

# Incidence of Hospital-Acquired and Ventilator-Associated Pneumonia in Patients with Severe COVID 19 on High Flow Oxygen

A. Papamanoli, J. Nakamura, J. Fung, J. Abata, N. Karkala, T. Tsui, J. Yoo, P. Grewal, A. Mojahedi, S. Dhaliwal, J. Robin, J. Hotelling, S. Rawal, A. Coritsidis, G. Psevdos, A. Kalogeropoulos, L. Marcos

### Introduction

- Hospital-acquired pneumonia (HAP) and ventilator-associated pneumonia (VAP) can be serious complications of coronavirus disease 19 (COVID-19).
- Co-infections may worsen outcomes and prolong hospitalization.
- This risk may be exacerbated by systemic corticosteroids and other adjunctive therapies.

# Methods

- We reviewed the records of all adults admitted to Stony Brook University Hospital, NY, from 3/1 to 4/15, 2020 with severe COVID-19 pneumonia, requiring high-flow O2 (non-rebreather mask, Venturi mask with FiO2 >50%, or high-flow nasal cannula).
- We excluded patients who received mechanical ventilation (MV) or died within 24h.
- Patients were followed until death or hospital discharge.
- We reviewed positive sputum cultures (PSC) for pathogenic microorganisms and calculated the incidence of HAP and VAP (nosocomial pneumonia, [NP]), rates of MV and impact on mortality. Fungi isolated from sputum were considered colonization unless associated with fungemia.
- We also examined the impact of adjunctive therapies with immunosuppressive potential (steroids and tocilizumab), on HAP or VAP.

# Results

- Among 469 patients (Table 1); 199 (42.4%) required intensive care and 172 (36.7%) MV.
- Median length of stay was 13 days (8-22).
- 105 (22.4%) patients had PSC; 59 were considered true pathogens (HAP: 11, VAP: 48), with predominance of *S. aureus (MSSA) 38.9%, Enterobacteriaceae 33.8% and Pseudomonas* species 18.6%, and 39 isolates were considered colonization (Table 2).
- Patients with PSC <48h (N=7) from admission, were not considered NP.
- The incidence of NP was 7.0 per 1000 patient-days (95%CI 5.5-8.5). Of 11 patients with HAP, 9 needed MV.
- NP was more frequent among patients receiving steroids (9.0 vs 5.7 per 1000 patient-days; P=0.023). Use of tocilizumab was not associated with NP (6.2 vs 8.4; P=0.11).
- Mortality was non-significantly higher in patients with (20/59, 33.9%) vs. without (103/410, 25.1%) NP (P=0.16).
- Intubation and length of stay were the strongest predictors of NP in multivariable models.

# Stony Brook Renaissance School of Medicine, Department of Medicine

### Table 1. Baseline Patient Characteristics (N=469)

Table 1. Baseline Patient Characte	eristics (IN=469)
Characteristic	Value
Age, years	61 (50-73)
Female	166 (35.4%)
White	249 (53.1%)
Black	31 (6.6%)
Asian	29 (6.2%)
Hispanic	158 (33.7%)
Body mass index, kg/m <sup>2</sup>	29.3 (26.1, 33.9)
Duration of symptoms, days	7.0 (3.5, 9.0)
O <sub>2</sub> saturation, %	91 (87, 93)
Temperature, °C	38.1 (37.5 <i>,</i> 39.0)
Hypertension	265 (56.5%)
Diabetes	155 (33.1%)
Coronary artery disease	71 (15.1%)
Atrial fibrillation	58 (12.4%)
Chronic lung disease	49 (10.4%)
Chronic kidney disease	48 (10.2%)
Congestive heart failure	45 (9.6%)
Asthma	36 (7.7%)
Immunocompromised	35 (7.5%)
Statins	180 (38.4%)
Angiotensin-converting enzyme inhibitors	74 (15.8%)
Angiotensin receptor blockers	73 (15.6%)
NT-proBNP pg/mL	205 (56, 991)
Troponin, ng/mL	0.01 (0.01, 0.01)
Creatine phosphokinase, IU/L	163 (80 <i>,</i> 375)
Erythrocyte sedimentation rate, mm/h	54 (31, 80)
C-reactive protein, mg/dL	11.9 (6.4, 19.3)
D-Dimer, ng/mL	362 (241, 747)
Procalcitonin, ng/mL	0.21 (0.13, 0.49)
Ferritin, ng/ml	919 (489, 1534)
Lactate dehydrogenase, IU/L	407 (305, 538)
Interleukin-6, pg/mL	63 (30, 112)
Lymphocyte count, K/uL	0.8 (0.6, 1.1)
Creatinine, mg/dL	1.0 (0.8, 1.3)
Alanine transaminase, IU/L	34 (21, 55)
Aspartate aminotransferase, IU/L	46 (32, 70)
International normalized ratio	1.2 (1.1, 1.3)
Corrected QT interval on ECG, ms	437 (418, 460)
Values are N (%) or median (25 <sup>th</sup> , 75 <sup>th</sup> per	centile)

Values are N (%) or median (25", 75" percentile)

**Correspondence:** Aikaterini Papamanoli, MD Aikaterini.Papamanoli@stonybrookmedicine.edu Stony Brook University Medical Center, 101 Nicolls Road, HSC, T-15, Rm 020 Stony Brook, NY 11794-8153 Tel #: +1 (631) 444-3490

## Table 2. Distribution of Microorganisms In Positive Sputum Cultures

	-
True pathogens	
VAP N= 48	HAP N=11
Staphylococcus aureus (MSSA)	21 Staphylococcus aureus (MSSA)
Pseudomonas aeruginosa	8 Pseudomonas aeruginosa
Klebsiella (enterobacter) aerogenes	6 Staphylococcus aureus (MRSA)
Klebsiella pneumoniae	5 Aspergillus fumigatus
Stenotrophomonas (Xanthomonas) maltophilia	3 Klebsiella pneumoniae
Klebsiella pneumoniae MDR	2 Klebsiella (enterobacter) aerogenes
Staphylococcus aureus (MRSA)	2 Candida albicans
Candida tropicalis	2
Escherichia coli	2
Candida albicans	1
Streptococcus agalactiae (group B) beta hemolytic	1
Streptococci (group C) beta hemolytic	1
Burkholderia cepacia complex	1
Acinetobacter baumannii complex	1
Proteus mirabilis	1
Klebiella oxytoca MDR	1
Escherichia coli MDR	1
Citrobacter farmeri MDR	1
Citrobacter koseri (Citrobacter diversus)	1
Candida dubliniensis	1
Candida parapsilosis	1
Streptococcus pneumoniae	1
Pseudomonas putida	1
Colonization	
Candida albicans	34
Candida tropicalis	6
Candida parapsilosis	5
Candida krusei	1
Candida dubliniensis	1
L	

MDR: Multidrug resistant; MRSA: Methicillin-resistant *Staphylococcus aureus;* MSSA: Methicillin-sensitive *Staphylococcus aureus* 

but not tocilizumab.



### Conclusion

 Among high risk COVID-19 patients, NP is a common complication. MSSA and *Enterobacteriaceae* were the most frequent isolates.

• The risk increases with intubation, longer hospital stay and use of steroids