

Treating COVID-19+ Patients in Place: Clinical Deep Dive into the Research



FMDA Journal Club

April 22, 2020

Diane Sanders-Cepeda, DO, CMD – Presenter

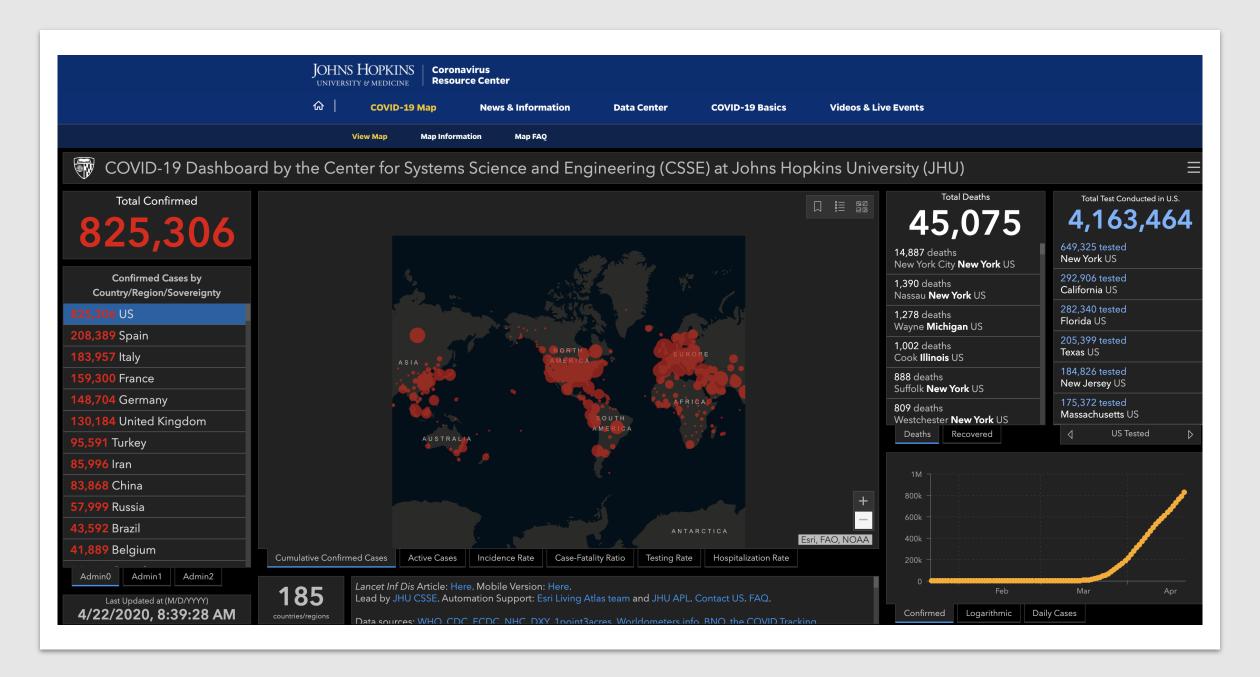
COVID 19 state of the state

Clinical Updates

COVID 19 literature review

Open Discussion

Agenda



Cases by County

Cases by Zip Code

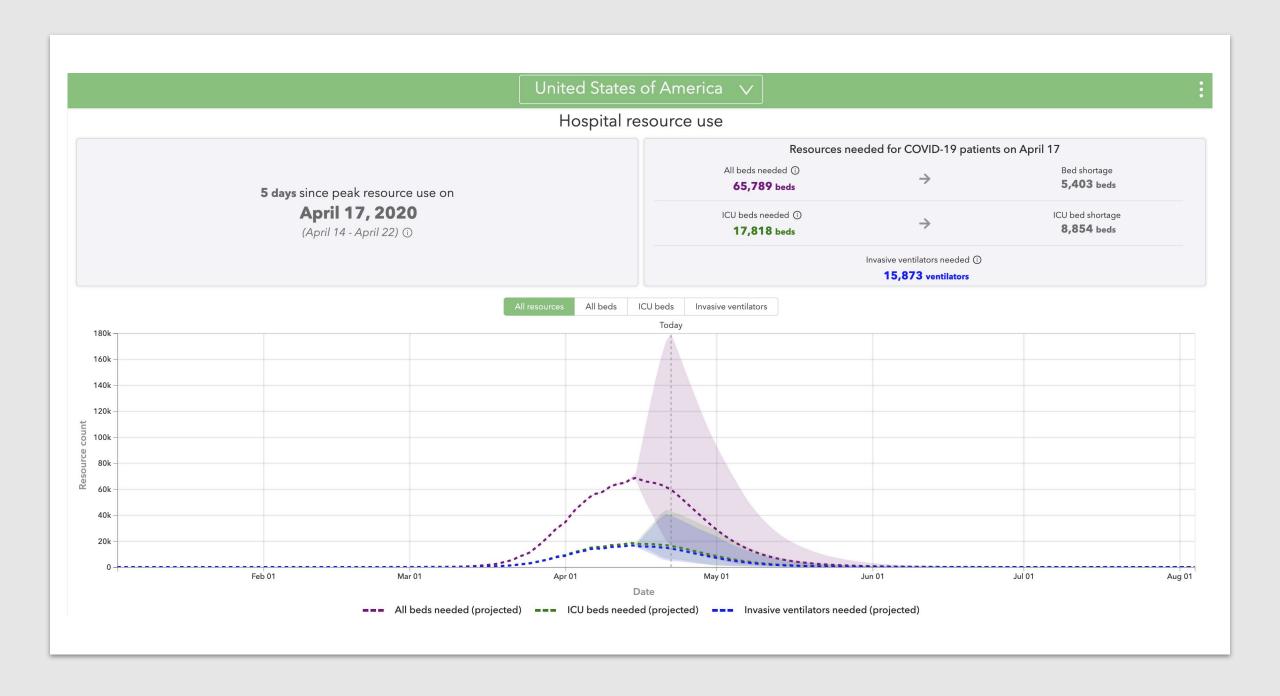
USA and World

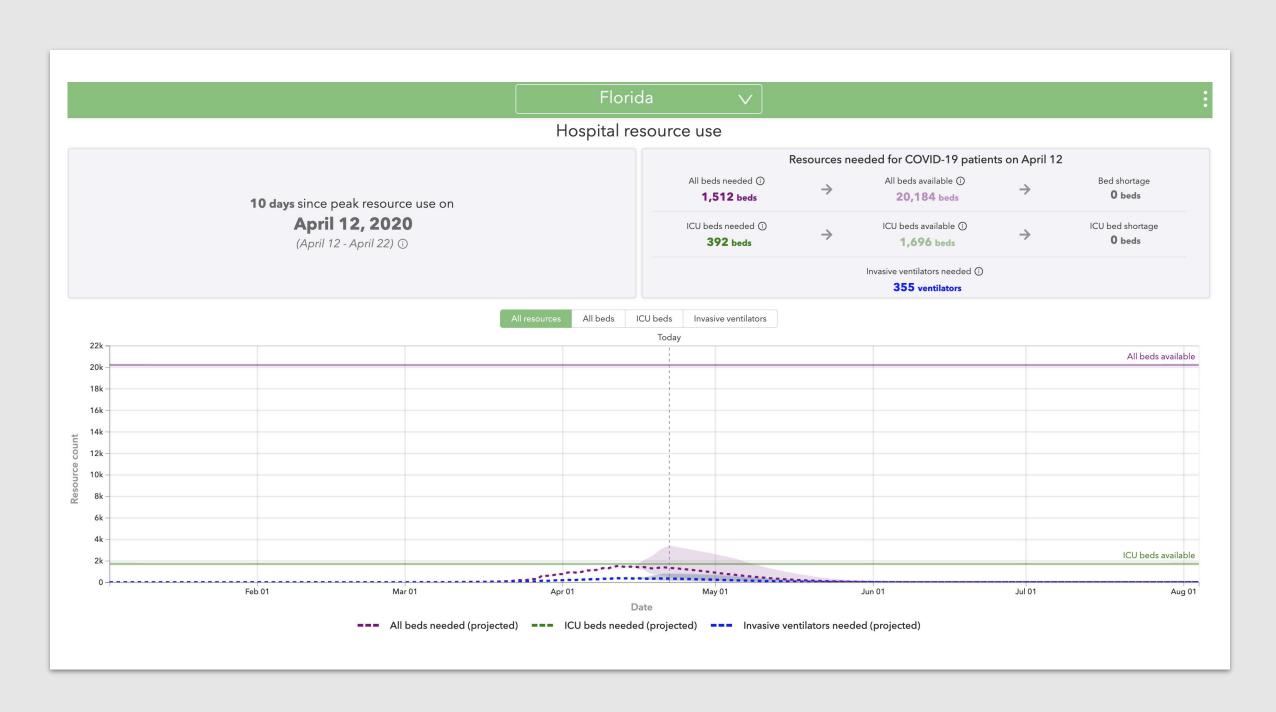
Recent Cases

Deaths by Day

Florida Cases

Florida Testing





DEPARTMENT OF HEALTH & HUMAN SERVICES Centers for Medicare & Medicaid Services 7500 Security Boulevard, Mail Stop C2-21-16 Baltimore, Maryland 21244-1850



Center for Clinical Standards and Quality/Quality, Safety & Oversight Group

Ref: QSO-20-26-NH

DATE: April 19, 2020

TO: State Survey Agency Directors

FROM: Director

Quality, Safety & Oversight Group

SUBJECT: Upcoming Requirements for Notification of Confirmed COVID-19 (or COVID-

19 Persons under Investigation) Among Residents and Staff in Nursing Homes

Memorandum Summary

- *CMS is committed* to taking critical steps to ensure America's health care facilities are prepared to respond to the 2019 Novel Coronavirus (COVID-19) Public Health Emergency (PHE).
- Communicable Disease Reporting Requirements: To ensure appropriate tracking, response, and mitigation of COVID-19 in nursing homes, CMS is reinforcing an existing requirement that nursing homes must report communicable diseases, healthcare-associated infections, and potential outbreaks to State and Local health departments. In rulemaking that will follow, CMS is requiring facilities to report this data to the Centers for Disease Control and Prevention (CDC) in a standardized format and frequency defined by CMS and CDC. Failure to report cases of residents or staff who have confirmed COVID -19 and Persons under Investigation (PUI) could result in an enforcement action. This memorandum summarizes new requirements which will be put in place very soon.
- *Transparency:* CMS will also be previewing a new requirement for facilities to notify residents' and their representatives to keep them up to date on the conditions inside the facility, such as when new cases of COVID-19 occur.





Nebulizer Treatment for COVID-19 Infected Residents with Severe Respiratory Symptoms

COVID-19 infections cause a range of symptoms but often impact the respiratory tract. These symptoms range from sore throat or cough to more severe pulmonary symptoms. More severe symptoms can include reactive airway disease with wheezing and shortness of breath or extreme respiratory compromise, sometimes due to acute respiratory distress syndrome (ARDS), that may require mechanical ventilation. The onset of the severe respiratory symptoms can occur at any time in the course of the disease and can often come on very quickly.

This document deals with COVID-19 positive residents with severe respiratory symptoms who need nebulizer treatments. CDC explains the risk:

CDC considers nebulizer treatments may be an aerosol generating procedure. While "aerosols generated by nebulizers are derived from medication in the nebulizer. It is uncertain whether potential associations between performing this common procedure and increased risk of infection might be due to aerosols generated by the procedure or due to increased contact between those administering the nebulized medication and infected patients" as CDC describes in the FAQs on their website.

COVID 19 Literature Review

Case Reports

The NEW ENGLAND JOURNAL of MEDICINE

CORRESPONDENCE

COVID-19 CASES

To rapidly communicate information on the global clinical effort against Covid-19, the Journal has initiated a series of case reports that offer important teaching points or novel findings. The case reports should be viewed as observations rather than as recommendations for evaluation or treatment. In the interest of timeliness, these reports are evaluated by in-house editors, with peer review reserved for key points as needed.

Immune Thrombocytopenic Purpura in a Patient with Covid-19



Figure 1. Purpuric Lesions on the Patient's Lower Extremity.

The NEW ENGLAND JOURNAL of MEDICINE

CORRESPONDENCE

COVID-19 CASES

To rapidly communicate information on the global clinical effort against Covid-19, the Journal has initiated a series of case reports that offer important teaching points or novel findings. The case reports should be viewed as observations rather than as recommendations for evaluation or treatment. In the interest of timeliness, these reports are evaluated by in-house editors, with peer review reserved for key points as needed.

Coagulopathy and Antiphospholipid Antibodies in Patients with Covid-19

Table 1. Demographic and Clinical Characteristics and Laboratory Findings.*					
Characteristic	Patient 1	Patient 2	Patient 3		
Demographic characteristics					
Age — yr	69	65	70		
Sex	Male	Female	Male		
Initial findings					
Medical history	Hypertension, diabetes, stroke	Hypertension, diabetes, coronary artery disease, no history of thrombosis	Hypertension, emphysema, nasopharyngeal carcinoma, stroke		
Symptoms at disease onset	Fever, cough, dyspnea, diarrhea, headache	Fever, cough, dyspnea	Fever, fatigue, dyspnea, headache		
Imaging features	Ground-glass opacity, bilateral pulmonary infiltrates	Ground-glass opacity, bilateral pulmonary infiltrates	Bilateral pulmonary infiltrates		
Treatment before admission to ICU	Oseltamivir, intravenous immune globulin	Antibiotics	Antibiotics, ribavirin, rosuvastatin		
Days from disease onset to thrombotic event	18	33	10		
Findings on admission to ICU					
Days since disease onset	24	21	24		
Disease severity	Critical	Critical	Critical		

Characteristic	Patient 1	Patient 2	Patient 3
Laboratory findings			
White-cell count (per mm³)	17,790	6730	8710
Differential count (per mm³)			
Total neutrophils	16,290	6230	7090
Total lymphocytes	430	290	790
Total monocytes	800	170	430
Platelet count (per mm³)	78,000	79,000	180,000
Hemoglobin (g/liter)	111	99	92
Albumin (g/liter)	26.3	32.6	24.4
Alanine aminotransferase (U/liter)	15	11	8
Aspartate aminotransferase (U/liter)	23	20	20
Lactate dehydrogenase (U/liter)	632	233	417
Creatinine (µmol/liter)	80	58	86
Creatine kinase (U/liter)	63	335	16
EGFR (ml/min/1.73 m²)	86.6	93.2	78.5
High-sensitivity cardiac troponin I (pg/ml)	3876.8	14.3	125.4
Prothrombin time (sec)	17.0	17.2	15.1
Activated partial-thromboplastin time (sec)	43.7	45.3	47.6
Fibrinogen (g/liter)	4.15	4.42	6.42
Fibrin degradation products (mg/liter)	85.5	8.1	7.3
D-dimer (mg/liter)	>21.00	2.84	3.23
Serum ferritin (µg/liter)	ND	2207.8	ND
Procalcitonin (ng/ml)	0.11	0.18	0.40
High-sensitivity C-reactive protein (mg/liter)	112.0	56.0	125.4
Antiphospholipid antibodies	Anticardiolipin IgA, anti $-eta_2$ -glycoprotein I IgA and IgG	Anticardiolipin IgA, anti $-oldsymbol{eta}_2$ -glycoprotein I IgA and IgG	Anticardiolipin IgA, anti $-eta_2$ -glycoprotein I IgA and IgG
Imaging features	Multiple cerebral infarctions in bilateral frontal parietal occipital lobe and bilateral basal ganglia, brain stem, and bilateral cerebellar hemispheres	Multiple cerebral infarc- tions in right frontal and bilateral parietal lobe	Multiple cerebral infarction in frontal lobe, right fro tal parietal temporal oc cipital lobe, and bilatera cerebellar hemispheres

* EGFR denotes estimated glomerular filtration rate, ICU intensive care unit, and ND not determined.

THE LANCET

CORRESPONDENCE | ONLINE FIRST

Hospital-based use of thromboprophylaxis in patients with COVID-19

Alex C Spyropoulos ○ • Walter Ageno • Elliot S Barnathan

Published: April 21, 2020 • DOI: https://doi.org/10.1016/S0140-6736(20)30926-0

Investigating COVID 19

Research Letter



February 27, 2020

Positive RT-PCR Test Results in Patients Recovered From COVID-19

Lan Lan, MD¹; Dan Xu, MD¹; Guangming Ye, MD²; et al

≫ Author Affiliations | Article Information

JAMA. 2020;323(15):1502-1503. doi:10.1001/jama.2020.2783



Research Letter

ONLINE FIRST

FREE

April 15, 2020

Rates of Co-infection Between SARS-CoV-2 and Other Respiratory Pathogens

David Kim, MD, PhD¹; James Quinn, MD, MS¹; Benjamin Pinsky, MD, PhD²; et al.

Author Affiliations | Article Information

JAMA. Published online April 15, 2020. doi:10.1001/jama.2020.6266

Table 1. Patient Characteristics and Sites of Specimen Collection, by SARS-CoV-2 and Non-SARS-CoV-2 Pathogen Status

Table 1. Patient Characteristics and Sites of Specimen Collection, by SARS-CoV-2 and Non-SARS-CoV-2 Pathogen Status

	SARS-CoV-2 status, No. (%)						
Characteristic	Negative (n = 1101)	40	Positive (n = 116)				
	Positive for other respiratory pathogen	Negative for other respiratory pathogen	Positive for other respiratory pathogen	Negative for other respiratory pathogen			
No. of samples	294	807	24	92			
No. of patients ^a	292	800	23	92			
Age, mean (range), y ^b	48.8 (7-82)	43.8 (1-100)	50.8 (9-88)	43.3 (1-98)			
Female, No./total (%)b	161/292 (55.1)	443/800 (55.4)	12/23 (52.2)	52/92 (56.5)			
Site of specimen collection, No./total (%) ^c							
Outpatient clinic	115/294 (39.1)	347/807 (43.0)	11/24 (45.8)	39/92 (42.4)			
Emergency department							
Discharged	122/294 (41.5)	301/807 (37.3)	12/24 (50.0)	38/92 (41.3)			
Admitted ^d	28/294 (9.5)	109/807 (13.5)	1/24 (4.2)	15/92 (16.3)			
Inpatient	29/294 (9.9)	50/807 (6.2)	0/24	0/92			

Abbreviation: SARS-CoV-2, severe acute respiratory syndrome coronavirus 2.

- ^a Row sum (1207) is greater than the total number of unique patients (1206) because 1 patient was tested twice, 11 days apart, with different results for non-SARS-CoV-2 pathogens, and so appears in the first 2 columns.
- b Mean age and proportion female are calculated with respect to unique patients.
- ^c Proportions of samples collected at different sites are calculated with respect to numbers of samples.
- Denotes patients tested in the emergency department and admitted to an inpatient ward from the emergency department.

Table 2. Proportions of Specimens Positive for Non-SARS-CoV-2 Respiratory Pathogens and Mean Patient Ages for Each Subgroup, by SARS-CoV-2 Result^{a,b}

Table 2. Proportions of Specimens Positive for Non-SARS-CoV-2 Respiratory Pathogens and Mean Patient Ages for Each Subgroup, by SARS-CoV-2 Result^{a,b}

	SARS-CoV-2 status						
Pathogen	Negative (n = 1101)		Positive (n = 116)				
	Proportion positive for other respiratory pathogen, No. (%) ^b	Mean age of positive patients, y	Proportion positive for other respiratory pathogen, No. (%) ^b	Mean age of positive patients, y			
Influenza							
A	29/1101 (2.6)	45.9	1/116 (0.9)	74.0			
В	8/1101 (0.7)	21.6	0/116 (0)				
RSV	32/1101 (2.9)	26.0	6/116 (5.2)	52.3			
Parainfluenza							
1	1/1101 (0.1)	71.0	1/116 (0.9)	43.0			
2	0/1101 (0)		0/116 (0)				
3	2/1101 (0.2)	40.0	1/116 (0.9)	45.0			
4	5/1101 (0.5)	26.6	1/116 (0.9)	36.0			
Metapneumovirus	47/1101 (4.3)	41.1	2/116 (1.7)	67.0			
Rhinovirus/enterovirus	133/1101 (12.1)	32.6	8/116 (6.9)	42.1			

Abbreviations: RSV, respiratory syncytial virus; SARS-CoV-2, severe acute respiratory syndrome coronavirus 2.

^a Positive results for non-SARS-CoV-2 pathogens may in some cases represent the detection of residual virus in resolved cases, rather than clinical co-infection as such.

The Search for a Treatment

Clinical Review & Education

JAMA | Review

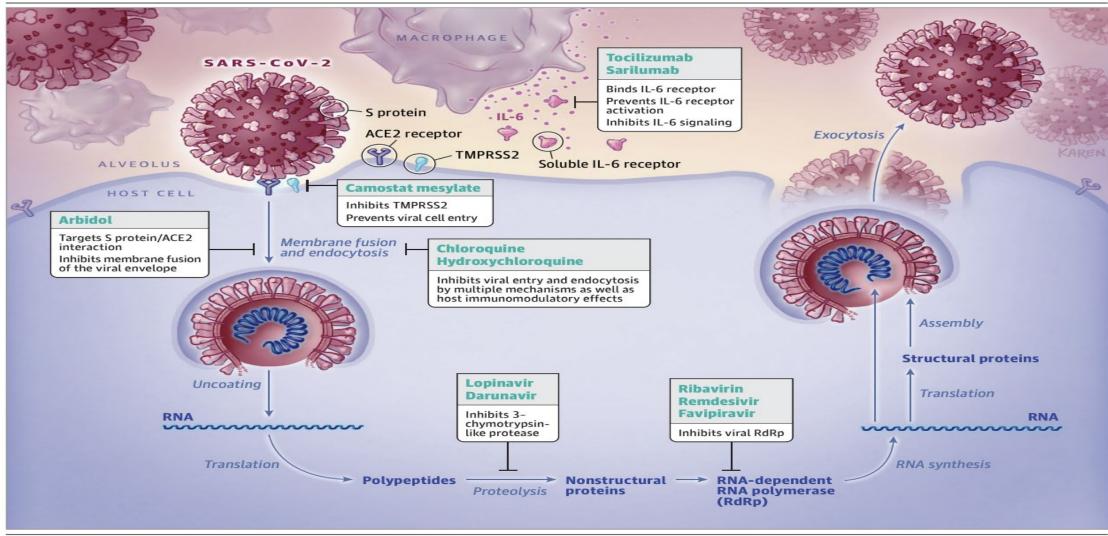
Pharmacologic Treatments for Coronavirus Disease 2019 (COVID-19) A Review

James M. Sanders, PhD, PharmD; Marguerite L. Monogue, PharmD; Tomasz Z. Jodlowski, PharmD; James B. Cutrell, MD

IMPORTANCE The pandemic of coronavirus disease 2019 (COVID-19) caused by the novel severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) presents an unprecedented challenge to identify effective drugs for prevention and treatment. Given the rapid pace of scientific discovery and clinical data generated by the large number of people rapidly infected by SARS-CoV-2, clinicians need accurate evidence regarding effective medical treatments for this infection.

- Viewpoint
- Related article

Figure. Simplified Representation of Severe Acute Respiratory Syndrome Coronavirus 2 (SARS-CoV-2) Viral Lifecycle and Potential Drug Targets



Schematic represents virus-induced host immune system response and viral processing within target cells. Proposed targets of select repurposed and investigational products are noted. ACE2, angiotensin-converting enzyme 2; S protein, spike protein; and TMPRSS2, type 2 transmembrane serine protease.

Table 2. Summary of Treatment and Clinical Outcomes From Early COVID-19 Clinical Series

Source	Huang et al, 2020 ⁹¹	Chen et al, 2020 ⁹²	Wang et al, 2020 ⁵¹	Yang et al, 2020 ⁹³	Young et al, 2020 ⁹⁴	Kujawski et al, 2020 ⁶⁶	Guan et al, 2020 ⁹⁵
Study setting and region	Wuhan Jinyintan Hospital, China (12/16/19-1/2/20)	Wuhan Jinyintan Hospital, China (1/1/20-1/20/20)	Zhongnan Hospital, Wuhan, China (1/1/20-1/28/20)	Wuhan Jinyintan Hospital, China (12/24/19-1/26/20))	4 Singapore hospitals (1/23/20-2/3/20)	US-confirmed cases (1/20/20-2/5/20)	National Chinese cases (12/19/19-1/29/20)
No. of patients	41 Hospitalized	99 Hospitalized	138 Hospitalized	52 (All ICU)	18 Hospitalized	12 (Only 7 hospitalized)	1096 Hospitalized
Age, median (IQR), y	49 (41-58)	Mean (SD), 55.5 (13.1)	56 (42-68)	Mean (SD), 59.7 (13.3)	47 (31-73)	53 (21-68)	47 (35-58)
Sex, No. (%)							
Male	30 (73)	67 (68)	75 (54)	35 (67)	9 (50)	8 (67)	637 (58)
Female	11 (27)	32 (32)	63 (46)	17 (33)	9 (50)	4 (33)	459 (42)
ICU status/ complications, No. (%)	ICU: 13 (32); ARDS: 12 (29); MI: 5 (12); AKI: 3 (7); shock: 3 (7); secondary infection: 4 (10)	ICU: 23 (23); ARDS: 17 (17); AKI: 3 (3); shock: 4 (4); VAP: 1 (1)	ICU: 36 (26); ARDS: 27 (20); MI: 10 (7.2); arrhythmia: 23 (17); AKI: 5 (3.6); shock: 12 (8.7)	ICU: 52 (100); ARDS: 35 (67); MI: 12 (23); AKI: 15 (29); bacterial infection: 8 (15)	ICU: 2 (11); ARDS: 0 (0); secondary bacterial infection: (0)	ICU: 1 (8); culture-positive secondary bacterial infection: 0 (0)	ICU: 55 (5); ARDS: 37 (3.4); AKI: 6 (0.5); shock: 12 (1.1)
Treatments, No. (%)							
Supportive care	NIV/HFNC: 10 (24); MV: 2 (5); ECMO: 2 (5); KRT: 3 (7)	NIV: 13 (13); MV: 4 (4); ECMO: 3 (3); KRT: 9 (9)	NIV: 15 (10.9); MV: 17 (12); ECMO: 4 (2.9); KRT: 2 (1.5)	NIV: 29 (56); MV: 22 (42); ECMO: 6 (12); KRT: 9 (17)	Supplemental oxygen: 6 (33); MV: 1(6)	Supplemental oxygen: 4 (33)	Oxygen: 454 (41); NIV: 56 (5); MV: 25 (2); ECMO: 5 (0.5); KRT: 9 (0.8)
Specific agents	Antivirals (oseltamivir): 38 (99); antibacterials: 41 (100); corticosteroids: 9 (22)	Antivirals (oseltamivir, ganciclovir, or lopinavir/ ritonavir): 75 (76); antibacterials: 70 (71); antifungals: 15 (15); corticosteroids: 19 (19); IVIG: 27 (27)	Antivirals (oseltamivir): 124 (90); antibacterials: moxifloxacin: 89 (64), ceftriaxone: 34 (23), azithromycin: 25 (18); corticosteroids: 62 (45)	Antivirals: 23 (44); antibacterials: 49 (94); corticosteroids: 30 (58); IVIG: 28 (54)	Antivirals (lopinavir/ ritonavir): 5 (42); other antivisor antibacterials: NR	Antivirals (remdesivir): 3 (25); antibacterials: 5 (42); corticosteroids: 2 (17)	Antivirals (oseltamivir): 393 (36); antibacterials: 637 (58); antifungals: 31 (2.8); corticosteroids: 204 (19); IVIG: 144 (13)
Discharged alive, No. (%)	28 (68)	31 (31)	47 (34)	NR	8 (75)	100 (100)	55 (5)
Deaths, No. (%)	6 (15)	11 (11)	6 (4.3)	32 (62)	0	0	15 (1.4)

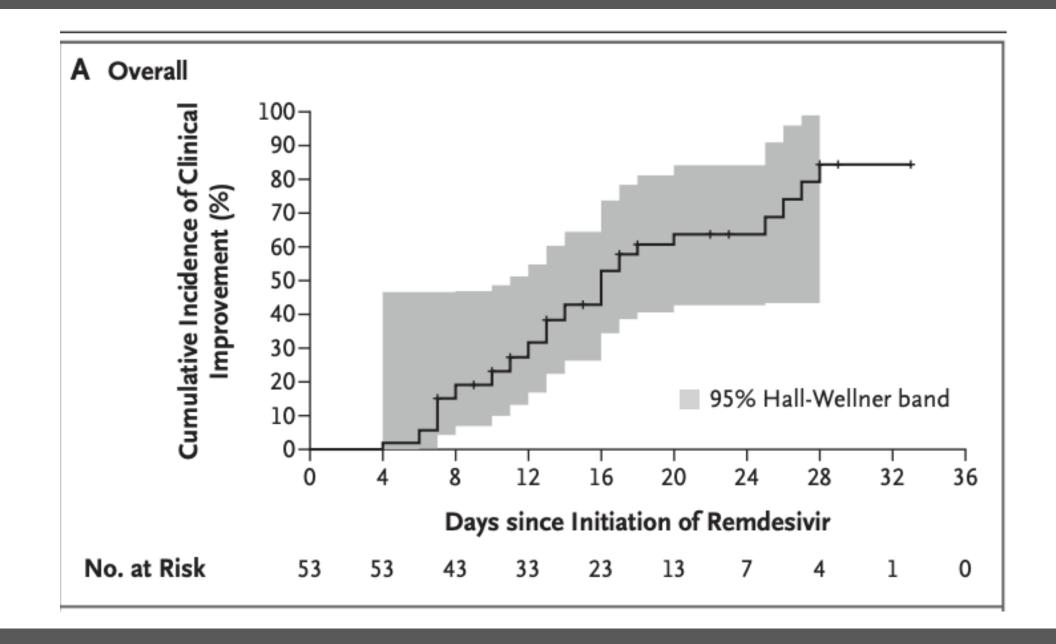
Abbreviations: AKI, acute kidney injury; ARDS, acute respiratory distress syndrome; COVID-19, coronavirus disease 2019; ECMO, extracorporeal membrane oxygenation; HFNC, high-flow nasal cannula; ICU, intensive care unit; IQR, interquartile range; IVIG, intravenous immunoglobulins;

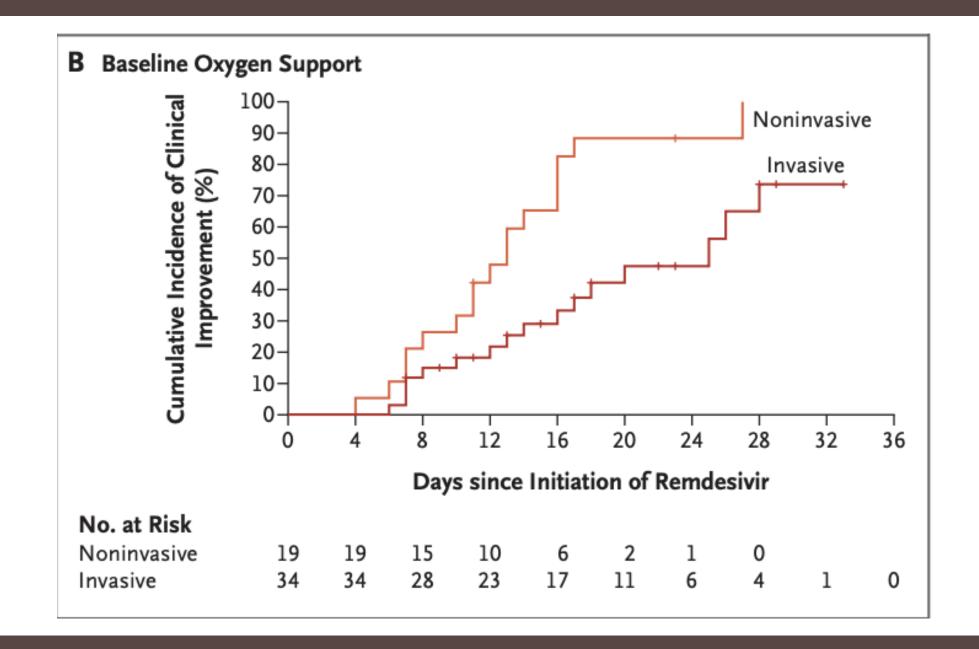
MI, myocardial infarction; MV, invasive mechanical ventilation; KRT, kidney replacement therapy; NIV, noninvasive ventilation; NR, not reported; VAP, ventilator-associated pneumonia.

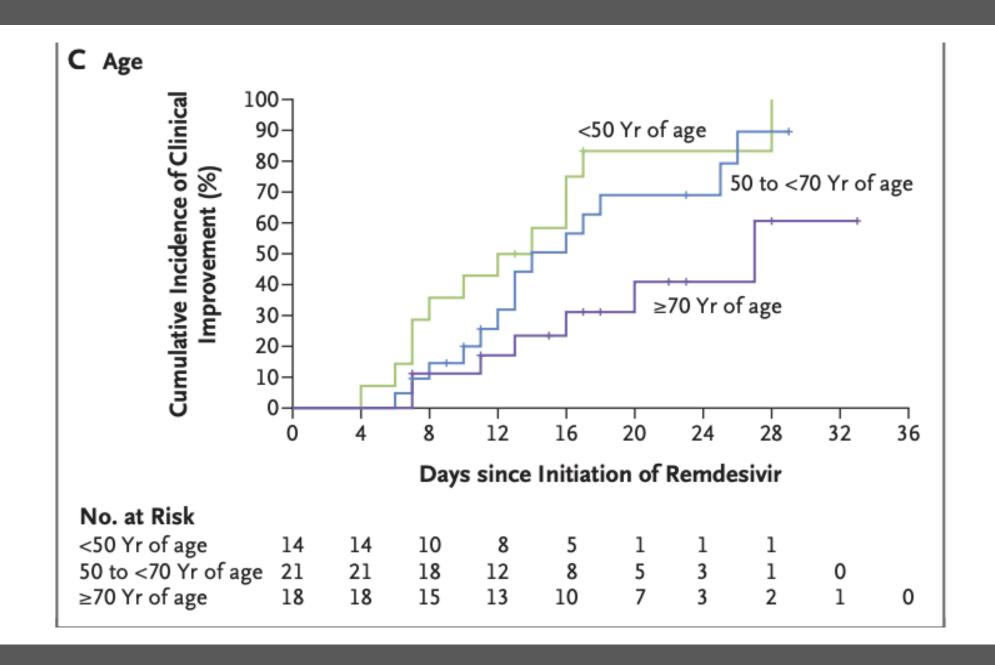
ORIGINAL ARTICLE

Compassionate Use of Remdesivir for Patients with Severe Covid-19

J. Grein, N. Ohmagari, D. Shin, G. Diaz, E. Asperges, A. Castagna, T. Feldt,
G. Green, M.L. Green, F.-X. Lescure, E. Nicastri, R. Oda, K. Yo, E. Quiros-Roldan,
A. Studemeister, J. Redinski, S. Ahmed, J. Bernett, D. Chelliah, D. Chen,
S. Chihara, S.H. Cohen, J. Cunningham, A. D'Arminio Monforte, S. Ismail,
H. Kato, G. Lapadula, E. L'Her, T. Maeno, S. Majumder, M. Massari,
M. Mora-Rillo, Y. Mutoh, D. Nguyen, E. Verweij, A. Zoufaly, A.O. Osinusi,
A. DeZure, Y. Zhao, L. Zhong, A. Chokkalingam, E. Elboudwarej, L. Telep,
L. Timbs, I. Henne, S. Sellers, H. Cao, S.K. Tan, L. Winterbourne, P. Desai,
R. Mera, A. Gaggar, R.P. Myers, D.M. Brainard, R. Childs, and T. Flanigan







Event	Invasive Ventilation (N = 34)	Noninvasive Oxygen Support (N=19)	Total (N = 53)
	nı	ımber of patients (percent	t)
Any adverse event	22 (65)	10 (53)	32 (60)
Adverse events occurring in 2 or more patients			
Hepatic enzyme increased*	8 (24)	4 (21)	12 (23)
Diarrhea	1 (3)	4 (21)	5 (9)
Rash	3 (9)	1 (5)	4 (8)
Renal impairment	4 (12)	0	4 (8)
Hypotension	3 (9)	1 (5)	4 (8)
Acute kidney injury	2 (6)	1 (5)	3 (6)
Atrial fibrillation	2 (6)	1 (5)	3 (6)
Multiple-organ-dysfunction syndrome	3 (9)	0	3 (6)
Hypernatremia	3 (9)	0	3 (6)
Deep-vein thrombosis	3 (9)	0	3 (6)
Acute respiratory distress syndrome	1 (3)	1 (5)	2 (4)
Pneumothorax	2 (6)	0	2 (4)
Hematuria	2 (6)	0	2 (4)
Delirium	1 (3)	1 (5)	2 (4)
Septic shock	2 (6)	0	2 (4)
Pyrexia	1 (3)	1 (5)	2 (4)
Any serious adverse event	9 (26)	3 (16)	12 (23)
Serious events occurring in 2 or more patients			
Multiple-organ-dysfunction syndrome	2 (6)	0	2 (4)
Septic shock	2 (6)	0	2 (4)
Acute kidney injury	2 (6)	0	2 (4)
Hypotension	2 (6)	О	2 (4)

^{*} Adverse-event terms are based on the *Medical Dictionary for Regulatory Activities*, version 22.1. Hepatic enzyme increased includes the following terms: hepatic enzyme increased, alanine aminotransferase increased, aspartate aminotransferase increased, and transaminases increased. Elevated hepatic enzymes resulted in discontinuation of remdesivir therapy in 2 patients.

Management Guidelines



Expert U.S. panel develops NIH treatment guidelines for COVID-19

"Living document" expected to be updated often as new clinical data accrue.

Overview and Spectrum of COVID-19

Summary Recommendations

- The COVID-19 Treatment Guidelines Panel (the Panel) **does not recommend** the use of any agents for pre-exposure prophylaxis (PrEP) against severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) outside of the setting of a clinical trial (AIII).
- The Panel **does not recommend** the use of any agents for post-exposure prophylaxis (PEP) against SARS-CoV-2 infection outside of the setting of a clinical trial (AIII).
- The Panel recommends no additional laboratory testing and no specific treatment for persons with suspected or confirmed asymptomatic or presymptomatic SARS-CoV-2 infection (AIII).
- At present, no drug has been proven to be safe and effective for treating COVID-19. There
 are insufficient data to recommend either for or against the use of any antiviral or
 immunomodulatory therapy in patients with COVID-19 who have mild, moderate, severe,
 or critical illness (AIII).

Infectious Diseases Society of America Guidelines on the Treatment and Management of Patients with COVID-19

Published by IDSA, 4/11/2020

COVID-19 Guideline, Part 2: Infection Prevention - Coming Soon

COVID-19 Guideline, Part 3: Diagnostics - Coming Soon

Adarsh Bhimraj*, Rebecca L. Morgan**, Amy Hirsch Shumaker, Valery Lavergne**, Lindsey Baden, Vincent Chi-Chung Cheng, Kathryn M. Edwards, Rajesh Gandhi, William J. Muller, John C. O'Horo, Shmuel Shoham, M. Hassan Murad**, Reem A. Mustafa**, Shahnaz Sultan**, Yngve Falck-Ytter**

- Recommendation 1. Among patients who have been admitted to the hospital with COVID-19, the IDSA guideline panel recommends hydroxychloroquine/chloroquine in the context of a clinical trial. (Knowledge gap)
- Recommendation 2. Among patients who have been admitted to the hospital with COVID-19, the IDSA guideline panel recommends hydroxychloroquine/chloroquine plus azithromycin only in the context of a clinical trial. (Knowledge gap)
- Recommendation 3. Among patients who have been admitted to the hospital with COVID-19, the IDSA guideline panel recommends the combination of lopinavir/ritonavir only in the context of a clinical trial. (Knowledge gap)
- Recommendation 4. Among patients who have been admitted to the hospital with COVID-19 pneumonia, the IDSA guideline panel suggests against the use of corticosteroids. (Conditional recommendation, very low certainty of evidence)
- Recommendation 5. Among patients who have been admitted to the hospital with ARDS due to COVID-19, the IDSA guideline panel recommends the use of corticosteroids in the context of a clinical trial. (Knowledge gap)
- Recommendation 6. Among patients who have been admitted to the hospital with COVID-19, the IDSA guideline panel recommends tocilizumab only in the context of a clinical trial. (Knowledge gap)
- Recommendation 7. Among patients who have been admitted to the hospital with COVID-19, the IDSA guideline panel recommends COVID-19 convalescent plasma in the context of a clinical trial. (Knowledge gap)

What will Tomorrow Bring?

Emerging Research



For Journalists

Podcasts & Rad

By Dana Sparks

Convalescent Plasma: A Therapy for COVID-19?

April 6, 2020







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Bacille Calmette-Guérin (BCG)vaccination and COVID-19

Scientific Brief

Lessons Learned

ORIGINAL ARTICLE

Epidemiology of Covid-19 in a Long-Term Care Facility in King County, Washington

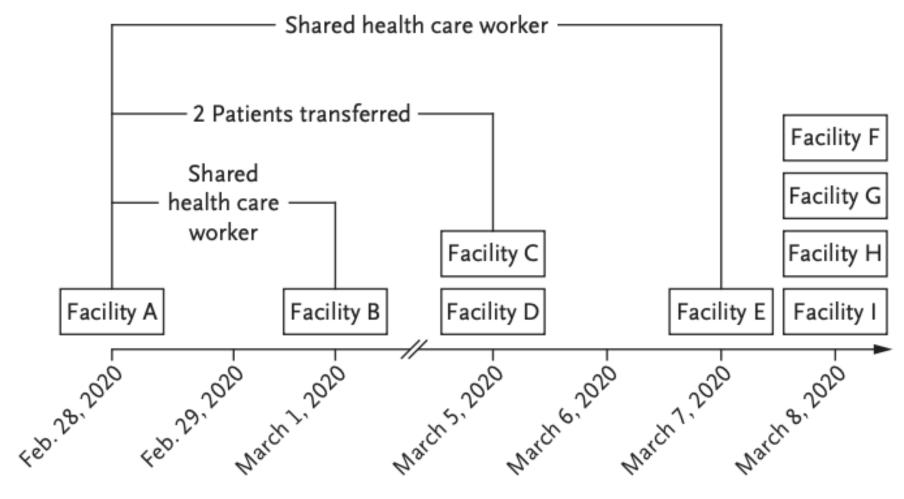
Temet M. McMichael, Ph.D., Dustin W. Currie, Ph.D., Shauna Clark, R.N., Sargis Pogosjans, M.P.H., Meagan Kay, D.V.M., Noah G. Schwartz, M.D., James Lewis, M.D., Atar Baer, Ph.D., Vance Kawakami, D.V.M., Margaret D. Lukoff, M.D., Jessica Ferro, M.P.H., Claire Brostrom-Smith, M.S.N., Thomas D. Rea, M.D., Michael R. Sayre, M.D., Francis X. Riedo, M.D., Denny Russell, B.S., Brian Hiatt, B.S., Patricia Montgomery, M.P.H., Agam K. Rao, M.D., Eric J. Chow, M.D., Farrell Tobolowsky, D.O., Michael J. Hughes, M.P.H., Ana C. Bardossy, M.D., Lisa P. Oakley, Ph.D., Jesica R. Jacobs, Ph.D., Nimalie D. Stone, M.D., Sujan C. Reddy, M.D., John A. Jernigan, M.D., Margaret A. Honein, Ph.D., Thomas A. Clark, M.D., and Jeffrey S. Duchin, M.D., for the Public Health–Seattle and King County, EvergreenHealth, and CDC COVID-19 Investigation Team*

Table 1. Demographic and Clinical Characteristics of Persons with Confirmed Covid-19 Linked to Facility A.* Health Care Residents Personnel Visitors Total (N = 101)Characteristic (N = 16)(N = 167)(N = 50)Median age (range) - yr 83 (51-100) 43.5 (21–79) 62.5 (52-88) 72 (21–100) Sex — no. (%) Male 32 (31.7) 12 (24.0) 55 (32.9) 11 (68.8) Female 69 (68.3) 38 (76.0) 5 (31.2) 112 (67.1) Hospitalized — no. (%) 55 (54.5) 8 (50.0) 66 (39.5) Yes 3 (6.0) 9 (8.9) 44 (88.0) 8 (50.0) 61 (36.5) Nο Unknown 37 (36.6) 3 (6.0) 40 (24.0) 0 Died — no. (%) 35 (21.0) Yes 34 (33.7) 0 1 (6.2) 67 (66.3) 50 (100.0) 132 (79.0) No 15 (93.8) Chronic underlying conditions — no. (%)† 68 (67.3) 2 (12.5) 74 (44.3) Hypertension: 4 (8.0) Cardiac disease 61 (60.4) 68 (40.7) 4 (8.0) 3 (18.8) 41 (40.6) 2 (12.5) 43 (25.7) Renal disease 0 Diabetes mellitus 32 (31.7) 1 (6.2) 38 (22.8) 5 (10.0) Obesity 37 (22.2) 31 (30.7) 3 (6.0) 3 (18.8) Pulmonary disease 32 (31.7) 2 (4.0) 2 (12.5) 36 (21.6) Cancer 15 (14.9) 15 (9.0) Compromised immune system 9 (8.9) 0 0 9 (5.4) Liver disease 6 (5.9) 0 6 (3.6) 0

^{*} Data are for persons with confirmed Covid-19 that was epidemiologically linked to Facility A, including residents of King County and Snohomish County, from February 27 through March 18, 2020.

[†] For chronic underlying conditions, "no" and "unknown" are combined. Percentages represent the number of cases with information on the coexisting condition, irrespective of missing data. Data on chronic underlying conditions were missing for 1 resident, 5 health care personnel, and 1 visitor.

[‡] Hypertension was the only chronic underlying condition in 7 residents, 2 health care personnel, and 1 visitor.



Date of First Covid-19 Case Confirmed in Facility





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This meeting has been recorded and will be available at www.fmda.org/journalclub.php