

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



# Adult Respiratory Syncytial Virus (RSV) Session

**Camille Kotton, MD**

**Chair, Adult RSV Work Group**

Advisory Committee on Immunization Practices (ACIP)

June 26, 2024

# Adult RSV Work Group Membership

## ACIP Voting Members

Camille Kotton (Chair)  
Keipp Talbot  
Sarah Long  
Albert Shaw

## Ex Officio Members

Rachel Zhang (FDA)  
Nicholas Geagan (FDA)  
Nadine Peart Akindele (FDA)  
Sonnie Kim (NIH/NIAID)  
Michelle Juaneza (HRSA)  
Uzo Chukwuma (IHS)

## CDC Co-Leads

Michael Melgar  
Amadea Britton

## Liaisons

Kenneth Schmader (AGS)  
Vidya Sundareshan (ACP)  
Gretchen LaSalle (AAFP)  
April Killikelly (NACI, PHAC)  
Winnie Siu (NACI, PHAC)  
Katherine Williams (APTR)  
Ruth Lynfield (NFID)  
Bindy Crouch (AIM)  
Steven Pergam (IDSA)  
Elizabeth Skoy (APhA)

## Consultants

Robert Atmar (Baylor College of Medicine)  
Doug Campos-Outcalt (University of Arizona)  
Helen Chu (University of Washington)  
Peter Donofrio (Vanderbilt University)  
Marie Griffin (Vanderbilt University)  
Rebecca Morgan (Case Western Reserve University)  
Cynthia Lucero-Obusan (Veterans Health Administration)  
Tracy Ruckwardt (NIH/NIAID)  
Jonathan Temte (University of Wisconsin)

# Transition in Work Group Chair

- Dr. Albert Shaw will be the incoming work group chair for the adult RSV work group starting in July 2024.



# CDC Contributors

## Coronavirus and Other Respiratory Viruses Division

Melissa Coughlin	Danielle Moulia
Fatima Dawood	Ismael Ortega-Sanchez
Katherine Fleming-Dutra	Lakshmi Panagiotakopoulos
Kristen Folsom	Pragna Patel
Jarrett Gartin	Monica Patton
Monica Godfrey	Amanda Payne
Aron Hall	Derrell Powers
Fiona Havers	Mila Prill
Michele Hlavsa	Lauren Roper
Jefferson Jones	Diya Surie
Ruth Link-Gelles	Natalie Thornburg
Agustin Lopez	Raigan Wheeler
Josephine Mak	Megan Wallace
Meredith McMorrow	Trang Wisard
Noelle-Angelique Molinari	

## Immunization Safety Office

Anne Hause  
Pedro Moro  
Christine Olson  
David Shay  
Karen Broder  
John Su  
Eric Weintraub  
Michael McNeil  
Julianne Gee

## Influenza Division

Jill Ferdinands  
Lisa Grohskopf

## Immunization Services Division

Carla Black  
Kayla Calhoun  
Jennifer Kriss  
James Singleton  
Nicole Dowling  
Andrew Leidner  
Jamison Pike  
James Singleton  
Patricia Wodi

## NCIRD Office of the Director

Hannah Rosenblum  
Melinda Wharton  
Jessica MacNeil

# CDC and ACIP currently recommend that adults aged 60 years and older may receive a single dose of RSV vaccination, using shared clinical decision-making.

- ACIP voted on this recommendation at the June 2023 meeting
- Recommendation is not product-specific; administer whichever vaccine is available



# Recap of Adult RSV session, February 2023

- Moderna presented safety and efficacy data from the first 9 months of follow up in their phase 3 trial in adults aged  $\geq 60$  years
- CDC presented risk-stratified rates of RSV-associated hospitalization in U.S. adults aged 50 years and older
- CDC gave an update on uptake and implementation of RSV vaccine in U.S. adults aged 60 years and older during the first season following the recommendation
- CDC and FDA presented on post-marketing safety of the protein subunit RSV vaccines (GSK's AREXVY and Pfizer's ABRYYSVO) in adults aged 60 years and older
- CDC presented an analysis comparing the estimated magnitude of public health benefit and potential risk of Guillain-Barre syndrome (GBS) associated with protein subunit RSV vaccination in adults aged 60 years and older
- For adults aged 60 years and older who remain unvaccinated, ACIP agreed with encouraging timing of RSV vaccination in the late summer and early fall to optimize public health benefits

# Recent work group discussion

- Moderna mRESVIA for use in adults aged 60 years and older (received FDA approval May 31, 2024)
- GSK AREXVY for use in adults aged 50-59 years at increased risk of severe RSV disease (received FDA approval June 7, 2024)
- Continued discussion of safety, including risk of GBS, following RSV vaccination
- Shift from a shared clinical decision-making recommendation to:
  - Potential universal recommendation in adults 75 and older
  - Potential risk-based recommendation in adults 60–74
  - Potential risk-based recommendation in adults 50–59



# Re-evaluating Shared Clinical Decision-Making (SCDM) recommendation

- In June 2023, ACIP recommended RSV vaccination in adults 60 and older using shared clinical decision-making
  - In addition to benefits of vaccination, the shared clinical decision-making discussion, as intended by ACIP, is also meant to include discussion of the potential risk of vaccine-associated adverse events associated with RSV vaccine, specifically Guillain Barre Syndrome.
- In the first year following this recommendation, we have learned:
  - Feedback from healthcare providers that having SCDM conversations is not simple in practice.
  - Unlike a universal recommendation where there's a clear call to action to vaccinate, with SCDM the call to action is to discuss with a healthcare provider, a less clear message.
- A risk-based recommendation for adults 60–74 and a universal recommendation for adults  $\geq 75$  years would potentially highlight more clearly for providers and other public health officials which adults are likely to benefit from an RSV vaccine and provide a clearer recommendation for patients.



# Agenda: Wednesday June 26, 2024

- Manufacturer presentation: ABRYOVO (Pfizer) safety and immunogenicity in non-pregnant adults aged 18-59 years
- Manufacturer presentation: AREXVY (GSK) immunogenicity with a 24-month revaccination interval
- Manufacturer presentation: mRESVIA (Moderna) season 2 safety and efficacy update
- Postmarketing safety updates: Vaccine Safety Datalink
- Evaluation of Guillain-Barre Syndrome (GBS) following RSV vaccination among adults 65 years and older
- Observational RSV vaccine effectiveness
- Economic analysis of adult RSV vaccination
- Update to benefits and risks discussion
- Comparison of economic analyses of adult RSV vaccination
- Evidence to Recommendations
- Clinical Considerations
- Dr. Iona Munjal (Pfizer)
- Dr. Susan Gerber (GSK)
- Dr. Rituparna Das (Moderna)
- Dr. James Donahue (Marshfield Clinic Research Institute)
- Dr. Patricia Lloyd (FDA)
- Dr. Diya Surie (CDC, NCIRD)
- Dr. David Hutton (University of Michigan)
- Dr. David Hutton (University of Michigan)
- Dr. Ismael Ortega-Sanchez (CDC/NCIRD)
- Dr. Michael Melgar (CDC/NCIRD), Lauren Roper (CDC/NCIRD), Dr. Amadea Britton (CDC/NCIRD)
- Dr. Michael Melgar (CDC/NCIRD)

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

Photographs and images included in this presentation are licensed solely for CDC/NCIRD online and presentation use. No rights are implied or extended for use in printing or any use by other CDC CIOs or any external audiences.

