



Tiny Shots, Big Protection: Childhood Immunizations

Continuing Pharmacy Education

Jennifer Goodwin, PharmD, PGY1, Advocate Illinois Masonic Medical Center

Delilah Velez, PharmD, PGY1, Advocate Illinois Masonic Medical Center

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Disclosures

The planner(s) and speaker(s) have indicated that there are no relevant financial relationships with any ineligible companies to disclose.

Learning Objectives

At the end of this session, learners should be able to:

- Outline the role of childhood immunizations in preventing disease and protecting public health
- Apply updated childhood immunization recommendations to patient care
- Recognize current misconceptions and misinformation regarding childhood immunization
- Identify effective communication strategies for addressing vaccine hesitancy in parents and caregivers

Outline



Immunization for Individual and Public Health

Current Childhood Vaccine Recommendations

Vaccine Misinformation

Pharmacist and Technician Role

Abbreviation Key

- ACIP: Advisory Committee on Immunization Practices
- AAP: American Academy of Pediatrics
- APC: Antigen Presenting Cell
- CDC: Center for Disease Control and Prevention
- HHS: Health and Human Services

Abbreviation Key

- DTaP/TDaP: Diphtheria, Tetanus, Pertussis
- Hib: Haemophilus Influenzae Type B
- Hep A: Hepatitis A
- Hep B: Hepatitis B
- HPV: Human Papillomavirus
- MMR: Measles, Mumps, Rubella
- PCV: Pneumococcal Conjugate Vaccine (PCV13, PCV15)
- PPSV: Pneumococcal Polysaccharide Vaccine (PPSV23)
- RSV: Respiratory Syncytial Virus

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Prenatal Development

Immune System Formation

- Fetal immune system is exposed to maternal antigens
- Fetus develops immune tolerance

Maternal IgG Antibody Transfer

- Cross the placenta at ~20 weeks and peak at birth
- Recognize and neutralize pathogens
- Provide passive immunity during first few months of life

Maternal Vaccination

- Increases neonatal protection by boosting antibody transfer
- Provides passive immunity until infant vaccine series begins

Pediatric Development

Birth

- Maternal IgG antibodies provide passive immunity
- Maternal antibodies can interfere with early vaccine response

Infancy

- Maternal antibodies decline, reducing passive protection
- Infant adaptive immunity matures, allowing initiation of the primary vaccine series

Childhood

- Series and booster doses spaced strategically
- Builds immunologic memory

Vaccine Timing in Early Childhood

What Parents Often Notice

- "My baby looks healthy"
- "There aren't many disease around anymore"
- "Why do we need vaccines early?"

What the Immune System is Actually Doing

- Early immunity is temporary
- Protection builds step-by-step as the immune system develops
- Vaccines are timed to protect during periods of highest vulnerability

Why Community Protection Matters for Public Health

- Infants cannot rely on individual immunity alone
- When vaccination rates decline, these immune gaps become exposed
- Protection early in life depends on reduced disease circulation

Herd Immunity

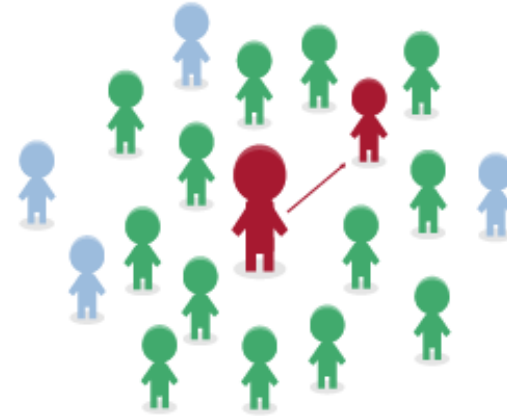
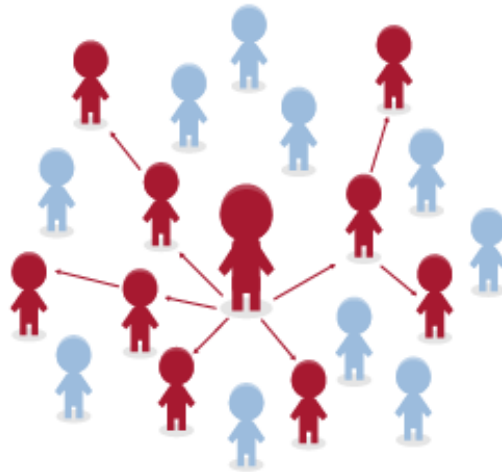
Occurs when a certain proportion of the population is immune to a disease via vaccination or past infection

Limits disease transmission, preventing outbreaks

Protects vulnerable individuals who cannot be vaccinated

Immunity thresholds vary by disease

Herd Immunity



No herd immunity

Herd immunity achieved

● Susceptible ● Infected ● Immune → Disease transmission

Source: GAO adaptation of NIH graphic. | GAO-20-646SP

Herd Immunity and Pediatrics

Neonates and infants rely on herd immunity for protection

School-aged children are major transmission reservoirs

High vaccination rates in this age group strongly supports public health

Small declines in pediatric coverage can lead to disease resurgence

Herd Immunity and Pediatrics

Neonatal Infection Protection

Herd immunity is built through consistent, widespread vaccination

High vaccine

health

Small declines in coverage can lead to disease resurgence

Vaccination Rates

78% of U.S. counties report decline in childhood vaccination rates

Average county-level coverage decreased from 93.9% → 91.3%

Consequences of Declining Rates:

Measles

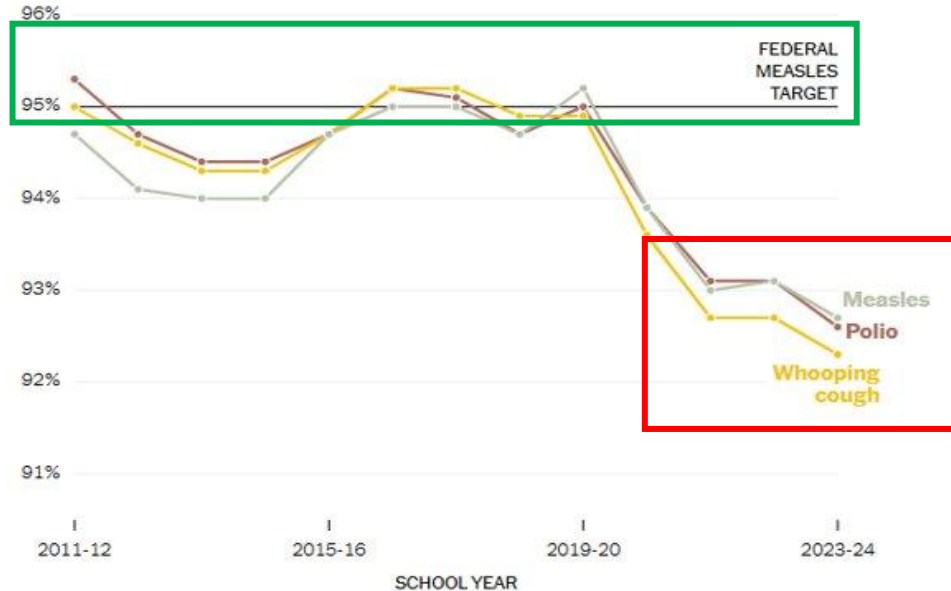
- 2,144 U.S. cases in 2025 (>30 year high)

Influenza

- 2024-25 season hospitalization peak (>10 year high)

Vaccination Rates

Share of U.S. kindergartners vaccinated against ...



Source: Centers for Disease Control and Prevention.

Vaccination Rates

Share of U.S. kindergartners vaccinated ages 5-16

When vaccination rates decline
preventable diseases return

| SCHOOL YEAR | Share of U.S. kindergartners vaccinated ages 5-16 |
|-------------|---------------------------------------------------|
| 2011-12 | 92% |
| 2012-13 | 91% |
| 2013-14 | 90% |
| 2014-15 | 89% |
| 2015-16 | 88% |
| 2016-17 | 87% |
| 2017-18 | 86% |
| 2018-19 | 85% |
| 2019-20 | 84% |
| 2020-21 | 83% |
| 2021-22 | 82% |
| 2022-23 | 81% |
| 2023-24 | 80% |

SOURCE: CENTERS FOR DISEASE CONTROL AND PREVENTION

Source: Centers for Disease Control and Prevention

Knowledge Check #1

Which statement best describes the role of childhood immunizations in public health?

- A. They primarily protect only the vaccinated individual
- B. They limit disease spread and protect vulnerable populations
- C. They reduce disease severity but not transmission
- D. They eliminate the need for infection-related precautions

Outline



Immunization for Individual and Public Health

Current Childhood Vaccine Recommendations

Vaccine Misinformation

Pharmacist and Technician Role

Who Makes Vaccine Recommendations

Vaccine Recommendation Bodies

ACIP

Provides
recommendations
to CDC

CDC

Implements
recommendations
into official policy

AAP

Publishes
recommendations
independently

Vaccine Recommendation Bodies

ACIP Update: 30 professional organization representatives no longer participate in workgroups to contribute their expertise

ACIP
Professional
representatives
to CDC

ACIP
recommendations
independently

Current CDC Vaccine Recommendations

Early Childhood Vaccines

| VACCINE OR PREVENTIVE ANTIBODY | BIRTH | 1 MONTH | 2 MONTHS | 4 MONTHS | 6 MONTHS | 7 MONTHS | 8 MONTHS | 12 MONTHS | 15 MONTHS | 18 MONTHS | 19 MONTHS | 20-23 MONTHS | 2-3 YEARS | 4-6 YEARS |
|--------------------------------|-------------------------------------------------|---------|----------|----------|----------|----------|----------------------------------|-----------|-----------|-----------|-----------|--------------|-----------|-----------|
| RSV antibody | Depends on mother's RSV vaccine status | | | | | | Depends on child's health status | | | | | | | |
| Hepatitis B | Dose 1 | Dose 2 | | | Dose 3 | | | | | | | | | |
| Rotavirus | | Dose 1 | Dose 2 | Dose 3 | | | | | | | | | | |
| DTaP | | Dose 1 | Dose 2 | Dose 3 | | | | Dose 4 | | | | | | Dose 5 |
| Hib | | Dose 1 | Dose 2 | Dose 3 | | | Dose 4 | | | | | | | |
| Pneumococcal | | Dose 1 | Dose 2 | Dose 3 | | | Dose 4 | | | | | | | |
| Polio | | Dose 1 | Dose 2 | Dose 3 | | | | | | | | Dose 4 | | |
| COVID-19 | At least 1 dose of the current COVID-19 vaccine | | | | | | | | | | | | | |
| Influenza/Flu | Every year. Two doses for some children | | | | | | | | | | | | | |
| MMR | | | | | | | | Dose 1 | | | | | | Dose 2 |
| Chickenpox | | | | | | | | Dose 1 | | | | | | Dose 2 |
| Hepatitis A | | | | | | | 2 doses separated by 6 months | | | | | | | |



ALL children should be immunized at this age



SOME children should get this dose of vaccine or preventive antibody at this age



Parents/caregivers should talk to their health care provider to decide if this vaccine is right for their child



Early Childhood Vaccine Changes

| Vaccines | CDC 2024 | CDC 2025/2026 |
|-----------------|------------------|-------------------------------------------------|
| Hepatitis B | + | high-risk group or shared decision-making |
| Hepatitis A | + | |
| Rotavirus | + | shared decision-making |
| COVID-19 | + | shared decision-making |
| Influenza | + | shared decision-making |
| MMR + Varicella | single injection | two split injections |
| RSV | palivizumab | nirsevimab |

Early Childhood Vaccine Changes

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| Rotavirus | + | shared decision-making |
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| Influenza | + | shared decision-making |
| MMR + Varicella | single injection | two split injections |
| RSV | palivizumab | nirsevimab |

Recent Updates to Vaccine Recommendations

RSV

Background

- Palivizumab (Synagis): Previous standard of care, discontinued 12/31/2025
- Abrysvo: Maternal RSV vaccine, given at 32-36 weeks gestation Sep-Jan

AAP Recommendation

- **Nirsevimab or Clesrovimab** for infants **<8 months entering first RSV season** if:
 - Mother unvaccinated or status unknown
 - Infant born <14 days after maternal vaccination
- **Nirsevimab** for **high-risk children 8-19 months entering second RSV season**

CDC Recommendation

- Follows same recommendations as AAP

Wilkins, D., Wählby Hamrén, U. et al. 2024.

Study Design

- Randomized, palivizumab-controlled phase 2/3 study

Population

- High-risk infants: preterm, CLD, or CHD
- N=925

Treatment Arms

- Nirsevimab (N=614): 1 dose + 4 monthly placebo injections
- Palivizumab (N=304): 5 monthly doses

Outcomes

- RSV nAbs at days 31, 151, and 361 post dose

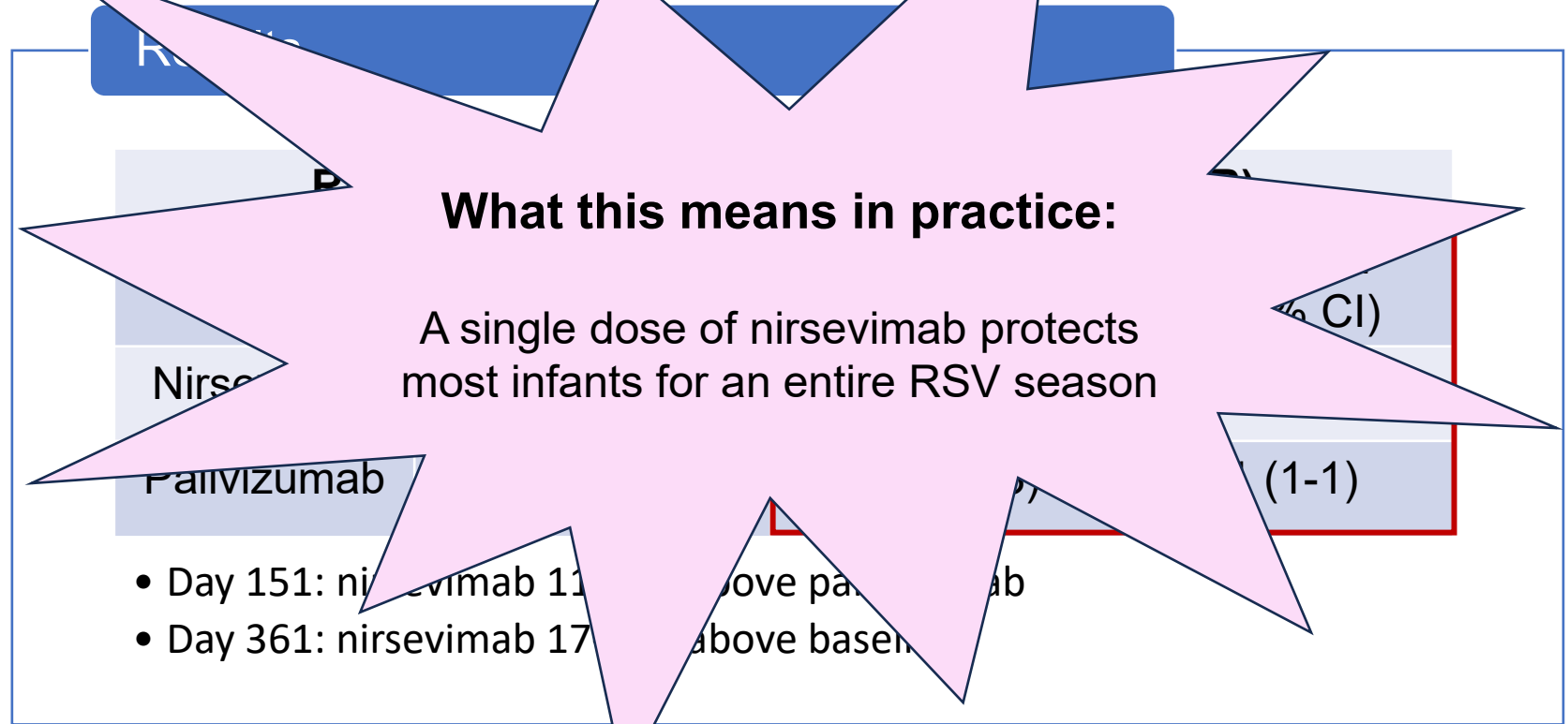
Wilkins, D., Wählby Hamrén, U. et al. 2024.

Results

| RSV nAb Geometric Mean Fold Rise (GMFR) | | | |
|-----------------------------------------|--------------------|---------------------|---------------------|
| | Day 31 (95% CI) | Day 151 (95% CI) | Day 361 (95% CI) |
| Nirsevimab | 334 (282-395) | 116 (103-130) | 17 (15-19) |
| Palivizumab | 7 (5-9) | 11 (9-13) | 1 (1-1) |

- Day 151: nirsevimab 11-fold above palivizumab
- Day 361: nirsevimab 17-fold above baseline

Wilkins, D., Wählby Hamré U. et al. 2024.



Hepatitis B Vaccine

Background

- Universal Hep B birth dosing since 1991 reduced pediatric infections by 99%
- Infants are highly vulnerable: ~90% infected at birth develop chronic infection

AAP Recommendation

- Dose 1: At birth
- Dose 2: ≥ 4 weeks after dose 1
- Dose 3: ≥ 8 weeks after dose 2 and ≥ 16 weeks after dose 1

CDC Recommendation

- Infants born to hepatitis B-negative mothers:
 - Shared decision making; initial dose >2 months old
- Infants born to hepatitis B-positive or unknown-status mothers:
 - Vaccine within 12 hours of birth

Lee, et al. 2006.

Study Design

- Systematic review and meta-analysis

Population

- Newborns of HBsAg-positive mothers who received the vaccine within 1 month of life
- 5 trials (N = 403)

Treatment Arms

- Plasma derived or recombinant vaccine (N = 252)
- Placebo (N = 151)

Outcomes

- Occurrence of hepatitis B infection

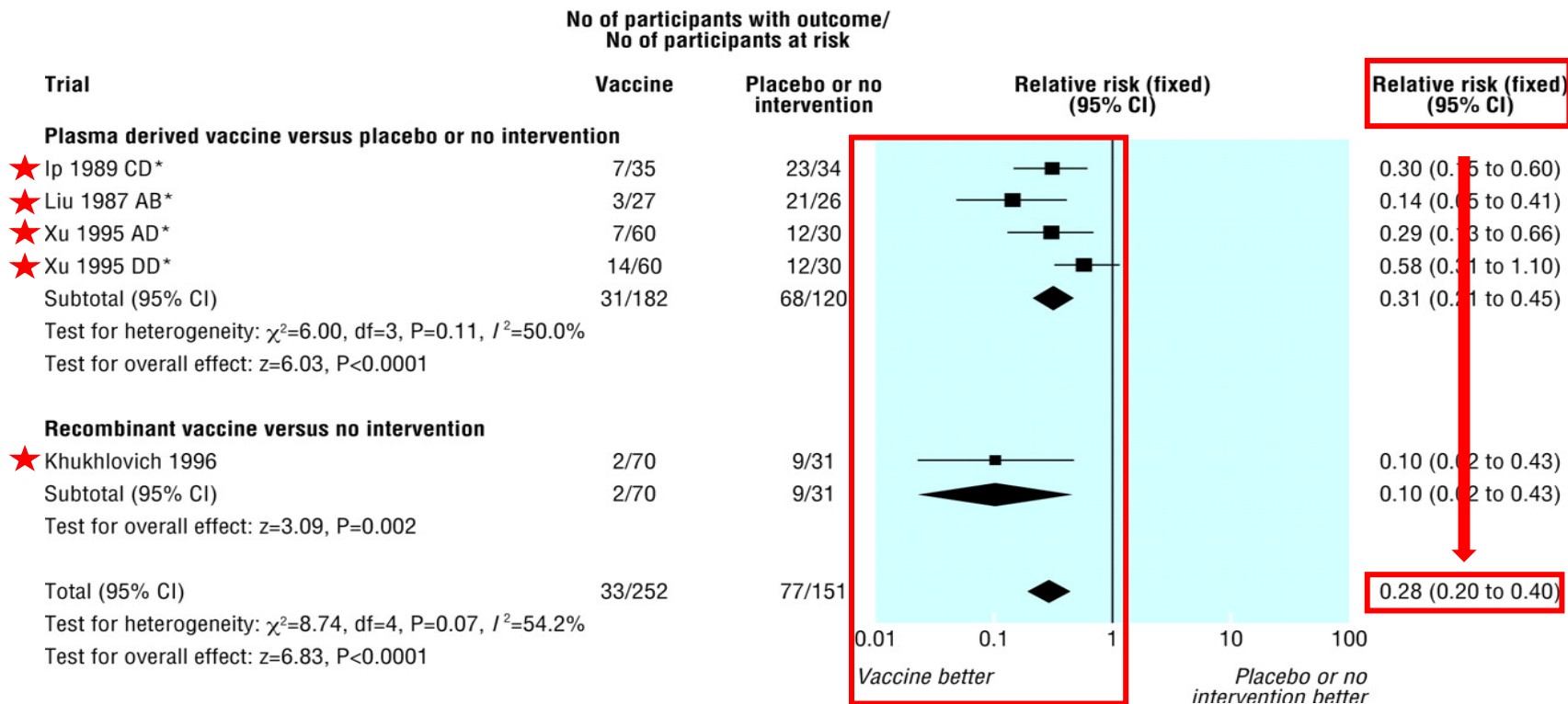
Lee, et al. 2006.

Results

Hepatitis B Occurrence

| | Vaccine (%) | Placebo (%) | Relative Risk (95% CI) |
|-----------------------------------|-------------|-------------|------------------------|
| Plasma Derived Vaccine (4 trials) | 31/182 (17) | 68/120 (57) | 0.31 (0.21-0.45) |
| Recombinant Vaccine (1 trial) | 2/70 (2) | 9/31 (29) | 0.10 (0.02-0.43) |
| All Trials Combined | 33/252 (13) | 77/151 (51) | 0.28 (0.20-0.40) |

Lee, et al. 2006.



Lee, et al. 2006.

No... with outco...
at ris...

Trial

Plasma derived vaccine versus

- ★ Ip 1989 CD*
- ★ Li...
- ★ Xu 1995
- ★ Xu 1995 DD

Subtotal (95% CI)

Test for heterogeneity: $\chi^2=$

Test for overall effect: $z=$

Recomb...

- ★ Kh...

Subtotal (95% CI)

Test for overall effect: $z=3.09, P=0.002$

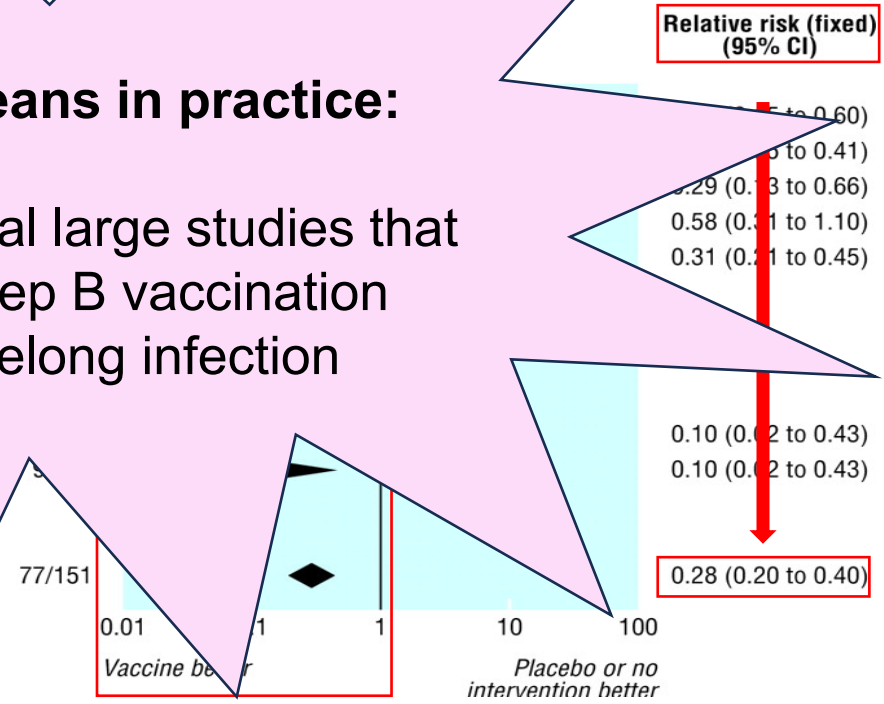
Total (95% CI)

Test for heterogeneity: $\chi^2=8.74, df=4, P=0.07, I^2=54.2\%$

Test for overall effect: $z=6.83, P<0.0001$

What this means in practice:

There are several large studies that show early Hep B vaccination prevents lifelong infection



COVID-19 Vaccine

Background

- COVID-19 continues to circulate with seasonal surges
- Vaccination reduces risk of severe disease and hospitalization

AAP Recommendation

- All children 6–23 months
- Children 2–18 years in the following groups: high risk/living with someone high risk for severe infection, long-term care settings, previously unvaccinated

CDC Recommendation

- Children ≥ 6 months based on shared clinical decision-making

Muñoz FM, et al. 2023.

Study Design

- Randomized, placebo-controlled, phase 2/3 trial

Population

- Healthy children stratified into 6-23 months and 2-4 years
- N = 1254

Treatment Arms

- COVID-19 Vaccine (N=873): 3 dose series
- Placebo (N = 381): 3 placebo injections

Outcomes

- First occurrence of COVID-19 after Dose 3

Muñoz FM, et al. 2023.

Results

| COVID-19 Infections | | | |
|----------------------|--------------------|--------------------|--------------------------------|
| | Vaccine (N=873) | Placebo (N=381) | Vaccine Efficacy % (95% CI) |
| 6 months to 4 years | 13 | 21 | 73.2 (43.8-87.6) |
| 6 months to <2 years | 4 | 8 | 75.8 (9.7-94.7) |
| 2 to 4 years | 9 | 13 | 71.8 (28.6-89.4) |

- Most common adverse events: mild-moderate fever/fatigue
- Few participants withdrew owing to events
- No deaths occurred

Muñoz FM, et al 2023.

Res

What this means in practice:

Vaccination against COVID-19 reduces symptomatic infection with acceptable short-term safety profile

6 months

6 m

2 to 4 years

- Most common adverse events: mild to moderate fever, fatigue
- Few participants with symptoms following to events
- No deaths occurred

MMR/Varicella

Background

- Available as a combined measles, mumps, rubella, and varicella or separate
- Recent discussion regarding separation due to evidence relating to febrile seizures

AAP Recommendation

- Continues to recommend giving families the choice between MMRV and MMR+V

CDC Recommendation

- Varicella as standalone rather than combination for children ages <4 years

Klein, NP. et al. 2010.

Study Design

- Comparative study using Vaccine Safety Datalink data from 2000-2008

Population

- Children aged 12-23 months that received first dose of either MMRV or MMR+V
- N=459,461

Treatment Arms

- MMRV (N=83,107)
- MMR+V, both vaccines given in same visit (N=376,354)

Outcomes

- Seizure occurrence, days 7-10 after vaccination

Klein, NP. et al. 2010.

Results

| Post-Vaccination Seizures, days 7-10 | | | | |
|---------------------------------------|--------------------|----------------------|----------------------|---------|
| | MMRV (n=83,107) | MMR+V (n=376,354) | RR (95% CI) | P-value |
| Seizures | 77 (0.093%) | 174 (0.046%) | | <0.001 |
| Seizure Risk After Vaccination* | | | 1.98 (1.43- 2.73) | <0.001 |

Klein, NP. et al 2010.

What this means in practice:

Both MMRV and MMR+V are effective; separation slightly lowers seizure risk

Seiz

.001

After
Vaccination*

<0.001

Considerations of MMRV Data

Seizure Risk

- Febrile seizures occur in ~2-5% of children (CDC)
- Data included children with concurrent febrile illnesses
- Seizures occurred in <1% of children in either group

Vaccines for Children (VFC) Program

- ~50% of US children qualify for the VFC program
- Due to seizure risk, CDC suggests MMR+V for those <4 years old
- CDC voted to continue MMRV use in the same age group within the VFC program

Evidence & Practice Change

| Vaccines | Clinical Evidence Supports | Implement Into Practice |
|-------------------|-----------------------------------------------------------------------|-----------------------------------|
| RSV | Single dose vaccination protects for whole season | Prevention for most infants |
| Hep B | Meta-analysis showed decrease in HBV and near elimination in children | Early-life vaccination supported |
| COVID-19 | Showed efficacy and favorable safety profiles | Routine, recommended immunization |
| MMR/ Varicella | Combination vaccines studies indicated increased febrile seizure risk | Shared-decision making |

Knowledge Check #2

Which of the following statements is incorrect?

- A. A single dose of RSV immunization provides protection for the RSV season in most infants
- B. Early hepatitis B vaccination is supported by evidence, showing decreased infection and near elimination in children
- C. COVID-19 vaccines have demonstrated efficacy and favorable safety profiles
- D. Combination MMR/Varicella vaccines have not been associated with any increased risk of febrile seizures and do not require shared decision-making

Outline



Immunization for Individual and Public Health

Current Childhood Vaccine Recommendations

Vaccine Misinformation

Pharmacist and Technician Role

Vaccine Misinformation

Children receive too many vaccines

Vaccines cause autism

Children do not need Hep B vaccination before age 12 since it is transferred sexually

Vaccine Misinformation

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Number of Pediatric Vaccines

Misinformation:

- U.S. Department of Health and Human Services (HHS) claim children receive **~92 vaccines** to be compliant with CDC schedule

Fact:

- Children who follow the CDC's recommended immunization schedule get **~30 vaccines**

Extended Spacing of Vaccines

Benefits:

- Easier assessment of adverse reactions
- Willingness of parents to vaccinate their children
- Some immunity is better than none

Risks:

- Lack of adequate immunity
- More appointments/higher costs
- Repeated stress for additional vaccine visits
- Mixed messages regarding importance of scheduling

Vaccine Misinformation

Children receive too many vaccines

Vaccines cause autism

Children do not need Hep B vaccination before age 12 since it is transferred sexually

Vaccines and Autism

Misinformation:

- Stemmed from a now-discredited 1998 study alleging a MMR—Autism link

Fact:

- 2019 study found no link between vaccines and autism
- Study included > 650,000 children born from 1999-2010

Disproval of 1998 MMR-Autism Trial

Diagnoses

- 3 of 9 children reported with regressive autism never received autism diagnosis

Pre-existing conditions

- All 12 children were reported "previously normal"
- 5 children had pre-existing behavioral concerns

Recruitment and funding

- Case series of only 12 children
- Participants were recruited through anti-MMR campaigners
- Commissioned and funded for planned litigation

Hviid, A. et al. 2019.

Study Design

- Nationwide cohort study

Population

- Children born in Denmark from 1999—2010
- N=657,461

Treatment Arms

- MMR-vaccinated children (N=625,842)
- MMR-unvaccinated children (N=31,619)

Outcomes

- Diagnosis of autism

Hviid, A. et al. 2019.

Results

| Diagnosis of Autism | | | | |
|---------------------|---------------------------|----------------------------|--------------------------|---------|
| | Vaccinated (N=625,842) | Unvaccinated (N=31,619) | Hazard Ratio (95% CI) | P-value |
| All children | 5992 (0.96%) | 525 (1.66%) | 0.93 (0.85-1.02) | 0.085 |
| Male | 4844 | 416 | 0.97 (0.87-1.08) | |
| Female | 1148 | 109 | 0.79 (0.64-0.97) | |

Vaccines and Autism

Autism and Vaccines

QUESTIONS AND CONCERNS | PAGE 2 OF 9 | ALL PAGES ↓



For Everyone

NOV. 19, 2025

CDC updates from
11/19/25

KEY POINTS

- The claim "vaccines do not cause autism" is not an evidence-based claim because studies have not ruled out the possibility that infant vaccines cause autism.
- Studies supporting a link have been ignored by health authorities
- HHS has launched a comprehensive assessment of the causes of autism, including investigations on plausible biologic mechanisms and potential causal links.

Vaccine Misinformation

Children receive too many vaccines

Vaccines cause autism

Children do not need Hep B vaccination before age 12 since it is transferred sexually

Hepatitis B

Misinformation:

- Hep B is only sexually transmitted
- Hep B vaccination should be delayed to 12 years of age

Fact:

- Hep B can be transmitted through bodily fluids, blood, sexually, or perinatally
- Crucial to vaccinate in infancy
- Can be transmitted during birth or by other positive contacts

Hepatitis B



"Silent Killer"— 50-90% of adults with chronic infection are unaware of status until serious liver damage develops



Children are particularly susceptible and more likely to develop chronic infection that leads to life-threatening complications



CDC's update states that birth dose should only be given if mother is Hep B positive or status unknown

Hepatitis B



are

CDC's update from 1/6/26:

infection
develops



**If mother Hep B negative: shared-
decision making & first dose delayed
until ≥ 2 months old**

likely to



**CDC's update states that birth dose should only be given
if mother is Hep B positive or status unknown**

Knowledge Check #3

Which of the following statements is true?

- A. Children receive 92 vaccines to complete the age 0-18 CDC immunization schedule
- B. Hepatitis B infections are exclusively sexually transmitted
- C. Extended spacing of vaccinations does not affect the body's mounted immune response
- D. A large trial from 1998 linking the MMR vaccine to autism was discredited and proven fraudulent

Outline



Immunization for Individual and Public Health

Current Childhood Vaccine Recommendations

Vaccine Misinformation

Pharmacist and Technician Role

Reframing the Narrative

- The AAP and FrameWorks Institute recommend vaccine messaging that emphasizes collective health and prevention
- Key framing strategies:

Community protection (schools, childcare, parks)

Vaccination supports long-term health and prevention

Immunization is a routine public health practice

Vaccine Hesitancy

Differentiate between hesitancy and access issues

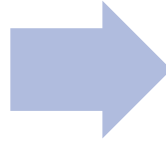


Target interventions based on barrier

Use Intentional Language

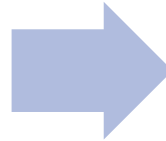
- Use language that emphasizes prevention and preparedness

Fight disease



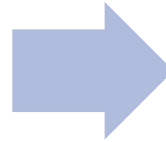
Prepare the immune system

Decision to vaccinate



Manage routine vaccinations

Herd immunity



Community immunity

Pharmacist and Technician Role

- Vaccine promotion and misinformation prevention are public health responsibilities
- Key roles:

Assess
immunization status

Provide evidence-
based
recommendations

Serve as a trusted
source of vaccine
information

Knowledge Check #4

Which approach is most likely to reduce misunderstanding during vaccine discussions?

- A. Emphasizing disease severity using fear-based messaging
- B. Correcting misinformation by labeling it as disinformation
- C. Focusing only on individual vaccine decisions
- D. Using neutral, preparation-oriented language

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Questions?

Jennifer Goodwin
jennifer.morrow@aah.org

Delilah Velez
delilah.velez@aah.org