Clinical Laboratory COVID-19 Response Call Monday, May 3, 2021 at 3:00 PM EDT

Welcome

- Jasmine Chaitram, CDC Division of Laboratory Systems (DLS)
- Surveillance Testing of Wastewater for COVID-19
 - Amy Kirby, CDC Division of Foodborne, Waterborne, and Environmental Diseases (DFWED)
- Investigation of Linked Clusters of SARS-CoV-2 Variant B.1.351
 - Kenneth Feder, Maryland Department of Health
- FDA Update
 - Tim Stenzel, U.S. Food and Drug Administration (FDA)

CDC Preparedness Portal

https://www.cdc.gov/csels/dls/preparedlabs/covid-19-clinical-calls.html

Find CLCR call information, transcripts, and audio recordings on the CDC Preparedness Portal



Schedule for Clinical Laboratory COVID-19 Response Calls

The next call will be on **Monday, May 17** from 3:00 PM to 4:00 PM EDT



We Want to Hear from You!

Training and Workforce Development

Questions about education and training?

Contact LabTrainingNeeds@cdc.gov



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How to Ask a Question

- Using the Zoom Webinar System
 - Click the Q&A button in the Zoom webinar system
 - Type your question in the Q&A box and submit it
 - Please do not submit a question using the chat button





If you are a patient, please direct any questions to your healthcare provider



Slide decks may contain presentation material from panelists who are not affiliated with CDC. Presentation content from external panelists may not necessarily reflect CDC's official position on the topic(s) covered.

Center for Surveillance, Epidemiology, and Laboratory Services

Surveillance Testing of Wastewater for COVID-19

CDC Division of Foodborne, Waterborne, and Environmental Diseases (DFWED)



Center for Surveillance, Epidemiology, and Laboratory Services

Investigation of Linked Clusters of SARS-CoV-2 Variant B.1.351

Kenneth Feder Maryland Department of Health



Center for Surveillance, Epidemiology, and Laboratory Services

FDA Update

Tim Stenzel

U.S. Food and Drug Administration (FDA)



U.S. Food and Drug Administration (FDA)

COVID-19 Emergency Use Authorization (EUA)
Information for Medical Devices

https://www.fda.gov/medical-devices/emergencysituations-medical-devices/emergency-useauthorizations

COVID-19 In Vitro Diagnostic EUAs

https://www.fda.gov/medical-devices/coronavirus-disease-2019-covid-19-emergency-use-authorizations-medical-devices/vitro-diagnostics-euas

COVID-19 Frequently Asked Questions

https://www.fda.gov/emergency-preparedness-and-response/coronavirus-disease-2019-covid-19/coronavirus-disease-2019-covid-19-frequently-asked-questions

COVID-19 Updates

https://www.fda.gov/emergency-preparedness-and-response/mcm-legal-regulatory-and-policy-framework/emergency-use-authorization#2019-ncov

FDA Townhall Meetings

https://www.fda.gov/medical-devices/workshopsconferences-medical-devices/virtual-town-hall-seriesimmediately-effect-guidance-coronavirus-covid-19diagnostic-tests-06032020

 Independent Evaluations of COVID-19 Serological Tests

https://open.fda.gov/apis/device/covid19serology/



U.S. Food and Drug Administration (FDA)

COVID-19 Diagnostic Development
CDRH-EUA-Templates@fda.hhs.gov

- Spot Shortages of Testing Supplies: 24-Hour Support Available
 - 1. Call 1-888-INFO-FDA (1-888-463-6332)
 - 2. Then press star (*)
- FDA MedWatch

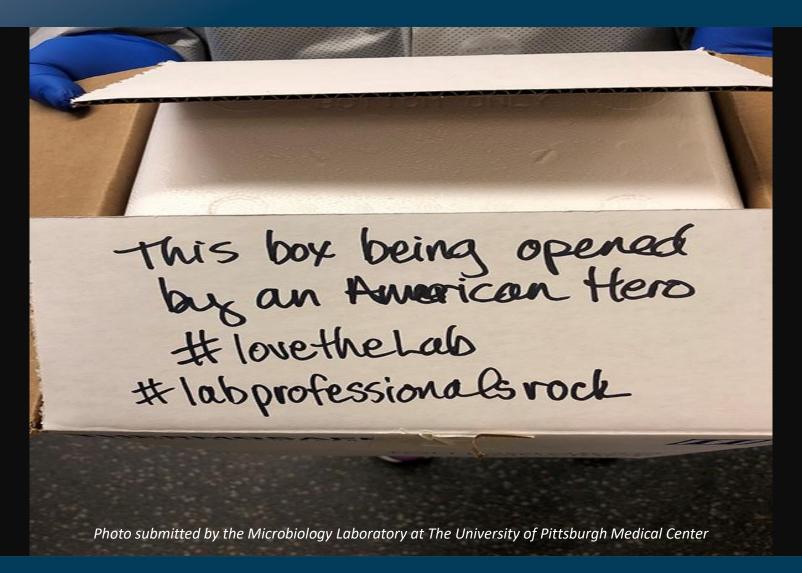
https://www.fda.gov/safety/medwatch-fda-safety-information-and-adverse-event-reporting-program



CDC Social Media



Thank You For Your Time!



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13