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The impact of COVID-19 pandemic on conducting emergency medicine clinical research



The COVID-19 pandemic challenges the status quo of conducting emergency medicine (EM) research [1]. Research generates new knowledge for the advancement of the medical field, abiding by ethical parameters [1–3]. As EM is on the frontlines of this global crisis, the aim of this correspondence is to consider the impacts of the COVID-19 pandemic on EM clinical research.

Financial burdens for EM research have been an ongoing concern prior to the COVID-19 pandemic. For example, NIH funding has proved a challenge for EM researchers, garnering only 1.7% of funds [4–6]. Additional safety considerations in EM Departments due to COVID-19 drastically increased [7]. Researchers have the added obstacle of devising pathways to safely continue EM research. There has also been a drop in non-COVID-19 patients. The 31%–45% reduction in ED visits across the US, raises concern for patients not seeking care [8]. For instance, there had been a decrease in patients presenting with strokes and STEMIs [8]. This patient population is a crucial element for research; thus, the potential number of patients that could contribute to EM trials is decreased. On the other hand, COVID-19, though disruptive, has opened up new research opportunities [4,9]. For example, Stanford's EM Department is funding a study evaluating clinical characteristics of COVID-19 patients [10,11]. Other avenues include exploring topics regarding domestic safety during a pandemic, patient and physician well-being, and alternative communication avenues [3]. Though most institutions around the country have begun to proceed past the COVID-19 crisis, the new pathways devised to maintain research during such unsteady times must be incorporated into future practices, cultivating sustainable research methods that are resilient to crises.

During this crisis, the price of patient-safety must be weighed against the cost of halting clinical trials. However, some clinical trials have adapted to the current circumstances and are carrying out their trials via telemedicine, producing sustainable research methods [12]. Though not all clinical research can be fully conducted via telemedicine, such as interventional research, it can be incorporated through its use in participant virtual consenting and follow-up [13–15]. This spares patients traveling, as well as reduces potential COVID-19 exposure [1]. For non-interventional research, virtual resources offer web-based platforms to execute survey-based studies as well as data processing components of various types of studies. There is a compelling applicability of utilizing telemedicine in a post-COVID-19 era to conduct clinical research. Continued use of telemedicine could allow clinical trials to address patient barriers to care. Addressing financial and geographical barriers, (although technical barriers could be created) could allow for a larger patient population in clinical trials, increasing the statistical power of trials.

Another future application of telehealth is through its ability to continue the exchange of ideas, despite social distancing hindrances [9,13–15]. Virtual technology maintains education; for example, residents can continue their research education and collaborations through online lectures, meetings, and web conferences [9,15]. Previously, virtual resources have been underutilized in research, though in light of the COVID-19 pandemic, its functionality in clinical research is beginning to reveal its efficiency and efficacy. The applicability of telemedicine has to be fully explored [8,13–15].

In addition to expanding the use of telemedicine into research where applicable, the mental health of EM researchers should be considered moving forward. Throughout this pandemic, researchers should remain cognizant of symptoms of burnout and depression, which include anxiety, fatigue, and an overwhelming sense of sadness, and seek psychosocial support if this occurs [16,17]. To prevent negative psychological effects, clinical researchers should perform regular self-check-ins, engage in physical activity, and stay connected with family and friends [16]. Continuing to part take in these discussions is a productive approach to support one's mental well-being and should be continued even after the end of this global crisis [8,16,17].

The future recommendation for EM research productivity should be focused on building and maintaining sustainable practices throughout the rest of the COVID-19 pandemic and into the post-COVID-19 era. We have several recommendations moving forward. For conducting research virtually, non-interventional or observational studies should be adaptable to virtual secure platforms to create alternative efficient, portable methods that can expand participant populations. From this, we can draw on lessons to evaluate how funding may change in the future and implement strategies to maximize EM clinical research. In regard to maintaining good mental health practices, we recommend optimizing proactive measures such as regular physical activity and time with family and friends.

Research, an integral aspect of the EM community, has been affected in many ways by this global pandemic. Safety issues have to be considered to properly decide how best to proceed or temporarily halt research studies. According to the Declaration of Helsinki, “While the primary purpose of medical research is to generate new knowledge, this goal can never take precedence over the rights and interests of individual research subjects.” [2] Although these are trying times, the creative avenues designed by researchers to continue working ethically fosters inspiration for the future.

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