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Online Article IN THIS ISSUE Resistance to Bebtelovimab

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COVID-19 Update

Resistance to Bebtelovimab

 $\mbox{Revised 12/16/22:}$ The EUA for bebtelovimab has been withdrawn. Click here for more information.

The FDA has warned that the investigational anti-SARS-CoV-2 monoclonal antibody bebtelovimab is not expected to retain activity against the Omicron variants BQ.1 and BQ.1.1.¹ Bebtelovimab (LY-CoV1404 – Lilly) is available under an FDA Emergency Use Authorization (EUA) for IV treatment of mild to moderate COVID-19 in high-risk patients \geq 12 years old who weigh \geq 40 kg for whom alternative treatment options are unavailable or inappropriate.^{2,3} The drug remains authorized for use in all regions of the US.¹

The relative prevalence of SARS-CoV-2 variants BQ.1 and BQ.1.1 has increased in recent weeks. In the week ending November 12, 2022, they were estimated to have caused ~44% of COVID-19 cases in the US, up from ~9% of cases 4 weeks earlier.⁴

The NIH recommends that high-risk nonhospitalized adults with COVID-19 be treated with either oral ritonavir-boosted nirmatrelvir (*Paxlovid*) or IV remdesivir (*Veklury*); ritonavir-boosted nirmatrelvir is preferred.⁵ Both of these therapies decreased the risk of hospitalization or death significantly more than placebo in large, randomized, double-blind trials.^{6,7} If these drugs are inappropriate or unavailable, use of molnupiravir (*Lagevrio*; available under an EUA) or bebtelovimab (only if the majority of circulating SARS-CoV-2 strains in the region are susceptible to bebtelovimab) is recommended.^{5,8} Ritonavirboosted nirmatrelvir, remdesivir, and molnupiravir are expected to retain activity against SARS-CoV-2 variants BQ.1 and BQ.1.1.¹

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