



HAP, VAP & All of That

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Objectives

1

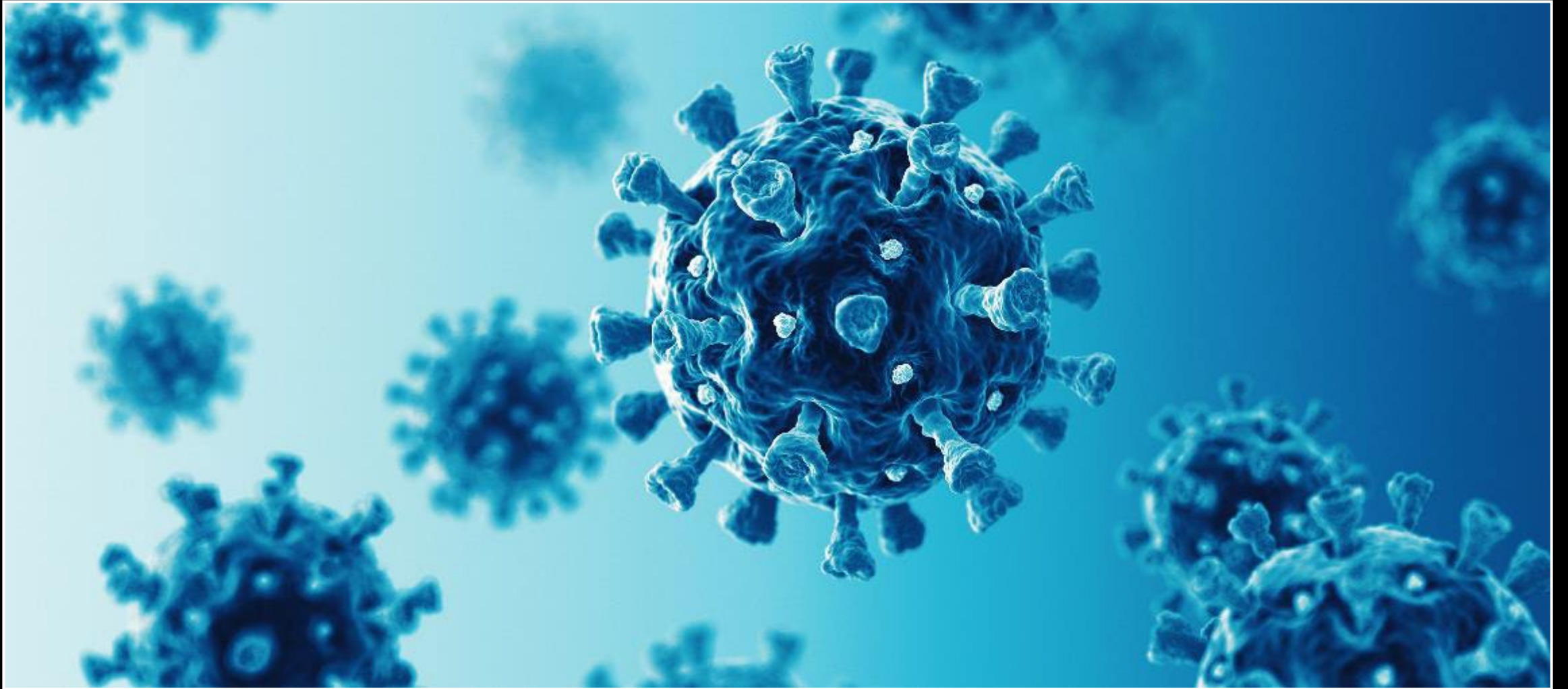
Recognize the definitions of NV-HAP, VAE, and VAP

2

Discuss the epidemiology and pathophysiology of all types of HAP

3

Review recommendations for prevention of all types of HAP

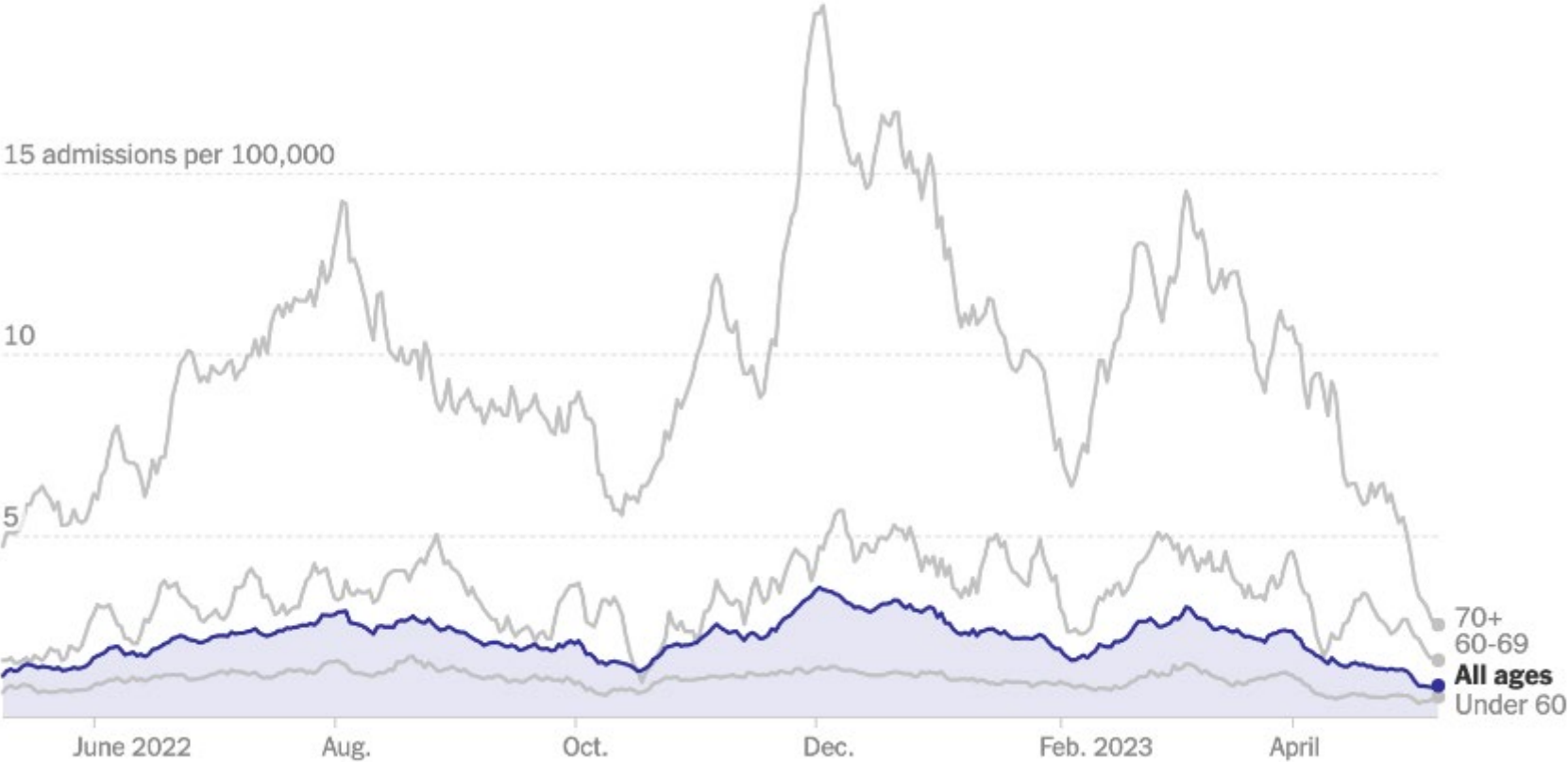


But First, COVID Update

Nebraska COVID Hospital Cases – May 8, 2023

Daily Covid hospital admissions

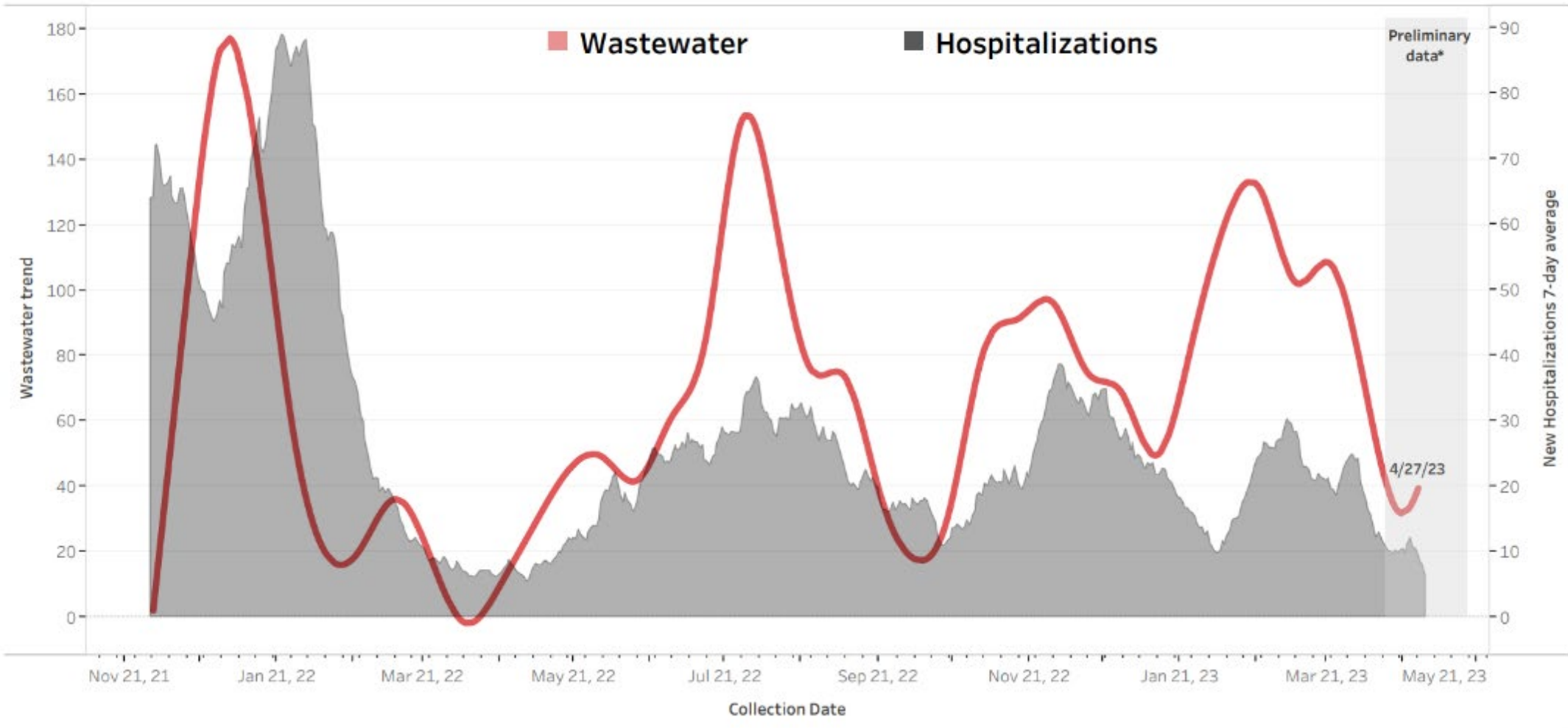
Avg. on May 8 14-day change
16 **-35%**



Nebraska SARS-CoV-2 Wastewater Surveillance Report

Nebraska Statewide SARS-CoV-2 Wastewater Levels and COVID-19 Hospitalizations

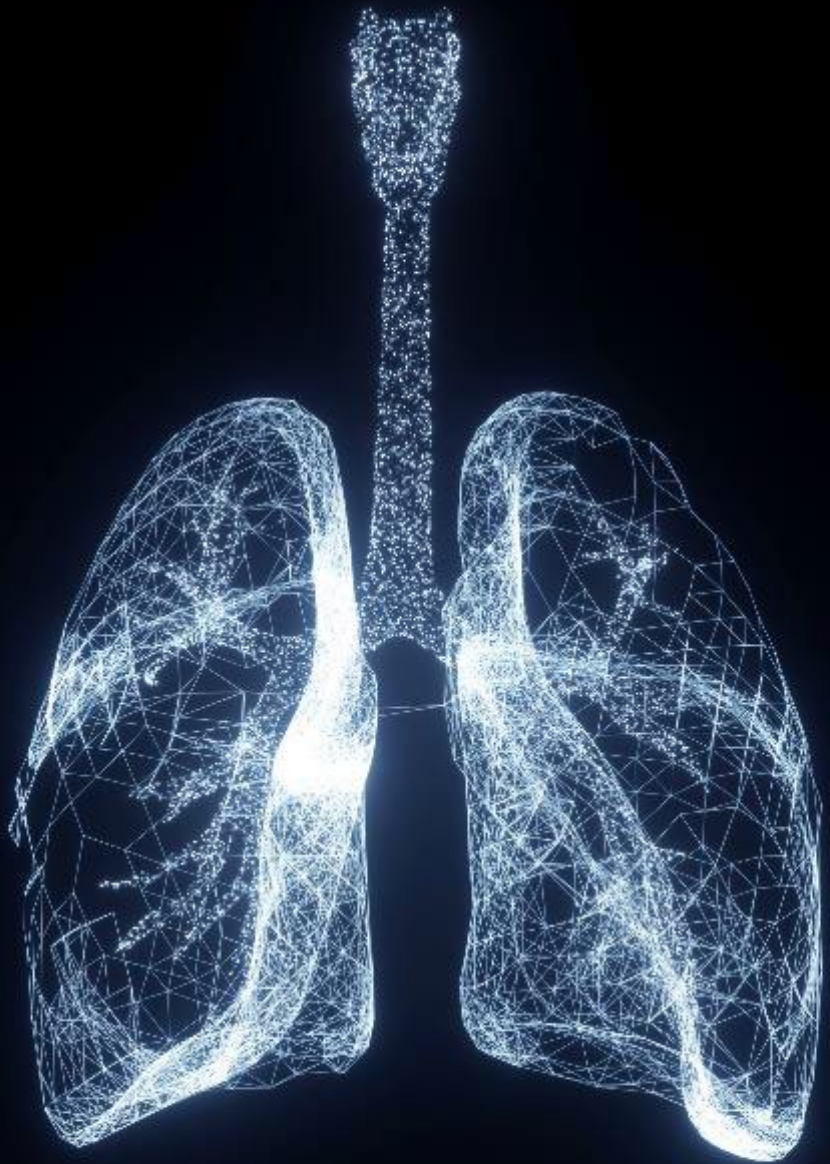
Updated 5/2/2023





Arcturus – Newest COVID Variant

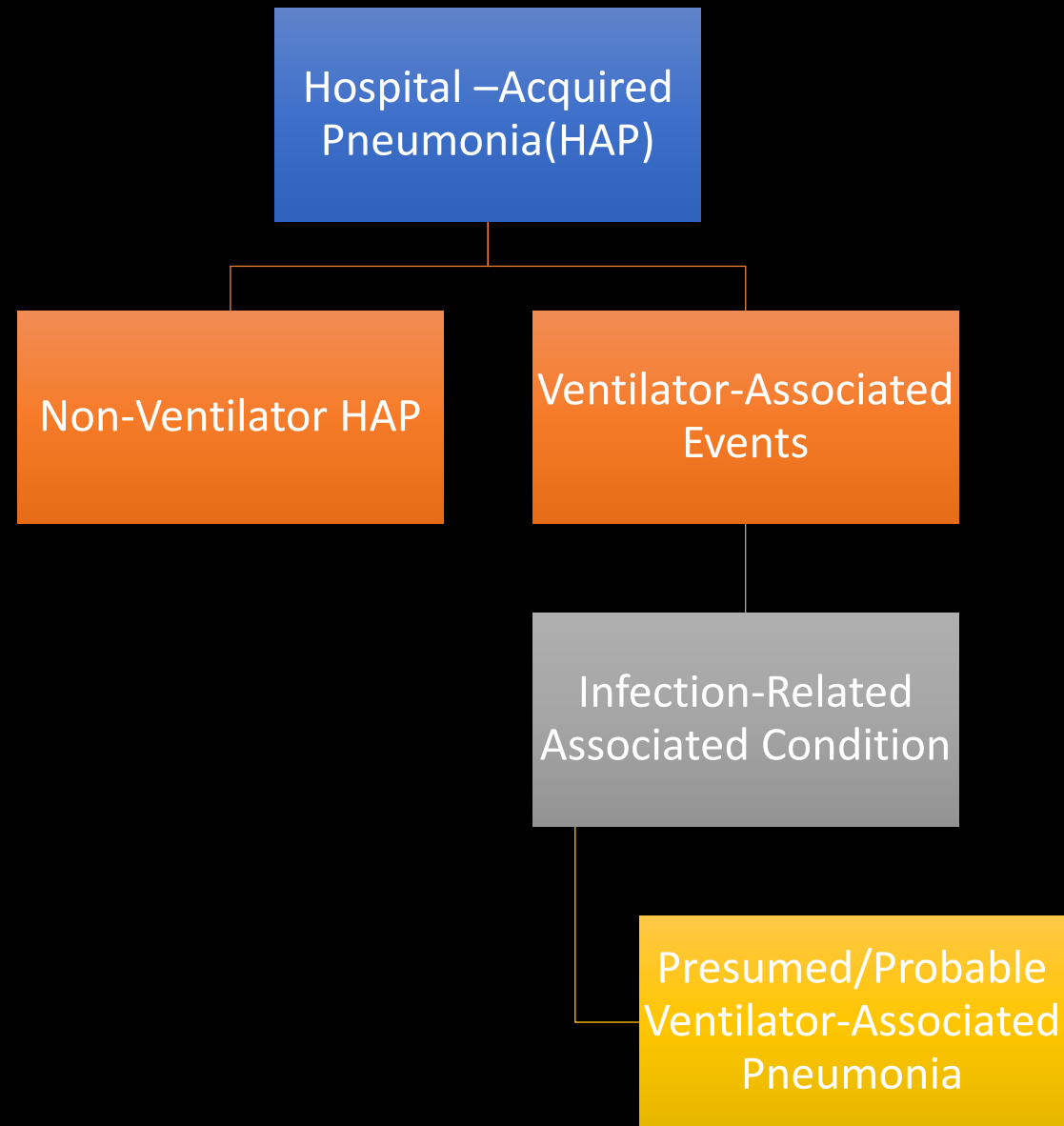
- Reported in > 30 countries and growing in the US
- Anticipated to take over current strains
- Possible summer surge; but not a 'variant of concern'
- Flu-like symptoms, more pink-eye/itchy eye symptoms
- Vaccine still effective



Community-Acquired
to Hospital-Acquired

The Different Pathophysiology of Pneumonias







Burden of HAP


- HAP is the most common nosocomial infection
- Impacting ~1 in 100 patients
- NV-HAP = 60-70% of all HAPs
- Approximately 5-10% of patients on a ventilator develop VAEs
- NV-HAP, VAE and VAP all carry increased morbidity, mortality, LOS and cost
- Attributable mortality from VAP is ~10% but varies based on type and severity of illness

VAE vs VAP – Why Does It Matter?

VAE is NOT synonymous with VAP



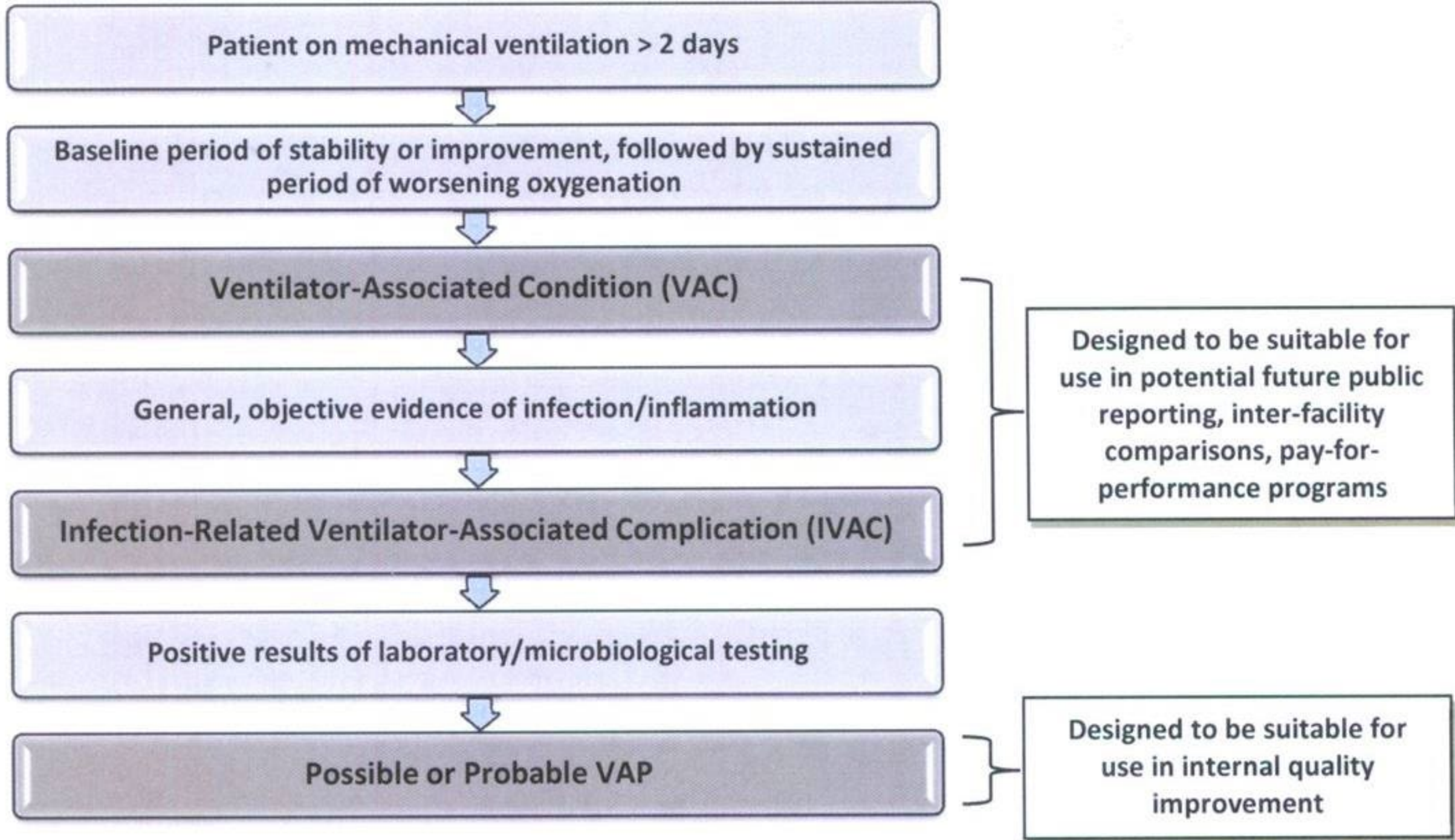
Only 25-33% of VAEs are due to pneumonia; 'mild' pneumonia may not meet ventilator requirements



VAE is considered a quality metric given associated poor outcomes with VAE



VAE is increasingly demonstratable to be preventable with implementation of best practices



Patient on mechanical ventilation > 2 days

Baseline period of stability or improvement, followed by sustained period of worsening oxygenation

Ventilator-Associated Condition (VAC)

General, objective evidence of infection/inflammation

Infection-Related Ventilator-Associated Complication (IVAC)

Positive results of laboratory/microbiological testing

Possible or Probable VAP

Designed to be suitable for use in potential future public reporting, inter-facility comparisons, pay-for-performance programs

Designed to be suitable for use in internal quality improvement

Patient has a baseline period of stability or improvement on the ventilator, defined by ≥ 2 calendar days of stable or decreasing daily minimum FiO_2 or PEEP values. The baseline period is defined as the two calendar days immediately preceding the first day of increased daily minimum PEEP or FiO_2 .



After a period of stability or improvement on the ventilator, the patient has at least one of the following indicators of worsening oxygenation:

- 1) Increase in daily minimum FiO_2 of ≥ 0.20 (20 points) over the daily minimum FiO_2 in the baseline period, sustained for ≥ 2 calendar days.
- 2) Increase in daily minimum PEEP values of ≥ 3 cmH_2O over the daily minimum PEEP in the baseline period, sustained for ≥ 2 calendar days.

Example

Date	PEEP (min)	FiO2 (min)
Jan 1	10	100
Jan 2	5	50
Jan 3	5	40
Jan 4	5	40
Jan 5	8	60
Jan 6	8	50
Jan 7	8	40
Jan 8	5	40
Jan 9	5	40

VAE

Patient meets criteria for VAC



On or after calendar day 3 of mechanical ventilation and within 2 calendar days before or after the onset of worsening oxygenation, the patient meets both of the following criteria:

- 1) Temperature $> 38^{\circ}\text{C}$ or $< 36^{\circ}\text{C}$, **OR** white blood cell count $\geq 12,000$ cells/mm³ or $\leq 4,000$ cells/mm³.

AND

- 2) A new antimicrobial agent(s)* is started, and is continued for ≥ 4 calendar days.

*See [Appendix](#) for eligible agents.

Patient meets criteria for VAC and IVAC

AND

On or after calendar day 3 of mechanical ventilation and within 2 calendar days before or after the onset of worsening oxygenation, ONE of the following criteria is met:

- 1) Purulent respiratory secretions (from one or more specimen collections)
 - Defined as secretions from the lungs, bronchi, or trachea that contain \geq 25 neutrophils and \leq 10 squamous epithelial cells per low power field [lpf, x100].
 - If the laboratory reports semi-quantitative results, those results must be equivalent to the above quantitative thresholds.

OR

- 2) Positive culture (qualitative, semi-quantitative or quantitative) of sputum*, endotracheal aspirate*, bronchoalveolar lavage*, lung tissue, or protected specimen brushing*

**Excludes the following:*

- Normal respiratory/oral flora, mixed respiratory/oral flora or equivalent
- *Candida* species or yeast not otherwise specified
- Coagulase-negative *Staphylococcus* species
- *Enterococcus* species

Compendium Updates

Infection Control & Hospital Epidemiology (2022), **43**, 687–713
doi:[10.1017/ice.2022.88](https://doi.org/10.1017/ice.2022.88)



SHEA/IDSA/APIC Practice Recommendation

Strategies to prevent ventilator-associated pneumonia, ventilator-associated events, and nonventilator hospital-acquired pneumonia in acute-care hospitals: 2022 Update

Michael Klompas MD, MPH^{1,2} , Richard Branson MSc, RRT³ , Kelly Cawcutt MD, MS⁴ , Matthew Crist MD⁵ ,
Eric C. Eichenwald MD^{6,7}, Linda R. Greene RN, MPS, CIC⁸, Grace Lee MD⁹, Lisa L. Maragakis MD, MPH¹⁰,
Krista Powell MD, MPH⁵ , Gregory P. Priebe MD¹¹ , Kathleen Speck MPH¹², Deborah S. Yokoe MD, MPH¹³ and
Sean M. Berenholtz MD, MHS^{12,14,15}

Essential Practices

- Good evidence that the intervention decreases the average duration of mechanical ventilation, length of stay, mortality, and/or costs. Benefits likely outweigh risks,





Essential Practices

Intervention	Quality of Evidence
Avoid intubation & prevent reintubation Use HF O2 or NIPPV when safe & feasible	High
Provide early enteral vs parenteral nutrition	High
Change the ventilator circuit only if visibly soiled or malfunctioning	High
Maintain & improve physical conditioning	Moderate
Minimize Sedation Avoid benzos in favor of other agents Use a protocol to minimize sedation Implement a ventilator liberation protocol	Moderate
Provide oral care with toothbrushing but <i>without</i> chlorhexidine	Moderate
Elevate head of bed to 30-45	Low

Additional Approaches

- Good evidence that the intervention improves outcomes in some populations, but may confer some risk in others.
- May lower VAP rates but insufficient data to determine impact on duration of mechanical ventilation, LOS or mortality




Additional Approaches

Intervention	Quality of Evidence
Use selective oral or digestive decontamination in countries and ICUs with low prevalence of antibiotic resistant organisms	High
Utilize endotracheal tubes with subglottic secretion drainage ports for patients expected to require >48-72 hours of mechanical ventilation	Moderate
Consider early tracheostomy	Moderate
Consider post-pyloric rather than gastric feeding for patients with gastric intolerance or at high risk for aspiration	Moderate



Generally Not
Recommended


- Inconsistently associated with lower VAP rates and no impact or negative impact on duration of mechanical ventilation, LOS or mortality
 - No impact on VAP rates, average duration of mechanical ventilation, LOS or mortality
- 

Generally Not Recommended

Intervention	Quality of Evidence
Oral care with chlorhexidine	Moderate
Probiotics	Moderate
Ultrathin polyurethane ET cuffs	Moderate
Tapered ET cuffs	Moderate
Automated control of ET cuff pressure	Moderate
Frequent cuff-pressure monitoring	Moderate
Silver-coated ETs	Moderate
Kinetic beds	Moderate
Prone positioning	Moderate
Chlorhexidine bathing	Moderate
Stress-ulcer prophylaxis	Moderate
Monitoring residual gastric volumes	Moderate
Early parenteral nutrition	Moderate



No
Recommendation

- No impact on VAP rates or other patient outcomes, unclear impact on costs
 - Closed endotracheal suctioning systems
- 

Burden of NV-HAP

- Most Common HAI
- ~50% require ICU
- ~20% need mechanical ventilation
- Length-of-stay longer by up to ~15 days
- Up to 30% crude mortality



A photograph of a hospital room, showing a patient lying in a bed with medical equipment and a nurse's station in the background. The room is dimly lit, with a focus on the patient's bed and the surrounding medical environment.

Risk Factors for NV-HAP

- Age > 65
- Depressed CNS
- Post-surgical patients
- Limited mobility
- Enteral feedings*
- Malnutrition
- Use of gastric acid suppressants
- Hyperglycemia



2020 Call To Action on NV-HAP



Surveillance & Prevention Barriers

- Definitions
- Surveillance
- Not YET reportable
- Leadership buy-in
- Evidence-based preventative strategies



Defining NV-HAP

Factor	Definition
Clinical features of NV-HAP	
Inclusion criteria	≥ 3 days hospitalization Not receiving mechanical ventilation
Worsening oxygenation sustained for ≥2 calendar days	Drop in pulse oximetry from ≥95% on ambient air to <95% on ambient air, or Initiation of supplemental oxygen, or Escalation of supplemental oxygen (flow rate or device)
Fever or Abnormal WBC	Temperature ≤36 or ≥38 °C, or WBC <4,000 or ≥12,000 cells/mm ³
Recognition/response by clinical team	
Performance of chest imaging	Evidence of order or procedure code for chest radiograph or computerized tomography
Initiation of new antibiotics	Administration of selected antimicrobials (supplement B) Not previously administered in past 2 days ≥3 days of administration

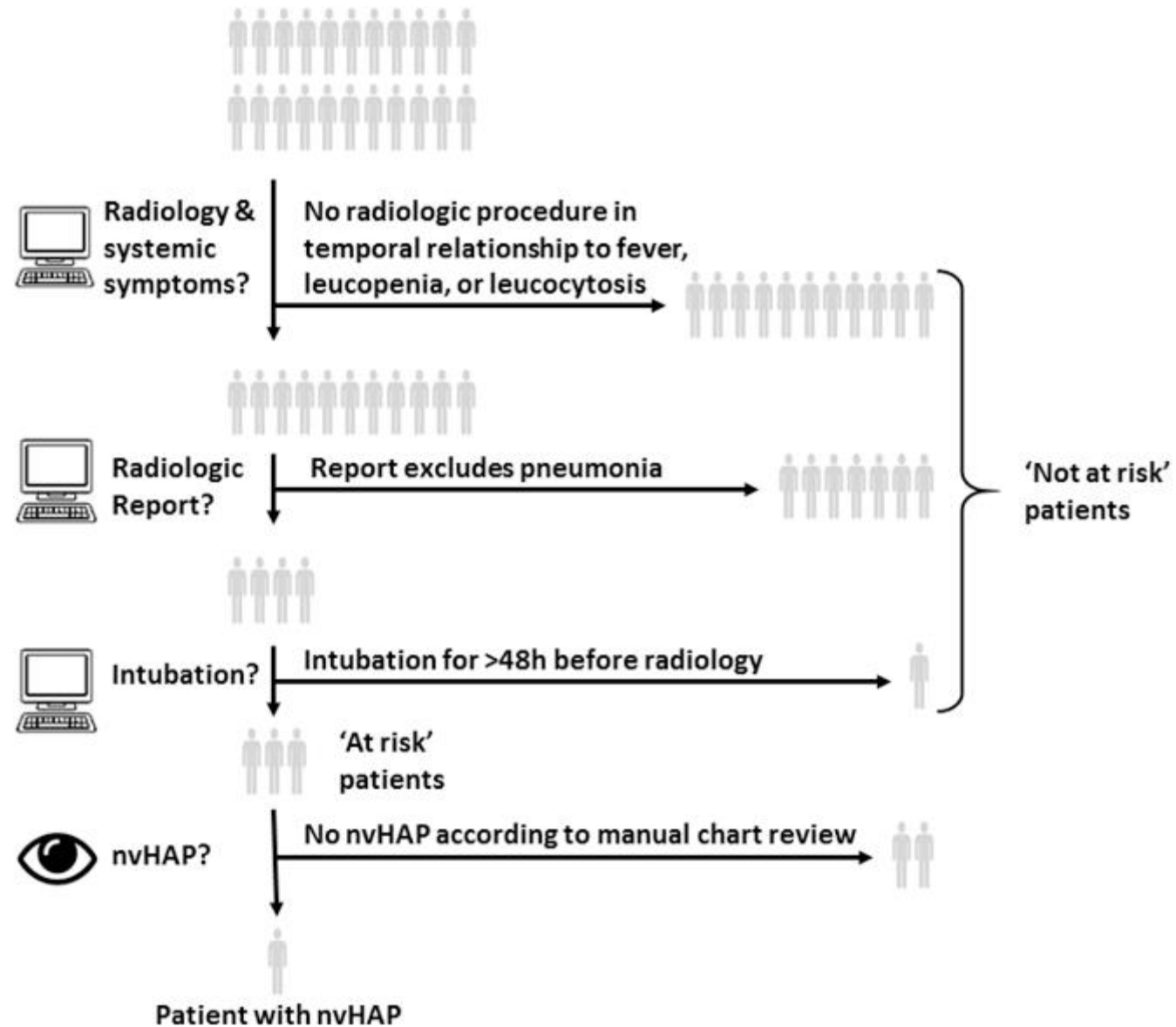


TWO Reasons Clinical Assessments May Not Be Adequate



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Surveillance for NV-HAP



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NV-HAP

Paucity of data, but increasing frequency

Recommendations:

- Oral care – toothbrushing
- Diagnose and manage dysphagia
- Early mobility
- Prevent nosocomial viral spread

The Joint Commission – Prevent NV-HAP Via:

Maintaining regular oral care

Maintaining patient mobility

Elevating the head of the patient's bed

Reducing the use of acid-suppressing medications

Minimizing sedation

Performing dysphagia screening in high-risk patients

Using modified diets and feeding strategies for patients with abnormal swallowing

Following standardized processes to place and manage feeding tubes

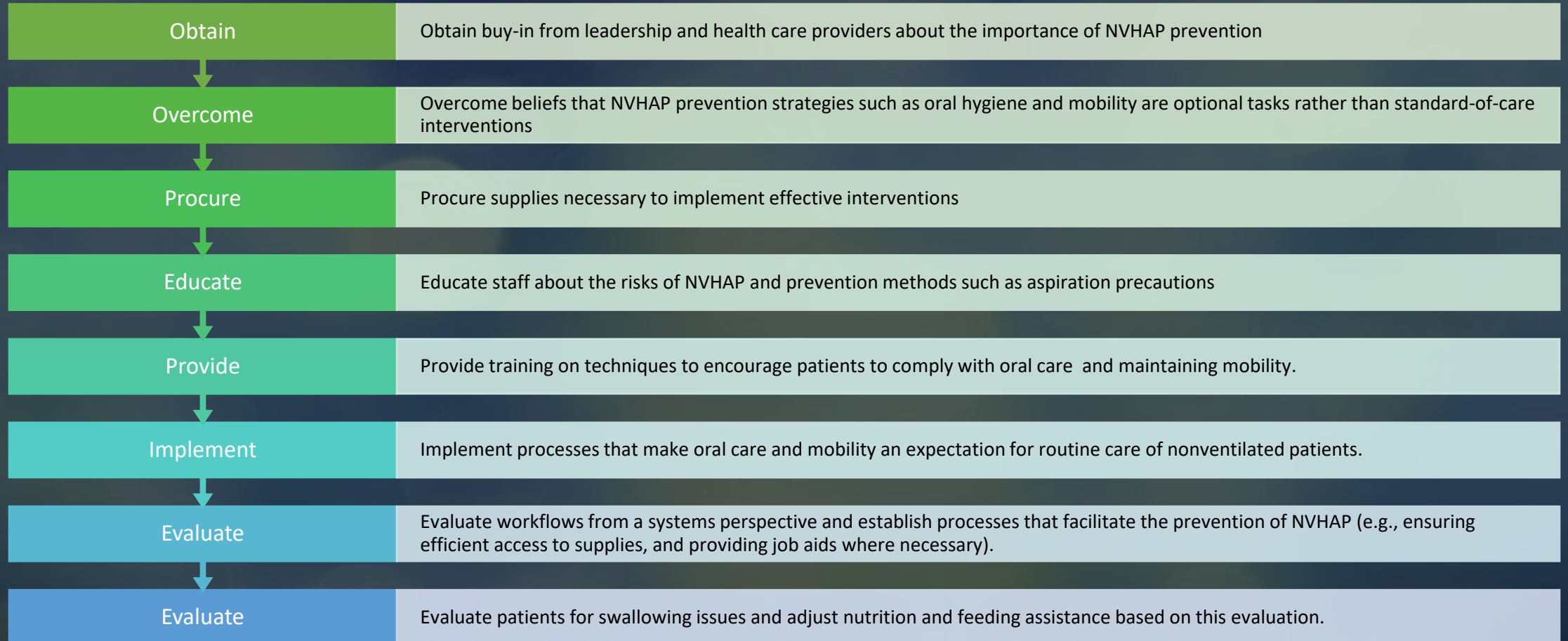
Breathing exercises

Using chest physiotherapy

Using incentive spirometry

Educating the patient and family about NVHAP prevention



The Joint Commission – Additional Actions



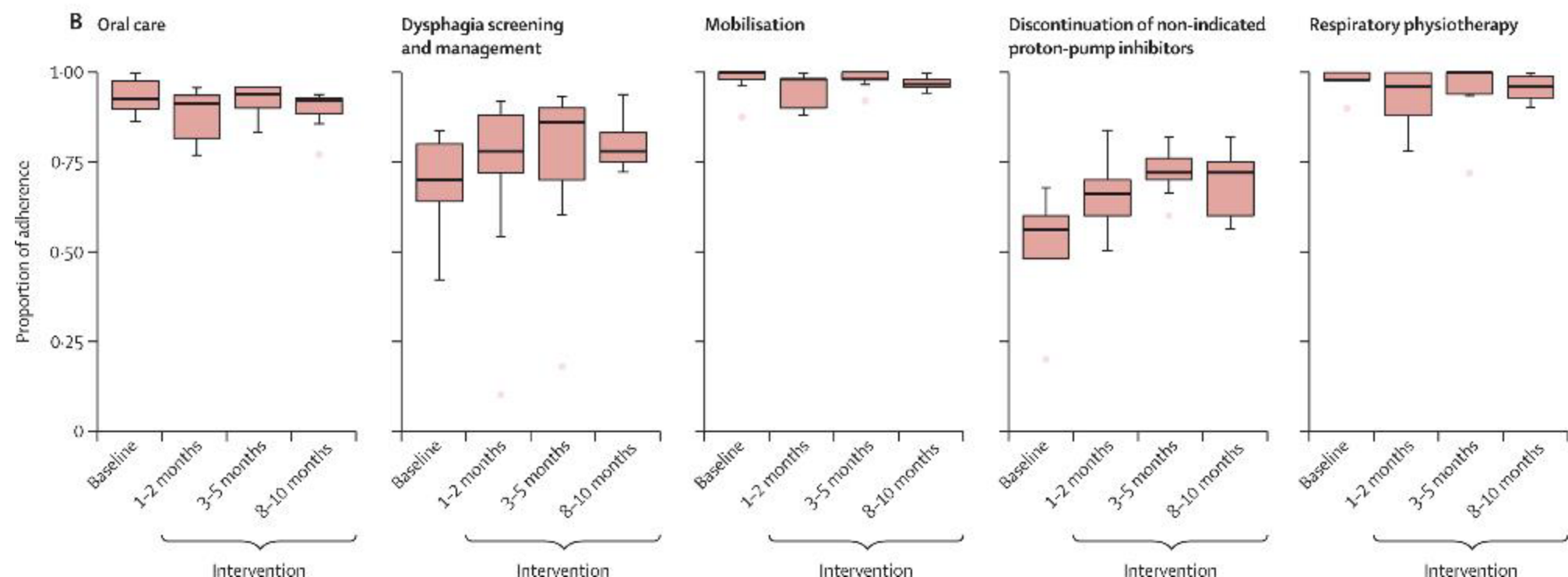
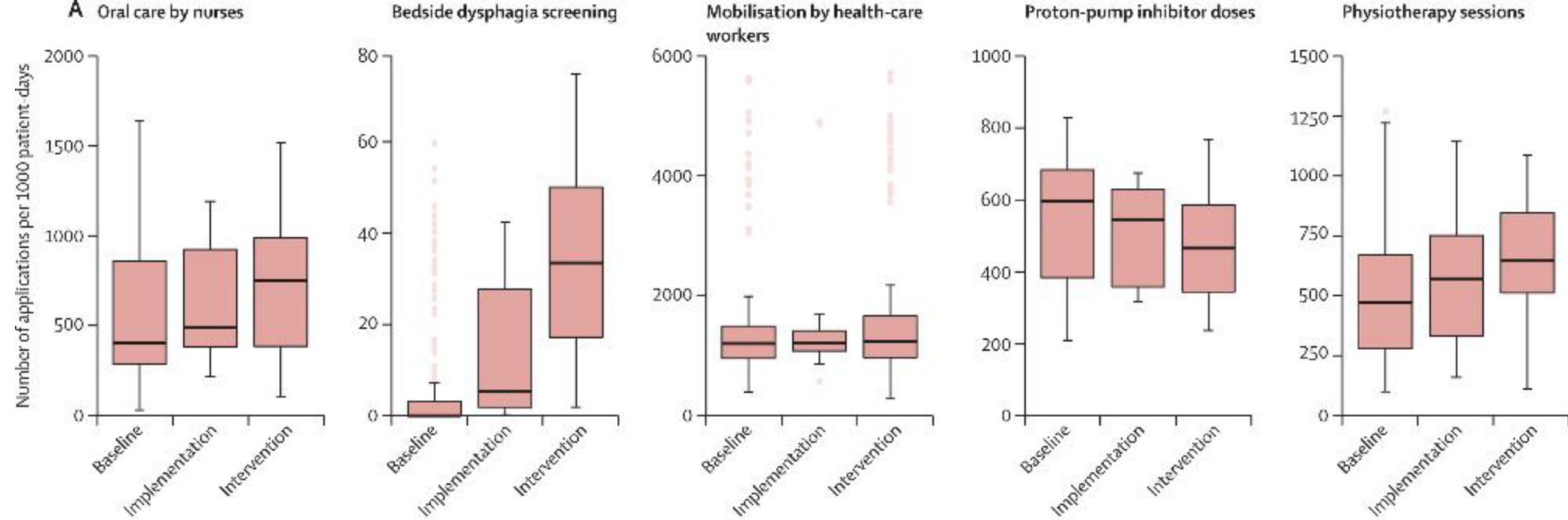
The Joint Commission – Additional Actions

Educate	Educate patients on risks and prevention methods that are implemented in the health care setting and should be continued after discharge.
Empower	Empower patients and family members to ask for assistance with oral care, feeding, and mobility and obtaining any needed supplies.
Perform	Perform outcome and/or process surveillance to determine rates of infection and compliance with processes to prevent NVHAP
Consider	Consider using standardized definitions such as those promulgated by the CDC to allow for aggregation of data and comparison infection rates with other organizations.
Develop	Develop operational NVHAP tracking systems to assess the impact of prevention initiatives
Implement	Implement processes to sustain NVHAP prevention for the long term

Prevention of non-ventilator-associated hospital-acquired pneumonia in Switzerland: a type 2 hybrid effectiveness–implementation trial

Aline Wolfensberger MD^{a †}  , Lauren Clack PhD^{a b †}, Stefanie von Felten PhD^c,
Mirjam Faes Hesse MA^a, Dirk Saleschus MA^a, Marie-Theres Meier RN^a, Katharina Kusejko PhD^a,
Prof Roger Kouyos PhD^a, Prof Leonhard Held PhD^c, Hugo Sax MD^{a d}

- NVHAP bundle implementation
 - Reduced NVHAP by 31%



Key Points



HAP is the most common Healthcare-Associated Infection; led by NV-HAP



Prevention of VAE/VAP improves patient outcomes, costs & length-of-stay



Call to action to start monitoring, and actively working to prevent, NV-HAP

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