



Do No Harm: Mitigating Risk Factors Vent and Non-Ventilator Pneumonia



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Disclosures

- △ Consultant-Michigan Hospital Association Keystone Center
- △ Subject matter expert on CAUTI, CLABSI, HAPI, Sepsis, Safety culture for HRET/AHA
- △ Consultant and speaker bureau
 - △ Stryker's Sage business
 - △ LaJolla Pharmaceutical
 - △ Potrero Medical
- ▲ Baxter Advisory Board

Session Objectives

- △ Define key fundamental evidence-based nursing care practices that reduce vent and non-vent HAP
- Discuss strategies to overcome barriers

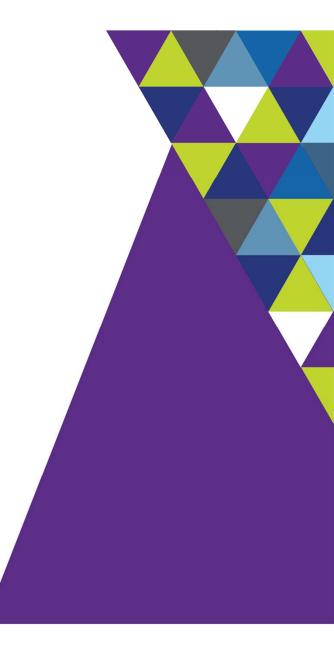


Notes on Hospitals: 1859

"It may seem a strange principle to enunciate as the very first requirement in a Hospital that it should do the sick no harm."

- Florence Nightingale

Advocacy = Safety



The Why

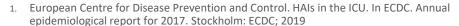




VAP

- 8.3% of ICU patient in Europe develop an HAI¹
- 6% is pneumonia with 97% associated with mechanical ventilation (9.5 VAP per 1000 intubation days)¹
- Most frequent organism pseudomonas aeruginosa¹
- △ VAP is associated with ↑ MV days and ↑ ICU & hospital LOS²
- △ Attributable mortality estimated to be 4.0– 13.5% ²
- Financial cost of a VAP episode has been estimated as approximately 15,000 to 40,000 US²







Building Blocks to Best Practice in Caring for Mechanically Ventilated Patients



Ventilator Bundle: HOB 30, Deep Vein Thrombosis (DVT) prophylaxis, Peptic Ulcer Disease (PUD) prophylaxis, Sedation interruption, Spontaneous breathing trial, daily care with chlorhexidine



VAP Bundle: HOB 30, Sedation interruption, Spontaneous breathing trial, oral care 6x per day, CHG rinse 2x per day, subglottic secretions drainage if expected to be ventilated > 72hrs

Risk Factor Categories for Hospital Acquired Pneumonia

Factors that increase bacterial burden or colonization

Factors that increase risk of aspiration



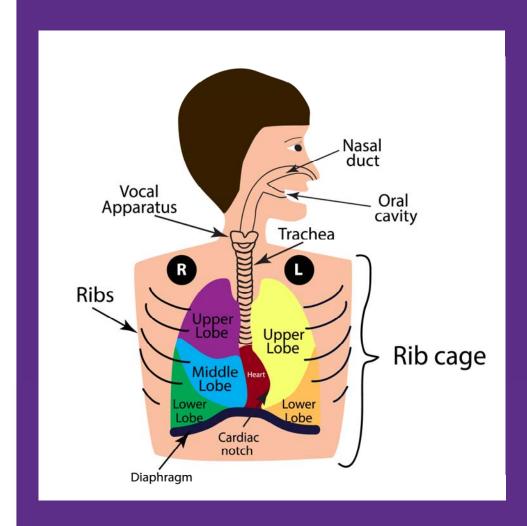
Single Ecosystem

- Entire respiratory tract is one ecosystem¹
 - △ Upper-nasal and oral cavities
 - △ Lower-alveoli
- ▲ Not sterile environment¹
- Oral flora changes in hospitalized patients²
- Relationship between dental plaque and pulmonary lavage fluid³



Johanson WG, et al. N Engl J Med. 1969 Nov 20;281(21):1137-40

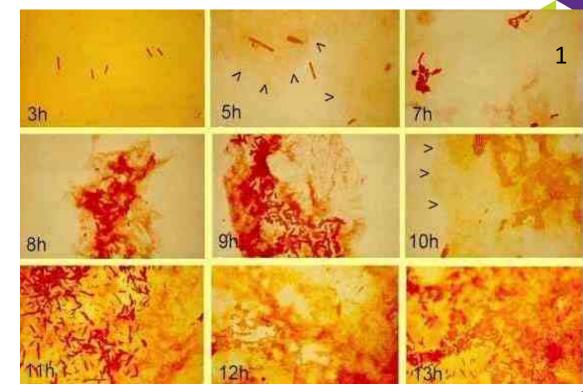




Where does Pneumonia Start: Oral Bacteria during Hospitalization & Illness



- Oral cavity¹
 - > 1 billion oral microbes
 - 700-1000 species
 - Replicate's 5 x in 24hr period
- Disruption of Microbiome²
 - Plaque, gingivitis, tooth decay
 - Reduced salivary flow/change in pH
- △ 24-48 hours for HAP pathogens in mouth³
- If aspirated =100,000,000 bacteria/ml saliva into lungs⁴



- Scannapieco FA, Stewart EM, Mylotte JM.. Crit Care Med. 1992;20:740-745.



Oral Cavity & VAP

- ▲ 89 critically ill patients
- Examined microbial colonization of the oropharynx through out ICU stay
- △ Used pulse field gel electrophoresis to compare chromosomal DNA
- Results:
 - △ Diagnosed 31 VAPs
 - △ 28 of 31 VAPs the causative organism was identical via DNA analysis



- △ 49 elderly nursing home residents admitted to the hospital
- Examined baseline dental plaque scores & microorganism within dental plaque
- Used pulse field gel electrophoresis to compare chromosomal DNA
- Results
 - △ 14/49 adults developed pneumonia
 - △ 10 of 14 pneumonias, the causative organism was identical via DNA analysis

Risk Factor Categories for Hospital Acquired Pneumonia

Factors that increase bacterial burden or colonization

Factors that increase risk of aspiration



Micro Aspiration during Sleep in Healthy Subjects

- Prospective duplicate full-night studies
- △ 10 normal male's 22-55 years of age
- Methods:
 - Radioactive 99 mTc tracer inserted into the nasopharynx
 - Lung scans following final awakening
 - No difference in sleep efficacy between 2 study nights

A Results:

50%

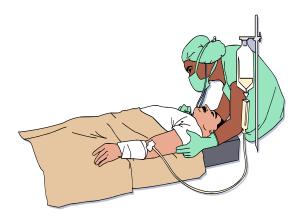
In the lung parenchyma

Body Position: Supine versus Semi-recumbent (30-45 degrees)



Methodology

- ▲ 19 mechanically ventilated patients
- 2 period crossover trial
- Study supine and semirecumbent positions over 2 days
- △ Labeled gastric contents (Tc 99m sulphur colloid)
- △ Measured q 30 min content of gastric secretions in endobronchial tree in each position
- Sampled ET secretions, gastric juice & pharyngeal contents for bacteria



Body Position: Supine versus Semi-recumbent

Results:

A Radioactive contents higher in endobronchial secretions in supine patients

Time dependent:

- Supine: 298cpm/30min vs. 2592cpm/300min
- HOB: 103cpm/30min vs.216cpm/300min



Same microbes cultured in all 3 areas

• HOB: 32%

• Supine: 68%

Oral Hygeine





Polling Question

- ▲ What is your current oral care regime at your facility?
 - △ CHG alone
 - △ Toothbrushing
 - △ Toothbrushing with CHG
 - △ Toothbrushing, CHG, cleansing swabs(Comprehensive kit)
 - △ Nothing

What Does the Evidence Tell Us?



Brush
CHG rinse alone
CHG rinse in Combination
Swab/Clean/Moisturize
Suction

All of the above

Comprehensive Oral Care Program

Literature Review: Oral Care Impact of VAP

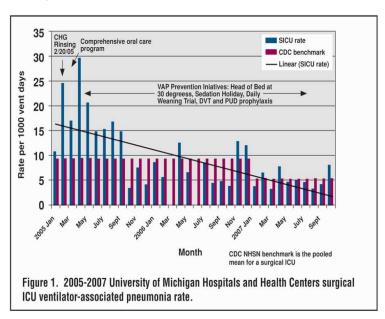
Comprehensive Oral Care:

- Reduction in VAP from 5.6 to 2.2¹
- Reduction in VAP from 4.10 (2005) to (2.15) in 2006 with addition of CPC & comprehensive oral care. Vent bundle & rotational therapy already being performed²
- Reduction in VAP from 12.0 to 8.0 (p=.060) with 80% compliance, vent bundle already being preformed, 1538 patients randomized to control or study group, Additional outcomes; ↓ vent days (p=.05), ↓ ICU LOS (p=.05) ↓ time to VAP (p=<.001) & reduction in mortality (p=.05) ³</p>
 - 1. Schleder B. et al. J Advocate Health 2002;4(1):27-30)
 - Powers J, et al. J Nurs Care Qual. 2007 Oct-Dec;22(4):316-21
 - 3. Garcia R et al AJCC, 2009;18:523-534)

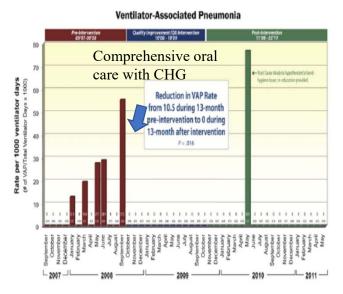
Literature Review: Oral Care Impact of VAP

Comprehensive Oral Care & CHG:

• Reduction in VAP to zero for 2 years, vent bundle, mobility, oral care & CHG with comprehensive education preformed (Murray TM et al. AACN Advanced Critical Care. 2007;18(2):190-199)



Dickinson S et al. SCCM Critical Connections, 02/2008



Heck K, et al. American Journal of Infection Control 40 (2012) 877-9





▲ Klompas Study-Retrospective review

- △ Single center
- △ Impact of vent bundle (5536 patients)
- △ Connection of CHG with increase mortality on patients vented > 3 days

△ Deschepper study: Retrospective Review

- △ Hospital wide retrospective cohort (82,274 patients)
- △ 11,133 patients received CHG oral care
- △ Divided into low exposure-cumulative dose < 300 mg (8080 pts)
- \triangle High exposure > 300 mg (3053 pts)
- △ 300 mg CHG is equivalent to 1 bottle of 250ml of oral care soln at .12%-covers 5-6 days at 3 times a day)
- In the sickest group CHG low or high exposure was not a risk for increased mortality
- Showed improvement on mortality in ICU patients ventilated < 96hrs and not harm if vented > 96 hrs
- Greatest risk for mortality increase is use in non-ICU patients.

Cochrane Meta-Analysis 2020 of RCT's



Analysis 1.1. Comparison 1: Chlorhexidine versus placebo/usual care, Outcome 1: Incidence of VAP

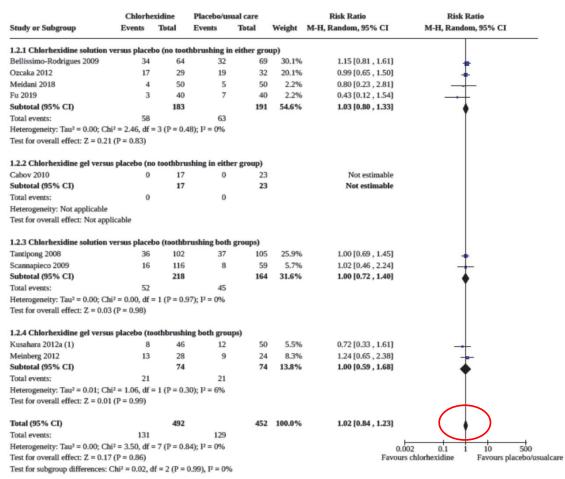
	Chlorhexidine Placebo			ual care		Risk Ratio	Risk Ratio	
Study or Subgroup	Events	Total	Events	Total	Weight	M-H, Random, 95% CI	M-H, Random, 95% CI	
1.1.1 Chlorhexidine solution	versus place	bo (no too	thbrushing in	either gro	up)			
Fu 2019	7	40	37	40	9.0%	0.19 [0.10, 0.37]		
Meidani 2018	6	50	15	50	7.6%	0.40 [0.17, 0.95]		
Grap 2011 (1)	7	21	10	18	8.6%	0.60 [0.29 , 1.25]		
Ozcaka 2012	12	29	22	32	10.6%	0.60 [0.37, 0.98]	_	
Bellissimo-Rodrigues 2009	16	64	17	69	9.7%	1.01 [0.56 , 1.83]		
Quon 2017	4	8	2	8	4.5%	2.00 [0.50 , 8.00]		
Subtotal (95% CI)		212		217	50.0%	0.57 [0.33 , 1.00]		
Total events:	52		103					
Heterogeneity: Tau ² = 0.33; Ch		f = 5 (P = 0		%			- 1	
est for overall effect: Z = 1.9		-5(1-0	.005),1 - 72					
.1.2 Chlorhexidine gel versu	u placeho (n	a taathbru	ching in sith	er group)				
Cabov 2010	1 piacebo	17	6	23	2.6%	0.23 [0.03 , 1.70]		
Koeman 2006	13	127	23	130				
Subtotal (95% CI)	10	144	20	153		0.53 [0.29 , 0.97]		
Total events:	14	144	29	155	12.070	0.35 [0.25 , 0.57]	•	
Heterogeneity: Tau ² = 0.00; Ch		- 1 /B - 0 °					- 1	
Test for overall effect: Z = 2.04		- I (P - 0.3	30), 1 - 076				- 1	
rest for overall effect. 2 - 2.0	f (F = 0.04)							
1.1.3 Chlorhexidine solution	versus place	bo (toothb	rushing both	groups)				
Tantipong 2008	5	58	10	52	6.6%	0.45 [0.16 , 1.23]		
Scannapieco 2009 (2)	14	97	12	49	8.9%	0.59 [0.30 , 1.18]		
Berry 2011 (3)	4	33	1	43	2.4%	5.21 [0.61 , 44.47]		_
Subtotal (95% CI)		188		144	17.8%	0.74 [0.29 , 1.89]		
Total events:	23		23					
Heterogeneity: Tau ² = 0.36; Ch	ni ² = 4.30. df	= 2 (P = 0.1	12): F = 53%				- 1	
Test for overall effect: Z = 0.64		-(-	-,, -					
1.1.4 Chlorhexidine gel versu	s placeho (te	anthbrushi	ng both grou	ns)				
Kusahara 2012a (4)	15	46	16	50	9.8%	1.02 [0.57 , 1.82]		
Meinberg 2012	18	28	11	24			T.	
Subtotal (95% CI)	10	74	- 11	74		1.22 [0.83 , 1.79]	K	
Total events:	33	/4	27	74	20.270	1.22 [0.03 , 1./3]	•	
rotar events: Heterogeneity: Tau² = 0.00; Ch		= 1 (P = 0						NINIT 4
		- 1 (F = 0.4	1, 1 - 076					NNT 1
	J (P = 0.32)							· - ·
Test for overall effect: Z = 1.00		618		588	100.0%	0.67 [0.47, 0.97]		
		919						
Total (95% CI)	122	918	182					
otal (95% CI) otal events:				66%			0.02 0.1 10	50
Fest for overall effect: Z = 1.00 Fotal (95% CI) Fotal events: Heterogeneity: Tau ² = 0.26; Ci Fest for overall effect: Z = 2.14	ni² = 35.29, di			66%			0.02 0.1 10 s chlorhexidine Favours pla	

Analysis 3.1. Comparison 3: Toothbrushing versus no toothbrushing, Outcome 1: Incidence of VAP

	Toothbrushing		No toothbrushing		Risk Ratio Weight M-H, Random, 95% CI		Risk Ratio M-H, Random, 95% CI	
Study or Subgroup	Events Total		Events Total					
3.1.1 Powered toothbru	ısh + usual	care (± CI	IX) versus u	ısual care (± CHX)			
Pobo 2009 (1)	15	74	18	73	23.5%	0.82 [0.45 , 1.50]	-	
Yao 2011 (2)	4	28	14	25	12.7%	0.26 [0.10, 0.67]		
Subtotal (95% CI)		102		98	36.2%	0.49 [0.16, 1.53]		
Total events:	19		32					
Heterogeneity: $Tau^2 = 0$.	52; Chi ² = 4	4.05, df = 1	(P = 0.04); I	$^{2} = 75\%$				
Test for overall effect: Z	= 1.23 (P =	0.22)						
3.1.2 Toothbrush + CH	X versus C	HX alone						
Lorente 2012	21	217	24	219	25.7%	0.88 [0.51, 1.54]	_	
De Lacerda 2017	17	105	28	108	26.4%	0.62 [0.36, 1.07]	-	
Subtotal (95% CI)		322		327	52.1%	0.74 [0.50, 1.09]	A	
Total events:	38		52				•	
Heterogeneity: Tau ² = 0.	.00; Chi ² = 0	0.77, df = 1	(P = 0.38); I	$^{2} = 0\%$				
Test for overall effect: Z	= 1.53 (P =	0.13)						
3.1.3 Toothbrush + pov	idone iodir	ie versus p	ovidone iodi	ine alone				
Long 2012	4	31	11	30	11.6%	0.35 [0.13, 0.98]		
Subtotal (95% CI)		31		30	11.6%	0.35 [0.13, 0.98]		
Total events:	4		11				~	
Heterogeneity: Not appl	icable							
Test for overall effect: Z	= 1.99 (P =	0.05)						
Total (95% CI)		455		455	100.0%	0.61 [0.41, 0.91]		
Total events:	61		95					
Heterogeneity: $Tau^2 = 0$.	.08; Chi ² = 6	5.71, df = 4	(P = 0.15); I	$^{2} = 40\%$		0.0	01 0.1 1 10	
Test for overall effect: Z	= 2.44 (P =	0.01)					Toothbrushing No toothb	

Impact on Mortality

Analysis 1.2. Comparison 1: Chlorhexidine versus placebo/usual care, Outcome 2: Mortality



Zhao T, et al. Cochrane Database Syst Rev. 2020 Dec 24;12:CD008367

It is More than CHG

- △ .12% CHG application 2x daily is a small part of the oral care equation
- △ It is the comprehensive and frequent delivery of oral hygiene, including toothbrushing and cleansing



Ventilated Patients

- △ Assessment completed within 6hrs of admission & then q 12hrs
- △ Standardized oral assessment tool
- △ Toothbrushing should occur X2 daily additionally oral cleansing with swabs, suctioning and moisturization of the mouth q 2-4hrs

△ Tools to use

- Pediatric toothbrush followed by suctioning or Suction toothbrush (consider using a single use)
- · Swab for cleaning and moisturizing/suction swab if available to suction debris with cleaning
- Consider using oral care tools & supplies that can be kept at the bedside

△ Oral care cleansing solutions

- Use of an oral antiseptic rinse like CHG or CPC after brushing or in combination with comprehensive oral care
- Advise caution with routine use of CHG/consult team
- With swab cleaning use CPC, 1.5% H2O2 or sterile water



Does Compliance Make A Difference?

Oral care compliance & use of the ventilator bundle resulted in a 89.7% reduction in VAP

VAP rates for the years of the study



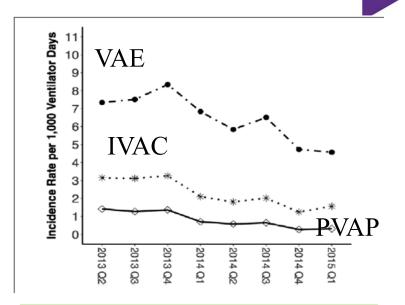
Compliance rates for the years of the study





Impact of a New Bundle/2 State Collaborative

- △ 38 hospitals, 56 ICU's in 2 states from October 2012 to March 2015
- Evidence based interventions, teamwork & safety culture
- Head-of-bed elevation, use of subglottic secretion drainage endotracheal tubes, oral care, chlorhexidine mouth care, and daily spontaneous awakening and breathing trials.



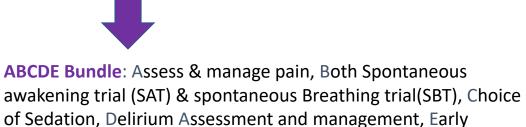
- VAE: 7.34 to 4.58 cases per 1,000 ventilator-days (p = 0.007)
- IVAC 3.15 to 1.56 per 1,000 ventilator days (p = 0.018)
- PVAP 1.41 to 0.31 cases per 1,000 ventilator-days (p = 0.012)

Building Blocks to Best Practice in Caring for Mechanically Ventilated Patients

Ventilator Bundle: HOB 30, Deep Vein Thrombosis (DVT) prophylaxis, Peptic Ulcer Disease (PUD) prophylaxis, Sedation interruption, Spontaneous breathing trial, daily care with chlorhexidine¹



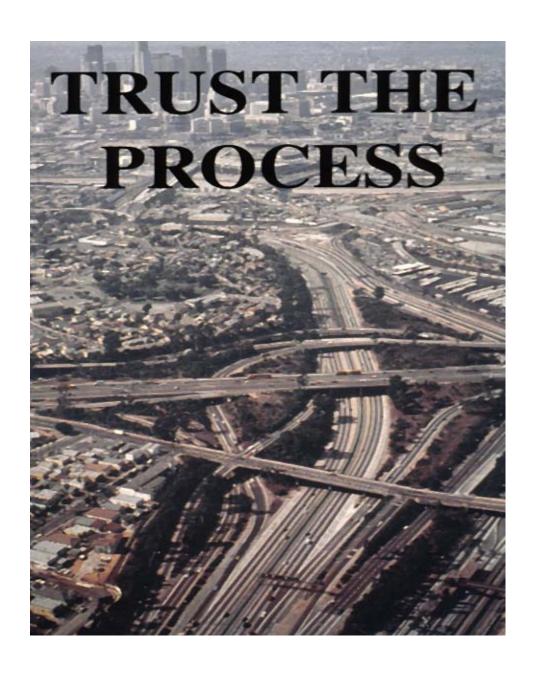
VAP Bundle: HOB 30, Sedation interruption, Spontaneous breathing trial, oral care 6x per day, CHG rinse 2x per day, subglottic secretions drainage if expected to be ventilated > 72hrs²



- 1. http://www.ihi.org/resources/Pages/Tools/HowtoGuidePreventVAP.aspx
- 2. Rawat N, et al. Crit Care Med, 2017;45:1208-1215
- 3. www.ICÚliberation.org

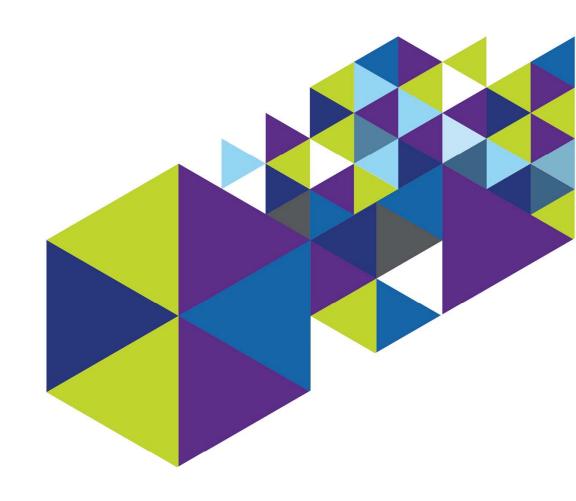
Mobility, Family and Patient Engagement³







Non-Vent Pneumonia: Addressing Risk Factors



Build the Will: NV-HAP Causes Harm

- △ HAP 1st most common HAI in U.S.^{1,2}
- △ 1 in every 4 hospital infections are pneumonia¹
 - △ 60% non-ventilator
- △ Increased mortality →15.5%-30.9%³
 - △ 8½ x more likely to die than equally sick patients who did not get non-vent HAP⁴
- △ Increased morbidity \rightarrow 50% are not discharged home^{5,6,7}
 - \triangle Extended LOS \rightarrow 7-9 days^{5,6,7}
 - \triangle Increased Cost \rightarrow \$36K to \$54K per case⁶
 - \triangle 2x likely for readmission <30 day^{5,6}
 - △ 46% ↑ ICU utilization^{5,6}
 - △ Increase antibiotic utilization⁸



- 2. Strassle PD, et al. Infect Control Hosp Epidemiol. 2020 Jan;41(1):73-79.
- . Giuliano K, et al. Am J of Infect Control. 2018;46:322-327
- Micek ST, et al. Chest. 2016 Nov;150(5):1008-1014.
- 5. Baker D, Quinn B et al. J Nurs Care Qual, 2019 1-7
- 6. Giuliano K, et al. Am J of Infect Control. 2018;46:322-327
- 7. Davis J et al. Pa Patient Safety Advisory, 2018;15(3)
- 8. Lacerna CC, et al. Infec control & Hosp Epidemiology 2020;41, 547-552



United Kingdom- Non-Ventilator HAP

- ▲ 1.5% of hospital inpatients in England have a hospital-acquired respiratory infection
- △ Over half are hospital-acquired pneumonia and are not associated with intubation.
- A Hospital-acquired pneumonia is estimated to increase hospital stay by about 8 days
- Reported mortality rate that ranges from 30–70%.
- Variations in clinical management and outcome occur across the UK.



Hospital-Acquired Pneumonia:

Non-Ventilated versus Ventilated Patients in Pennsylvania

Purpose:

△ Compare VAP and NV-HAP incidence, outcomes

Methods:

- Pennsylvania Database queried
- △ All nosocomial pneumonia data sets (2009-2016)



Results:

Table 1. Penns	Table 1. Pennsylvania Nosocomial Pneumonia Incidence and Number of Patients with NV-HAP or VAP Who Died									
Year	Number of NV-HAP Patients	Number of NV-HAP Patients Who Died	Percentage of Patients with NV-HAP Who Died (Confidence Interval)	Number of VAP Patients	Number of VAP Patients Who Died	Percentage of Patients with VAP Who Died (Confidence Limit)				
2009	1,977	364	18.41 (16.52–20.3)	922	163	17.68 (14.96–20.39)				
2010	1,848	366	19.81 (17.78–21.83)	737	144	19.54 (16.35–22.73)				
2011	1,780	318	17.87 (15.9–19.83)	643	127	19.75 (16.32–23.19)				
2012	1,620	307	18.95 (16.83–21.07)	571	112	19.61 (15.98–23.25)				
2013	1,528	285	18.65 (16.49–20.82)	767	160	20.86 (17.63–24.09)				
2014	1,419	256	18.04 (15.83–20.25)	901	199	22.09 (19.02–25.16)				
2015	1,427	277	19.41 (17.13–21.7)	912	218	23.90 (20.73–27.08)				
2016	1,380	280	20.29 (17.91–22.67)	980	221	22.55 (19.58–25.52)				
Total	12,979	2453	18.89%	6433	1344	20.89%				

- Mortality
- Incidence
- ▲ Total deaths
- Total cost
- Wide-spread



NV-HAP SMCS Research Findings: 2010

Incidence:

- △ 115 adults
- △ 62% non-ICU
- △ 50% surgical
- △ Average age 66
- Common comorbidities:
 - CAD, COPD, DM, GERD
- Common Risk Factors:
 - Dependent for ADLs (80%)
 - CNS depressant meds (79%)

24,482 patients and **94,247** pt days

Cost:

- \$4.6 million
- △ 23 deaths
- △ Mean Extended LOS 9 days
- △ 1,035 extra days



Quinn, B. et al. Journal of Nursing Scholarship, 2014. 46(1):11-19

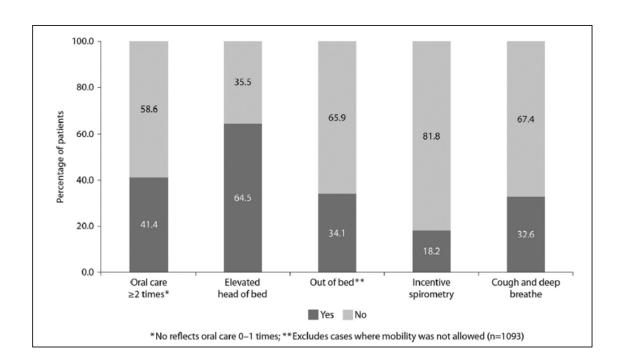
HAPPI-2 Incidence of Non-Ventilator Hospital-Acquired Pneumonia



- ▲ Multicenter retrospective chart review
- △ Extracted NV-HAP cases per the 2014 ICD-9-CM codes for pneumonia not POA and the 2013 CDC case definition
- 21 hospitals completed data collection
- △ Measured nursing care missed 24hrs before diagnosis
- Non-vent HAP occurred on every unit

HAPPI-2 Incidence of Non-Ventilator Hospital-Acquired Pneumonia

Missed nursing care 24 hours prior to Non-Vent HAP dx.





HAPPI-2 Incidence of Non-Vent Hospital-Acquired Pneumonia

Results:

- 1,300 NV-HAP (0.12-2.28 per 1,000 pt days)
 - △ 15.8% mortality
 - \triangle 50% < 66 yrs old
 - △ 63% non-surgical
 - △ 70.8% outside the ICU
 - △ 27.3 % in ICU
 - △ 18.8% transferred to ICU
 - △ 37.3% LOS >20 days
 - \triangle 57.7% LOS > 15 days
 - △ 40.6% admitted from home were discharged back to home
 - △ 19.3% readmitted within 30 days
 - \triangle \$36.4 -\$52.56 million in extra costs

- Med-Surg (43.1%; n = 560)
- Telemetry (8.5%; n = 111)
- Progressive (7.2%; n = 93)
- Oncology (4.9%; n = 64)
- Orthopedic (2.8%; n = 37)
- Neurology (1.5%; n = 19)
- Obstetric (0.2%; n = 3)

Is Pneumonia Part of the Sepsis Picture?

30-50% of sepsis cases may initiate with pneumonia¹

Site of infection	Frequency %		Mortality %	
	Male	Female	Male	Female
Respiratory	41.8	35.8	22.0	22.0
Bacteremia	21.0	20.0	33.5	34.9
Genitourinary	10.3	18.0	8.6	7.8
Abdominal	8.6	8.1	9.8	10.6
Device related	1.2	1.0	9.5	9.5
Wound/ soft tissue	9.0	7.5	9.4	11.7
Central nervous system	0.7	0.5	17.3	17.5
Endocarditis	0.9	0.5	23.8	28.1
Other/ unspecified	6.7	8.6	7.6	6.5

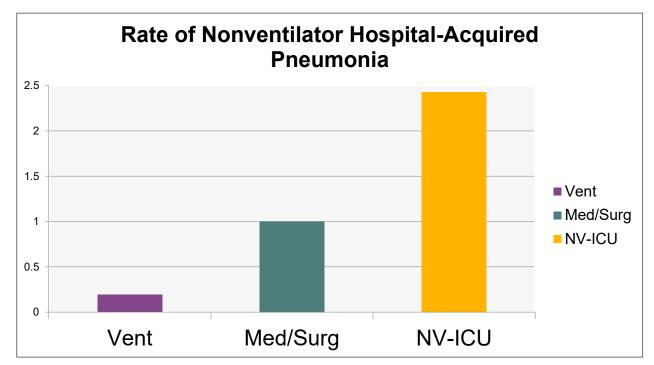
Risk of developing sepsis 28x greater with NVHAP than with pneumonia on admission²

^{1.} Angus DC, et al. N Engl J Med. 2013 Aug 29;369(9):840-51.

^{2.} Giuliano K, et al. Am J of Infect Control. 2018;46:322-327



Where is the Highest Risk for NV-HAP?



NV-HAP per 1000 patient days

Baker D, Quinn B, Amer J of Infect Control, 2018;46:2-7 Slide courtesy of Barb Quinn Addressing the risk-factors associated with NV-HAP through evidence based fundamental nursing care strategies



Risk Factors for Pneumonia



Pathogens

- Hospital environment
- Healthcare workers
- Disruption of normal oral flora

Aspiration

- Supine position
- CNS depressant medications
- Invasive tubes

Weak Host

- Surgery
- Immobility
- Co-morbid conditions



Weak Host: Who is at Highest Risk?

- ▲ Male
- Elderly
- Surgical
- **△** ICU
- Chronic disease
 - △ DM, CHF, CKD, COPD, alcoholism

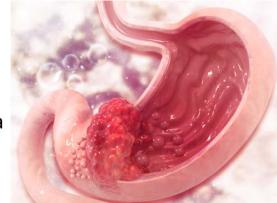
- ▲ Immunocompromised
- △ More than 6 medications
- △ Low albumin
- △ On antibiotics
- △ Dependent for ADLs
- **&** Smokers



Slide courtesy of Barb Quinn

Stewardship of Stress Ulcer Prophylaxis (SUP)

- △ The most common complication of SUP is pneumonia¹
- ▲ ICU enteral fed patients¹
 - △ no benefit & may increase risk for pneumonia Avoid unnecessary use
- △ Acute Stroke patients (Systematic Review & Meta-Analysis)^{2,3}
 - △ Acid suppressive medications are an important contributor to pneumonia development, especially PPIs
- △ May lead to loss of protective bacteriostatic effect of gastric acid^{1,3}
- △ Higher risk of Clostridium difficile infection when combined with antibiotics¹



- 1. Huang et. al (2018). Critical Care 22(20), 1-9.
- 2. Marchina et al (2019). J of the Neurological Sciences, 400;122-128.
- 3. Herzig SJ. et. Al (2014) Ann Neurol. 76(5): 712-178.

Systematic Review of Inpatient Mobilization

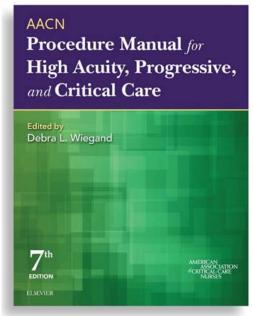
- △ Literature review of research studies that provides evidence to the consequences of mobilizing or not mobilizing hospitalized adult patients
- 36 studies were included
- Findings in four theme areas:
 - △ Physical outcomes include pain relief, reduced deep vein thrombosis, less fatigue, less delirium, less pneumonia, improved physical function (no relationship to falls)
 - \triangle Psychological outcomes include less anxiety, \downarrow depressive mood, \downarrow distress symptoms, \uparrow comfort and \uparrow satisfaction
 - △ Social outcomes include ↑quality of life and more independence
 - \triangle Organizational outcomes include \checkmark length of stay, \checkmark mortality and \checkmark cost













Procedure 4: Endotracheal Tube Care and Oral Care

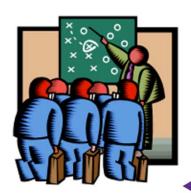
Authors:

Kathleen M Vollman Mary Lou Sole Barbara Quinn

SMCS HAP Prevention Plan

Phase 1: Oral Care

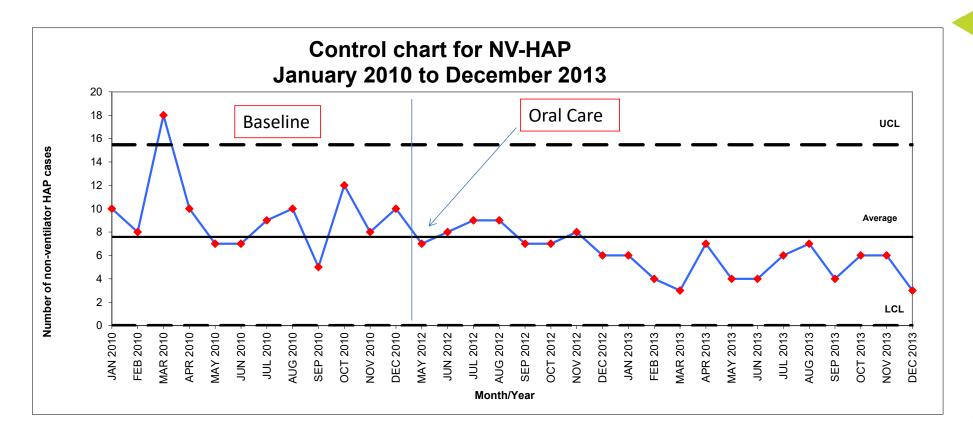
- △ Formation of new quality team: Hospital-Acquired Pneumonia Prevention Initiative (HAPPI)
- △ New oral care protocol to include non-ventilated patients
- △ New oral care products and equipment for all patients
- △ Staff education and in-services on products
- △ Ongoing monitoring and measurement
 - △ Monthly audits



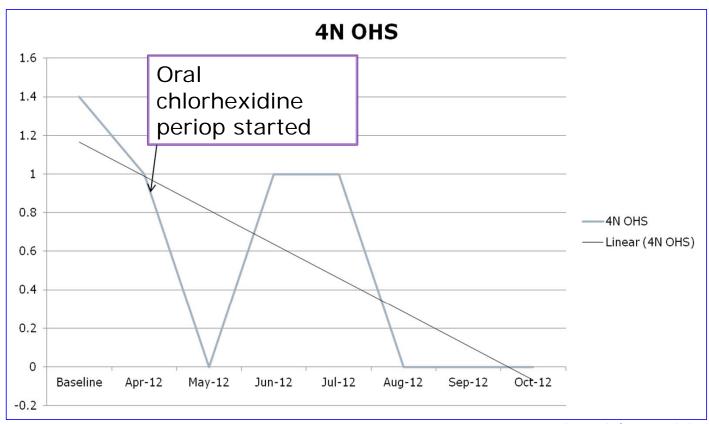
Protocol – Plain & Simple

Patient Type	Tools	Procedure	Frequency
Self Care / Assist	 Brush, paste, rinse, moisturizer Soft-bristled toothbrush Toothpaste with dentifrice Antiseptic mouth rinse (alcohol-free) Moisturizer (Petroleum-free) 	Provide tools Brush 1-2 minutes Rinse	4X / day
Dependent / Aspiration Risk	Suction toothbrush kit (4)	Package instructions	4X / day
Dependent / Vent	ICU Suction toothbrush kit (6)CHG for vent & cardiac surgery patients	Package instructions	6X / day
Dentures	Denture cup, brush Cleanser Adhesive	Remove dentures & soak Brush gums, mouth Rinse	4X / day

NV-HAP Incidence 50 % Decrease from Baseline



Open Heart Surgery Patients: NV-HAP Reduced 75%





- 60 NV-HAP avoided Jan 1 − Dec. 31 2013
- ♠ \$2,400,000 cost avoided
- 117,600 cost increase for supplies
- \$2,282,400 return on investment

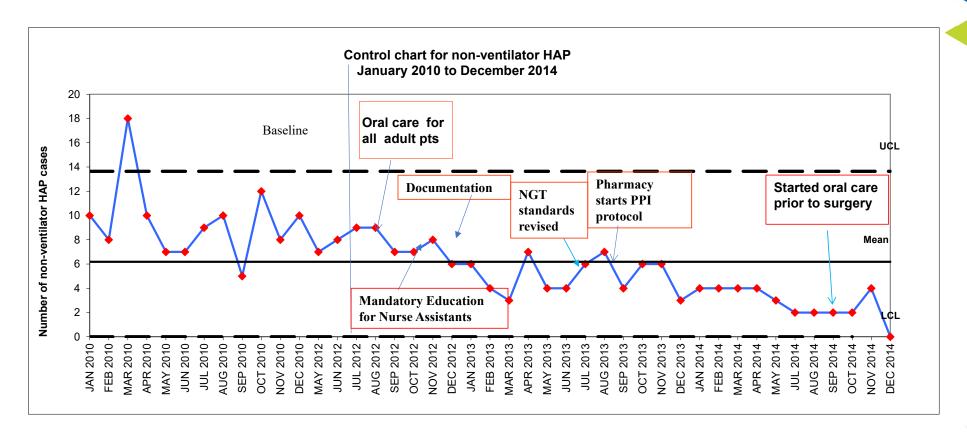
8 lives saved

PRICELESS



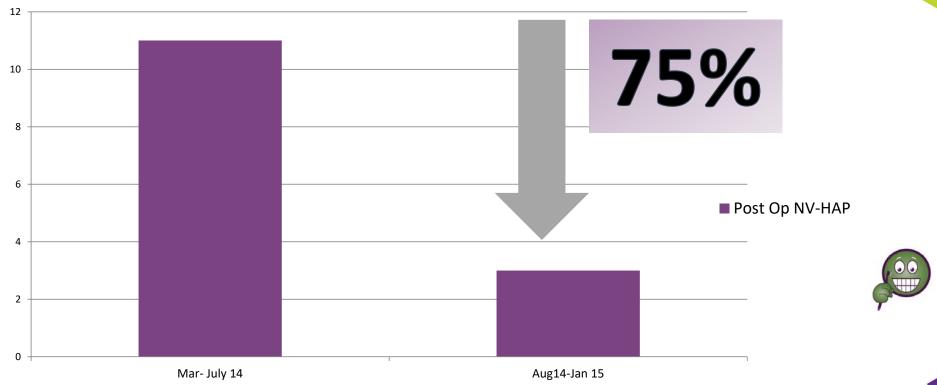
NV-HAP ↓ 70% from baseline!





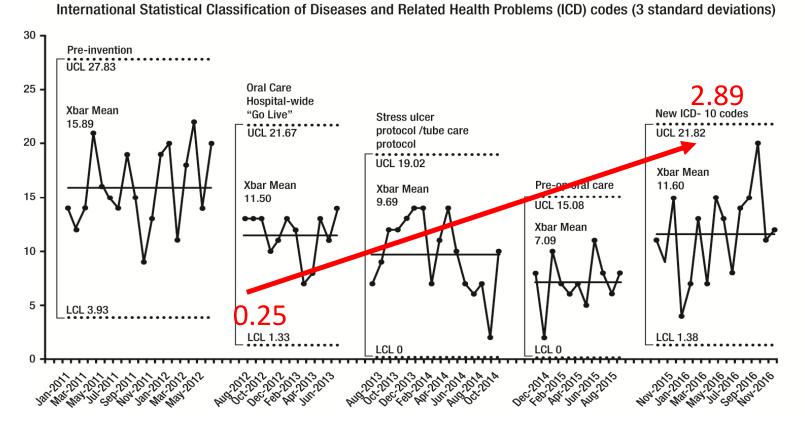
Post-Operative NV-HAP (all adult inpatient surgery) Incidence 6 months Pre-Oral Care vs. 6 Months After





Sustainability Hospital Wide Oral Care from .25 to 2.89 (almost 3x a day)

Figure 1: Statistical process control R and X-bar-charts:





Outcomes:

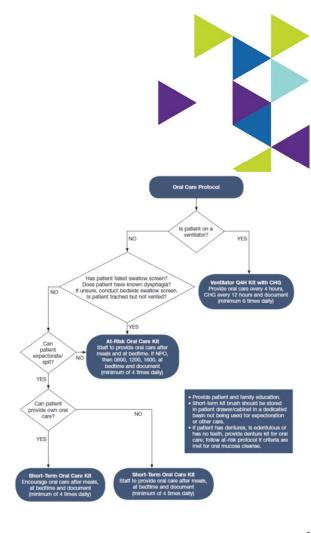
From the Beginning to 2014

- △ Between May 2012 and December 2014
- △ Sutter Medical Center avoided 164 cases of NV-HAP:
 - △ \$5.9 million
 - △ 31 lives
 - △ 656-1476 extra days in the hospital



Nurse Driven Oral Care Protocol to Improve NV-HAP

- △ QI project, 650 bed level 1 trauma center
- △ Data measure retrospectively/prospectively using ICD 9 & 10 codes not POA for NV-HAP and VAP
- △ 7 months baseline, 7 months intervention
- ▲ Method:
 - △ Evaluated current practice, the literature and oral care supplies
 - △ Pilot program with new oral care protocols/supplies for self care, assisted oral care and ventilator oral care
 - △ Expanded to whole hospital post pilot area



Results

Staff adherence to protocol 76% (36%-100%)

▲ NV-HAP

△ Baseline: 202 charts/52 NV-HAP's-20 deaths

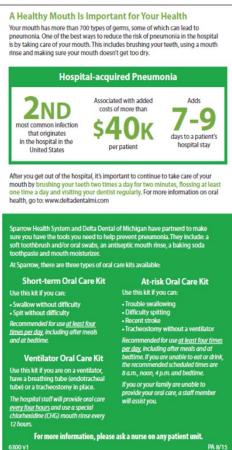
△ Post: 215 charts/26 NV-HAP's (p< 0.0001)-4 deaths

△ Baseline: 56 VAE's/ 12 VAP's (2.87 per 1000 vent days)

△ Post: 49 VAE's/3 VAP's (1.26 per 1000 vent days

50% reduction in NV-HAP, avoided 16 deaths & 1.4 million dollars

Figure 2. Patient Education Information Sheet



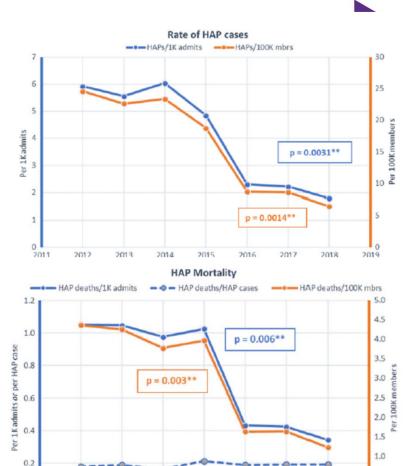
A Successful Program to \downarrow NVHAP in a Large Hospital System

0.0

2011

2012

- 21 hospital system
- Longitudinal observational design
- Intervention
 - △ Upright for meals, mobilization, swallow evaluation, sedation restrictions, rigorous oral care, feeding tube care (ROUTE)
- Additional results
 - △ Reduction in antibiotic days
 - Carbapenem, quinolone, aminoglycoside & vancomycin
 - △ ↓ Benzodiazepine use



0.5

0.0

2019

p = 0.439

2017

2018

Lacerna CC, et al. *Infection Control & Hospital Epidemiology*. 2020;41(5):547-552.



Non-Vent Patients

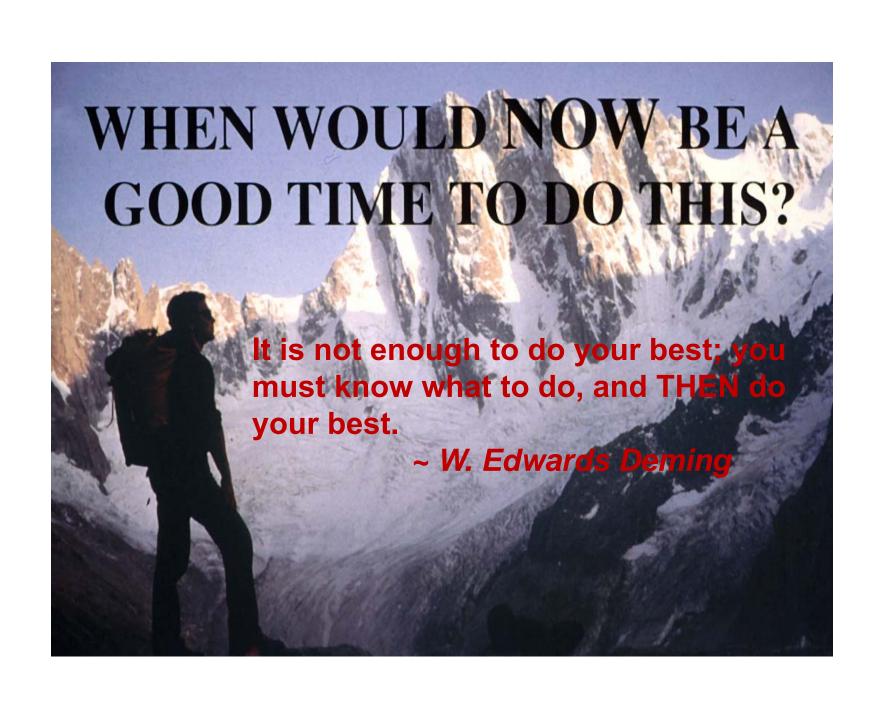
- △ Provided for patients who are unable to manage their own care or secretions safely
- △ Assessment completed within 6hrs of admission
- △ Standardized oral assessment tool
- △ Toothbrushing should occur X2 daily additionally oral cleansing with swabs, suctioning and moisturization of the mouth q 2-4hrs (schedule adjusted to accommodate patient condition/sleep

△ Tools to use

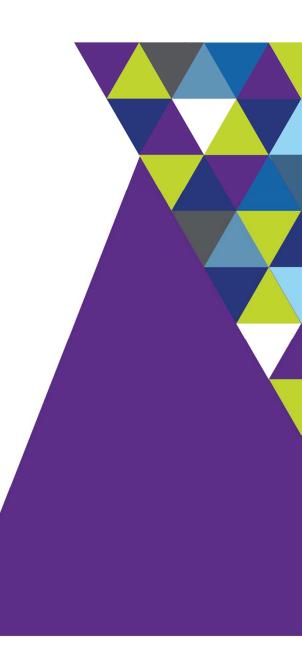
- Pediatric toothbrush followed by suctioning or Suction toothbrush (consider using a single use)
- · Swab for cleaning and moisturizing/suction swab if available to suction debris with cleaning
- Consider using oral care tools & supplies that can be kept at the bedside
- No recommendation on toothpaste for bacteria control

△ Oral care cleansing solutions

With swab cleaning use CPC, 1.5% H2O2 or sterile water



Forbid yourself to be deterred by poor odds just because your mind has calculated that the opposition is too great. If it were easy, everyone would do it.



HAI prevention courses by Kathleen Vollman

https://www.medbridgeeducation.com/advancing-nursing





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