COVID-19 outcomes of IHCA for patients with nonshockable in the setting of COVID-19 may not actually differ from pre–COVID-19. Nonetheless, this study found only 18 patients (1.3%) achieved return of spontaneous circulation, 4 (3%) survived to 30 days, and only 1 (<1%) achieved a favorable neurological outcome by 30 days. In this issue of JAMA Internal Medicine, Thapa et al² report what to our knowledge is the first US data on outcomes for IHCA among patients with COVID-19. In their case series of 54 patients, 52 (96%) had a nonshockable initial rhythm, 29 (54%) achieved return of spontaneous circulation, and 0 survived to hospital discharge (95% CI, 0%-6.6%). This very low hospital survival is likely driven by several factors, including critical illness in most patients at the time of arrest and the many patients with nonshockable initial rhythms. Additionally, presumed respiratory etiology of arrest for most patients, lack of therapies to effectively treat the underlying disease, and potential delays in response time for donning of personal protective equipment may have contributed to poor outcomes.

These small case series reporting hospital survival after IHCA among patients with COVID-19 must be interpreted with caution, as only 1 or 2 additional survivors would make important differences in the observed estimates. Outcomes in the setting of COVID-19 may not actually differ from pre–COVID-19 outcomes of IHCA for patients with nonshockable rhythms, for whom hospital survival is often less than 15%. Nonetheless, this article² represents important early evidence suggesting outcomes for IHCA in patients with COVID-19 pneumonia are likely poor, particularly among patients with respiratory failure. Improving outcomes for patients with severe illness with COVID-19 and IHCA will be challenging, as few of the likely drivers of poor outcomes (eg, nonshockable rhythms, respiratory etiologies of arrest, and underlying critical illness) are modifiable. While these early results should not warrant universal do-not-attempt-resuscitation (DNAR) orders for patients with COVID-19, they highlight the importance of conducting goals-of-care discussions early during the course of COVID-19 and revisiting those discussions with changes in clinical status (worsening or improvement). Moreover, the existing data may warrant clinician recommendations for DNAR, particularly in patients with severe respiratory failure who are at high risk of IHCA. An informed assent approach, in which the patient or family is invited to allow the clinician to assume responsibility for the DNAR decision, may be appropriate in select patients to help alleviate the psychological burden of decision-making on patients and families during this stressful time. Like traditional informed consent, this approach places substantial responsibility on clinicians to have open, respectful, and thoughtful communication with patients and families.

Although this study was not designed to examine racial disparities, it is notable that two-thirds of the patients were Black. Previous studies have reported that a larger minority of Black patients request CPR in the context of poor prognoses. Black persons also have lower rates of advance care planning documentation and report poorer quality communication during serious illness and greater mistrust in the health system that are associated with long-standing and ongoing disparities in health care. Finding ways to respect differences in preferences and eliminate disparities in high-quality communication during serious illness is critically important. Building trust with patients is crucial to effective communication, and clinician recommendations made without trust have potential for harm. In the context of COVID-19, Black persons and persons of color are more likely to contract COVID-19 or develop serious illness requiring hospitalization; this association is most likely because of disparities. As such, the urgency of eliminating racial disparities in health care has never been clearer.

The long-standing need to improve the conduct and timeliness of high-quality goals-of-care discussions for patients with serious illness has become even more important in the time of COVID-19. Promotion of early goals-of-care discussions should be a priority for patients, families, clinicians, health systems, and policy makers. Such a shared focus offers substantial opportunity for health system and public health interventions. Established programs, such as The Conversation Project (Institute for Healthcare Improvement; http://www.theconversationproject.org) and PRE-PARE For Your Care (The Regents of the University of California; http://www.prepareforyourcare.org), both of which offer new COVID-19–specific guidance, are important resources to help prepare patients and their families for in-the-moment decision-making should they be hospitalized with COVID-19. For selected patients with chronic life-limiting illness and preferences for limitations on life-sustaining treatments, completing the Physician Orders for Life-Sustaining Treatment may reduce unwanted
high-intensity care near the end of life. Although there are important limitations on current data regarding outcomes of IHCA for patients with COVID-19, we have enough data to conclude that it is important to implement programs to promote conversations about values and goals in the community and early goals-of-care discussions for patients hospitalized with COVID-19.

Matthew E. Modes, MD, MPP, MS
Robert Y. Lee, MD, MS
J. Randall Curtis, MD, MPH

Author Affiliations: Division of Pulmonary, Critical Care and Sleep Medicine, University of Washington, Seattle (Modes, Lee, Curtis); Cambia Palliative Care Center of Excellence, University of Washington, Seattle (Modes, Lee, Curtis).

Corresponding Author: J. Randall Curtis, MD, MPH, Division of Pulmonary, Critical Care, and Sleep Medicine, Box 359762, Harborview Medical Center, University of Washington, Seattle, WA 98104 (jrc@u.washington.edu).


Conflict of Interest Disclosures: Dr Modes reported grants from the National Institutes of Health (T32 # HL 125195; University of Washington Palliative Care T32) during the conduct of the study. Dr Lee reported grants from the National Institutes of Health during the conduct of the study. Dr Curtis reported grants from the National Institutes of Health, Cambia Health Foundation, and National Palliative Care Research Center outside the submitted work.


Characteristics and Reporting of Number Needed to Treat, Number Needed to Harm, and Absolute Risk Reduction in Controlled Clinical Trials, 2001-2019

Controlled clinical trials, which are used to guide the decisions made by patients, clinicians, and policy makers, often only report measures of relative effect. However, absolute measures, such as the absolute risk reduction (ARR), the number needed to treat (NNT), and the number needed to harm (NNH), which measure the difference in the observed risk of an event between 2 interventions and the number of patients who need to be treated to achieve 1 additional favorable or adverse outcome, respectively, can be easier to interpret, more clinically meaningful, and less likely to exaggerate differences when outcome risk is low. In part because only 5% of trials published in highly cited journals before 1998 reported NNT and/or ARR, the Consolidated Standards of Reporting Trials (CONSORT) statement recommended that trials with binary outcomes report both relative and absolute measures. We assessed the recent trends and characteristics of absolute measure reporting in highly cited medical journals to determine if there have been improvements over time.

Methods | We identified the 6 most-cited medical journals according to InCites Journal Citation Reports (Clarivate Analytics 2019) (Table 1). For each journal, we reviewed all issues published in 2001, 2007, 2013, and 2019 to identify all controlled clinical trials that reported analyses testing superiority of the intervention to control and abstract-level binary outcomes, including hazard ratios. For eligible trials, we identified key study characteristics and recorded whether at least 1 abstract-level positive (P < .05) binary efficacy and/or safety outcome was reported. Next, we determined whether any NNT, NNH, and/or ARR was reported in the abstract and/or full text. For each NNT/NNH, we recorded if reporting was for primary or secondary end points and whether 95% CIs, P values, and corresponding effect estimates were provided. Fisher exact and Mann-Whitney U tests were conducted in R, version 3.4.0 (R Foundation for Statistical Computing) (2-sided P < .05). Because publicly available data were used, this study did not require ethics approval or patient consent.

Results | We identified 875 controlled trials meeting the aforementioned criteria, of which 76 (8.7%) reported at least 1 NNT, 8 (0.9%) reported at least 1 NNH, and 249 (28.5%) reported at least 1 ARR (Table 1). In total, 292 trials (33.4%) reported at least 1 NNT, NNH, and/or ARR. A total of 80 (9.1%) reported at least 1 NNT and/or NNH, which remained relatively constant between 2001 and 2019; ARR reporting increased from 26 of 140 (18.6%) to 105 of 282 (37.2%); P < .001).

In the therapeutic area of oncology had the lowest rates of reporting NNT, NNH, and/or ARR, but there were no differences by intervention tested, patient follow-up, enrollment, or funding sources (Table 2). Trials with at least 1 statistically significant end point were more likely to report an NNT/NNH than those without (75 of 624 [12.0%] vs 5 of 251 [2.0%]; P < .001).

Among all 197 NNT/NNH reports, 95 (48.2%) were for primary end points and 76 (38.6%) had a 95% CI and/or P value. There were 114 NNT/NNH reports with a corresponding effect estimate reported anywhere in the text, of which 88 (77.2%) were statistically significant; 55 (48.2%) only had corresponding relative measures.

Discussion | Among 875 controlled trials with binary outcomes and/or hazard ratios published in highly cited general medical journals, fewer than one-tenth reported at least 1 NNT or NNH, but more than one-quarter reported at least 1 ARR. The majority of NNT/NNH reports were presented for statistically significant end points but without 95% CIs or P values. These findings raise concerns about persistent...