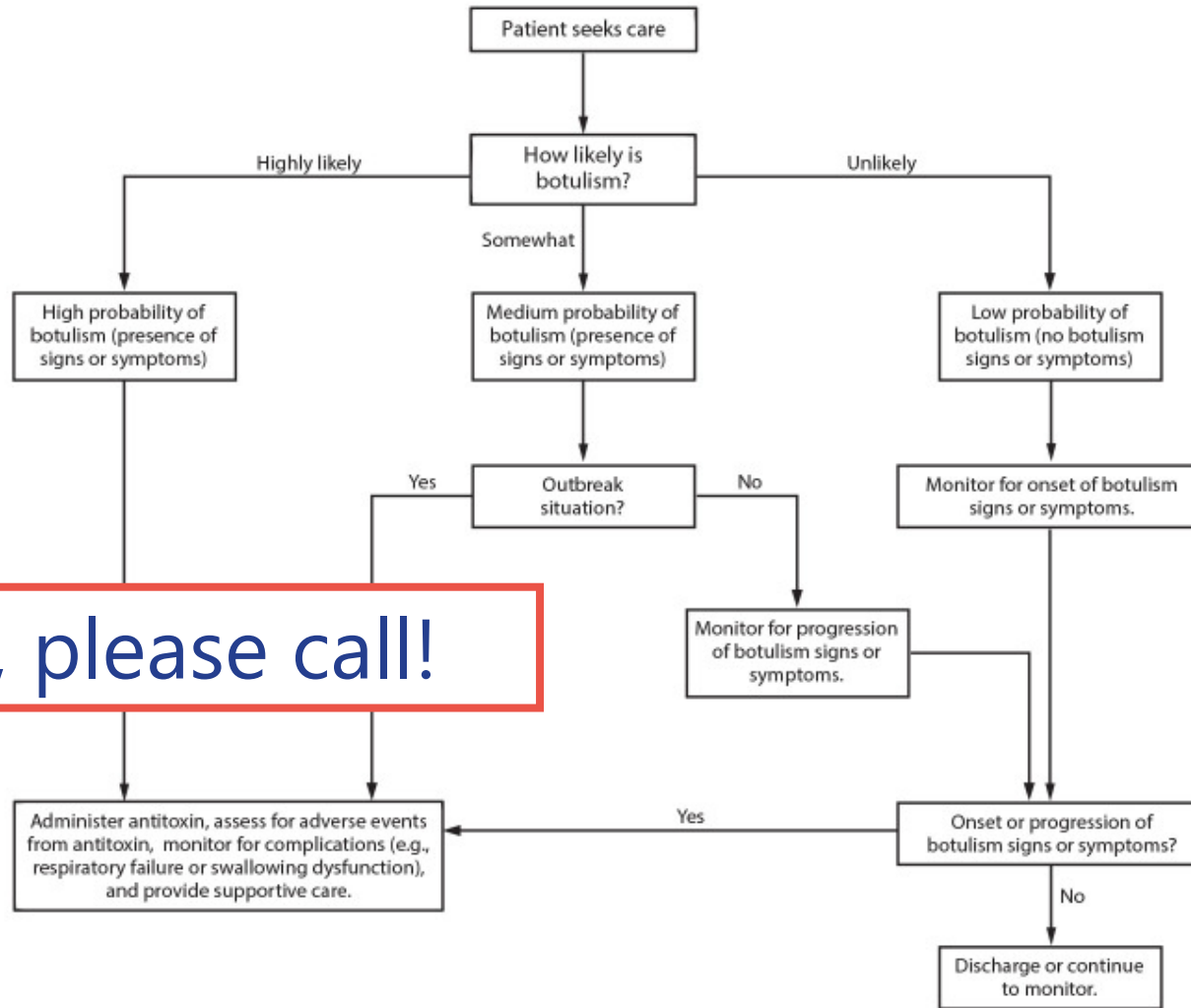
Reporting Suspect Botulism

- Botulism is **immediately reportable upon suspicion** per 410 IAC 1-2.5-75 & 76
 - First reports SHOULD NOT be through NBS
 - **Please call to report!**
- We cannot do rule-out testing for botulism. If we're testing, we're treating
- Decision to treat is based on clinical presentation and is made by the physician along with IDOH and CDC
- IDOH & CDC need to authorize release of antitoxin should a patient truly need it.

Whom to Contact

Contact Type	Name	Phone Number
Indiana Department of Health	Enteric Epidemiologist Epi on Call (After Hours)	(317) 233-7125 (317) 233-1325
Non-Infant Patients	CDC Emergency Operations Center	(770) 488-7100
Infant Patients	California Department of Health Infant Botulism Treatment & Prevention Program	(510) 231-7600

Medical Decision Making



When in doubt, please call!

Botulism Anti-Toxin (BAT) Logistics

Consult with CDC & IDOH



CDC releases antitoxin from nearest Quarantine Station (Chicago); coordinates shipment



Antitoxin arrives at healthcare facility to be administered by treatment team (clinician, pharmacist, etc.)

Anti-Toxin Administration and Actions

[Preparing and Administering Heptavalent Antitoxin for the Treatment of Botulism](#)



[▶ Low Resolution Video](#)

Centers for Disease Control and Prevention
1600 Clifton Road, MS C09
Atlanta, GA 30333

Clinical Outcome Report*

**Please include copy of discharge summary*

Please complete upon discharge or death and email to CDC at bot surveillance@cdc.gov or fax to 1-678-731-1542

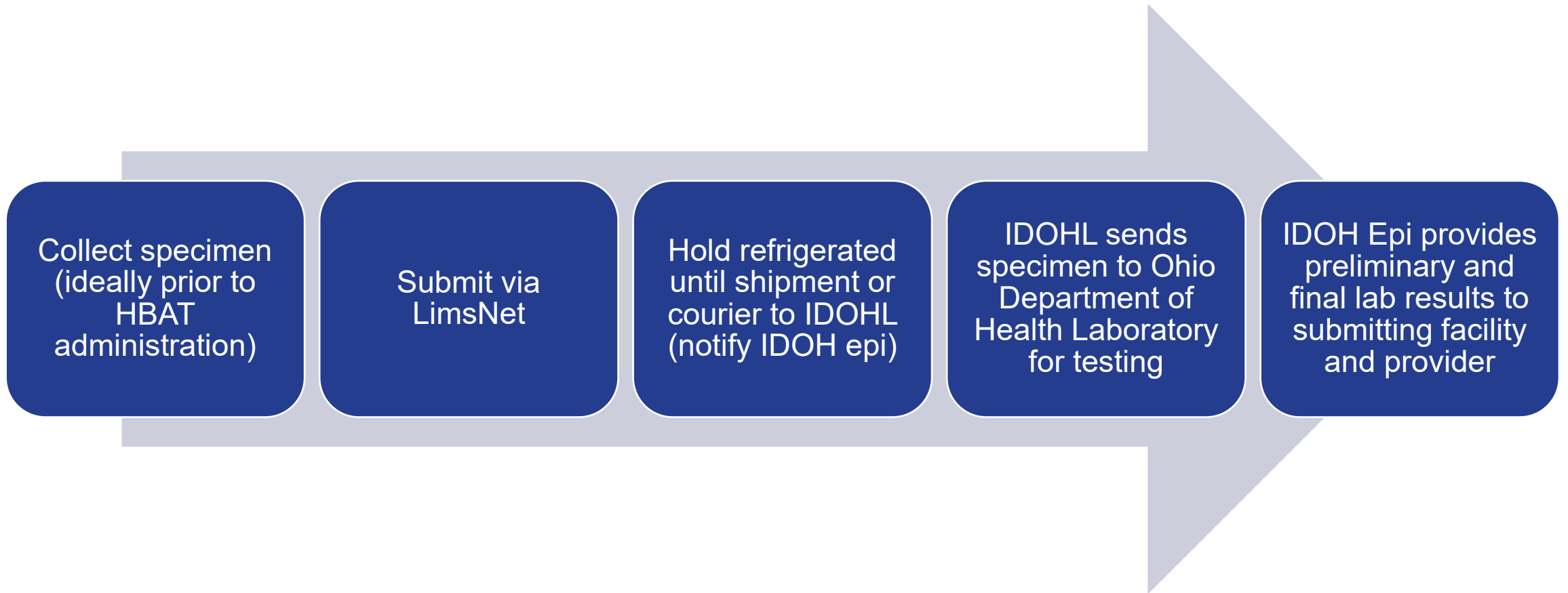
REPORTING AGENCY			
Treating Physician - Last Name, First Name	Telephone Number	Fax Number	Today's Date
Attending Physician Name - Last Name, First Name	Telephone Number	Fax Number	Specialty
Hospital Name	City	State	Zip Code
DEMOGRAPHIC INFORMATION			
Patient Name - Last Name, First Name, Middle Initial	City	State	Zip Code
Date of Birth	Sex <input type="checkbox"/> Male <input type="checkbox"/> Female		
CLINICAL OUTCOME INFORMATION			
How many days was patient hospitalized? _____ days			
How many days was patient in intensive care? _____ days			
Did patient require mechanical ventilation? <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unk			
If yes, how many days was patient on a ventilator? _____ days			
Did patient require a tracheostomy? <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unk			
If yes, when was the tracheostomy done? _____			
Did the patient develop pneumonia? <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unk			
What was the final diagnosis? (please check one)			
<input type="checkbox"/> Botulism <input type="checkbox"/> Tick paralysis <input type="checkbox"/> Paralytic shellfish poisoning			
<input type="checkbox"/> Myasthenia gravis <input type="checkbox"/> Eaton-Lambert syndrome <input type="checkbox"/> Other _____			
<input type="checkbox"/> Guillain-Barre syndrome <input type="checkbox"/> Stroke or central nervous system mass or lesion			
Was treatment given for any of the above diagnosis (even if it wasn't the final diagnosis)? <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unk			
If yes, specify type <input type="checkbox"/> Botulism Antitoxin <input type="checkbox"/> Plasmapheresis <input type="checkbox"/> Neostigmine/Physostigmine <input type="checkbox"/> Other Immunoglobulin therapy _____			
Did the patient develop an adverse event after botulism antitoxin administration? <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unk			
If yes, specify adverse event _____			
Did the patient die? <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unk			
If yes, When did patient die? _____			
What was the cause of death? _____			
If no, Where was patient discharged? <input type="checkbox"/> Home <input type="checkbox"/> Nursing home <input type="checkbox"/> Physical therapy/rehabilitation facility <input type="checkbox"/> Other (specify) _____			
Did patient have residual disability upon discharge? <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unk			
If yes, please specify types below (check as many as apply)			
<input type="checkbox"/> Proximal Upper Extremity Weakness <input type="checkbox"/> Diminished deep tendon reflexes <input type="checkbox"/> Other _____			
<input type="checkbox"/> Distal Upper Extremity Weakness <input type="checkbox"/> Fatigue <input type="checkbox"/> Other _____			
<input type="checkbox"/> Proximal Lower Extremity Weakness <input type="checkbox"/> Stroke or central nervous system mass or lesion			
<input type="checkbox"/> Distal Lower Extremity Weakness <input type="checkbox"/> Other _____			
ADDITIONAL INFORMATION			
Comments / Remarks: _____			

Testing for Botulism

- We **cannot** test an individual to rule-out the possibility of botulism because:
 - Delaying botulism treatment while waiting on testing results can cause severe, adverse consequences for the patient.
 - Testing is extensive
 - If the result is negative: Turnaround time is 9 days
 - If the result is positive: Turnaround time is 17 days
 - Botulism testing is subject to false positive results.

Alternatively for public health purposes we do require testing if we are releasing anti-toxin to be able to confirm the presence of toxigenic *C. botulinum*

Testing procedures



Clostridium botulinum & Botulinum Neurotoxin

Specimen Collection, Handling, and Shipping Guidelines

Disease/Agent	Specimen Selection					Shipping & Storage*	Specimen Handling
Botulism (botulinum toxin)	Specimen type	Clinical syndrome				Specimen(s) of choice for confirming botulism: 1. Serum 3. Stool 2. Wound/tissue 4. Incriminated food	
		Foodborne	Infant	Wound	Intentional release (airborne)		
	Enema fluid – 20 ml	X	X		X	2-8°C	Purge with a minimal amount of sterile non-bacteriostatic water to minimize dilution of toxin
	Food sample – 10-50g Liquid sample – 10-50ml	X	X		X	Ship at temperature as found when collected	Foods that support <i>C. botulinum</i> growth will have a pH of 3.5-7.0; most common pH is 5.5-6.5. Submit food in original container where possible or in leak-proof sealed transport devices. Botulinum toxin in commercial products is rare; contact the US FDA immediately if a commercial product is suspected of containing botulinum toxin.
	Gastric fluid – 20 ml	X, A				2-8°C	Collect up to 20 ml
	Intestinal fluid – 20 ml	A	A			2-8°C	Intestinal contents from various areas of the small and large intestines should be provided
	Nasal swab (anaerobic swab)				X	Room Temp	For aerosolized botulinum toxin exposure, obtain nasal cultures for <i>C. botulinum</i> and serum for mouse toxicity testing
	Serum – 15-20 mls	X, A		X	X	2-8°C	Serum should be obtained as soon as possible after the onset of symptoms and before antitoxin is given. Serum is required for mouse toxicity testing. In infants, serum is generally not useful, since the toxin is quickly absorbed before serum can be obtained.
	Stool ≥25 g	X	X	X	X	2-8°C	Botulism has been confirmed in infants with only “pea-size” stools. Please note: Anticholinesterase given orally, as in patients with myasthenia gravis, has been shown to interfere with toxin testing. <i>C. botulinum</i> has been isolated from stools following antitoxin treatment.
	Vomitus - 20 ml	X				2-8°C	Collect up to 20 ml
	Wound or Tissue			X		2-8°C	Anaerobic swab or transport system

*Refrigerated specimens that cannot be shipped to assure receipt within 72 hours of collection, freeze at ≤ -20°C

A= Autopsy specimens acceptable for certain specimen types

Revised 2012

Microbiology Client Services Manual



Submit Tests Packages Test Results Personalized Settings

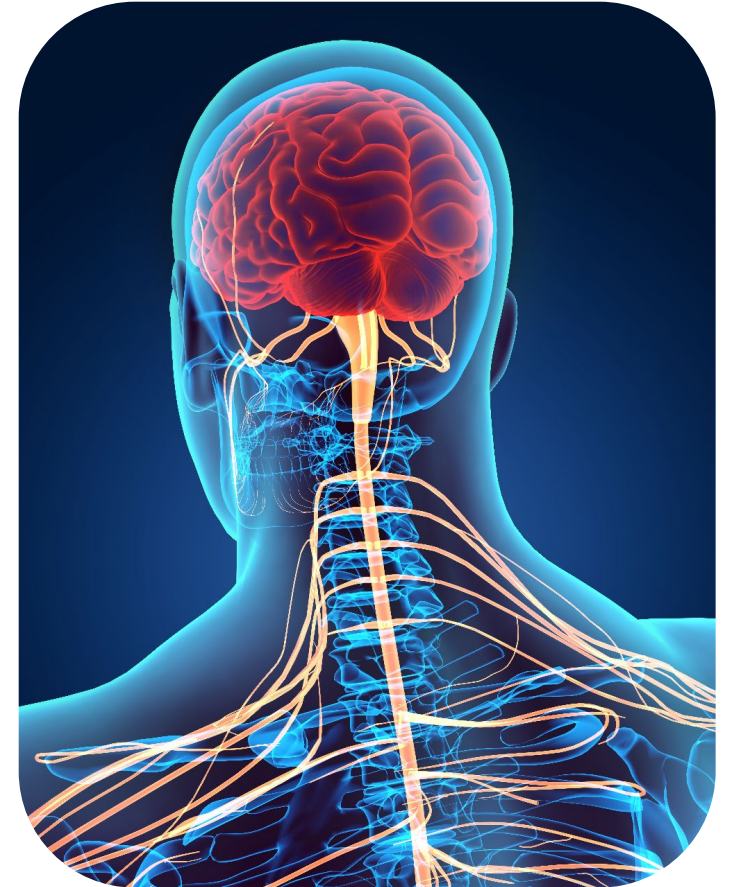
Patient Interview



- Please notify IDOH staff or CDC during the consultation if a high-risk exposure was identified during patient intake and evaluation.
- If deemed necessary, the LHD or an enteric epidemiologist will contact the patient or family to obtain exposures of concern. This is included, but not limited to foods, drug use, and environmental exposures.
- While talking with the patient, please notify them that someone may be reaching out to them for additional questions.

General Reminders

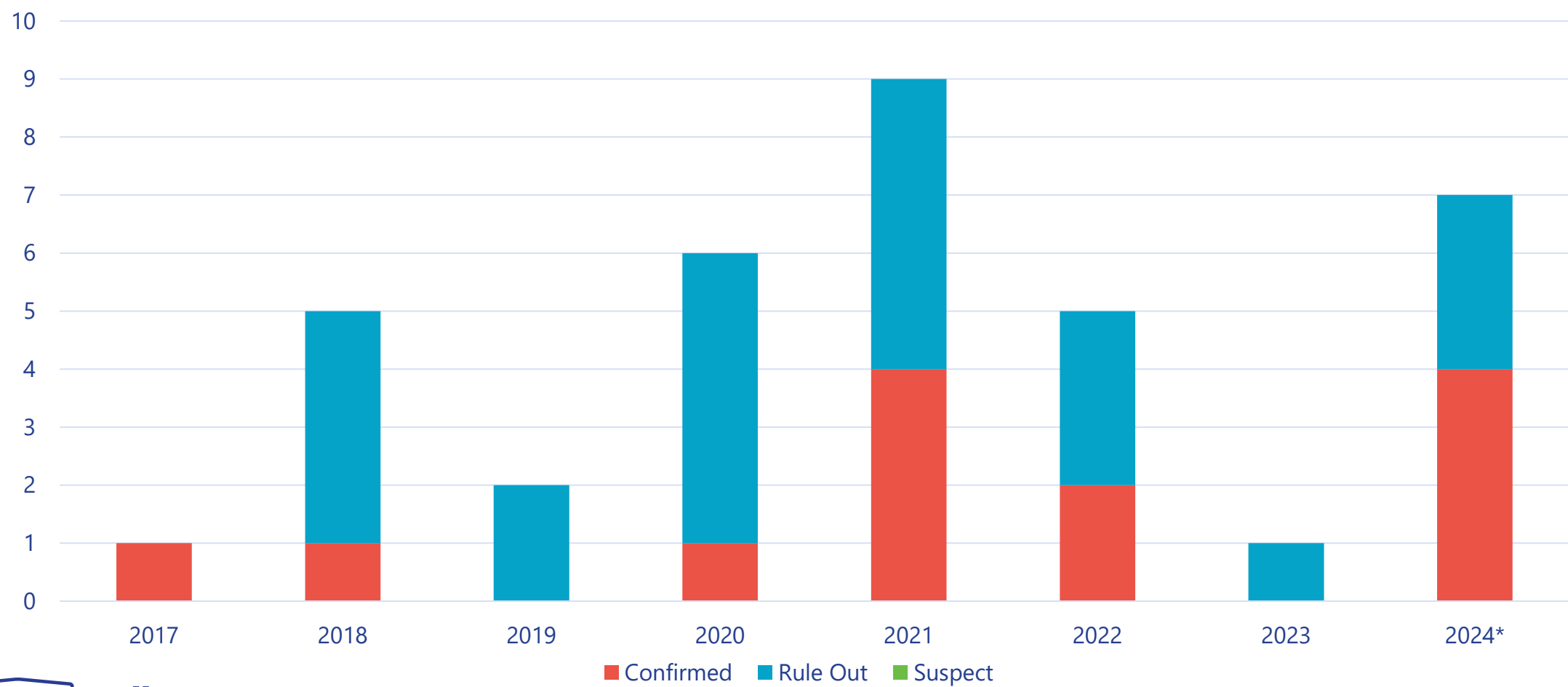
- Please give a clinical point contact to IDOH to help streamline ongoing process and they can also help with any questions directly between epi and clinician.
- Try to include and notify your hospital infection preventionist—they know how to get a hold of us!



Indiana Botulism Cases 2013-2024*

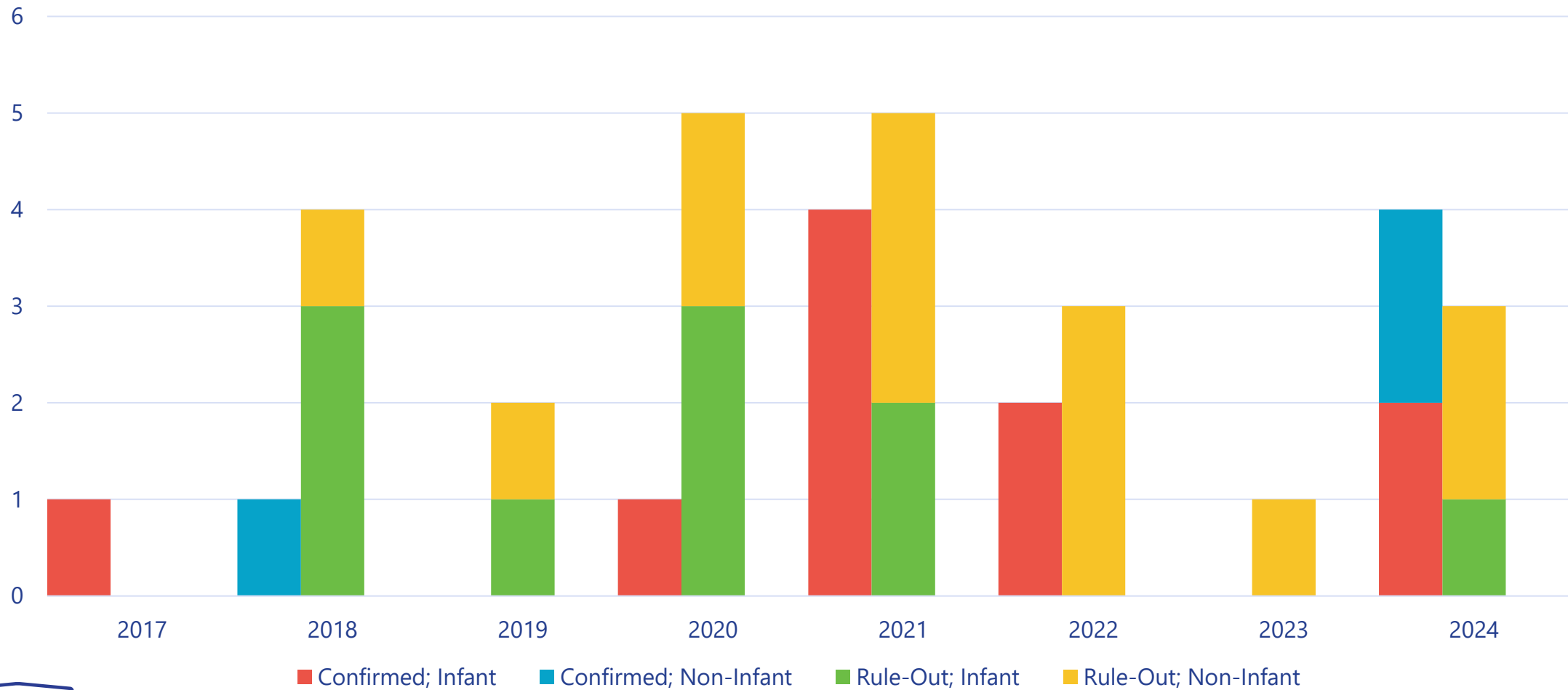


Botulism Rule-Outs per Year (2017-2024*)



*2024 pending

Botulism Rule-Outs per Year by Type (2017-2024*)



*2024 pending

Resources & Links

- CDC Clinician Resources for Botulism
 - <https://www.cdc.gov/botulism/hcp/clinician-resources/index.html>
- CA DPH Infant Botulism Treatment and Prevention Program
 - <https://www.cdph.ca.gov/Programs/CID/DCDC/Pages/InfantBotulism.aspx#>
- Rao AK, Sobel J, Chatham-Stephens K, Luquez C. Clinical Guidelines for Diagnosis and Treatment of Botulism, 2021. MMWR Recomm Rep 2021;70(No. RR-2):1–30. DOI: <http://dx.doi.org/10.15585/mmwr.rr7002a1>
 - <https://www.cdc.gov/mmwr/volumes/70/rr/rr7002a1.htm>

Questions?

Nicole Stone, MPH

Respiratory & Enteric Disease
Epidemiology Director

nstone2@health.in.gov





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UPDATES ON ACUTE FLACCID MYELITIS

MAKAYLA CULBERTSON

SENIOR VACCINE-PREVENTABLE
DISEASE EPIDEMIOLOGIST

09/27/2024

Acute Flaccid Myelitis Updates





- As of September 18, 2024, this year CDC has received 23 reports of suspected AFM, with 14 confirmed cases in 11 states
 - This includes one suspect case from Indiana
- As we enter the fall season, seasonal increases in the circulation of respiratory pathogens, including enteroviruses, is to be expected.
- Enterovirus D-68 (EV-D68) is believed to be the main enterovirus responsible for the increases in AFM cases observed during 2014, 2016, and 2018

Clinical Presentation of AFM

- Patients with acute flaccid myelitis can present with:
 - Sudden arm or leg weakness
 - Pain in the arms or legs
 - Pain in the neck or back
 - Difficulty moving the eyes or drooping eyelids
 - Facial droop
 - Difficulty swallowing or slurred speech
- These symptoms often happen after a viral infection

Reporting AFM

- Identify a patient under investigation (PUI); patient with onset of acute flaccid limb weakness and an MRI showing spinal cord lesions in at least some gray matter
- Contact the Indiana Department of Health (Business hours: 317-233-7125 After Hours: 317-233-1325)
- Collect CSF, whole stool, respiratory, and serum specimens to send to IDOHL for CDC forwarding
- MRI imaging will also be requested

SAMPLE	AMOUNT	TUBE TYPE	PROCESSING	STORAGE	SHIPPING
CSF	0.15 mL, 0.5-2 mL preferred (collect at same time or within 24hrs of serum if feasible)	Cryovial 	Spun and CSF removed to cryovial	Freeze at $\leq -20^{\circ}\text{C}$	Frozen on dry ice.
Respiratory Nasopharyngeal (NP)/Oropharyngeal (OP) swab	0.5 mL, 1 mL preferred (minimum amount)	N/A 	Store in vial transport medium	Freeze at $\leq -20^{\circ}\text{C}$	Frozen on dry ice.
Serum	0.5 mL, 1 mL preferred (collect at same time or within 24hrs of CSF if feasible)	Tiger/red top for collection; separate tube for shipping 	Spun and serum aliquot removed to separate tube	Freeze at $\leq -20^{\circ}\text{C}$	Frozen on dry ice.
Stool	1 gram, 10 – 20 grams preferred (2 samples collected 24hrs apart)	Sterile container 	N/A	Freeze at $\leq -20^{\circ}\text{C}$	Frozen on dry ice. Rectal swabs should not be sent in place of stool.

Please, always include whole stool specimens to help identify pathogens and rule out poliovirus.

AFM Resources

- [CDC AFM Cases](#)
- [CDC Tools and Resources for Providers](#)
- [CDC AFM Job Aid for Clinicians](#)
- [CDC AFM Symptom Infographic](#)
- [CDC Clinical Guidance for the Acute Medical Treatment of AFM](#)
- [AFM Physician Consult and Support Portal](#)

CDC Healthcare provider tools and resources AFM

HEAD SHOULDERS KNEES & TOES

Unexplained proximal muscle weakness in children can occur in some neurologic conditions and can be easily missed during exams that only focus on distal strength.

When examining children with sudden limb, neck, or trunk weakness, remember **head, shoulders, knees, and toes**.

Raise KNEES

Muscle Group:

- * Hips

Ask:

- * Are they limping or dragging a leg?
- * Can they put on pants?
- * Can they do a squat and recover?



Reach down & touch TOES

Muscle Group:

- * Trunk

Ask:

- * Are they waddling or falling while walking?
- * Can they sit up and stand without support?
- * Can they get a toy off the ground while standing?



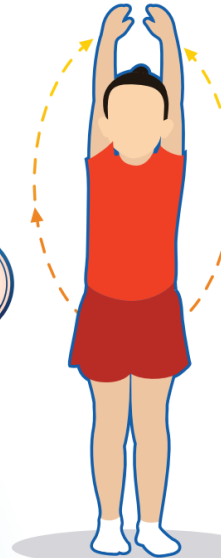
Lift both arms above the HEAD

Muscle Group:

- * Shoulder Girdle

Ask:

- * Are they using one limb less?
- * Can they put on a T-shirt?
- * Can they give a high-five with each hand?



Shrug the SHOULDERS

Muscle Group:

- * Neck/Shoulder Girdle

Ask:

- * Is one shoulder higher than the other?
- * Can they throw a ball overhead?
- * Can they hold up their head?



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Questions?

Makayla Culbertson

Senior VPD Epi

mculbertson@health.in.gov





Respiratory Disease Update (Flu, COVID, RSV)



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CDC respiratory season prediction

Upcoming 2024–25 respiratory season peak hospitalization burden likely similar to or lower than last year

Combined peak hospitalization burden of COVID-19, influenza, and RSV

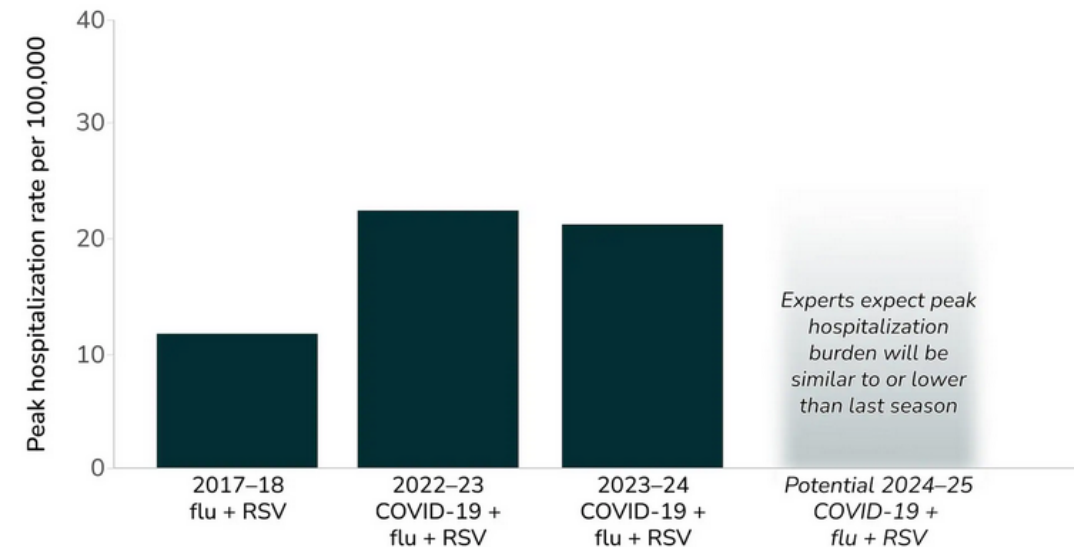


Figure 1: Experts believe there's a roughly 80% chance that the hospitalization burden for COVID-19, influenza, and RSV combined at their peak during the 2024–25 season will be similar to or lower than last season. The 2017–18 bar provides context for a severe season pre-pandemic; 2017–18 RSV data does not include pediatric hospitalizations. Data are from RESP-NET.

WHAT YOU NEED TO KNOW ABOUT FALL VACCINES 2024

Immunizations have been shown to lower risk of severe disease. Speak to your health care provider about the best timing for you.

Vaccine

Who

What

When



People 6 months of age and older

Updated 2024–2025 flu vaccine

During flu season. September and October remain the best times for most people to get vaccinated



Everyone aged 6 months and older should get 1 updated Moderna, Novavax, or Pfizer COVID-19 vaccine to be up to date.

Updated 2024–2025 COVID-19 vaccine

During fall and winter respiratory disease season



Adults over 75 and older and adults 60–74 at increased risk of severe RSV

NOT AN ANNUAL VACCINE

Eligible adults can get any time, best time is in late summer and early fall

WHAT YOU NEED TO KNOW ABOUT FALL VACCINES 2024

Immunizations have been shown to lower risk of severe disease. Speak to your health care provider about the best timing for you.

Vaccine

Who

What

When



Pregnant women at 32–36 weeks

Pfizer Abrysvo is the only RSV vaccine approved for pregnant women

September through January



Infants 19 months and younger

Monoclonal antibody shot

October through the end of March





Nirsevimab was **90% effective
at protecting infants from
RSV-associated hospitalization***

**Clinicians, talk to parents about
nirsevimab, a preventive antibody**

* Early estimates from the New Vaccine Surveillance Network, October 2023–February 2024



[bit.ly/mm7309a4](https://www.cdc.gov/mmwr/volumes/73/wr/mm7309a4.htm)

MARCH 7, 2024

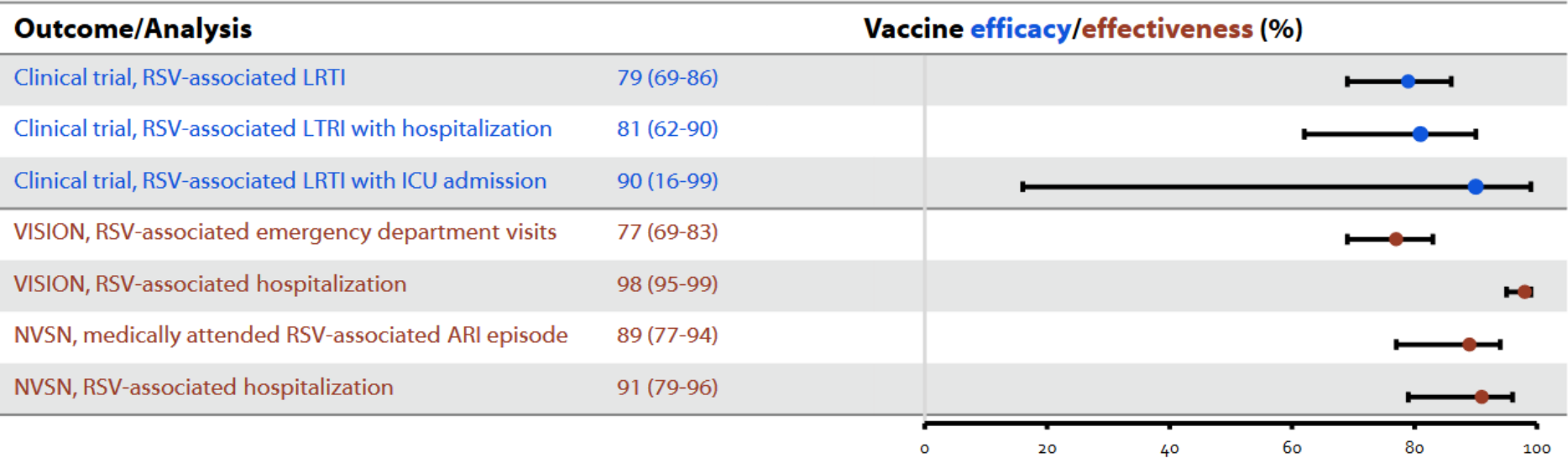
MMWR



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<https://www.cdc.gov/mmwr/volumes/73/wr/mm7309a4.htm>

Observational data indicate nirsevimab is working as expected (vs. RCT results) during the first RSV season after approval among infants in their first RSV season



Results may not be comparable across studies due to differences in outcome definitions, timing, and other factors.

<https://www.cdc.gov/mmwr/volumes/72/wr/mm7234a4.htm>
RCT = randomized clinical trial | ARI = acute respiratory illness

CDC RSV survey

- Survey done in March-April this year
- Coverage among those surveyed:
 - Maternal RSV vaccination among eligible pregnant women was 32.6%
 - Nirsevimab among infants was 44.6%
 - 55.8% of infants were protected by either or both products.
 - Receipt of a provider recommendation was strongly associated with both maternal and infant immunization.

RSV maternal vaccine and infant monoclonal antibody recommendations from CDC

Infants and young children

- To prevent severe RSV disease in infants, CDC recommends either maternal RSV vaccination or infant immunization with RSV monoclonal antibodies. **Most infants will not need both.**

Vaccination for pregnant women

- 1 dose of maternal RSV vaccine during weeks 32 through 36 of pregnancy, administered **September through January**. Pfizer Abrysvo is the only RSV vaccine recommended during pregnancy.

Immunization for infants and young children (monoclonal antibodies)

- 1 dose of nirsevimab is recommended for infants younger than 8 months of age who were born shortly before or are entering their first RSV season (**typically October through March**)
- 1 dose of nirsevimab for infants and children aged 8–19 months who are at increased risk for severe RSV disease and entering their second RSV season.
- *Note:* A different monoclonal antibody, palivizumab, is limited to children aged 24 months and younger with certain conditions that place them at high risk for severe RSV disease. It must be given once a month during RSV season.

Older Adult RSV Recommendation

- ACIP recommends adults 75 years and older receive a single dose of RSV vaccine
 - Removes shared clinical decision-making language
 - Includes the new Moderna RSV vaccine
- ACIP recommends adults 60-74 who are at increased risk of severe RSV disease receive a single dose of RSV vaccine
 - Shared clinical decision-making recommendation versus a risk-based recommendation
 - Adults that have already receive da dose of RSV do not need to receive another dose in the 2024-25 season

COVID-19 in Indiana



Indiana COVID-19 Home Dashboard

Below results are as of 9/24/2024, 11:59 PM. Dashboard updates by 5 p.m. on Wednesdays.

7-Day Average
COVID-19 Counts

COVID-19 Hospital
Admissions
22 (↓9)

Emergency
Department Visits
for
COVID-Like Illness
307 (↓39)

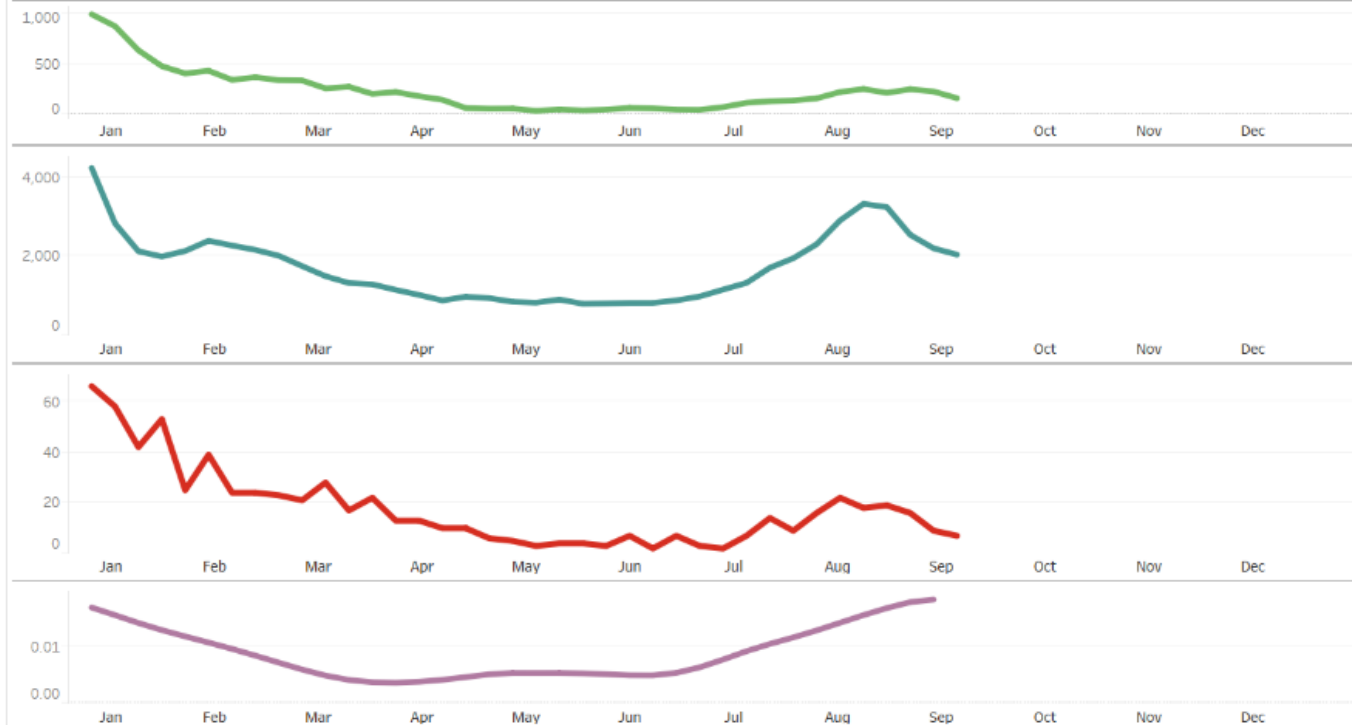
COVID-19 Deaths
1 (No Change)

SARS-CoV-2
Wastewater
Concentration
0.0177 (↑0.0010)
2,022,677 Total
Population Served

COVID-19 Trends

- 2024 COVID-19 Hospital Admissions
- 2024 Emergency Department Visits for COVID-Like Illness
- 2024 COVID-19 Deaths
- 2024 Concentration of SARS-CoV-2 in Wastewater
- 2020 2021 2022 2023

Year Selection
(filters Timeseries only)
2024



All numbers are provisional and reflect only those reported to IDOH. Numbers should not be characterized as a comprehensive total and may change as more data is reported.

COVID-19 Update for the United States

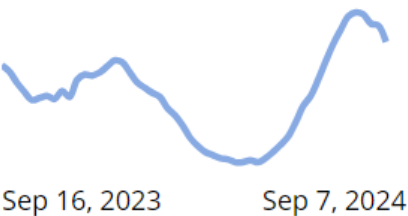
Early Indicators

Test Positivity >

% Test Positivity

14.9%

Week ending September 7, 2024
Previous week 16.5%

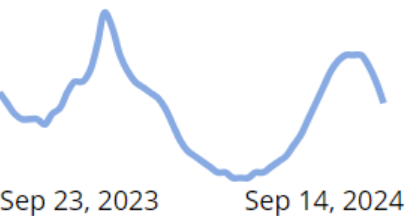


Emergency Department Visits >

% Diagnosed as COVID-19

1.7%

Week ending September 14, 2024
Previous week 2.1%



These early indicators represent a portion of national COVID-19 tests and emergency department visits. [Wastewater](#) information also provides early indicators of spread.

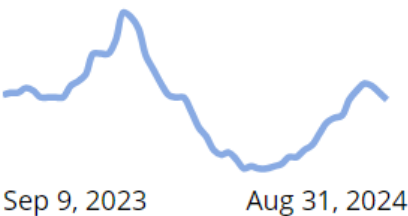
Severity Indicators

Hospitalizations >

Rate per 100,000 population

4.1

Week ending August 31, 2024
Previous week 4.4

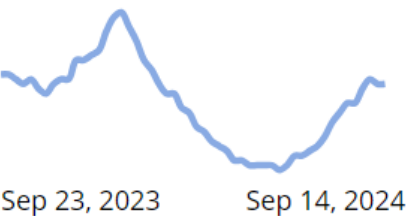


Deaths >

% of All Deaths in U.S. Due to COVID-19

2.3%

Week ending September 14, 2024
Previous week 2.3%



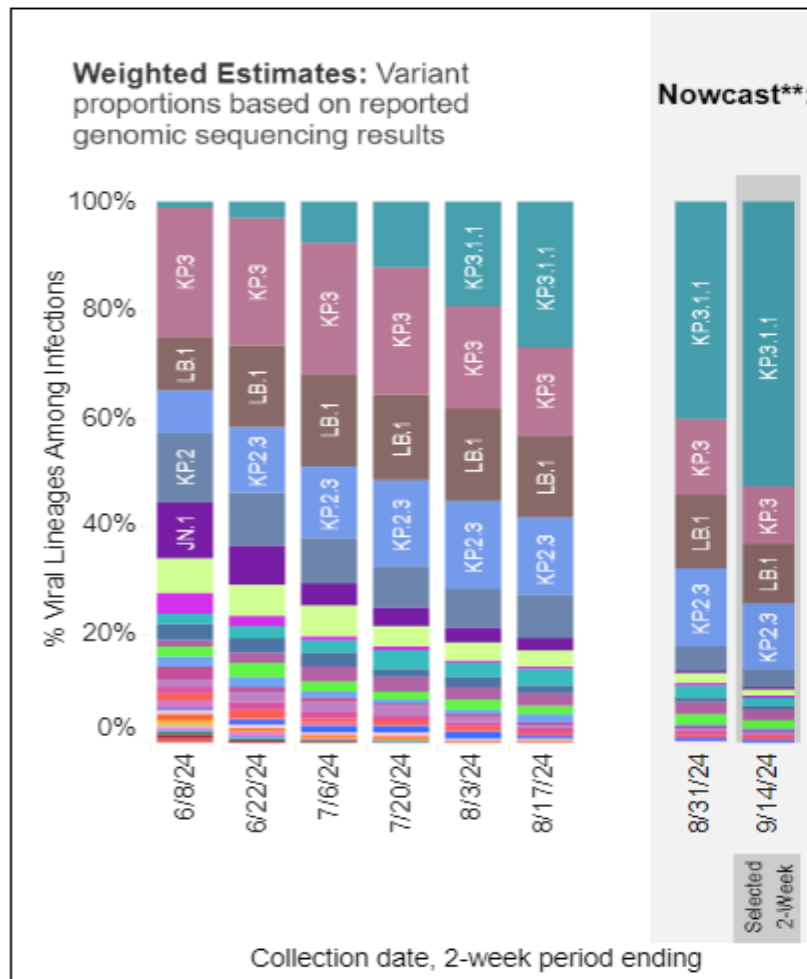
CDC | Test Positivity data through: September 7, 2024; Emergency Department Visit data through: September 14, 2024; Hospitalization data through: August 31, 2024; Death data through: September 14, 2024.
Posted: September 23, 2024 2:52 PM ET

Weighted and Nowcast Estimates in United States for 2-Week Periods in 5..

Nowcast Estimates in United States for 9/1/2024 – 9/14/2024



Hover over (or tap in mobile) any lineage of interest to see the amount of uncertainty in that lineage's estimate.



USA

WHO label	Lineage #	US Class	%Total	95%PI	
Omicron	KP.3.1.1	52.7%	48.6-56.8%		
	KP.2.3	12.2%	10.8-13.8%		
	LB.1	10.9%	9.4-12.6%		
	KP.3	10.6%	9.3-12.1%		
	KP.2	3.1%	2.2-4.2%		
	LP.1	2.1%	1.4-3.0%		
	KP.1.1.3	1.9%	1.4-2.8%		
	JN.1.18	1.7%	0.6-4.0%		
	KP.1.1	1.5%	1.2-1.9%		
	KS.1	0.7%	0.4-1.0%		
	KP.2.15	0.7%	0.4-1.0%		
	LF.3.1	0.6%	0.4-0.9%		
	JN.1.16.1	0.6%	0.4-0.8%		
	KP.4.1	0.2%	0.1-0.4%		
	JN.1.11.1	0.2%	0.1-0.3%		
	JN.1	0.2%	0.1-0.3%		
	KW.1.1	0.0%	0.0-0.1%		
	XDV.1	0.0%	0.0-0.1%		
	JN.1.16	0.0%	0.0-0.0%		
	JN.1.7	0.0%	0.0-0.0%		
	KP.1.2	0.0%	0.0-0.0%		
	KQ.1	0.0%	0.0-0.0%		
	JN.1.8.1	0.0%	0.0-0.0%		
	JN.1.32	0.0%	0.0-0.0%		

COVID-19 Recommendation


- ACIP recommends 2024-2025 COVID-19 formulation for individuals 6 months of age and older
- COVID-19 Hospitalizations
 - 50% of hospitalized children and adolescents have no underlying conditions
 - 2/3 of all COVID-19 associated hospitalizations are in adults 65+
 - 19% of adults 65+ who were hospitalized were residents of a long-term care facility
- COVID-19 Vaccine Effectiveness
 - 53% effective against symptomatic infections in adults
 - 66-71% effective against ED/UC visits among children 9 mo -17 yrs



COVID-19 vaccine recs MMWR 9/19/24

**CDC recommends the 2024-25
COVID-19 vaccine for everyone 6 months and older**

**An updated vaccine protects
against:**

- ☒ COVID-19 variants spreading now
- ☒ Severe illness, hospitalization,
and death



 bit.ly/mm7337e2
SEPTEMBER 10, 2024 

FDA Approves updated COVID-19 vaccines

FDA NEWS RELEASE

- **Based on KP.2**
- **Monovalent**

FDA Approves and Authorizes Updated mRNA COVID-19 Vaccines to Better Protect Against Currently Circulating Variants

Unvaccinated individuals 6 months through 4 years of age are eligible to receive three doses of the updated, authorized Pfizer-BioNTech COVID-19 Vaccine or two doses of the updated, authorized Moderna COVID-19 Vaccine.

Individuals 6 months through 4 years of age who have previously been vaccinated against COVID-19 are eligible to receive one or two doses of the updated, authorized Moderna or Pfizer-BioNTech COVID-19 vaccines (timing and number of doses to administer depends on the previous COVID-19 vaccine received).

Individuals 5 years through 11 years of age regardless of previous vaccination are eligible to receive a single dose of the updated, authorized Moderna or Pfizer-BioNTech COVID-19 vaccines; if previously vaccinated, the dose is administered at least 2 months after the last dose of any COVID-19 vaccine.

Individuals 12 years of age and older are eligible to receive a single dose of the updated, approved Comirnaty or the updated, approved Spikevax; if previously vaccinated, the dose is administered at least 2 months since the last dose of any COVID-19 vaccine.

Additional doses are authorized for certain immunocompromised individuals ages 6 months through 11 years of age as described in the Moderna COVID-19 Vaccine and Pfizer-BioNTech COVID-19 Vaccine fact sheets.



<https://www.fda.gov/news-events/press-announcements/fda-approves-and-authorizes-updated-mrna-covid-19-vaccines-better-protect-against-currently#:~:text=The%20updated%20mRNA%20COVID%2D19,through%2011%20years%20of%20age.>

FDA Approves Novavax COVID-19 vaccine

FDA NEWS RELEASE

JN.1

Emergency Use Authorization

What You Need to Know

- Individuals 12 years of age and older who have never been vaccinated with any COVID-19 vaccine are eligible to receive two doses of this updated vaccine, 3 weeks apart.
- Individuals who have been vaccinated only with one dose of any Novavax COVID-19 vaccine are eligible to receive one dose of the updated Novavax COVID-19 vaccine at least 3 weeks after the previous dose.
- Those who have been vaccinated with a prior formula of a COVID-19 vaccine from another manufacturer or with two or more doses of a prior formula of the Novavax COVID-19 vaccine are eligible to receive a single dose of the updated Novavax COVID-19 vaccine at least 2 months after the last dose of a COVID-19 vaccine.

FDA Authorizes Updated Novavax COVID-19 Vaccine to Better Protect Against Currently Circulating Variants

Updated CDC COVID vaccine recommendations

Ages 12 years and older

COVID-19 vaccination history ⁶	2024–2025 vaccine	Number of 2024–2025 doses indicated	Dosage (mL/ug)	Interval between doses
Unvaccinated	Moderna	1	0.5 mL/50 ug	—
	OR			
	Novavax	2	0.5 mL/5 ug rS protein and 50 ug Matrix-M adjuvant	Dose 1: Day 0 Dose 2: 3–8 weeks after Dose 1*
	OR			
	Pfizer-BioNTech	1	0.3 mL/30 ug	—
1 or more doses any mRNA, NOT including 1 dose any 2024–2025 COVID-19 vaccine	Moderna	1	0.5 mL/50 ug	At least 8 weeks after last dose
	OR			
	Novavax	1	0.5 mL/5 ug rS protein and 50 ug Matrix-M adjuvant	At least 8 weeks after last dose
	OR			
	Pfizer-BioNTech	1	0.3 mL/30 ug	At least 8 weeks after last dose

1 dose any Novavax	Novavax	1	0.5 mL/5 ug rS protein and 50 ug Matrix-M adjuvant	Dose 2: 3–8 weeks after Dose 1**
2 or more doses any Novavax, NOT including 1 dose any 2024–2025 COVID-19 vaccine	Moderna	1	0.5 mL/50 ug	At least 8 weeks after last dose
	OR			
	Novavax	1	0.5 mL/5 ug rS protein and 50 ug Matrix-M adjuvant	At least 8 weeks after last dose
	OR			
	Pfizer-BioNTech	1	0.3 mL/30 ug	At least 8 weeks after last dose
2 or more doses any Novavax, INCLUDING 1 dose any 2024–2025 COVID-19 vaccine	No further doses indicated			

So what about those less than 6 mo old?

CDC MMWR 9/26:

- COVID-19–associated hospitalization rates among infants aged <6 months remain higher than those among any other age group except adults aged ≥ 75 years
 - Rates were comparable to hospitalization rates in adults aged 65–74 years.
 - Among approximately 1,000 hospitalized infants with COVID-19, 22% were admitted to an intensive care unit, and nine died while hospitalized.
 - The percentage of hospitalized infants whose mothers had been vaccinated during pregnancy was 18% during October 2022–September 2023 and decreased to <5% during October 2023–April 2024.
- CDC recommends COVID vaccination for pregnant women to protect infants in this age group.

Free COVID-19 Tests



COVID-19 Testing

Order Your 4 Free At-home COVID-19 Tests

Every U.S. household is eligible to order 4 free at-home tests.

Need help placing an order for your at-home tests?
Call [1-800-232-0233](tel:1-800-232-0233) (TTY [1-888-720-7489](tel:1-888-720-7489)).

Seasonal Influenza Vaccine

FLU PREVENTION TIPS



- ACIP affirmed the recommendation for a routine annual influenza vaccination for individuals 6 months and older who do not have contraindications
- No shortages or delays in shipment expected

FluMist now available for self or caregiver administration

FDA NEWS RELEASE

FDA Approves Nasal Spray Influenza Vaccine for Self- or Caregiver-Administration

*First Influenza Vaccine That Does Not Need to be Administered by a Health
Care Provider*

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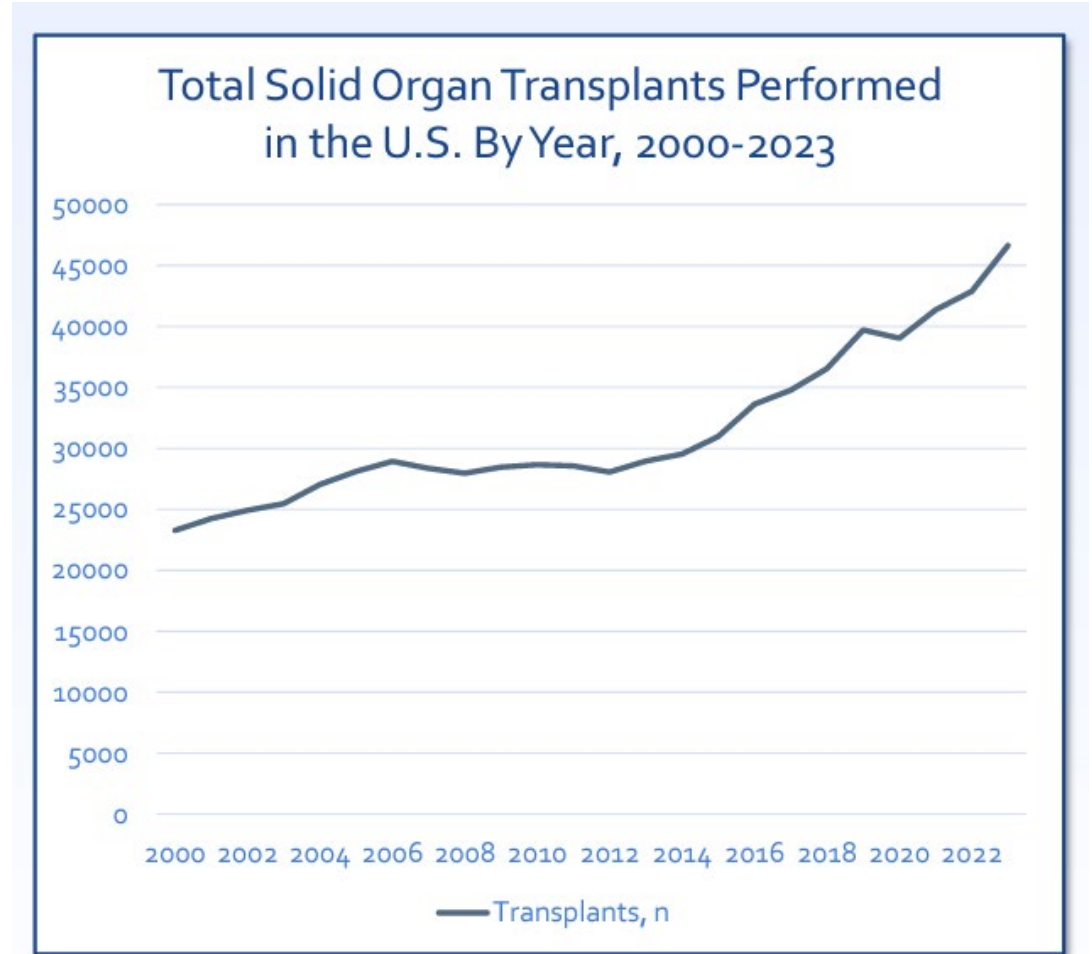
[✉ Email](#)

[🖨 Print](#)

High Dose/Adjuvanted Flu Vaccines for Solid Organ Transplant Recipients Recommendation

- New recommendation for high dose or adjuvanted flu vaccines for solid organ transplant in adults 18+
 - Previously only approved for 65+

[Link](#) to review of studies



CDC Study – Early Flu Antiviral Treatment Decreases Risk of Death

- FluSurv-NET 2012 - 2019
- Delayed initiation of antiviral treatment in patients hospitalized with influenza-associated pneumonia was associated with higher risk of death
- 30-day mortality
 - 7.5% - Day 0
 - 8.5% - Day 1
 - 10.2% - Days 2-5
- [CDC Link](#)

Clinical Infectious Diseases

MAJOR ARTICLE

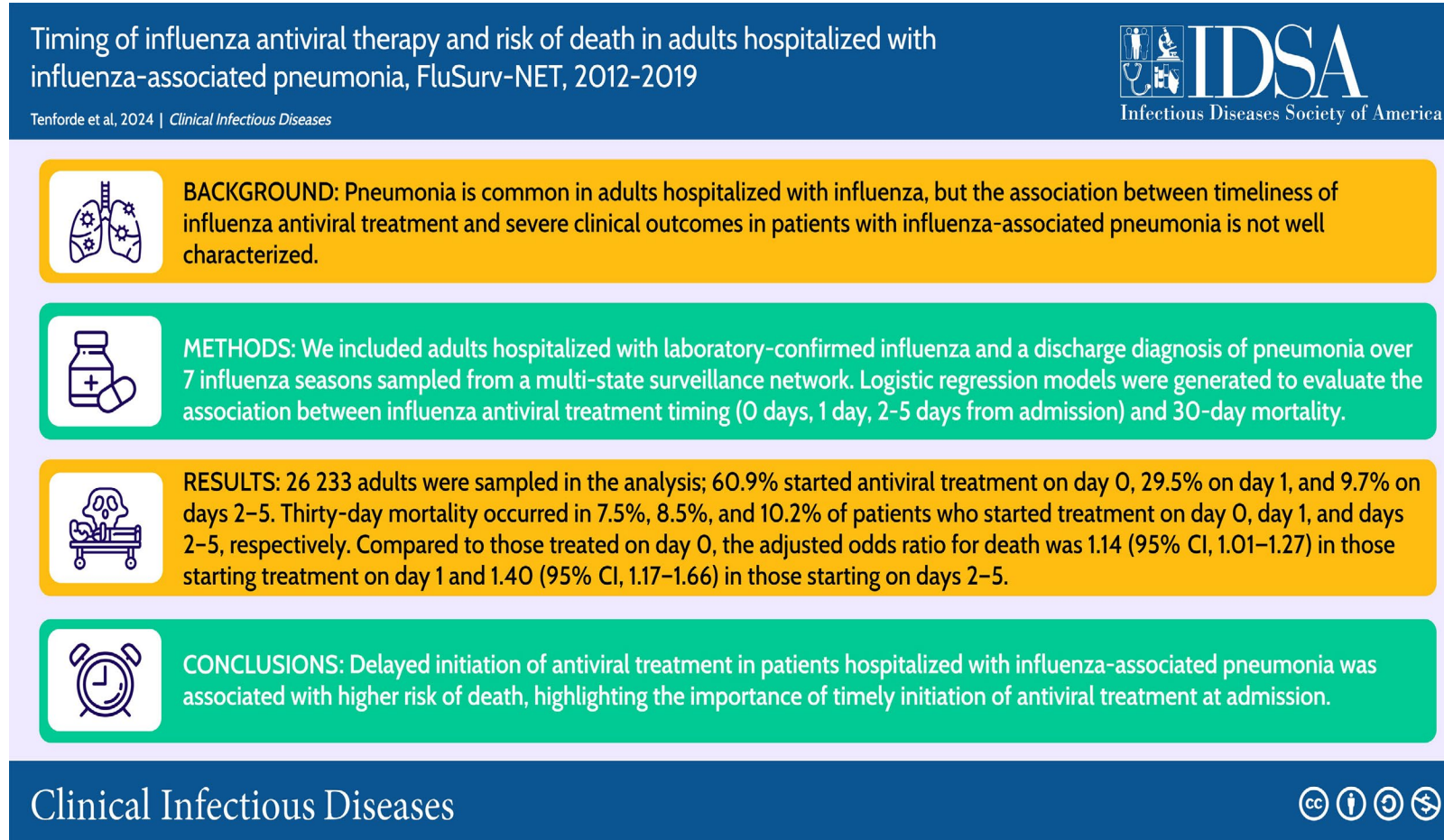


Timing of Influenza Antiviral Therapy and Risk of Death in Adults Hospitalized With Influenza-Associated Pneumonia, Influenza Hospitalization Surveillance Network (FluSurv-NET), 2012–2019

Mark W. Tenforde,^{1,6} Kameela P. Noah,¹ Alissa C. O'Halloran,¹ Pam Daily Kirley,² Cora Hoover,³ Nisha B. Alden,⁴ Isaac Armistead,⁴ James Meek,⁵ Kimberly Yousey-Hindes,⁵ Kyle P. Openo,^{6,7,8} Lucy S. Witt,^{6,7} Maya L. Monroe,⁹ Patricia A. Ryan,⁹ Anna Falkowski,¹⁰ Libby Reeg,¹⁰ Ruth Lynfield,¹¹ Melissa McMahon,¹¹ Emily B. Hancock,¹² Marisa R. Hoffman,¹² Suzanne McGuire,¹³ Nancy L. Spina,¹³ Christina B. Felsen,¹⁴ Maria A. Gaitan,¹⁴ Krista Lung,¹⁵ Eli Shiltz,¹⁵ Ann Thomas,¹⁶ William Schaffner,¹⁷ H. Keipp Talbot,¹⁷ Melanie T. Crossland,¹⁸ Andrea Price,¹⁸ Svetlana Masalovich,¹ Katherine Adams,¹ Rachel Holstein,¹ Devi Sundaresan,¹ Timothy M. Uyeki,¹ Carrie Reed,¹ Catherine H. Bozio,¹ and Shikha Garg¹

¹Influenza Division, Centers for Disease Control and Prevention, Atlanta, Georgia, USA; ²California Emerging Infections Program, Oakland, California, USA; ³California Department of Public Health, Richmond, Virginia, USA; ⁴Colorado Department of Public Health and Environment, Denver, Colorado, USA; ⁵Connecticut Emerging Infections Program, Yale School of Public Health, New Haven, Connecticut, USA; ⁶Georgia Department of Public Health, Georgia Emerging Infections Program, Atlanta, Georgia, USA; ⁷Division of Infectious Diseases, Emory University School of Medicine, Atlanta, Georgia, USA; ⁸Research, Atlanta Veterans Affairs Medical Center, Decatur, Georgia, USA; ⁹Maryland Department of Health, Emerging Infections Program, Baltimore, Maryland, USA; ¹⁰Michigan Department of Health and Human Services, Lansing, Michigan, USA; ¹¹Health Protection Bureau, Minnesota Department of Health, St Paul, Minnesota, USA; ¹²New Mexico Department of Health, New Mexico Emerging Infections Program, Santa Fe, New Mexico, USA; ¹³New York State Department of Health, Albany, New York, USA; ¹⁴University of Rochester School of Medicine and Dentistry, Rochester, New York, USA; ¹⁵Ohio Department of Health, Columbus, Ohio, USA; ¹⁶Public Health Division, Oregon Health Authority, Salem, Oregon, USA; ¹⁷Division of Infectious Diseases, Vanderbilt University Medical Center, Nashville, Tennessee, USA; and ¹⁸Salt Lake County Health Department, Salt Lake City, Utah, USA

Graphical abstract This graphical abstract is also available at Tidbit: ...





Infectious Diseases of Public Health Importance

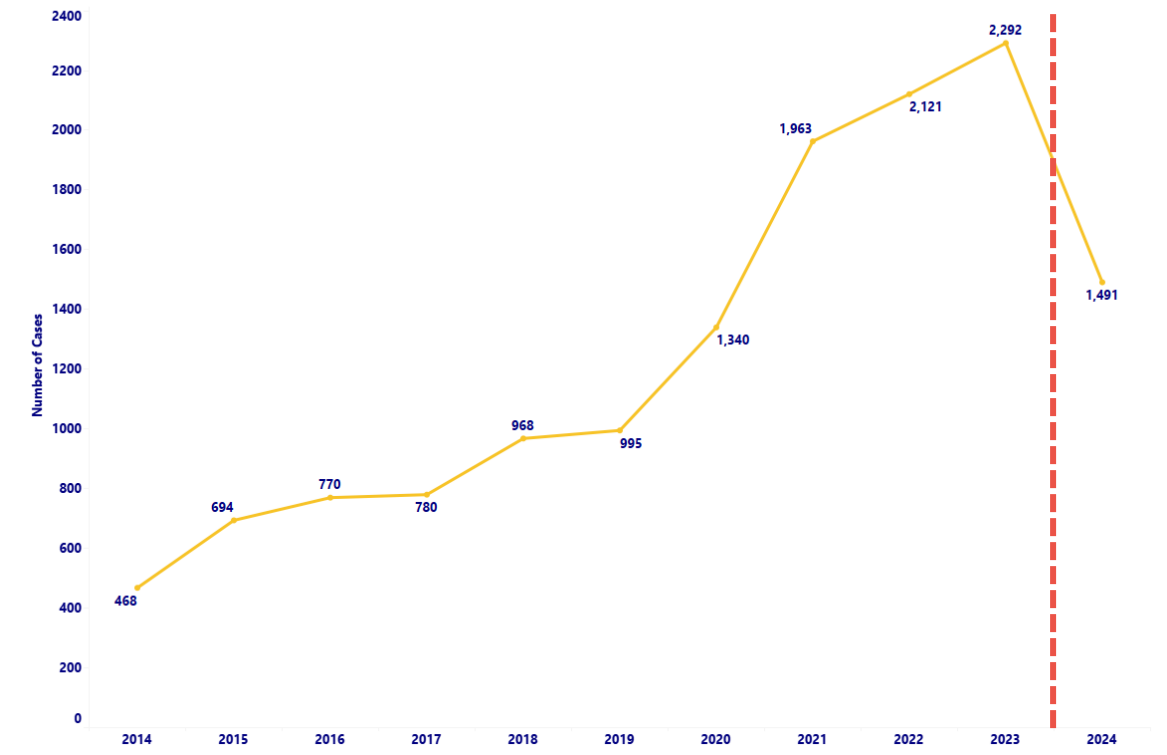


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Adult Syphilis Morbidity

- Rates of adult syphilis have been on the rise since 2014 in Indiana, reaching 33.9 (per 100,000) in 2023. Year to date there have been 1,491 cases of adult syphilis reported in 2024*.
- Black/African American Hoosiers have the highest rates of syphilis when compared to other races
- While rates are still highest among men, women (including those of childbearing age) have had the highest percent increase over time in recent years
 - From 2019-2023 there was a 283% increase in syphilis cases among females of childbearing age (15-44 years old)

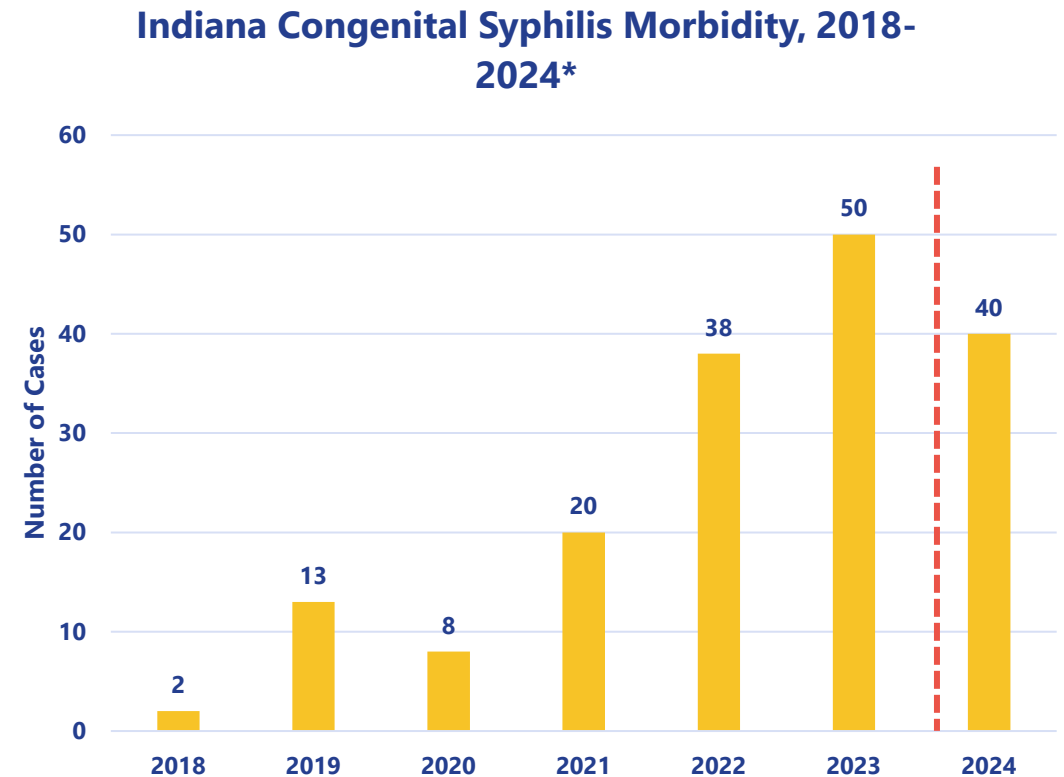
Adult Syphilis Morbidity, Indiana, 2014-2024*



*2024 STI data are preliminary.

Congenital Syphilis Morbidity

- From 2018-2023 there has been a 2,400% increase in congenital syphilis (CS) cases, with 50 cases reported in 2023 and 40 cases reported year to date in 2024*.
 - 31 counties have reported at least one CS case since 2018.
- Of the 40 CS cases reported this year in Indiana, **2 were stillbirths**



*2024 STI data are preliminary.

Importance of ED screening

- Study of about 300,000 people in Chicago
- Prior to study, about 3.6% of patients screened for syphilis. After implementation, about 24.4%.
- Pregnant women:
 - Pre-intervention testing was 5.9% (272 of 4,579), post-intervention was 49.9% (2,061 of 4,129)
 - Cases went from 2 to 15

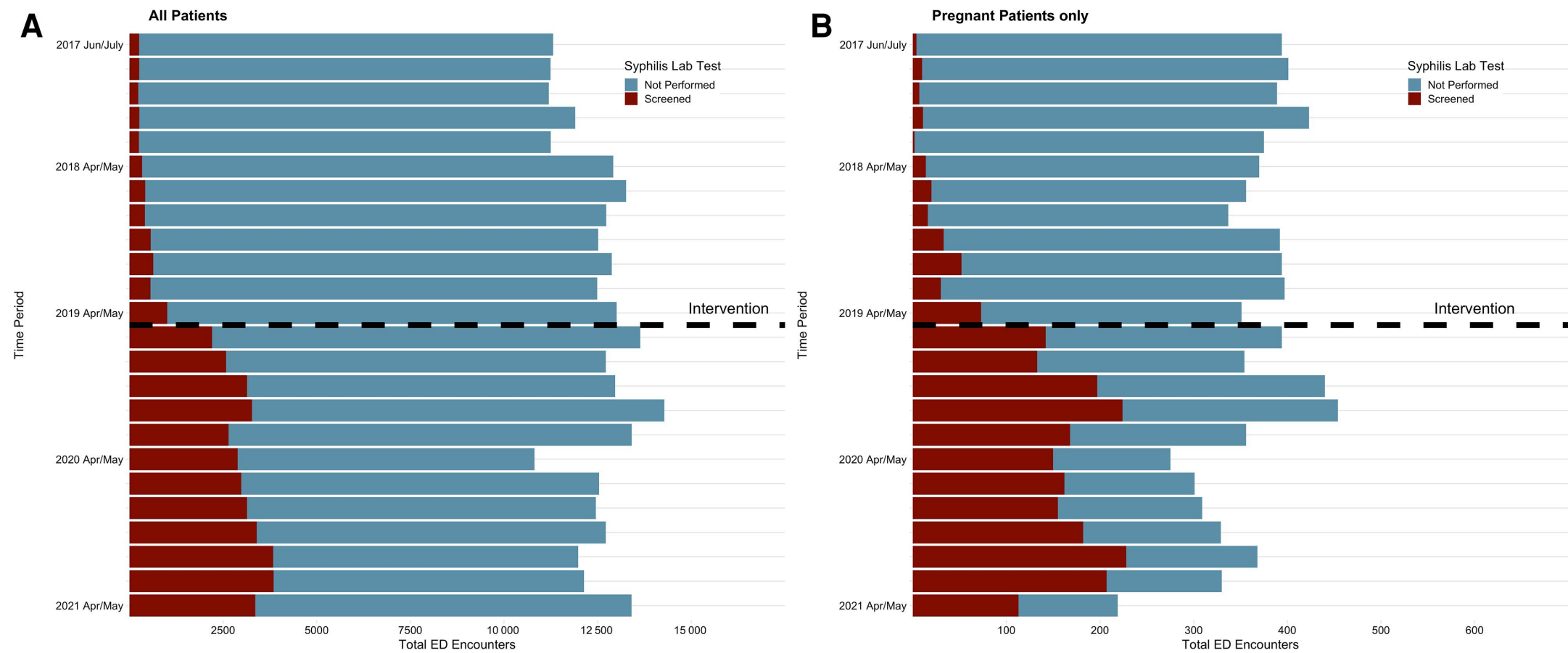
Media Advisory

Tuesday, September 10, 2024

Emergency department screening more than doubles detection of syphilis cases

NIH-supported study shows potential of strategy to reach people with and without symptoms.

Importance of ED screening



Recommendations

- Perform syphilis testing on all patients upon finding a positive pregnancy test
- Test all pregnant patients three times during pregnancy (at initial prenatal visit, again at 28-32 weeks of gestation, and then at delivery)
- Meet people where they are with syphilis testing and treatment outside of settings in which pregnant patients are typically encountered.
 - This could include emergency departments, urgent cares, primary care visits, jail/prison intake, local health departments, community programs, and other addiction services.
- Perform screening and treatment of all sexually active women and their partners for syphilis in counties with [high syphilis rates](#)
- Perform screening and appropriate treatment for those with other risk factors for syphilis (have unprotected sex and do not use condoms or do not use them correctly, have multiple sex partners, have a sex partner who has syphilis and have sex with a partner who has multiple sex partners)
- Treat all pregnant women who are infected with syphilis immediately upon diagnosis, according to their clinical stage of infection. Treatment must be with penicillin G benzathine (Bicillin LA).

Congenital Syphilis is Preventable

Toolkit can be found here:

<https://www.in.gov/health/audiences/clinicians/clinical-guidelines-and-references/congenital-syphilis-clinician-toolkit/>

Includes:

- Dashboards (Adult and Congenital Syphilis)
- Case definitions
- Treatment algorithm
- Clinical staging
- Treatment information



TB INTENSIVE TRAINING COURSE

OCTOBER 2-4

**The Welborn Conference
Center**

412 Mulberry St. Evansville, IN 47713

DETAILS

This three day training, for healthcare providers, will cover TB and LTBI diagnosis and treatment along with other special topics including epidemiology, laboratory testing, and TB in pregnancy. CMEs and CNEs will be provided.

Two options for attendance

- 1 day only, focus on Latent TB Infection
- 3 day comprehensive workshop on TB

REGISTER HERE

<https://www.in.gov/health/idepd/tuberculosis/#Upcoming> On Demand Training

QUESTIONS

Contact alg278@njms.rutgers.edu and
shorrocks@health.in.gov

RUTGERS
New Jersey Medical School
GLOBAL TUBERCULOSIS INSTITUTE

**Indiana
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Health**

**TB Prevention
and Care
Program**

Increase in *Mycoplasma pneumoniae*

- *M. pneumoniae* usually peaks every 3 to 7 years, with variation of strain types contributing to this pattern.
- During fall 2023, started to see a re-emergence globally and since that time, there has been an increase in cases in the U.S.
- *M. pneumoniae* infections are common bacterial respiratory infections that are most common in young adults and school-aged children.
- In 2024 CDC has seen an increase in *M. pneumoniae* infections, including in young children.
 - Keep Mp on the differential diagnosis for respiratory infections, especially if the infection does not respond to beta-lactams
 - Testing can include serology or molecular, such as RVP, if available. IgM can have false positives but if pre-test probability is high it is likely accurate.
- So far macrolide resistance has been low and the patients have been responsive to Azithro

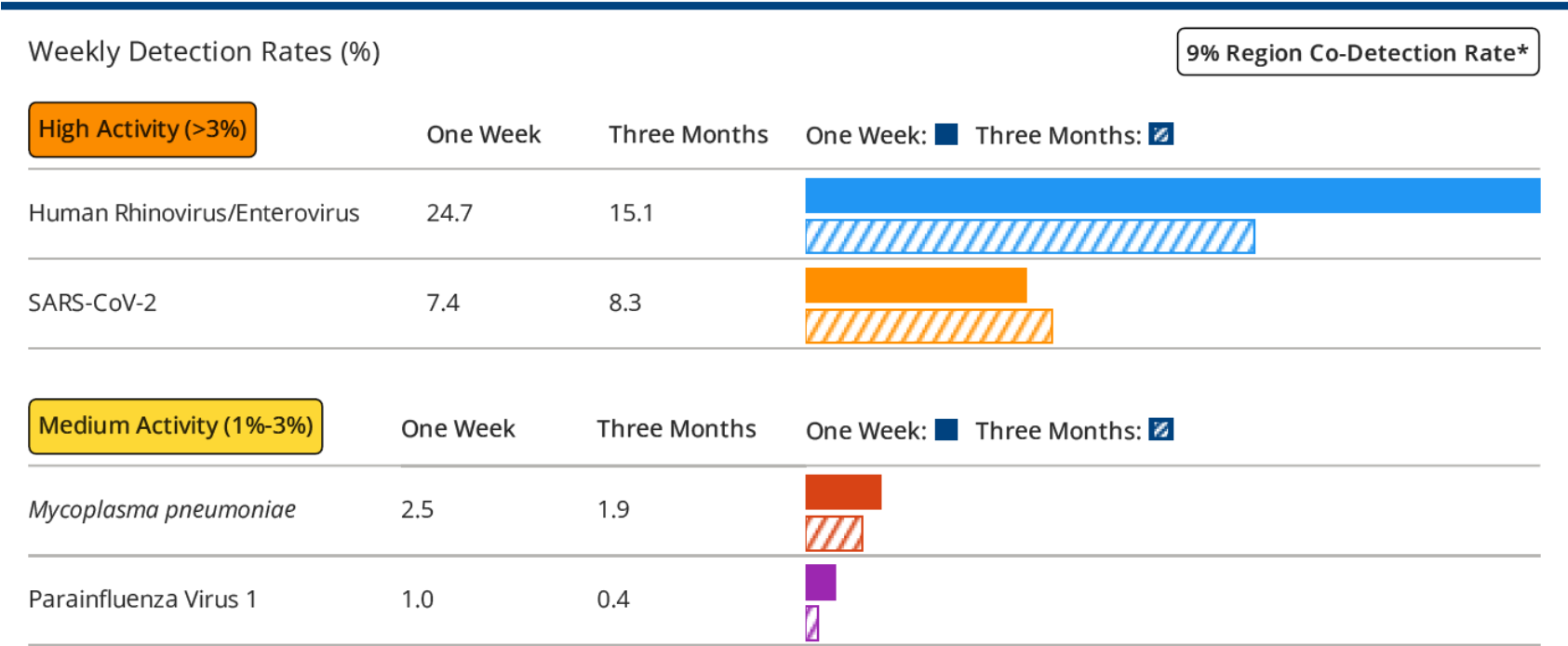
Biofire Respiratory Viral Panel



BIOFIRE® Syndromic Trends

Midwest Region

Respiratory Report
RP2.1



Resources about Mp

1. CDC *Mycoplasma pneumoniae* Infection Surveillance and Trends - <https://www.cdc.gov/mycoplasma/php/surveillance/index.html>
2. Clinical Care of *Mycoplasma pneumoniae* Infection - <https://www.cdc.gov/mycoplasma/hcp/clinical-care/index.html>
3. Laboratory Testing for *Mycoplasma pneumoniae* - <https://www.cdc.gov/mycoplasma/php/laboratories/index.html>
4. Submitting Specimens for *Mycoplasma pneumoniae* Testing - <https://www.cdc.gov/mycoplasma/php/laboratories/specimen-packing.html>
5. MMWR (Notes from the Field): Reemergence of *Mycoplasma pneumoniae* Infections in Children and Adolescents After the COVID-19 Pandemic, United States, 2018-2024 - https://www.cdc.gov/mmwr/volumes/73/wr/mm7307a3.htm?s_cid=mm7307a3_w

**What's
new?**

**Recent Updates from
the CDC**



**Indiana
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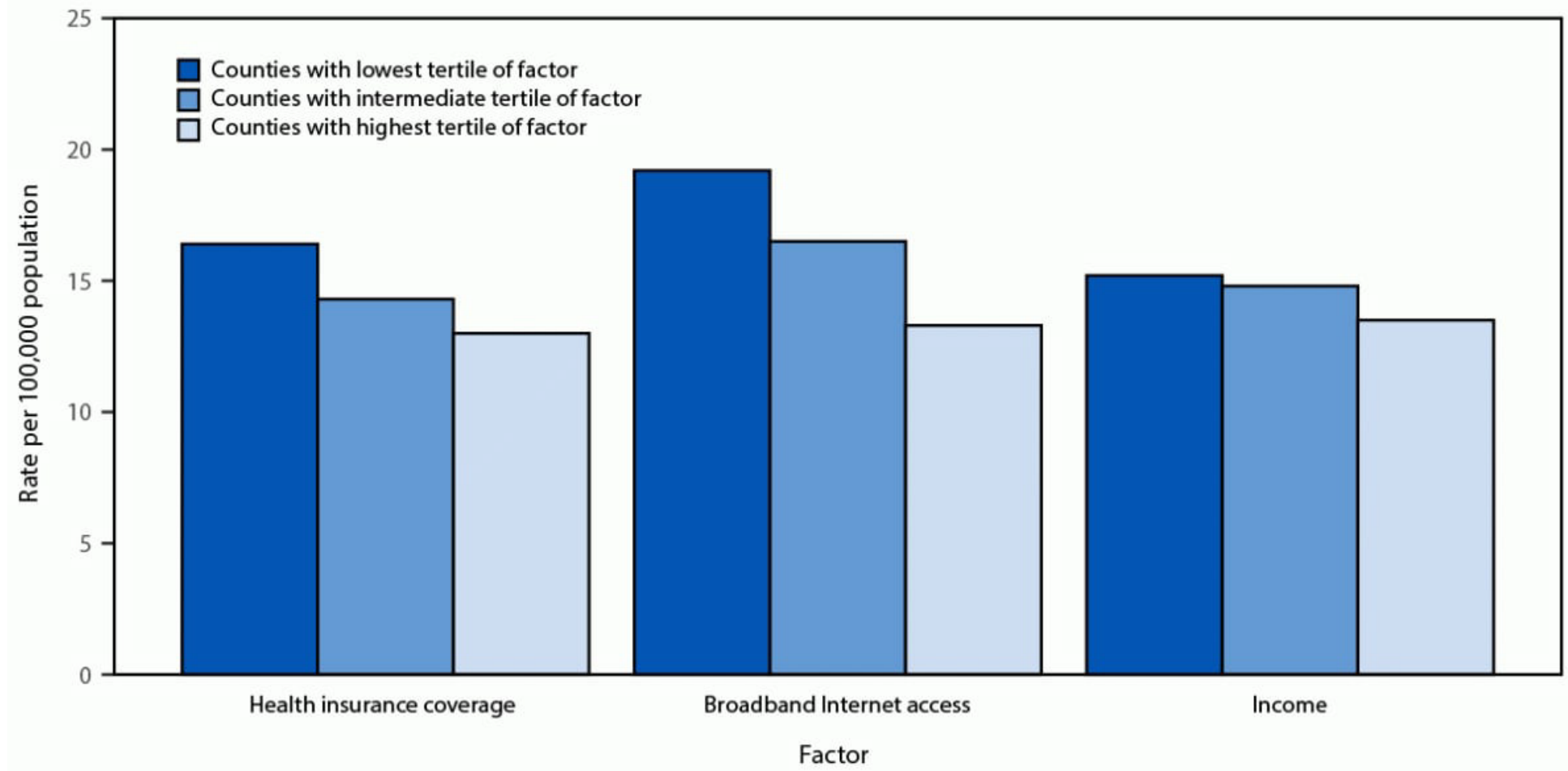
Suicide Rates and County Level Factors



**Indiana
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https://www.cdc.gov/mmwr/volumes/73/wr/mm7337e1.htm?s_cid=mm7337e1_w

Suicide Rates and County Level Factors





Other Public Health Updates



Indiana
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of
Health

Niemann-Pick Disease Type C Treatment

- Miplyffa (arimoclomol)
- Oral medication
- Approved to treat neurological symptoms associated with NPC in adults and children 2 years and older
- NPC is a rare genetic disease that results in progressive neurological symptoms and organ dysfunction

FDA NEWS RELEASE

FDA Approves First Treatment for Niemann-Pick Disease, Type C

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ements

For Immediate Release: September 20, 2024

[FDA approves NPC treatment](#)

Youth E-Cigarette Use Drops

FDA NEWS RELEASE

Youth E-Cigarette Use Drops to Lowest Level in a Decade

Youth Use of Nicotine Pouches Remains Low

[LinkedIn](#) [Email](#) [Print](#)

Among U.S. middle and high school students:

E-cigarette use declined from **7.7%** in 2023 to **5.9%** in 2024



Nicotine pouch use remained low (**1.8%**) in 2024



Youth use of any tobacco product is unsafe

Talk with youth about the harms of tobacco product use



bit.ly/mm7335a3

SEPTEMBER 5, 2024

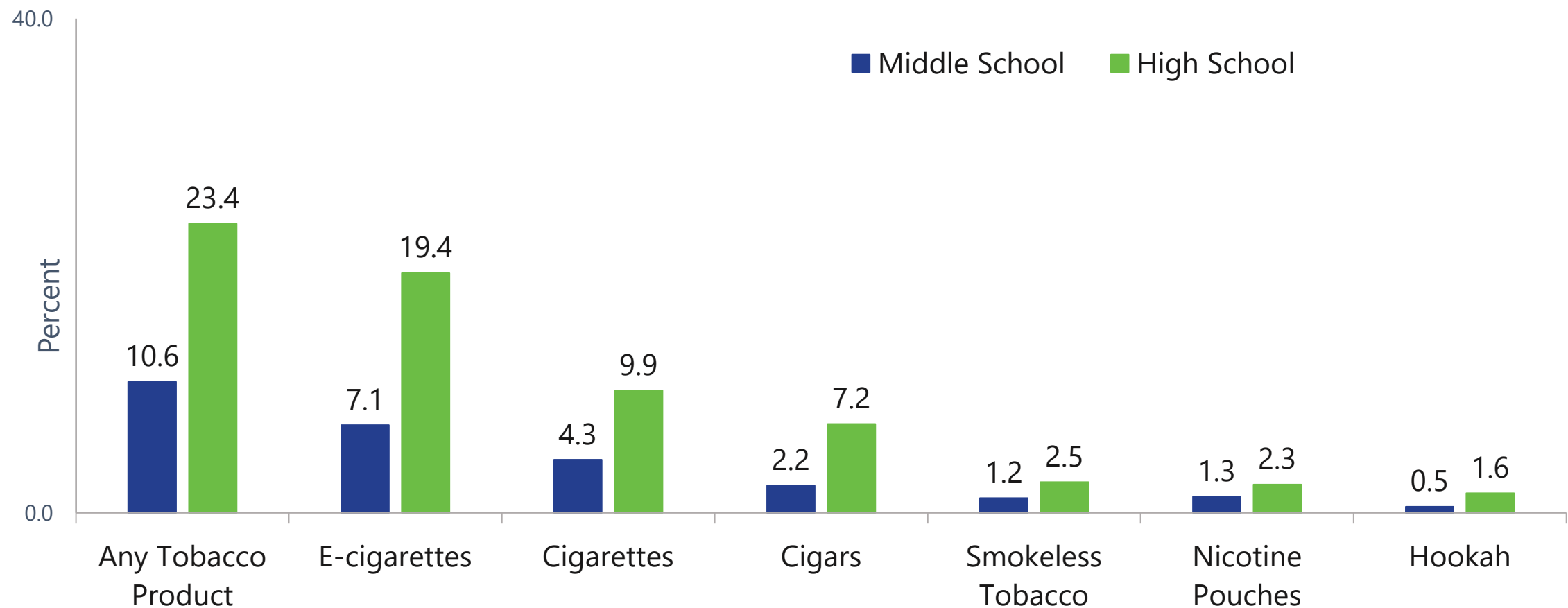
* Data from the 2024 National Youth Tobacco Survey

MMWR

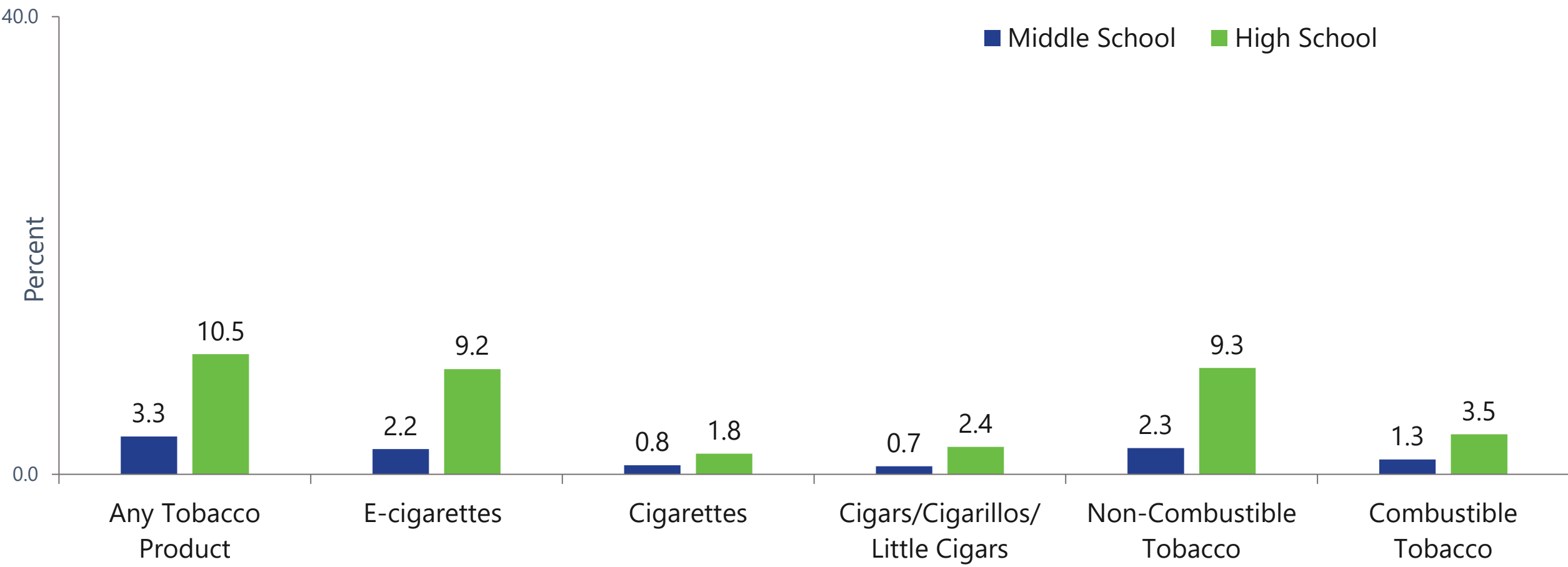


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Ever Use of Tobacco Products Among Middle & High School Students, IYTS 2022



Current Use of Tobacco Products Among Middle & High School Students, IYTS 2022



High School: Current use of cigarettes, cigars, smokeless tobacco, pipe, hookah, snus, dissolvable tobacco, nicotine pouches, e-cigarettes or heated tobacco products.
Middle School: Current use of cigarettes, cigars, smokeless tobacco, pipe, hookah, snus, dissolvable tobacco, nicotine pouches, or e-cigarettes

A state investment in local public health



Your Community Info

Health First Indiana



Indiana
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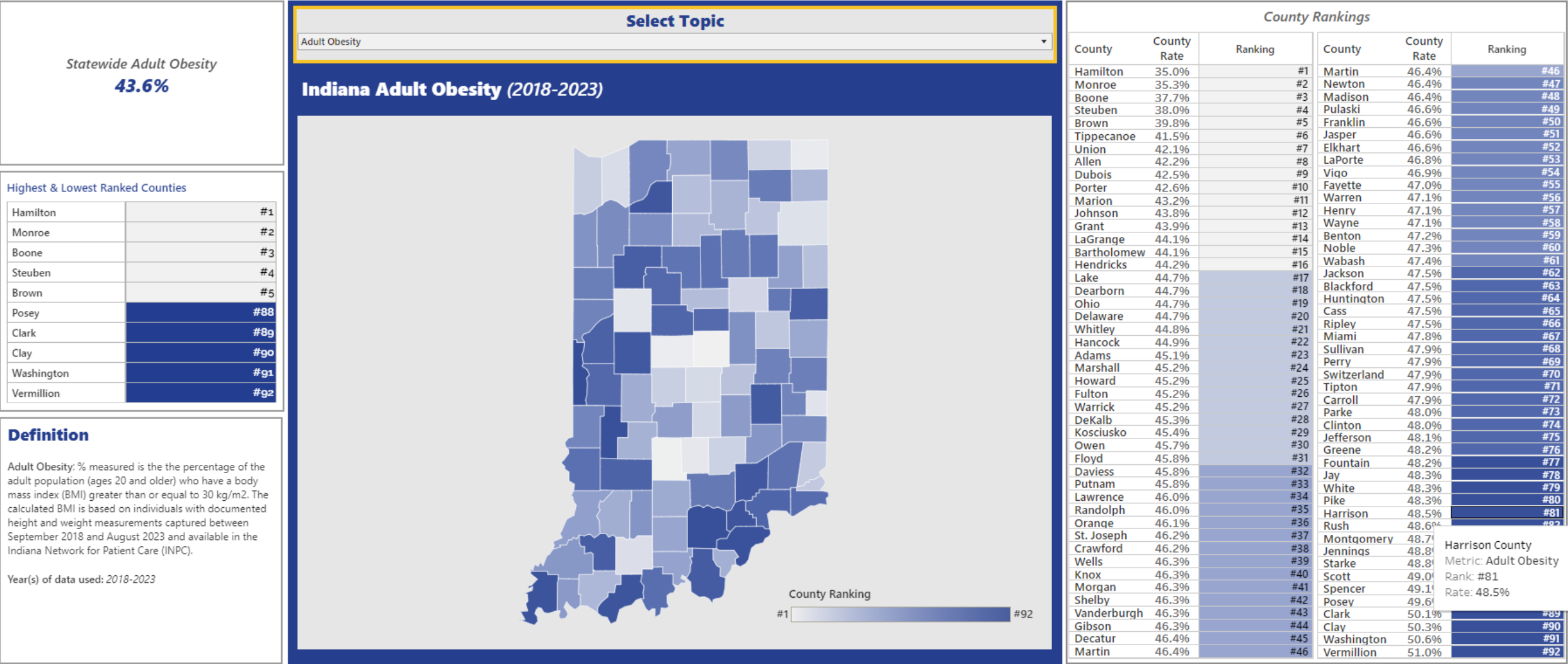
Funding Status

92 local health departments have opted-in for 2025

100% of the State of Indiana will receive
Health First Funding in 2025

County Health Scorecard

<https://www.in.gov/healthfirstindiana/county-health-scorecard/>



Activity Tracker - Statewide

Core Service	Number of Services Provided
Access & Linkage to Clinical Care	23,367
Child & Adult Immunizations	68,953
Childhood Lead Screening & Case Management	11,517
Chronic Disease Prevention & Reduction	40,998
Emergency Preparedness	8,344
Fatality Review (Child, Infant, Fetal, Suicide, Overdose)	16,083
Infectious Disease Prevention and Control	77,971
Maternal and Child Health	70,262
Student Health/School Health Liaison	140,040
Tobacco Prevention and Cessation	32,750
Trauma and Injury Prevention and Education	59,549
Tuberculosis (TB) Prevention and Case Management	40,746
TOTAL NUMBER OF SERVICES PROVIDED	590,580

Pledge to Act

- Indiana Hospital Association (IHA) and Indiana Chamber of Commerce have committed to supporting public health efforts
- Pledge was created as a collaboration between healthcare organizations and businesses to help Hoosiers reach their optimal health
- **All Indiana Hospital Association members have taken the pledge**

Success Story: Lake County

- In partnership with Franciscan Health, Lake County Health Department (LCHD) kicked off a six-week event series called “Walk with a Doc”
- One-hour walking sessions begin with an educational discussion led by a physician or clinical staff member. Participants meet at Lake County Health Department and are eligible for fun incentives.
- First walking session took place on Sept. 12 and featured a kidney specialist. The walks provide an opportunity for attendees to participate in physical activity and ask clinicians questions.



Success Story: Wells County

- Wells CHD hosted free community baby shower on Sept. 5
- New and expecting mothers were eligible for prizes like play packs, strollers, car seats, highchairs, diapers, wipes and baby food. About 80 participants visited each vendor and went home with much-needed baby supplies and food.
- Brought several key partners together to better serve families in Wells County and share information on local resources



Public Health Day at the Statehouse

- Show support for public health by wearing blue and gold

Features:

- A celebration of an investment in public health
- Partnerships in action
- Local Health Department Awards
- Networking and light refreshments
- Save the date: March 12



Call to Action

- Join the conversations about public health funding in your county
- Remember, there is no one-size-fits-all approach
- Remember that better physical and mental health makes Indiana more attractive for families, businesses
- The bottom line: Health First Indiana is **all** of us!



Useful Resources



**Indiana
Department
of
Health**

Safe Sleep ASL videos

[Music/With Closed Captioning](#)
[Music/Without Closed Captioning](#)
[Fully Silent/Without Closed Captioning](#)



Ways to connect with us

- Access our [webpage](#) with resources for clinicians
- Please let us know what topics you'd like us to cover:
Email Gcrowder@health.in.gov
- Sign up for IHAN– Indiana Health Alert Network
<https://ihan-in.org>
- MARK YOUR CALENDARS - Clinician webinars for 2024:
Oct. 25, Nov. 22, Dec. 27

Questions?

CONTACT:

Guy Crowder, M.D., M.P.H.T.M.
Chief Medical Officer

GCrowder@health.in.gov

Next call: Noon, October 25

