

“Harmful laws can exacerbate stigma and discrimination, infringe on people’s rights and undermine public health responses,” according to Winnie Byanyima, Executive Director of UNAIDS. “To ensure responses to the pandemic are effective, humane and sustainable, governments must use the law as a tool to uphold the human rights and dignity of people affected by COVID-19.”

The COVID-19 Law Lab is a database of laws that countries have implemented in response to the pandemic. It includes state of emergency declarations, quarantine measures, disease surveillance, legal measures relating to mask-wearing, social distancing, and access to medication and vaccines. The database will continue to grow as more countries and themes are added.

It will also feature research on different legal frameworks for COVID-19. These analyses will focus on the human rights impacts of public health laws and help countries identify best practices to guide their immediate responses to COVID-19 and socioeconomic recovery efforts once the pandemic is under control. It builds off the work of the UHC Legal Solutions Network, which was established to help countries achieve universal health coverage through the implementation of rights-based legal frameworks.

“We need to track and evaluate how laws and policies are being used during the Pandemic to understand what works,” said Dr. Matthew M. Kavanagh, faculty in Georgetown University’s Department of International Health. Katie Gottschalk, Executive Director of the O’Neill Institute for National and Global Health Law at Georgetown University Law Center added, “We must learn lessons from the early stage of pandemic policies to implement the most effective laws going forward – the COVID-19 Law Lab allows us to do just that.”

Available from: <https://www.who.int/news-room/detail/22-07-2020-new-covid-19-law-lab-to-provide-vital-legal-information-and-support-for-the-global-covid-19-response>

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## WHO AND UNICEF WARN OF A DECLINE IN VACCINATIONS DURING COVID-19

**4 July 2020** - WHO today accepted the recommendation from the Solidarity Trial’s International Steering Committee to discontinue the trial’s hydroxychloroquine and lopinavir/ritonavir arms. The Solidarity Trial was established by WHO to find an effective COVID-19 treatment for hospitalized patients.

The International Steering Committee formulated the recommendation in light of the evidence for hydroxychloroquine vs standard-of-care and for lopinavir/ritonavir vs standard-of-care from the Solidarity trial interim results, and from a review of the evidence from all trials presented at the 1-2 July WHO Summit on COVID-19 research and innovation.

These interim trial results show that hydroxychloroquine and lopinavir/ritonavir produce little or no reduction in the mortality of hospitalized COVID-19 patients when compared to standard of care. Solidarity trial investigators will interrupt the trials with immediate effect.

For each of the drugs, the interim results do not provide solid evidence of increased mortality. There were, however, some associated safety signals in the clinical laboratory findings of the add-on Discovery trial, a participant in the Solidarity trial. These will also be reported in the peer-reviewed publication.

This decision applies only to the conduct of the Solidarity trial in hospitalized patients and does not affect the possible evaluation in other studies of hydroxychloroquine or lopinavir/ritonavir in non-hospitalized patients or as pre- or post-exposure prophylaxis for COVID-19. The interim Solidarity results are now being readied for peer-reviewed publication.

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