

Respiratory Syncytial Virus (RSV): Recognizing and Mitigating Risk in Vulnerable Adults

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Educational Objectives

By completing this educational activity, the participant should be better able to:

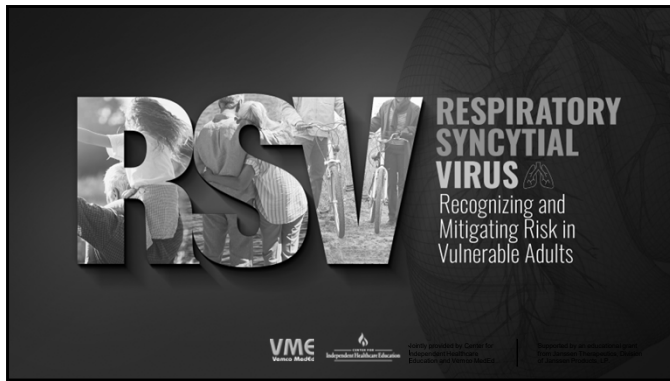
1. Discuss the burden of respiratory syncytial virus (RSV) infection in adults and list patient risk factors for severe infection and hospitalization.
2. Describe diagnostic approaches to differentiate RSV from other respiratory viral infections in adults.
3. Identify current and emerging approaches to prevent RSV in vulnerable adults.

Speakers' Disclosures

Dr. Gravenstein has disclosed that he has received research support from Pfizer, Sanofi, and Seqirus; he is on the speaker's bureau for Catapult, GlaxoSmithKline, Longeron, Merck, Novartis, Pfizer, Sanofi, and Seqirus; he is an independent contractor for Catapult Consultants and Healthcentric Advisors; and he is on the advisory board for Longeron, Merck, and Sanofi.

Supporter Disclosure

This educational activity is supported by an educational grant from Janssen Pharmaceuticals. It has been planned and produced by VemCo MedEd with Texas Academy of Family Physicians strictly as an accredited continuing medical education activity.



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Activity Description

Target Audience
This activity is intended to meet the needs of primary care providers including internists, family physicians, osteopathic physicians, physician assistants, and nurse practitioners. This program will target PCPs who are involved in the evaluation and management of adults at risk of serious RSV infection.

Learning Objectives
At the conclusion of the educational activity, the learner should be able to:

- Discuss the burden of respiratory syncytial virus (RSV) infection in adults and list patient risk factors for severe infection and hospitalization
- Describe diagnostic approaches to differentiate RSV from other respiratory viral infections in adults
- Identify current and emerging approaches to prevent RSV in vulnerable adults

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Faculty and Disclosure

Stefan Gravenstein, MD, MPH
Professor of Medicine
Director, Division of Geriatrics and Palliative Care
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Dr. Stefan Gravenstein reported the following relevant financial relationships with ineligible companies:

- Consulting: Sanofi, Merck & Co., Inc., Pfizer Inc., Novavax, VaxArt, Janssen, Moderna, GSK, Reviral
- Speakers Bureau: Seqirus, Sanofi, Janssen
- Research Support: Seqirus, Sanofi, Pfizer Inc.
- Advisory Board: Janssen

Dr. Gravenstein does not discuss off-label uses of any products.

All relevant financial relationships have been mitigated. No (other) speakers, authors, planners or content reviewers have any relevant financial relationships to disclose. Content review confirmed that the content was developed in a fair, balanced manner free from commercial bias. Disclosure of a relationship is not intended to suggest or condone commercial bias in any presentation, but it is made to provide participants with information that might be of potential importance to their evaluation of a presentation.

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Respiratory Syncytial Virus Basics: The Virus

Synonyms: Human RSV or hRSV; human orthopneumovirus

Negative sense, single stranded RNA virus

- 150nm diameter (some filamentous species can be several micrometers long)
- Codes for
 - Key internal structural proteins: Matrix protein [M], Nucleoprotein [N]
 - Proteins for the polymerase complex (P and L)
 - Nonstructural proteins [NS-1 and NS-2]: help evade innate immune response
 - Externally exposed transmembrane glycoproteins: small hydrophobic protein [SH], glycoprotein [G], **Fusion protein [F]**
 - Regulatory protein M2 proteins (M2-1 antitermination protein and M2-2, transcription/replication regulators)

RNA copying is error prone, allowing for rapid generation of single nucleotide polymorphisms...

Knipe DM, Howley PM, Cohen JI, Griffin DE, Lamb RA, Martin MA, Racaniello VR, Roizman B (ed). 2013. Fields virology, 6th ed. Lippincott Williams & Wilkins, Philadelphia, PA.
Griffiths C, Drews SJ, Marchant DJ. Clin Microbiol Rev. 2017;30(1):277-319.
Agoti CK, Otiemo JR, Gitahi CW, Camp PA, Nokes DJ. Emerg Infect Dis. 2014;20(6):950-9.
Mujase A, et al. Ann Allergy Asthma Immunol. 2020;125:36-46.

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RSV Basics: The Virus

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- M, N, P, L and nonstructural proteins [NS-1 and NS-2] that help evade innate immune response, externally exposed transmembrane glycoproteins (small hydrophobic protein [SH], glycoprotein [G], **fusion protein [F]**, and regulatory protein M2 proteins (M2-1 antitermination protein and M2-2, transcription/replication regulators)
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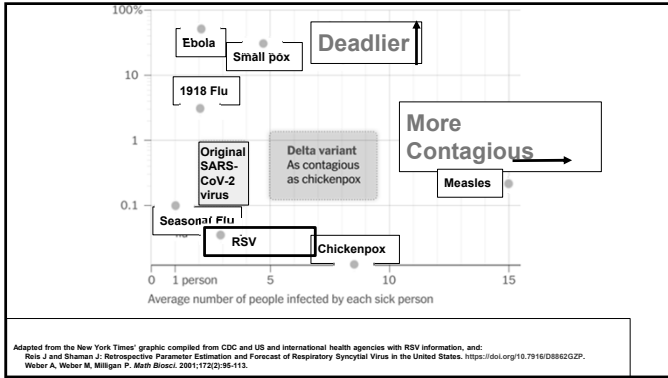
Infected cells fuse to form large cells, or syncytia

Spreads by air droplets or fomites

- Lands in eyes, nose or mouth

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 Infected cells fuse to form large cells, or syncytia

Spreads by droplets or fomites: it's pretty contagious!

- Lands on eyes, nose or mouth, transmits through the air and by fomite
- Binds to and infects airway epithelial cells
- R_0 has been estimated anywhere from 3 to 25 depending on model assumptions, but the value closer to 3 works well with predicting peak of outbreaks
- 70% of forecasts predict peak magnitude of RSV activity 4 weeks ahead of time
- CDC recommends "contact precautions" and contagious from 3 days to 4 weeks

Rein J, Shaman J (2016). *PLoS Comput Biol.* 2016;12(10): e1005133.
 Griffin C, Drews SJ, Marchant DJ. *Clin Microbiol Rev.* 2017;30(1):277-316.
 Weber A, Weber M, Milligan P. *Math Biosci.* 2005;172(2):95-113.

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Fun Fact

Infants have nearly all of the airways and alveoli they will have as adults

- This means a huge surface area to volume and especially tiny airways

This means that it takes less inflammation and bronchospasm to cause obstruction that results in wheezing and croup

It's **one of three reasons** children present differently from older adults with RSV infection

Hislop AA. *J Anat.* 2002;201(4):325-34.

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Risk Factors for Severe RSV Infection

- Age
- Overcrowding
- Smoke exposure (cooking, tobacco)
- Low SES
- Asthmatic mother (for risk in children)
- Co-morbidities** (and in older adults, multimorbidity)

SES, socio-economic status

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Susceptibility in Older Adults

RSV is among the top four causes of ILI (third before the advent of SARS-CoV-2), after enterovirus and influenza

- But RSV was the second most common cause of hospitalization
- Twice as likely as patients who had laboratory confirmation of influenza

95% of children have had RSV by age 2

- Essentially all adults have survived prior RSV, and will have some underlying immunity

Respiratory infections and related hospitalizations begin increasing around age 50 (P&I)

- Immune senescence
- In elderly, greater susceptibility with lower RSV-specific Ig and nasal IgA
- T-cell immunity declines with age

ILI, influenza-like illness
 Falsey AR, et al. *J Infect Dis.* 2014;209(12):1873-81.
 Falsey AR, Walsh EE. *J Infect Dis.* 1998;177(2):463-6.
 Walsh EE, Falsey AR. *J Infect Dis.* 2004;190(2):373-8.

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Reference:
 I. Hoshino K. *Immunol.* 1975;51(4):1659-1664.

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 Falsey AR, Walsh EE. *J Infect Dis.* 1998;177(2):463-6.
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Immune Senescence

T-cells change with age

- Reduced numbers of new T-cells and naive T-cells; reduced B-cell stimulation
 - B-cells make less antibody, reduced neutralizing antibody, isotype switched Ab repertoire
- Increased pool of memory T-cells
 - Memory T-cells have increasing dysfunction
 - Reduced IFN-gamma, cytokine production

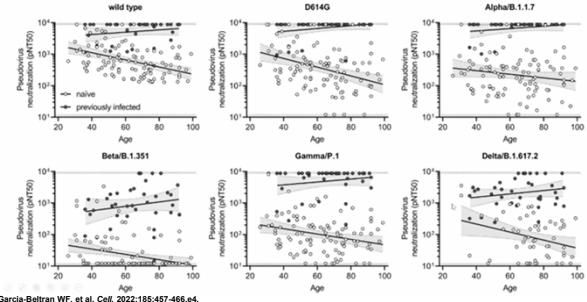
Dendritic cells (DC) present antigens to T-cells

- DC function is to present antigens to T-cells
- DC number and phenotype stable with age, but have declining function
 - Less able to process and present antigens, and to migrate to infected site (lung)
 - Increased level of pro-inflammatory cytokines on stimulation, and failure to recognize self (IL-6, TNF, INF-a)
 - Reduced TLR expression

Stephens LM, Varga SM. *Vaccines (Basel)*. 2021;9(6):624.
Agrawal A, Gupta S. *Ageing Res Rev*. 2011;10(3):336-65.

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Reduced Neutralizing Ab with Age after SARS-CoV-2 Vaccination



García-Betran WF, et al. *Cell*. 2022;185:457-466.e4.
Canaday DH, et al. *Clin Infect Dis*. 2021;73(2):212-215.

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Biologic Changes With Age Relate to Clinical Presentation

Biologic Change	Clinical effect
Reduced IL-6	Reduced fever, less efficient viral clearance
Impaired respiratory tract mucociliary function	Reduced cough, less efficient viral and mucous clearance
Delayed cytokine increase	Fewer symptoms at onset
Delayed cytokine normalization	Slower improvement and prolonged pro-inflammatory state
Reduced T-cell help	Reduced response to infection, vaccination; less durable
Reduced nutrition	Reduced physiologic reserve, more difficult rehabilitation
Brain Aging	Risk for delirium, sleep/appetite disturbance with cytokine storm

Gravenstein S, et al. *Med Health R I*. 2010;93(12):382-384.

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Fun Fact

Immune senescence is the second of three reasons why children present differently from older adults

- Children produce more cytokine faster (therefore faster and higher fever), and other cytokine-mediated symptoms
- Children may not have prior immunity, increasing peak viral shedding titers

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Most Clinicians Don't know that RSV is a Big Deal for Older Adults

Each year, up to 10% of older adults are infected with RSV in the US

- Closer to 10% in settings with close quarters (e.g., nursing homes, assisted living and senior housing)

Older adults more likely than younger adults to be hospitalized or die

Associated Risk Condition	Odds Ratio (95% CI)	P Value
Stroke, heart failure, chronic lung disease	~2 (1.02-4)	<0.05
Solid organ transplant	2.52 (0.88-7.22)	0.085
Chronic kidney disease	4.37 (2.74-6.98)	<0.001
Hematologic malignancy	5.17 (2.02-13.20)	0.001

Branche AR, et al. *Drugs Aging*. 2015;32(4):261-269.
Pastula ST, et al. *Open Forum Infect Dis*. 2017;4(1):ofw270.
Wyffels V, et al. *Adv Ther*. 2020;37:1203-1217.

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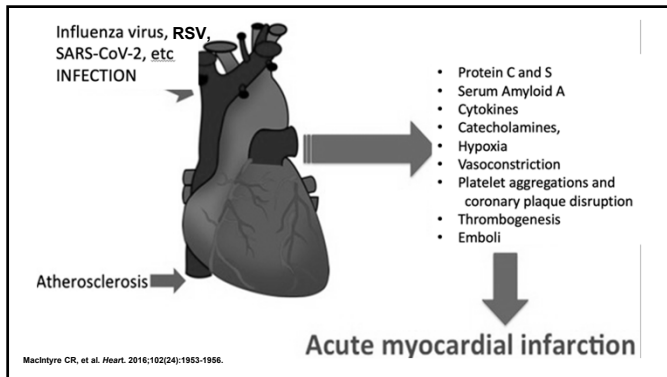
P&I begin increasing around age 50

- Immune senescence
 - In elderly, greater susceptibility with lower RSV-specific Ig and nasal IgA
 - T-cell immunity declines with age: reduced CD8 cytotoxic T-cell function; shift Th1 to Th2
- Decline in DC function

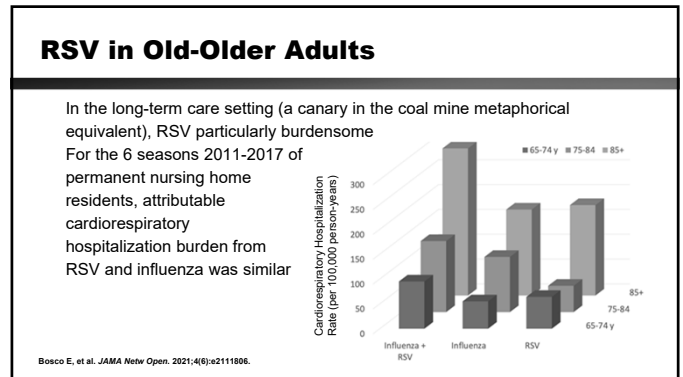
Older adults with severe RSV do show CD4 and CD8 T-cell responses *but unclear if severe disease is due to immunosenescence or "just" impaired T-cell responses and/or dysfunctional antibody*

Falsey AR, et al. *J Infect Dis*. 2014;209(12):1873-81.
Falsey AR, Walsh EE. *J Infect Dis*. 1998;177(2):463-6.
Walsh EE, Falsey AR. *J Infect Dis*. 2004;190(2):373-8.

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Fun Fact

Children have a better mucociliary escalator than older adults

- o With age, fewer cells and less efficient viral clearance on top of greater likelihood of polypharmacy—including drugs that dry secretions) change ability to clear virus
- o So early, wheezing, whooping more prominent with greater consequences from inflammation and earlier coughing
- o In older adults, productive coughing likely delayed a bit in course of illness and less wheezing

Children also don't typically have the other underlying conditions

- o So diagnostic confusion for other etiology (HF or COPD exacerbation) not as easily confounded by a diagnostic heuristic

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Rationale for RSV Testing in Older Adults

Antibiotic stewardship

- o As in Falsey study, antibiotic use high, and often potentially inappropriate

In the era of CoVID, diagnostic stewardship

- o Clinically, at onset RSV, SARS-CoV-2/CoVID, influenza, parainfluenza, etc. indistinguishable
- o Index of suspicion with one virus over another has context with diagnosed close contacts
- o Context: a negative SARS-CoV-2 test does not preclude SARS-CoV-2 infection
 - o PCR is highly sensitive when virus is present
 - o It can take days, sometimes weeks for SARS-CoV-2 PCR test to become positive
- o **Antigen test + @ higher titers** (it's contagious!), but **false - in up to half of infected**
- o Ruling in RSV makes a firm diagnosis and dual infection generally is uncommon
- o Multiplex testing identifies virus 40% or more often, can limit other tests
 - o Downside: more tests= more false positives, sample dependent

In the long-term care setting, having a diagnostic test has huge facility-level implications in these resource-poor environments: staffing, PPE, time and effort

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Management of RSV in Older Adults

Supportive

- o Bronchodilators (not FDA approved for this indication)
- o Steroids (especially with COPD) (not FDA approved for this indication)
- o O₂

Contact precautions

- o Frequent hand hygiene
- o Mask, ideally double mask (hook and loop) and properly fitted
- o Keep high-touch surfaces clean and disinfected
- o Isolate infected patients

Ribavirin, antibody treatment available not generally used in older adults, and not FDA approved for this indication in older adults

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The Path Forward

RSV is a big deal, but most clinicians caring for adult and older adult patients aren't aware of the implications

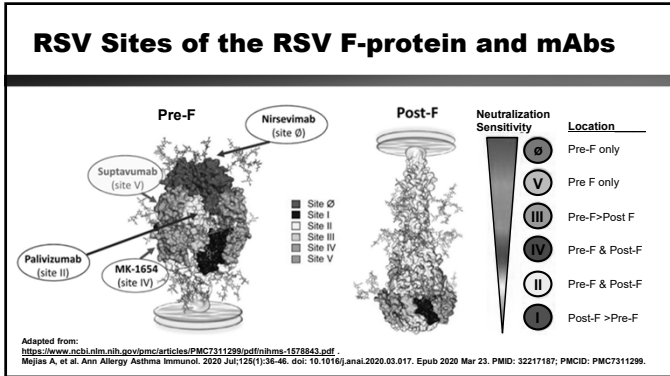
Because there are no meaningful approaches to prevention or intervention, there's little motivation to test or change awareness

Several vaccines in development, some in phase III now

- o Early failures with RSV vaccine (e.g., enhanced disease in vaccines studies in the 1960s) elevate the importance of safety signals of new vaccines

Neither monoclonal antibody or antivirals are likely to gain ground any time soon as a therapeutic options for older adults

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What's in Development: RSV Vaccines for the Elderly

Vaccine Type	Phase 1	Phase 2	Phase 3
Protein based (Inactivated, particle, subunit)	<ul style="list-style-type: none"> RSV SH Protein (Immunovaccine VIB) RSV F protein (NIH/NIAD/VRC) 	<ul style="list-style-type: none"> RSV G protein (Advaccine Biotechnology) RSV F protein (Pfizer) 	<ul style="list-style-type: none"> RSV F protein (RSVPreF3; GlaxoSmithKline)
Nucleic Acid		<ul style="list-style-type: none"> mRNA-1345 (Moderna) 	
Recombinant Vector		<ul style="list-style-type: none"> MVA (Bavarian Nordic) 	<ul style="list-style-type: none"> Adenovirus (Ad26.RSV.preF, Janssen)

Available at: https://path.azureedge.net/media/documents/RSV_snapshot-07APR2021_HighResolution_NonEditable_PDF_3KqK9PB.pdf

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Protein-based Vaccine Elicits Robust Immune Response in Elderly

RSVPreF3 contains recombinant subunit pre-fusion RSV antigen combined with adjuvant

Vaccine was well tolerated in Phase 1/2 studies in young and older adults

At one month post-immunization, elicited robust humoral and cellular immune response

- 10-fold increase in RSV-A neutralizing antibodies
- >12-fold increase in RSVPreF3 IgG antibodies

Phase 3 trial (AReSVi 004) started in 2021 to include up to 1650 adults ≥60 years with 3-year follow-up

- Interim results expected in second half of 2022

Schwarz TF, et al. *J Infect Dis.* 2021;jjab317. Presented at IDWeek 2020 (Abstract 119). <https://www.asnk.com/en/glob/media/press-releases/asnk-starts-phase-iii-rsv-candidate-vaccine-programme-for-older-adults/>.

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Early Signs of Success with Vector-based Vaccine in Elderly (CYPRESS Study)

Phase 2b study randomized 5782 individuals ≥65 years to receive Ad26.RSV.preF vaccine or placebo

Primary endpoint: First occurrence of RT PCR-confirmed RSV-mediated lower respiratory tract disease according to any of 3 case definitions:

- ≥3 symptoms of lower respiratory tract infection (LRTI)
- ≥2 symptoms of LRTI
- ≥2 symptoms of LRTI or ≥1 symptoms of LRTI plus ≥1 systemic symptom

Vaccine efficacy for each case definition was 80% (definition 1), 75% (definition 2), and 69.8% (definition 3)

Vaccine elicited a robust humoral and cellular immune response

A phase 3 trial (EVERGREEN) is underway

Presented at IDWeek 2021 (Abstract LB14). <https://www.contagionlive.com/view/cypress-trial-rsv-vaccine-older-adults-elicits-robust-immune-response>.

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Maximizing Protection in the Elderly: Co-Administration of RSV and Flu Vaccines

Phase 2a, double-blind, placebo-controlled study of 180 adults ≥60 years

Participants randomized to receive:

- Ad26.RSV.preF plus Fluorix on Day 1 and placebo on Day 29
- Placebo plus Fluorix on Day 1 and Ad26.RSV.preF on Day 29 (control)

Co-administration had an acceptable safety profile and showed no evidence of interference in immune response.

Results are compatible with simultaneous seasonal vaccination with both vaccines

Sadoff J, et al. *J Infect Dis.* 2021;223:699-708.

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Summary

RSV: is under-recognized for its importance in older adults

Altered presentation but indistinguishable clinically from other important causes of URI and LRTI

Awareness will increase as the clinical value of multiplex testing gains acceptance and is further established as a tool for:

- Antimicrobial stewardship
- Diagnostic stewardship
- Improving workforce and resource stability in under-resourced environments

We need an approach for primary prevention (vaccination!)

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Q&A

Evaluation Link:

<https://www.surveymonkey.com/r/RSV2022>

