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11-11-2020

Covid-19 Clinical Update 11/19/2020

George Washington University

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School of Medicine & Health Sciences

THE GEORGE WASHINGTON UNIVERSITY

Medicine Grand Rounds November 19, 2020

Critical Care Air Transport Karolyn Teufel, MD

Claiming CME Credit

TEXT the Unique Activity Code

PEXHOQ

to

202-902-9966



For questions regarding CME credits:

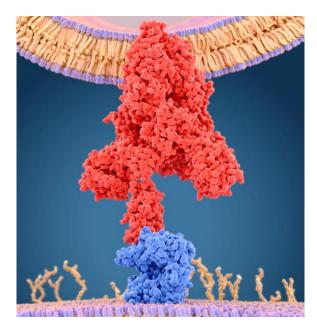
Contact the CEHP Office

Email: cehp@gwu.edu

Ph: 202-994-4285







1. EPIDEMIOLOGY

2. VACCINE

3. ANTIBODIES

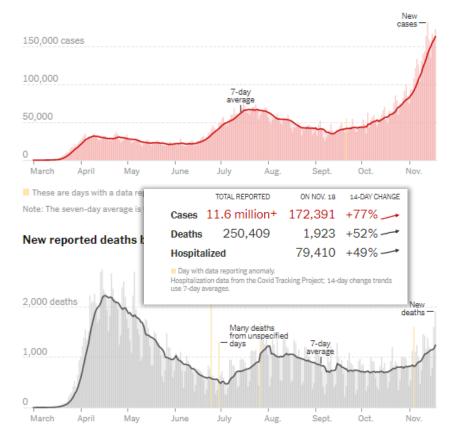
4. GW UPDATES

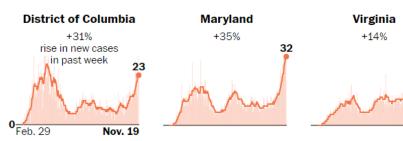
COVID-19 UPDATE

HANA AKSELROD, MD, MPH GW DIVISION OF INFECTIOUS DISEASES 11/19/2020

DISCLOSURES: NO FINANCIAL COI. INVESTIGATOR ON MODERNA VACCINE TRIAL. PRE-PRINT FINDINGS AND INVESTIGATIONAL USE MAY BE DISCUSSED.

New reported cases by day in the United States



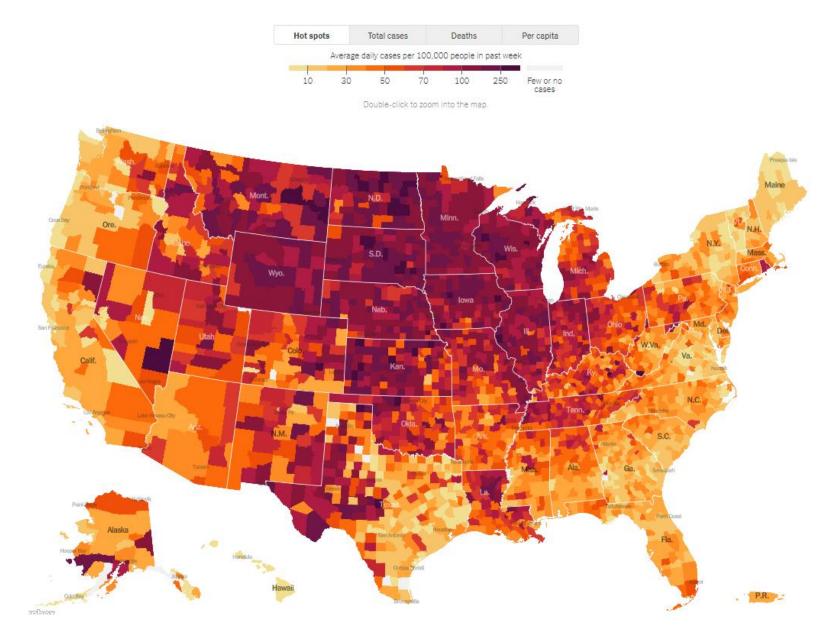


2,862 reported

per 100k

2,844 reported per 100k

2,482 reported per 100k



https://www.washingtonpost.com/graphics/local/dc-maryland-virginia-coronavirus-cases/ https://www.nytimes.com/interactive/2020/us/coronavirus-us-cases.html

NEWS RELEASES

Monday, November 16, 2020

Promising Interim Results from Clinical Trial of NIH-Moderna COVID-19 Vaccine Institute/Center

Contact

National Institute of Allergy and Infectious Diseases (NIAID)

Vaccine News

"PFIZER-BIONTECH VACCINE" BNT162B2

- 43,661 volunteers across 150 sites worldwide
- 170 cases of COVID-19
- 162 in placebo vs. 8 in vaccine group = 95% efficacy
- 10 cases of severe COVID-19, of which 9 (90%) occurred in placebo group
- 30% of US volunteers POC, 45% aged 56-85
- "Efficacy was consistent across . . . demographics; efficacy in adults over 65 years of age was over 94%"
- SAEs: HA (2% after dose #2), fatigue (4% after #2)
- Very cold (-80°C) storage and delivery chain required

"MODERNA-NIH VACCINE" MRNA-1273

- 30,000 volunteers across 100 US sites (25 CoVPN sites)
- 95 cases of symptomatic COVID-19
- 90 in placebo group vs. 5 in vaccine group = 94.5% efficacy
- 11 (12%) cases of severe COVID-19, all (100%) of which occurred in the placebo group
- 15 cases in older adults (ages 65+)
- ~37% of participants and 20 of the cases identified as people of color (12 Hispanic/LatinX, 4 Black or African Americans, 3 Asian American, 1 multiracial)
- SAEs: 2% after dose #1, 3% after dose #2 (myalgias, fatigue)
- More stable under standard cold chain conditions

https://www.statnews.com/2020/11/18/pfizer-biontech-covid19-vaccine-fda-data/; https://www.pfizer.com/news/press-release-detail/pfizer-and-biontech-conclude-phase-3-study-covid-19-vaccine; https://www.nih.gov/news-events/newsreleases/promising-interim-results-clinical-trial-nih-moderna-covid-19-vaccine; https://investors.modernatx.com/news-releases/news-release-details/modernas-covid-19-vaccine-candidate-meets-its-primary-efficacy

Coronavirus (COVID-19) Update: FDA Authorizes Monoclonal Antibody for Treatment of COVID-19



November 18, 2020

The COVID-19 Treatment Guidelines Panel's Statement on the Emergency Use Authorization of Bamlanivimab for the Treatment of COVID-19

On November 9, 2020, the Food and Drug Administration (FDA) issued an Emergency Use Authorization (EUA) for bamlanivimab (LY-CoV555) for the treatment of nonhospitalized patients who are at high risk of progressing to severe COVID-19 and/or hospitalization. Based on the available data, the Panel has determined the following:

At this t "insufficient data to recommend either for or against" the treatme "should not be considered the standard of care" Bamlan

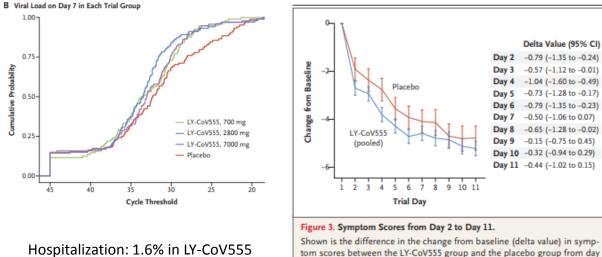
- More data are needed to assess the impact of bamlanivimab on the disease course of COVID-19 and to identify those people who are most likely to benefit from the drug. Health care providers are encouraged to discuss participation in bamlanivimab clinical trials with their patients.
- Patients who are hospitalized for COVID-19 should not receive bamlanivimab outside of a clinical trial.
- Given the possibility of a limited supply of bamlanivimab, as well as challenges distributing and
 administering the drug, patients at highest risk for COVID-19 progression should be prioritized for use of
 the drug through the EUA. In addition, efforts should be made to ensure that communities most affected
 by COVID-19 have equitable access to bamlanivimab.
- The Panel will continue to evaluate emerging clinical data on the use of bamlanivimab for the treatment of
 outpatients with mild to moderate COVID-19 and anticipates updating these recommendations as more
 information becomes available.

Antibody News

The NEW ENGLAND JOURNAL of MEDICINE

ORIGINAL ARTICLE

SARS-CoV-2 Neutralizing Antibody LY-CoV555 in Outpatients with Covid-19



Appendix.

Hospitalization: 1.6% in LY-CoV555 vs. 6.3% in placebo (NNT = 22)

> AE: GI mainly SAE: < 1%

https://pubmed.ncbi.nlm.nih.gov/33113295/; https://www.fda.gov/news-events/press-announcements/coronavirus-covid-19-update-fda-authorizes-monoclonal-antibody-treatment-covid-19 https://www.covid19treatmentguidelines.nih.gov/statement-on-bamlanivimab-eua

2 to day 11. The symptom scores ranged from 0 to 24 and included eight

about the symptom-scoring methods are provided in the Supplementary

domains, each of which was graded on a scale of 0 (no symptoms) to 3 (severe symptoms). The I bars represent 95% confidence intervals. Details





GW Updates

Article Info

THROMBOSIS Log in Register RESEARCH 2 LETTER TO THE EDITORS-IN-CHIEF | VOLUME 197, P65-68, JANUARY 01, 2021 PDF [545 KB] Figures The effect of anticoagulation on clinical outcomes in novel Coronavirus (COVID-19) pneumonia in a U.S. cohort Lei Lynn 😤 🖂 🛛 Juan A. Reyes 😤 🖾 🛛 Katrina Hawkins 🛛 ... Sneha Shah 🛛 Shant Ayanian 🖉 Karolyn Teufel 🛛 Show all authors Published: November 05, 2020 • DOI: https://doi.org/10.1016/j.thromres.2020.10.031 Check for updates **Highlights** Highlights Abstract Anticoagulation strategies for patients with COVID-19 infections remain unclear. CRediT authorship contribution statement · Outcomes of prophylactic versus therapeutic anticoagulation were compared Declaration of competing interest · In our cohort, therapeutic anticoagulation provided no mortality benefit over thromboprophylaxis. References

 Therapeutic anticoagulation was associated with increased adverse events.

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	FIND A DOCTOR	SPECIALTIES	PATIENTS	& VISITO	RS RESI	EARCH	LOCATIO	ONS CONTACT		۹
🖀 🕨 Specia	lties > Primary Care > CO	VID-19 Recovery Clinic	:						TI	0

COVID-19 Recovery Clinic

MULTI-DISCIPLINARY CLINIC OFFERING SPECIALIZED TREATMENT FOR PATIENTS RECOVERING FROM COVID-19

Our physicians at The GW Medical Faculty Associates have been caring for patients with COVID-19 since the early days of the pandemic in our region, and our researchers have been at the forefront of efforts to understand its pathogenesis, immunology, and treatment. As the COVID-19 pandemic continues, some patients who survive the disease experience long-lasting symptoms and negative effects on their health. These can include long-lasting effects on the heart, lung, immune system, mental health, and overall quality of life. Such persistent "long-haul" complications have been reported by our patients and described in the medical literature.

We are offering a new COVID-19 Recovery Clinic to support patients who struggle with long-term symptoms after COVID-19. The new clinic is co-directed by Dr. Monica Lypson and Dr. Aileen Chang from the Division of General Internal Medicine and Dr. Hana Akselrod from the Division of Infectious Diseases, with additional support from:

	Pulmonology	Cardiology				
Related Specialties	Geriatrics & Palliative Medicine	Psychiatry				
Family Practice	Memory Clinic	Rehabilitation Medicine				

Primary Care

Cuentos

Resident Clinic

COVID-19 Recovery Clinic

Visit Medicine Department

https://www.gwdocs.com/specialties/primary-care/covid-19-recovery-clinic/