Figure 2. FDA EUA criteria for the use of casirivimab/imdevimab for post-exposure prophylaxis of COVID-19 [1]

This EUA is for the use of the unapproved products casirivimab and imdevimab adult and pediatric individuals (12 years of age and older weighing at least 40 kg) for post-exposure prophylaxis of COVID-19 in individuals who are at high risk for progression to severe COVID-19, including hospitalization or death, and are:

- Not fully vaccinated **OR** who are not expected to mount an adequate immune response to complete SARS-CoV-2 vaccination (e.g., individuals with immunocompromising conditions including those taking immunosuppressive medications) **AND**
  - Have been exposed to an individual infected with SARS-CoV-2 consistent with close contact criteria per CDC criteria **OR**
  - Who are at high risk of exposure to an individual infected with SARS-CoV-2 because of occurrence of SARS-CoV-2 infection in other individuals in the same institutional setting (e.g., nursing homes, prisons).

Reference

1. U.S. Food and Drug Administration. Fact Sheet for Health Care Providers: Emergency Use Authorization (EUA) of Regen-CoV™ (casirivimab with imdevimab). Available at: https://www.fda.gov/media/145611/download. Accessed 9 April 2021.