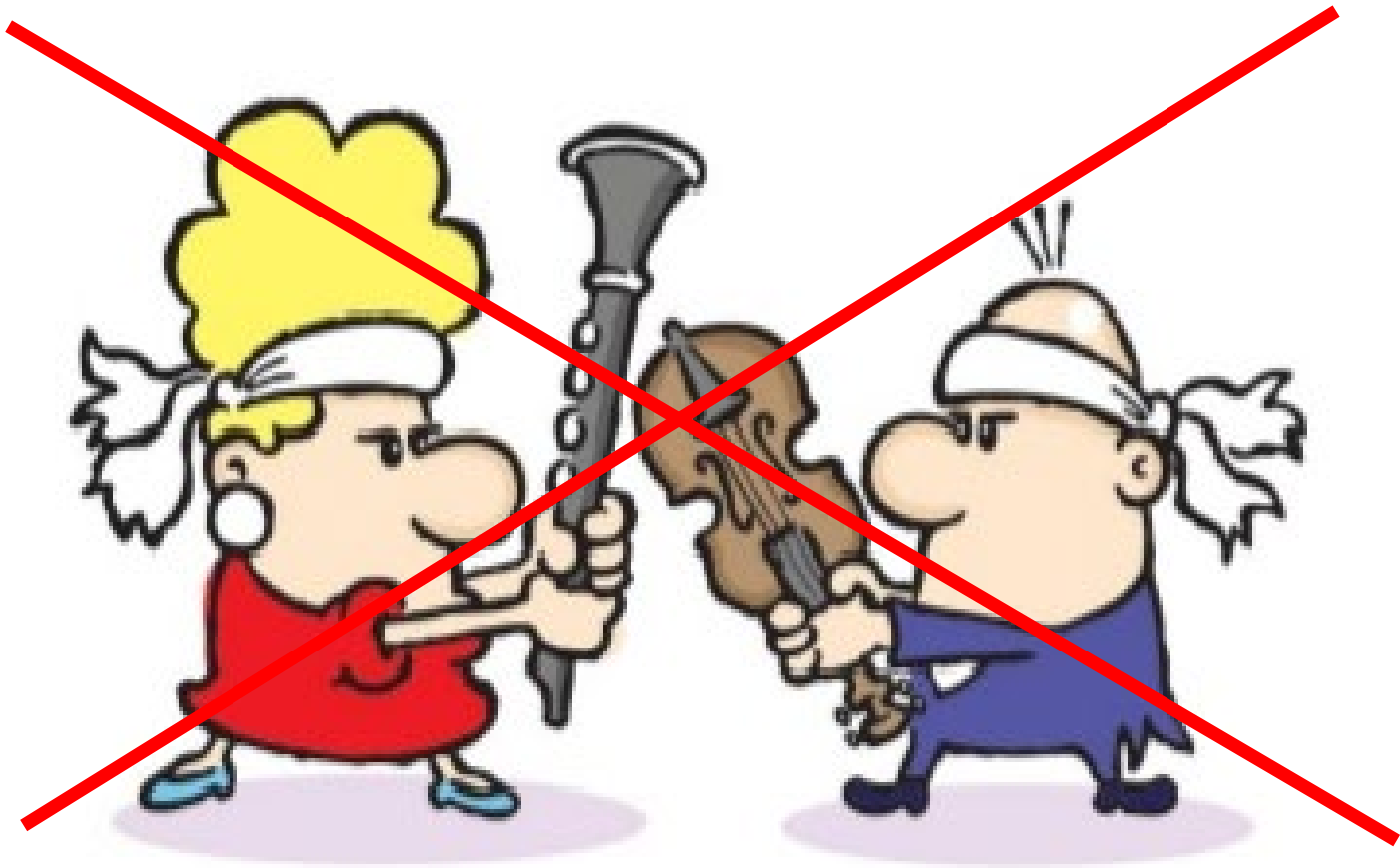




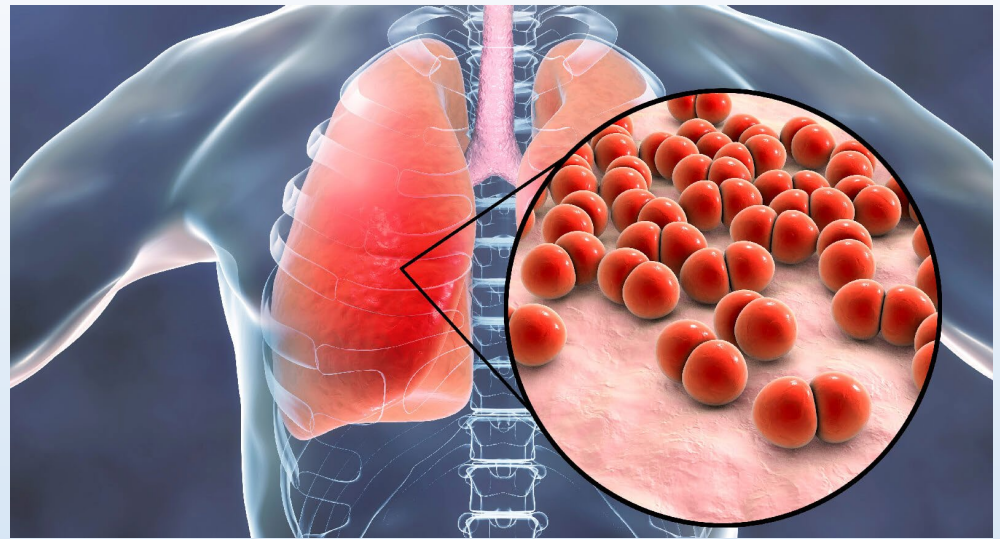
ventilátorová pneumonie na JIP

Jan Máca
KARIM FNO a LF OU

Conflict of Interest

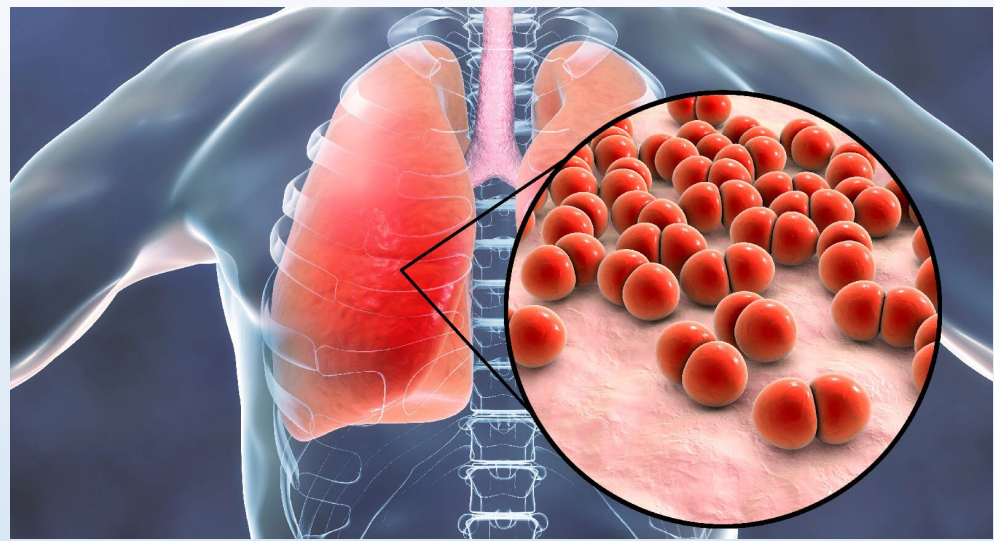


VAP



- definice - VAP vs. VAE
- etiopatogeneze
- diagnostika
- prevence
- ~~terapie~~

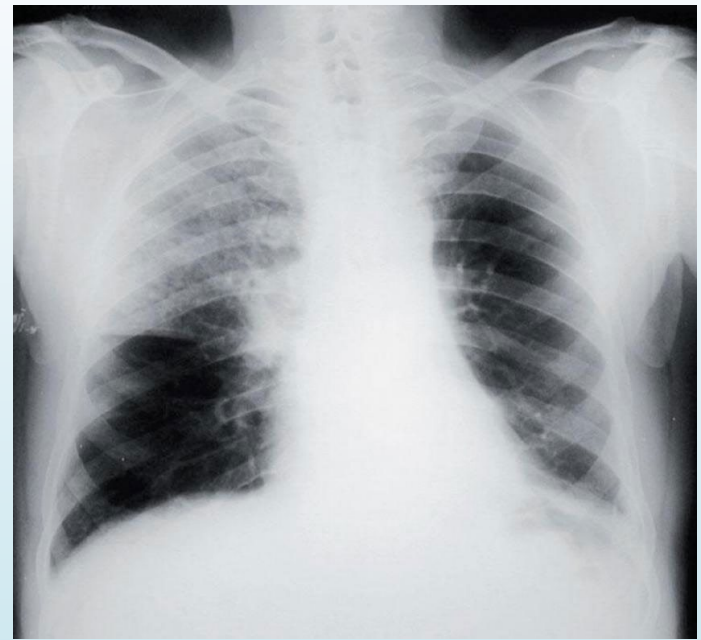
VAP



- **definice - VAP vs. VAE**
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VAP

klinické a epidemiologické dělení pneumonií



- **CAP** (community-acquired pneumonia)
- **HAP** (healthcare-associated pneumonia)
- **VAP** (ventilator-associated pneumonia)

healthcare-associated
infection

ventilator-associated pneumonia

VAP – časové dělení

1) časná

vznik v průběhu prvních **4 dnů** UPV

2) pozdní

vznik od **5. dne** UPV a dále

VAP

- výskyt až u **30%** pacientů na UPV
- mortalita **20-50%**
- **až 70%** - u rezistentních patogenů
- přítomnost VAP ↑ mortalitu o **30%**
- ↑ délku UPV
- ↑ délku hospitalizace
- ↑ náklady

VAE

VAEs – ventilator associated events



Centers for Disease Control and Prevention
CDC 24/7: Saving Lives, Protecting People™

reakce na **problematické** zikávání relevantních epidemiologických dat týkajících se **VAP**

2 důvody:

- radiologicko-diagnostický (rtg, dif. dg.)
- ekonomicko-společenský (zero VAP)

VAEs: definice

zhoršení oxygenace (**↑ potřeby FiO_2**) nebo **↑ PEEP** po **2 dnech** stabilního nastavení ventilačně/oxygenačních parametrů

VAE is **not synonymous** with VAP

Only **~25%–33%** of VAEs are due to pneumonia

most VAE are caused by **pneumonia, fluid overload, atelectasis, and/or ARDS**

VAEs – *ventilator associated events*



Centers for Disease Control and Prevention
CDC 24/7: Saving Lives, Protecting People™

filozofie VAEs

- nejde o klinickou definici
- není určena pro klinické použití

výhodou je snaha:

1. brát vážně ventilačně/oxygenační **změny**
2. o **včasné** nalezení jejich příčiny
3. zároveň se **vyvarovat** změn léčebné strategie při nevýznamných příčinách („ATB terapie atelektáz“)

problémem může být jejich interpretace poskytovatelem zdravotní péče, kontrolními institucemi

automated VAE detection (AI?)

VAEs in adults

1. ventilator-associated conditions (**VACs**)
2. infection-related ventilator-associated complications (**IVACs**)
3. possible VAP (**PVAP**)

VAEs and **VACs** are synonymous and are defined as:

- **↑** in the daily minimum **PEEP of ≥ 3 cmH₂O** sustained for ≥ 2 calendar days after ≥ 2 days of stable or decreasing daily minimum PEEP
- nebo*
- **↑ FiO₂ of ≥ 0.2** points sustained for ≥ 2 days after ≥ 2 days of stable or decreasing daily minimum FiO₂ levels

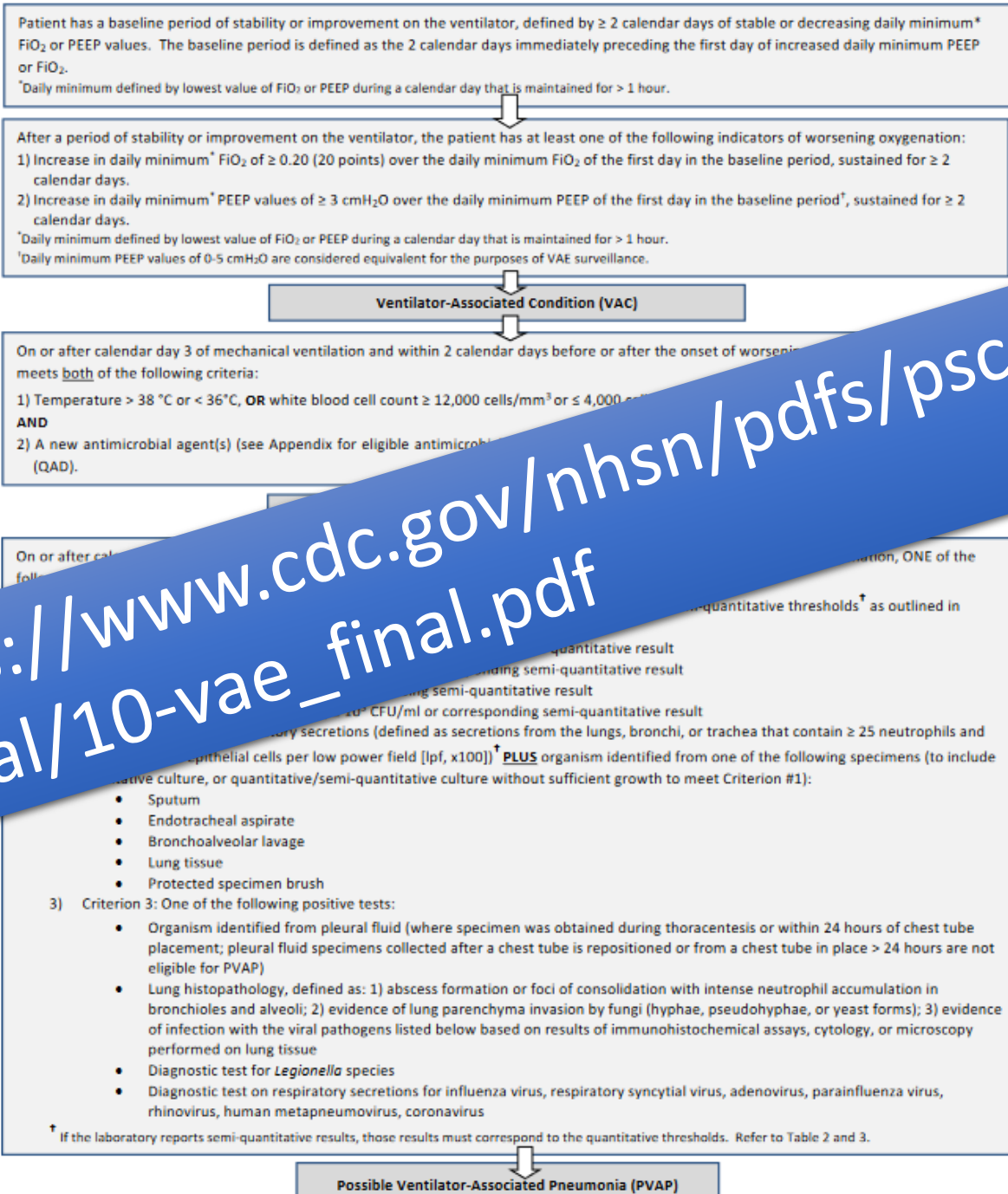
IVAC is defined as

- VAC with concurrent indications of **possible infection**, namely an abnormal temperature (**$< 36^{\circ}\text{C}$ or $> 38^{\circ}\text{C}$**) or white blood cell count (**$\leq 4,000$ or $\geq 12,000$ cells/mm³**)
- a*
- 1 or more **new antibiotic starts** that continue for ≥ 4 days, all beginning within 2 days before or 2 days after VAC onset

PVAP is defined as

- IVAC with indications that infection might be **localized to the lungs**. It requires **positive respiratory secretion** cultures, **pleural fluid**, positive assays for respiratory **viruses or Legionella**, or suggestive **histopathology** concurrent with the IVAC.
- The culture criterion can be fulfilled via **quantitative cultures** above various thresholds that vary depending upon specimen type or through **positive cultures** with any amount of growth if there is concurrent Gram-stain evidence of purulence

Figure 1: Ventilator-Associated Events (VAE) Surveillance Algorithm



https://www.cdc.gov/nhsn/pdfs/pscmanual/10-vae_final.pdf

potential risk factors for VAE

potential risk factors for VAE:

- **sedatives** (especially benzodiazepines and propofol)
- **opioids**
- **positive fluid balance**
- mandatory modes of mechanical ventilation with **high tidal volumes** and/or **high inspiratory driving pressures**
- **blood transfusions**
- **oral care** with **chlorhexidine**
- **stress ulcer prophylaxis**
- patient **transport**
- **gastric retention**
- **reintubation**
- **neuromuscular blockade**

interventions associated with ↓ VAE rates in interventional trials include:

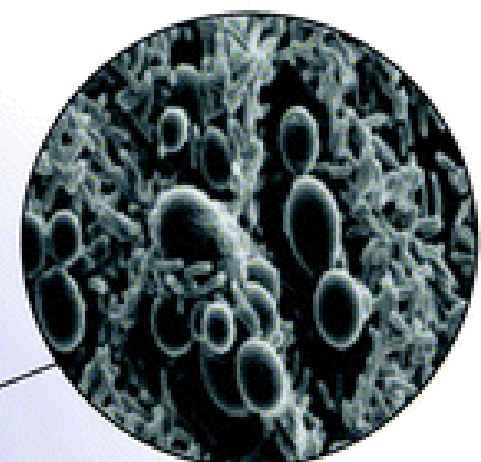
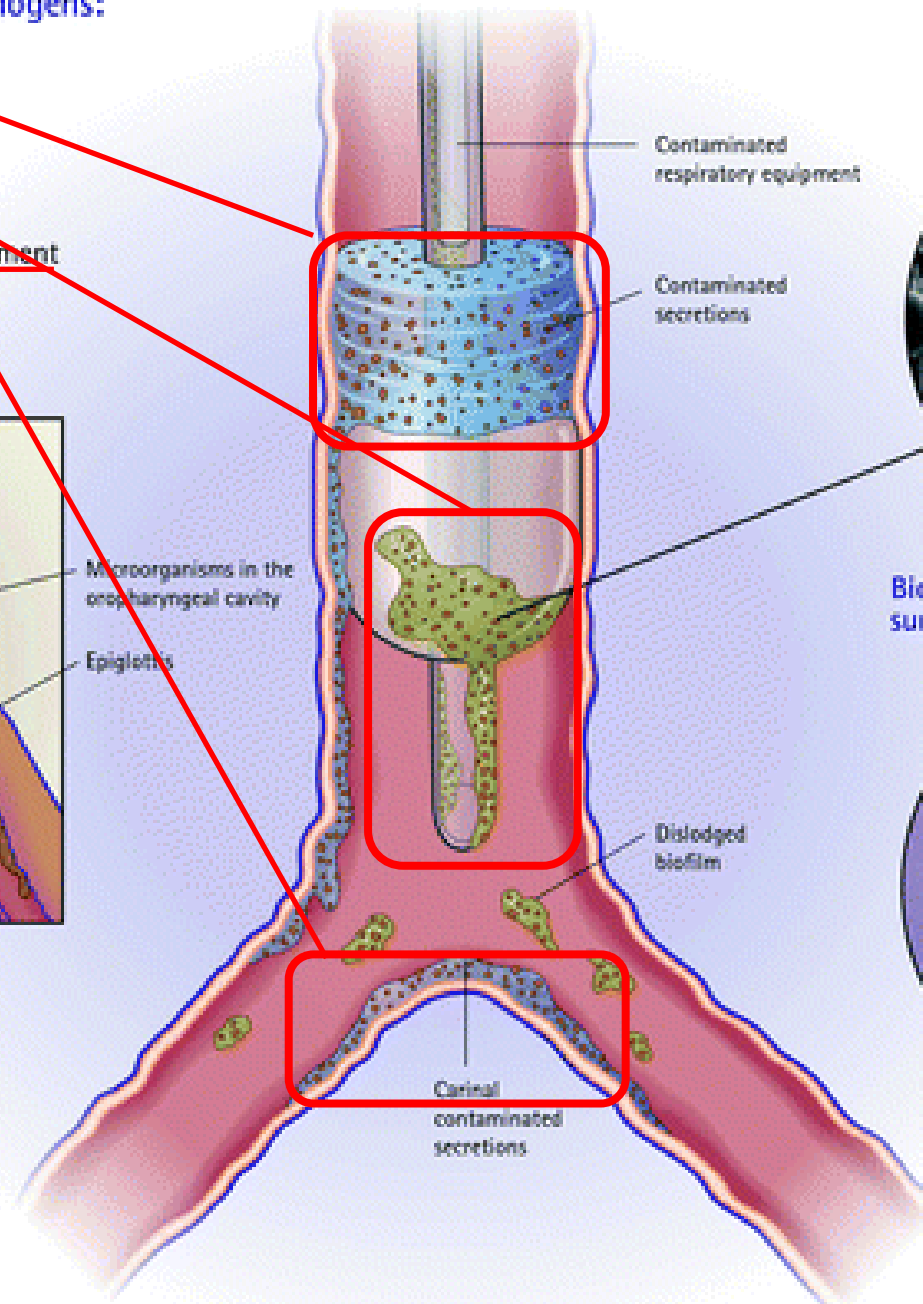
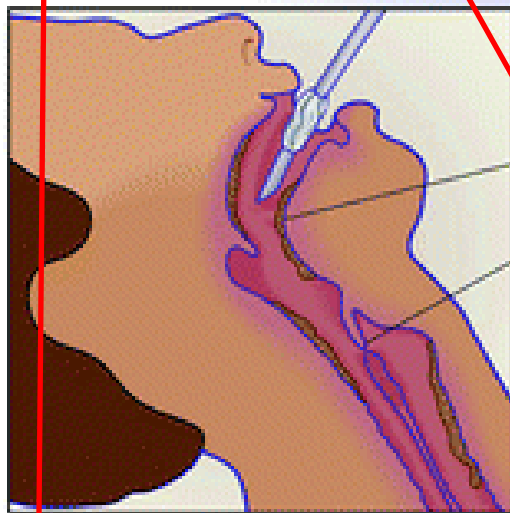
- spontaneous **awakening trials**
- spontaneous **breathing trials**
- **conservative fluid management**
- **dexmedetomidine**

VAP

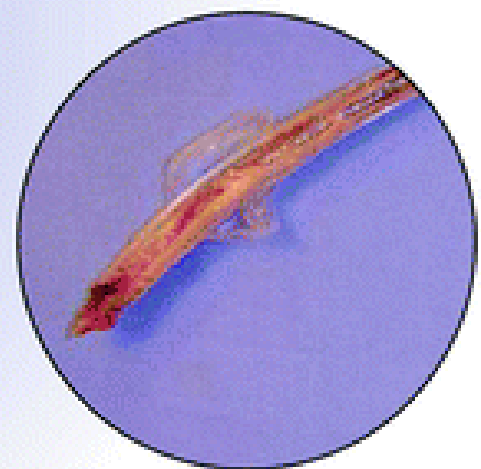
- definice - VAP vs. VAE
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- prevence

Common sources of VAP pathogens:

- Aspiration
- Intubation procedure
- Biofilm formation
- Contaminated secretions
- Contaminated respiratory equipment



Biofilm formation on inner and outer surface of the endotracheal (ET) tube



ET tube upon extubation

VAP

- definice - VAP vs. VAE
- etiopatogeneze
- **diagnostika**
- prevence

VAP - diagnostika

1. klinická manifestace
2. zobrazovací techniky
3. metody vyšetření a interpretace tracheálních a bronchoalveolárních vzorků
4. biomarkery

základní fyzikální vyš.

rtg, HRCT, UZ

invazivní vs. neinvazivní metody (TBA, BAL, PBS)

PCT, CRP, IL-6, sCD14-ST, sTREM-1

exhaled breath condensate?

benzen, cyklohexanon, pentanol a undecyl aldehyd

VAP – diagnostika (*Johanson*)

clinical, laboratory, radiographic, and microbiological criteria



- **nový nebo progredující infiltrát na rtg**

+ alespoň 2 další kritéria:

- **leukocytóza** $> 12\ 000$ nebo **leukopenie** $< 4\ 000 \times 10^9$
- **febrilie** $> 38^\circ\text{C}$
- změna **množství** anebo změna **charakteru** (*purulentní*) sekrece

ATS Consensus Conference (2005)

substantial interobserver variability

| CPIS points | 0 | 1 | 2 | |
|---|--------------------------------|---|--|---|
| 1. Tracheal secretions | Rare | | | |
| 2. Chest X-ray infiltrates | No infiltrate | | | |
| 3. Temperature, °C | ≥ 36.5 and ≤ 38.4 | | | |
| 4. Leukocytes count, per mm ³ | $\geq 4,000$ and $\leq 11,000$ | | | |
| 5. PaO ₂ /FiO ₂ , mmHg | > 240 or ARDS | | | |
| 6. Microbiology | Negative | | | |
| Footnote to Table 2. The modified CPIS | | | | |
| | | Oxygenation: PO ₂ /FiO ₂ ratio (mmHg) | Over 240 or existence of ARDS signs | 0 |
| | | | Less or equal to 240 and absence of ARDS signs | 2 |
| | | Chest X-ray | Existence of infiltration | 0 |
| | | | Disseminative infiltration | 1 |
| | | | Local infiltration | 2 |
| WBC: White blood cell | | | | |

Clinical Pulmonary Infection Score - CPIS

MCPIS

VAP - diagnostika

vitální požadavek je **rychlost** stanovení diagnózy

- u sepsy do **1 (-2) hod** od stanovení diagnózy je nutno nasadit ATB terapii
- jakékoliv prodlení významně negativně ovlivňuje outcome pacienta

VAP

- definice - VAP vs. VAE
- etiopatogeneze
- diagnostika
- **prevence**

prevention VAP - dříve

General measures

- Universal infection control measures
 - Hygiene
- Multidisciplinary team approach
 - Staff : Patient ratio

Prevention of aspiration

- Elevation of the head of the bed
- Endotracheal cuff pressure
- Avoiding circuit manipulation
- Drainage of subglottic secretions

Preventive measures for VAP

Decontamination

- Oral decontamination
- Selective GI decontamination
- Silver Endotracheal tube

Early extubation

- Early weaning protocol
- Daily sedation brakes

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







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2022

- Society for Healthcare Epidemiology (SHEA)
- Infectious Diseases Society of America (IDSA)
- American Hospital Association (AHA)
- Association for Professionals in Infection Control and Epidemiology (APIC)
- Centers for Disease Control and Prevention



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2022

strategies to **prevent**

- **ventilator-associated pneumonia (VAP)**
- ventilator-associated events (VAE)
- non-ventilator hospital-acquired pneumonia (NV-HAP)

in adults, children, and neonates

update of the Strategies to Prevent Ventilator-Associated Pneumonia in Acute Care Hospitals published in **2014**



I. kategorizace doporučení pro **prevenci VAP**

essential practices

- **should be** adopted by **all acute-care hospitals**
- good evidence that the intervention ↓ the average **duration of IMV, LOS, mortality, VAEs, antibiotic utilization, and/or costs** benefits likely outweigh **risks**

*in 2014 these were “basic practices,” renamed to **highlight their importance** as foundational for hospitals’ healthcare-associated infection (HAI) prevention programs*

essential practices

changes



- added a recommendation for **HFNO** or **NIPPV** as options to **avoid intubation, minimize duration of intubation, and prevent reintubations** (compared to COT)
- added a recommendation for **spontaneous awakening trials** or **sedation protocols** as effective strategies to **minimize sedation in adults**
- added a recommendation for **daily toothbrushing**
- reclassified endotracheal tubes with **subglottic secretion drainage** from an Essential Practice to an **additional approach**



1. avoid intubation and prevent reintubation

- ✓ use HFNO or NIPPV as appropriate whenever safe and feasible

2. minimize sedation

- ✓ avoid benzodiazepines in favor of other agents
- ✓ use a protocol to minimize sedation
- ✓ implement a ventilator liberation protocol

3. maintain and improve physical conditioning

4. elevate the head of the bed to 30–45°

5. provide oral care with toothbrushing but without chlorhexidine

6. provide early enteral vs. parenteral nutrition

7. change the ventilator circuit only if visibly soiled or malfunctioning (or per manufacturers' instructions)

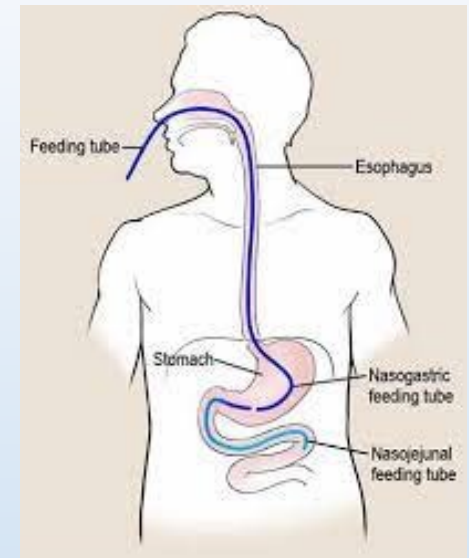
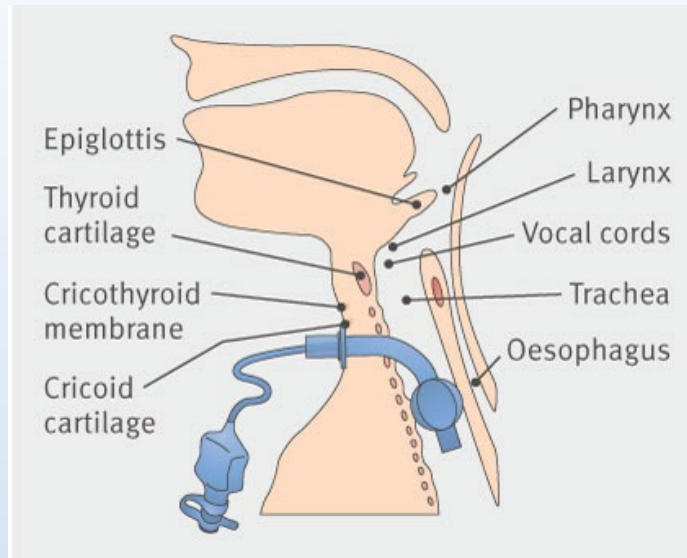


II. kategorizace doporučení pro **prevenci** VAP

additional approaches

- **considered** for use in locations and/or populations **within hospitals** when these **HAIs are not controlled after implementation of essential practices**
- 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 IMV, LOS, or mortality

in 2014 these were “special approaches”



additional approaches

changes

- **reclassified** endotracheal tubes with **subglottic secretion drainage** as an **additional approach** rather than an **essential practice** for adults and older children
- **added** a recommendation to consider **early tracheostomy**
- **added** a recommendation to consider **postpyloric** rather than **gastric feeding** in patients at high risk for aspiration

additional approaches

doporučení

1. use **selective oral** or **digestive decontamination** in countries and ICUs with **low prevalence of antibiotic-resistant organisms**
2. utilize endotracheal tubes with **subglottic secretion drainage** ports for patients expected to require **>48–72 hours of IMV**
3. consider **early tracheostomy**
4. consider **postpyloric** rather than **gastric feeding** for patients with **gastric intolerance** or at **high risk for aspiration**



III. kategorizace doporučení pro **prevenci VAP**

generally not recommended

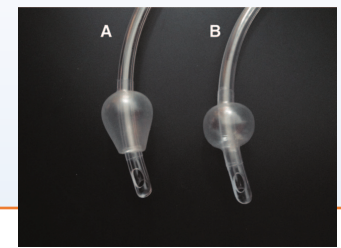
- **inconsistently** associated with **lower VAP rates**
- **no** or **negative** impact on duration of IMV, LOS, or mortality.
- **no impact** on **VAP rates**, **average impact** on duration of IMV, LOS, or mortality

not recommended

- **no** impact on **VAP rates** or other patient **outcomes**, **unclear** impact on **costs**

generally not recommended

Quality of Evidence:
MODERATE



1. tapered endotracheal tube cuffs
2. ultrathin polyurethane endotracheal tube cuffs
3. probiotics
4. frequent endotracheal cuff pressure monitoring
5. automated control of endotracheal cuff pressures
6. oral care with chlorhexidine, chlorhexidine bathing
7. silver-coated endotracheal tubes
8. kinetic beds
9. prone positioning
10. stress-ulcer prophylaxis
11. monitoring residual gastric volumes
12. early parenteral nutrition



not recommended

1. closed endotracheal suctioning systems



implementation of VAP, VAE, and NV-HAP prevention strategies

engagement

Develop a multidisciplinary team

at minimum: **unit directors, physicians, nurses, and respiratory therapists**
other: infection preventionists, pharmacists, nutritionists, physical therapists, occupational therapists, family members, and patient advocates.

Involve local champions

medical director, **nursing** director, **charge** nurses, director of **respiratory therapy**, engaged **frontline staff** (informal leaders)

Utilize peer networks

Horizontal networking of **peers** across hospitals

Comparing progress and benchmarks between ICUs can help units better understand their local strengths and weaknesses, learn from best practices, brainstorm solutions to common problems

implementation of VAP, VAE, and NV-HAP prevention strategies

education

Provide education sessions

introduction of evidence-based practices in the clinical setting

workshops, hands-on training, conferences, slide presentations, and/or interactive discussions, simulations

Provide educational materials

smartphone applications, interactive websites, pocket cards, brochures, posters, fact sheets, daily guides, guideline summaries, flow sheets and 1-page bulletins

implementation of VAP, VAE, and NV-HAP prevention strategies

execution

Standardize care processes

implementation of guidelines, bundles, protocols or pathways

establish **new care processes** as “normal behaviors”

daily **multidisciplinary rounds** are widely recommended

Create redundancy

reminders about best practice and can take the form of posters, bulletins, pens, stamps, pocket cards, 1-page signs, daily goal lists in patient rooms, checklists, and preprinted order sets, text messages, and screensavers on clinical computers

engage family members to assist with preventive care

implementation of VAP, VAE, and NV-HAP prevention strategies

evaluation

Measure performance

frequent **formal** and **informal audits** of clinical practice

measuring process and outcome measures

evaluating performance provides an ongoing, real-time image of actual implementation rates

analyze all or a representative sample of VAEs for **etiology and preventability**.
pneumonia, pulmonary edema, acute respiratory distress syndrome, and atelectasis are the precipitants for most VAEs

Provide feedback to staff

provide **regular feedback** on process and/or outcome data to staff

Feedback helps pinpoint **new areas for improvement** and marks successful transitions to new standards of care

VAP závěr



- VAP je **závažná** infekční komplikace v intenzivní péči
- diagnostika je **složitá** a **variabilní** (SE a SP nízká)
- je třeba na ni **myslet**
- diagnostika musí být **rychlá**
- nejasná budoucnost implementace **VAE**
- důraz na **důslednou aplikaci** preventivních opatření
- otázka **nezbytnosti** aktuálně **využívaných**, nicméně **nedoporučených preventivních opatření**
- důležitost **angažovanosti, edukace, vykonávání a vyhodnocování** prevence



děkuji za pozornost