# UC Irvine UC Irvine Previously Published Works

### Title

41. Assessing Past vs Present COVID-19 Infection: A Survey of Criteria for Discontinuing Precautions in Asymptomatic Patients

### Permalink

https://escholarship.org/uc/item/8bc671xm

**Journal** Open Forum Infectious Diseases, 8(Suppl 1)

### ISSN

2328-8957

### Authors

Gohil, Shruti Maurice, Annabelle Yokoe, Deborah <u>et al.</u>

### **Publication Date**

2021-11-01

Peer reviewed

*Methods.* We identified KTR with COVID-19 between 3/1/2020 and 4/30/2021. Patients were excluded if they had multiorgan transplant or hospital-acquired COVID-19. Data were analyzed by Cox regression with mAb administration as time-dependent variable, and the day of symptom onset as baseline.

**Results.** We studied 95 KTR; 20 received mAb. Comorbidities and immunosuppression were balanced between the two groups. mAb administration was associated with a significant decrease in hospitalizations or ER visits (15 vs. 76%, P< 0.001). This association remained significant after adjustment for confounders and by analyzing mAb administration as a time-dependent variable (Table: adj. HR 0.2, P=0.04). No KTR who received mAb died or required mechanical ventilation. Black or Hispanic KTR were less likely to receive mAb and more likely to be admitted to the hospital or visit the ER (Table).

#### Table

Analysis	Univariate			Multivariate		
Parameter	HR	95% CI	Р	HR	95% CI	Р
Age (years)	1.022	1.004-1.041	0.019	1.023	1.003-1.044	0.024
mAb	0.115	0.036-0.368	0.009	0.216	0.050-0.929	0.040
Chronic kidney disease	2.456	1.243-4.855	0.010	2.087	1.043-4.176	0.038
Black race	2.168	1.186-3.964	0.012	1.881	0.959-3.689	0.066
Hispanic ethnicity	1.701	1.003-2.883	0.049	2.029	1.111-3.703	0.021

#### Factors significantly associated with hospitalization or ER visit.

**Conclusion.** In our KTR population, mAb therapy for COVID-19 may have helped decrease hospitalizations and ER visits. Healthcare inequities, including access to investigational treatments, were exacerbated by the COVID-19 pandemic. Acknowledging the nonconcurrent control group as a limitation, we found a strong signal for benefit from mAb treatment. Antiviral mAb are a promising therapeutic modality for immunosuppressed patients.

Disclosures. Dimitrios Farmakiotis, M.D., Astellas (Grant/Research Support)Merck (Grant/Research Support)Viracor (Grant/Research Support)

40. Lenzilumab Efficacy and Safety in Newly Hospitalized COVID-19 Subjects: Results From a Phase 3 Randomized Double-Blind Placebo-Controlled Trial Zelalem Temesgem, MD<sup>1</sup>; Claudia R. Libertin, MD<sup>4</sup>; Gollen F. Kelley, MD, MPH<sup>5</sup>; Vincent Marconi, MD<sup>6</sup>; Robert Orenstein, DO<sup>7</sup>; Victoria Catterson, PhD<sup>8</sup>; William Aronstein, MD, PhD<sup>9</sup>; Cameron Durrant, MD<sup>10</sup>; Dale Chappell, MD<sup>10</sup>; Omar Ahmed, PharmD<sup>10</sup>; Gabrielle Chappell, MSc<sup>10</sup>; Andrew Badley, M.D.<sup>1</sup>; for the LIVE-AIR Study Group, n/a<sup>10</sup>; <sup>1</sup>Mayo Clinic, Rochester, MN; <sup>2</sup>Hennepin Healthcare Research Institute, Minneapolis, Minnesota; <sup>3</sup>Atrium Health, Charlotte, North Carolina; <sup>4</sup>Mayo Clinic, Jacksonville, Jacksonville, FL; <sup>5</sup>Emory University, Decatur, GA; <sup>6</sup>Atlanta VA, Atlanta, GA; <sup>7</sup>Mayo Clinic in Arizona, Phoenix, AZ; <sup>8</sup>BioSymetrics, Inc, New York, New York <sup>9</sup>CTI, Clinical Trial Services, Inc, Covington, Kentucky; <sup>10</sup>Humanigen, Inc, Burlingame, California

Drs. Meghan Lewis, Linda Sher, Michael Bowdish, Noah Wald-Dickler, Subarna Biswas, Lydia Lam, Khang Vo, Roy Poblete, May M. Lee, Douglass Hutcheon from the University of Southern California (USC) Keck and LAC Medical Centers; Drs. Zelalem Temesgen, and Andrew D. Badley, MD from the Mayo Clinic, Rochester MN; Drs. Charles D. Burger and Claudia R. Libertin from Mayo Clinic, Jacksonville FL; Dr. Jason Baker from Hennepin Healthcare Research Institute Minneapolis, MN; Dr. Victoria Catterson from BioSymetrics, Inc., New York, NY; Dr. William S. Aronstein from CTI, Covington, KY; Drs. Cameron Durrant, Dale Chappell, Omar Ahmed, and Gabrielle Chappell from Humanigen, Inc., Burlingame, CA; Drs. Robert Orenstein and Roberto Patron from Mayo Clinic Arizona, Drs. Colleen F. Kelley, John Gharbin, Caitlin Moran, Sheetal Kandiah, Valeria Cantos, Paulina Rebolledo, Carlos del Rio, Jeffrey Lennox, Carmen Polito, Paulina Rebolledo, Anandi Sheth from Emory University Medical Center and Grady Memorial Hospital; Drs. Anup Patel, Homero Paniagua from St. Barnabas Medical Center; Dr. Seife Yohannes from MedStar Washington Hospital Center; Drs. Alpesh Amin, Richard Lee, Miki Watanabe, Lanny Hsieh from the University of California-Irvine Medical Center; Drs. Martin Cearras, Amay Parikh, Jason Sniffen, Wilfred Onyia from AdventHealth Orlando; Drs. Vincent C. Marconi, Christopher Polk, Michael Boger, Lisa Davidson, Kiran Gajurel, Michael Leonard, Lewis McCurdy, Nestor Quezada, Mindy Sampson, Zainab Shahid, Stephanie Strollo, David Weinrib, Sara Zulfigar from Atrium Health; Drs. Cheryl McDonald, John Hollingsworth, John Burk, Joshua Berg, Daniel Barbaro, Andrew Miller, Lakshmi Sambathkumar, Stuart McDonald, Obinna Okoye from Texas Health Harris Methodist Hospital Fort Worth; Drs. Juan Pulido, Jennifer Fulton, William Gill from Baptist Health Research Institute Jacksonville; Drs. Richard Zuckerman, Lionel Lewis from Dartmouth-Hitchcock Medical Center; Dr. Chaitanya Mandapakala from St. Elizabeth Medical Center; Drs. Matthew Robinson, Brian Metzger from St. David's Medical Center; Drs. Maqsood Alam, Chrisoula Politis from Mercy Medical;

Drs. Anne Frosch, Linh Ngo from Hennepin Healthcare; Drs. Fernando Carvalho Neuenschwander, Estevão Figueiredo, Gualter Cançado, Gustavo Araujo, Lucas Guimarães from Hospital Vera Cruz (NUPEC) in Brazil; Drs. Ricardo Diaz, Natalia Bacellar, Celso Silva, Paulo Ferreira, from Escola Paulista de Medicina (UNIFESP) in Brazil; Dr. Marina Andrade Lima, Caroline Uber Ghisi, Camila Anton, Ricardo Albaneze from Hospital Dia do Pulmãoin Brazil; Dr. Daniel Wagner de Castro Lima Santos, Ana Caroline Iglessias, Marianna Lago, Paula Pietrobom, Maysa Alves from Hospital São Luiz do Jabaquara (IDOR) in Brazil; Drs. Juvencio José Duailibe Furtado, Leopoldo Trevelin, Valeria Telles, Francini Correa from Hospital Heliópolis in Brazil; Drs. Fabiano Ramos, Marina de A. R. Da Silva, Rebeca C. Lacerda Garcia, Ana Elizabeth G. Maldonado, Ana Carolina M. Beheregaray, Ana Maria T. Ortiz from Hospital São Lucas (PUCRS); Drs. Kleber Luz, Eveline Pipolo Milan, Janine Soares de Castro, Matheus José Barbosa Moreira, Renata Bezerra Onofre, Tácito do Nascimento Jácome, Victor Barreto Garcia, Victor Matheus Rolim de Souzafrom Centro de Pesquisas Clínicas de Natal (CPCLIN) in Brazil; Drs. Felipe Dal Pizzol, Cristiane Ritter, Marcelo B. Vinhas from Sociedade Literaria e Caritativa Santo Agostinho (SLCSA) in Brazil; Drs. Adilson Joaquim Westheimer Cavalcante, Julia Minghini, Loni Dorigo, Marina Salgado Miranda from Centro Multidisciplinar de Estudos Clínicos (CEMEC) in Brazil; Drs. Martti Anton Antila, Rebeca Brugnolli, Henrikki Antila from Consultoria Médica e Pesquisa Clínica (CMPC) in Brazil

#### Session: O-08. COVID-19 Treatment & Diagnostics

**Background.** Severe coronavirus disease 2019 (COVID-19) often results from the immune-mediated cytokine storm, triggered by granulocyte macrophage-colony stimulating factor (GM-CSF), potentially leading to respiratory failure and death. Lenzilumab, a novel anti-human GM-CSF monoclonal antibody, neutralizes GM-CSF and demonstrated potential to improve clinical outcomes in a matched case-cohort study of patients with severe COVID-19 pneumonia. This Phase 3 randomized, double-blind, placebo-controlled trial investigated the efficacy and safety of lenzilumab to improve the likelihood of survival without invasive mechanical ventilation (SWOV), bevond available treatments.

*Methods.* Hypoxic patients, hospitalized with COVID-19 (n=520), requiring supplemental oxygen, but not invasive mechanical ventilation, were randomized on Day 0 to receive lenzilumab (1800mg, n=261) or placebo (n=259), and available treatments, including remdesivir and/or corticosteroids; and were followed through Day 28.

**Results.** Baseline demographics were comparable between groups: male, 64.7%; mean age, 60.5 years; median CRP, 79.0 mg/L. Patients across both groups received steroids (93.7%), remdesivir (72.4%), or both (69.1%). Lenzilumab improved the primary endpoint, likelihood of SWOV in the mITT population, by 1.54-fold (HR: 1.54; 95%CI: 1.02-2.32, p=0.0403). Lenzilumab improved SWOV by 1.91-fold (nominal p=0.0073) and 1.92-fold (nominal p=0.0067) in patients receiving remdesivir or remdesivir and corticosteroids, respectively. A key secondary endpoint of incidence of IMV, ECMO or death was also improved in patients receiving remdesivir (p=0.0200) or remdesivir and corticosteroids (p=0.0180). Treatment-emergent serious adverse events were similar across both groups.

**Conclusion.** Lenzilumab significantly improved SWOV in hypoxic COVID-19 patients upon hospitalization, with the greatest benefit observed in patients receiving treatment with remdesivir and corticosteroids. NCT04351152

Disclosures. Zelalem Temesgem, MD, Humanigen, Inc (Grant/Research Support) Jason Baker, MD, Humanigen, Inc (Grant/Research Support) Christopher Polk, MD, Atea (Research Grant or Support)Gilead (Advisor or Review Panel member, Research Grant or Support)Humanigen (Research Grant or Support)Regeneron (Research Grant or Support) Claudia R. Libertin, MD, Gilead (Grant/Research Support) Colleen F. Kelley, MD, MPH, Gilead Sciences (Individual(s) Involved: Self): Grant/Research Support; Moderna (Individual(s) Involved: Self): Grant/Research Support; Novavax (Individual(s) Involved: Self): Grant/Research Support; Viiv (Individual(s) Involved: Self): Grant/Research Support Vincent Marconi, MD, Bayer (Consultant, Scientific Research Study Investigator)Eli Lilly (Consultant, Scientific Research Study Investigator)Gilead Sciences (Consultant, Scientific Research Study Investigator) ViiV (Consultant, Scientific Research Study Investigator) Victoria Catterson, PhD, Humanigen, Inc (Consultant) William Aronstein, MD, PhD, Humanigen, Inc (Consultant) Cameron Durrant, MD, Humanigen, Inc (Employee) Dale Chappell, MD, Humanigen, Inc (Employee) Omar Ahmed, PharmD, Humanigen, Inc (Employee) Gabrielle Chappell, MSc, Humanigen, Inc (Consultant) Andrew Badley, M.D., AbbVie (Consultant) for the LIVE-AIR Study Group, n/a, Humanigen, Inc (Grant/Research Support)

## 41. Assessing Past vs Present COVID-19 Infection: A Survey of Criteria for Discontinuing Precautions in Asymptomatic Patients

Shruti K. Gohil, MD, MPH<sup>1</sup>; Annabelle De St. Maurice, MD<sup>2</sup>; Deborah S. Yokoe, MD, MPH<sup>3</sup>; Deborah S. Yokoe, MD, MPH<sup>3</sup>; Stuart H. Cohen, MD<sup>4</sup>; Francesca J. Torriani, MD<sup>5</sup>; Jonathan Grein, MD<sup>6</sup>; Philip A. Robinson, MD<sup>7</sup>; Shannon C. Mabalot, MPH, CIC<sup>8</sup>; Paula Pedrani, BS<sup>1</sup>; Jessica Park, BS<sup>9</sup>; Richard Platt, MD, MSc<sup>10</sup>; Susan S. Huang, MD, MPH<sup>9</sup>; <sup>1</sup>UC Irvine School of Medicine, IRVINE, California; <sup>2</sup>University of California, Los Angeles, Los Angeles, CA; <sup>3</sup>University of California, San Francisco, San Francisco, CA; <sup>4</sup>University of California, Davis, Sacramento, CA; <sup>5</sup>University of California, San Diego, San Diego, CA; <sup>6</sup>Cedars-Sinai Medical Center, Los Angeles, CA; <sup>7</sup>Hoag Hospital, Irvine, California; <sup>8</sup>Sharp Memorial Hospital, San Diego, CA; <sup>9</sup>University of California, Irvine, Irvine, California;<sup>10</sup>Harvard Medical School, Boston, Massachusetts

### CDC Epicenters

Session: O-08. COVID-19 Treatment & Diagnostics

Background. COVID-19 patients can remain positive by PCR-testing for several months. Pre-admission or pre-procedure testing can identify recovered asymptomatic patients who may no longer be contagious but would require precautions according to current CDC recommendations (10 days). This can result in unintended consequences, including procedure delays or transfer to appropriate care (e.g., psychiatric or post-trauma patients requiring admission to COVID-19 units instead of psychiatric or rehabilitation facilities, respectively).

Methods. We conducted a structured survey of healthcare epidemiologists and infection prevention experts from the SHEA Research Network between March-April, 2021. The 14-question survey, presented a series of COVID-19 PCR+ asymptomatic patient case scenarios and asked respondents if (1) they would consider the case recovered and not infectious, (2) if they have cleared precautions in such cases, and if so, (3) how many transmission events occurred after discontinuing precautions. The survey used one or a combination of 5 criteria: history of COVID-19 symptoms, history of exposure to a household member with COVID-19, COVID-19 PCR cycle threshold (CT), and IgG serology. Percentages were calculated among respondents for each question.

Results. Among 60 respondents, 56 (93%) were physicians, 51 (86%) were hospital epidemiologists, and 46 (77%) had >10y infection prevention experience. They represented facilities that cumulatively cared for >29,000 COVID-19 cases; 46 (77%) were academic, and 42 (69%) were large ( >400 beds). One-third to one-half would consider an incidentally found PCR+ case as recovered based on solo criteria, particularly those with two consecutive high CTs or COVID IgG positivity recovered (53-55%) (Table 1). When combining two criteria, half to four-fifths of respondents deemed PCR+ cases to be recovered (Table 2). Half of those had used those criteria to clear precautions (45-64%) and few to none experienced a subsequent transmission event resulting from clearance.

Conclusion. The majority of healthcare epidemiologists consider a combination of clinical and diagnostic criteria as recovered and many have used these to clear precautions without high numbers of transmission.

#### Table 1: Assessment of COVID-19 recovery of asymptomatic case scenarios using solo criterion to identify potentially recovered patients for discontinuation of precautions.

SOLO CRITERION						
Asymptomatic Case Scenario	Percent That Believe Case Is Recovered and Not infectious	Percent That Have Used This Criterion to Clear Precautions <sup>1</sup>	Number of Transmission Events Among Those Released from Precautions Using This Criterion <sup>2</sup>			
HISTORY OF COVID-LIKE SYMPTOMS Clear history of COVID-like symptoms 1 month ago. Now COVID PCR CT* is 30. COVID IgG* negative.	42%	68%	0			
EXPOSURE TO COVID+ HOUSEHOLD/CLOSE CONTACT No history of symptoms. Known household COVID+ contact 5 weeks ago. Current COVID PCR CT is 30. COVID IgG negative.	33%	50%	0			
HIGH CYCLE THRESHOLD No history of symptoms or COVID+ contact. Current COVID PCR CT is >35. COVID IgG negative.	37%	45%	1			
TWO HIGH CYCLE THRESHOLDS No history of symptoms or COVID+ contact. Current COVID PCR CT is >35 on two separate days. COVID IgG negative.	53%	69%	1			
COVID IgG+ No history of symptoms or COVID+ contact. Current COVID PCR CT is 30. COVID IgG positive.	55%	52%	0			

Percent calculated among those who believe the case is recovered/not infectious

Abbreviations: CT = Cycle Threshold time, IgG – Immunoglobulin G

Table 2: Assessment of COVID-19 recovery of asymptomatic case scenarios using dual criteria to identify potentially recovered patients for discontinuation of precautions. 

DUA	CRITERIA		
Asymptomatic Case Scenario	Percent That Believe Case Is Recovered and Not infectious	Percent That Have Used This Criteria to Clear Precautions <sup>1</sup>	Number of Transmission Events Among Those Released from Precautions Using These Criteria <sup>2</sup>
HISTORY OF COVID-LIKE SYMPTOMS + EXPOSURE TO			
COVID+ HOUSEHOLD OR CLOSE CONTACT Patient with clear history of COVID-like symptoms 1 month ago, known household COVID+ contact 5 weeks ago.	52%	58%	O
HISTORY OF COVID-LIKE SYMPTOMS AND HIGH CYCLE			
THRESHOLD Patient with clear history of COVID-fike symptoms 1 month ago. No COVID+ contact. Current COVID PCR CT is >35. COVID IgG negative.	58%	60%	1
HISTORY OF COVID-LIKE SYMPTOMS AND TWO HIGH			
CYCLE THRESHOLDS Patient with clear history of COVID-like symptoms 1 month ago. No COVID+contact. Current COVID PCRCT is >35 on two separate days. COVID IgG negative.	65%	64%	0
HISTORY OF COVID-LIKE SYMPTOMS AND COVID IgG+ Patient with clear history of COVID-like symptoms 1 month			
ago. No known COVID+ contact. Current COVID PCR CT is 30. COVID IgG positive.	65%	56%	0
EXPOSURE TO COVID+ HOUSEHOLD OR CLOSE CONTACT			
AND HIGH CYCLE THRESHOLD Patient asymptomatic and no history of symptoms. Known household COVID+ contact 5 weeks ago, current COVID PCR CT is >35. COVID IgG negative.	50%	45%	O
EXPOSURE TO COVID+ HOUSEHOLD OR CLOSE CONTACT			
AND TWO HIGH CYCLE THRESHOLDS No history of symptoms. Known household COVID+ contact 5 weeks ago. Current COVID PCR CT is >35 on two separate days. COVID kgG negative.	60%	50%	1
EXPOSURE TO COVID+ HOUSEHOLD OR CLOSE CONTACT AND COVID IgG+ No history of symptoms. Known household COVID+ contact 5 weeks ago, current COVID PCR CT is 30. COVID IgG positive.	62%	46%	D
HIGH CYCLE THRESHOLD AND COVID IgG+ No history of symptoms. No known COVID+ contact. Current COVID PCR CT is>35, and COVID IgG positive.	63%	50%	1
TWO HIGH CYCLE THRESHOLDS AND COVID IgG+ No history of symptoms. No known COVID + contact. Current COVID PCR CT is ≥ 35 on two separate days. COVID IgG positive.	83%	48%	1

<sup>1</sup>Percent calculated among those who believe the case is recovered/not infectious. <sup>2</sup>Number of known transmission events

\*Abbreviations: CT = Cycle Threshold time, IgG – Immunoglobulin G

Disclosures. Shruti K. Gohil, MD, MPH, Medline (Other Financial or Material Support, Co-Investigator in studies in which participating hospitals and nursing homes received contributed antiseptic and cleaning products)Molnycke (Other Financial or Material Support, Co-Investigator in studies in which participating hospitals and nursing homes received contributed antiseptic and cleaning products)Stryker (Sage) (Other Financial or Material Support, Co-Investigator in studies in which participating hospitals and nursing homes received contributed antiseptic and cleaning products) Deborah S. Yokoe, MD, MPH, Nothing to disclose Stuart H. Cohen, MD, Seres (Research Grant or Support) Jonathan Grein, MD, Gilead (Other Financial or Material Support, Speakers fees) Richard Platt, MD, MSc, Medline (Research Grant or Support, Other Financial or Material Support, Conducted studies in which participating hospitals received contributed antiseptic product)Molnlycke (Other Financial or Material Support, Conducted studies in which participating hospitals received contributed antiseptic product) Susan S. Huang, MD, MPH, Medline (Other Financial or Material Support, Conducted studies in which participating hospitals and nursing homes received contributed antiseptic and cleaning products) Molnlycke (Other Financial or Material Support, Conducted studies in which participating hospitals and nursing homes received contributed antiseptic and cleaning products)Stryker (Sage) (Other Financial or Material Support, Conducted studies in which participating hospitals and nursing homes received contributed antiseptic and cleaning products)Xttrium (Other Financial or Material Support, Conducted studies in which participating hospitals and nursing homes received contributed antiseptic and cleaning products)