COVID-19 Therapeutics Authorized by FDAAt a Glance

| | | TESTING REQUIREMENTS | PRESCRIBING WINDOW | ELIGIBLE POPULATIONS |
|---------------------|-----------------------------------------------------------|---------------------------------------------|------------------------------------------|----------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|
| Monoclonal Antibody | REGEN-COV* (casirivimab plus imdevimab) | Positive direct SARS-CoV-2 viral test | Within 10 days after symptom onset | Adult and adolescent individuals (12 years of age and older who weigh at least 88 pounds [40 kg]) who are at high risk for severe COVID-19, including hospitalization or death. Please refer to FDA fact sheet for full list of Risks and Limitations: Regen-COV FDA |
| | BAMLANIVIMAB PLUS ETESEVIMAB* | Positive direct SARS-CoV-2 viral test | Within 10 days after symptom onset | Adult and pediatric individuals, including neonates, at high risk for progressing to severe COVID-19, including hospitalization or death. Please refer to FDA fact sheet for full list of Risks and Limitations: Bamlanivimab plus Etesevimab FDA |
| | SOTROVIMAB | Positive direct SARS-CoV-2 viral test | Within 10 days after symptom onset | Adult and pediatric patients (at least 12 years of age and older weighing at least 88 pounds [40 kg]) at high risk for progressing to severe COVID-19, including hospitalization or death. Please refer to FDA fact sheet for full list of Risks and Limitations: Sotrovimab FDA |
| | EVUSHELD (tixagevimab co-packaged with cilgavimab) | Positive direct SARS-CoV-2 viral test | Pre-exposure | Adult and adolescent individuals (12 years of age and older who weigh at least 88 pounds [40 kg]) for pre-exposure prophylaxis for prevention of COVID-19. Please refer to FDA fact sheet for full list of Risks and Limitations: Evusheld FDA |
| Oral Antiviral | PAXLOVID | Positive direct SARS-CoV-2 viral test | Within 5 days of symptom onset | Adults and pediatric patients (12 years of age and older weighing at least 88 pounds [40 kg]) at high risk for progressing to severe COVID-19, including hospitalization or death. Please refer to FDA fact sheet for full list of Risks and Limitations: Paxlovid FDA |
| | MOLNUPIRAVIR | Positive direct SARS-CoV-2 viral test | Within 5 days of symptom onset | Adults at high risk for progressing to severe COVID-19, including hospitalization or death, and for whom alternative COVID-19 treatment options authorized by FDA are not accessible or clinically appropriate. Please refer to FDA fact sheet for full list of Risks and Limitations: Molnupiravir FDA |

*Not expected to be effective against Omicron variant.

