Cepheid gets EUA for Xpert Xpress CoV-2/Flu/RSV Plus

October 2021—Cepheid received FDA emergency use authorization for its Xpert Xpress CoV-2/Flu/RSV Plus, a rapid molecular diagnostic test for qualitative detection of the viruses causing COVID-19, flu A, flu B, and respiratory syncytial virus infections from a single patient sample. The Plus version of the test provides three gene targets for SARS-CoV-2 detection—N2, E, and RdRP. The test is designed for use on any GeneXpert system and results are delivered in about 36 minutes.

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