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COVID-19 Therapeutics Information Brief

Changes to the document from the previous version are highlighted in yellow.

IMPORTANT/NEW COVID-19 Therapeutics Information

- Public Health Emergency Update
- Treatments Approved For Use
- Therapeutics Ordering Schedule
- Expiration Dating Extension Reminder
- COVID-19 Therapeutics Information Resources

Public Health Emergency Update

On January 30, 2023, the Biden Administration announced its intent to end the national emergency and public health emergency declarations on May 11, 2023, related to the COVID-19 pandemic. Importantly, the ending of the public health emergency declared by HHS under the Public Health Service Act will not impact FDA's ability to authorize devices (including tests), treatments or vaccines for emergency use. Existing emergency use authorizations (EUAs) for products will remain in effect and the agency may continue to issue new EUAs going forward when criteria for issuance are met. This means the process for obtaining, administering, and reporting COVID-19 therapeutics will remain unchanged. CMS reimbursement is also planned to continue.

Additional details and FAQs may be found here and here.

Detailed information and resources regarding Medicare and Medicaid Services may be found here.

Treatments Approved For Use

Details about treatment options that are expected to retain activity against COVID-19 can be found here and below:

Paxlovid is authorized for the treatment of mild-to-moderate COVID-19 in adults and pediatric patients (12 years of age and older weighing at least 40 kg) with positive results of direct SARS-CoV-2 viral testing, and who are at high risk for progression to severe COVID-19, including hospitalization or death.

Lagevrio is authorized for the treatment of mild-to-moderate COVID-19 in adults with positive results of direct SARS-CoV-2 viral testing who are at high risk for progression to severe COVID-19, including

hospitalization or death, and for whom alternative COVID-19 treatment options approved or authorized by FDA are not accessible or clinically appropriate.

Veklury is approved for the treatment of adults and pediatric patients (28 days of age and older and weighing at least 3 kg) with positive results of direct SARS-CoV-2 viral testing, who are not hospitalized and have mild-to-moderate COVID-19 and are at high risk for progression to severe COVID-19, including hospitalization or death.

COVID-19 convalescent plasma with high titers of anti-SARS-CoV-2 antibodies is authorized for the treatment of COVID-19 in patients with immunosuppressive disease or receiving immunosuppressive treatment, in inpatient or outpatient settings.

Therapeutics Ordering Schedule

Effective January 2023 Therapeutic Ordering transitioned to a monthly cadence. The planned schedule for upcoming orders are below:

Survey Sent	Survey Closed	Order Placed
2/13/2023	2/14/2023	2/15/2023
3/13/2023	3/14/2023	3/15/2023

- Iowa HHS encourages entities who do receive allocations of therapeutic products to notify and work with prescribers and LPHAs on the availability of therapeutic products in the community.
- The Department of Health and Human Services has released a COVID-19 Therapeutics locator.

Expiration Dating Extension Reminder

Recent shelf-life extensions have been issued for several products. Before wasting any product, be sure to check for expiration date extensions by using one or more of the following resources:

- FDA COVID-19 Therapeutics Extensions
- Call: 515-281-7317
- Email: C19Therapeutics@idph.iowa.gov

COVID-19 Therapeutics Information Resources

- COVID-19 Therapeutics Call Center: 515-281-7317.
- **COVID-19 Therapeutics Email**: Therapeutic questions from healthcare providers can be emailed to: c19Therapeutics@idph.iowa.gov
- <u>COVID-19 Therapeutics Table</u>: IDPH has developed a table of therapeutic products available for the treatment or prevention of COVID-19.
- Outpatient Therapeutics Decision Aid
- <u>Side-by-Side Overview Outpatient Therapeutics</u>
- NIH COVID-19 Treatment Guidelines