



FOR IMMEDIATE RELEASE
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TENNESSEE RECEIVES ORAL ANTIVIRAL COVID-19 TREATMENT SUPPLY
Molnupiravir and Paxlovid™ Available at Select Retail Pharmacies Across The State

NASHVILLE, Tenn. - Today, the Tennessee Department of Health (TDH) announced the state has received shipments of the Merck and Pfizer oral antiviral treatments for COVID-19.

The Food and Drug Administration (FDA) issued an [emergency use authorization](#) for molnupiravir by Merck and an [emergency use authorization](#) for Paxlovid™ by Pfizer as oral antiviral treatments of COVID-19. Early studies indicate these treatment options may reduce severe outcomes from COVID-19 including hospitalization or death. These treatments are recommended for individuals who are at high risk for progression to severe COVID-19 or have [underlying medical conditions](#).

	Molnupiravir	Paxlovid™
Approved age for use	18 years and older	12 years and older
When to start treatment	As soon as possible after testing positive, within 5 days of symptoms	As soon as possible after testing positive, within 3 days of symptoms
Typical regimen	40 pills over a 5-day period	30 pills over a 5-day period
Use in pregnancy or breastfeeding	No	Yes

Consult with your physician about your risk factors when taking these treatments. Both drugs require a prescription.

The Tennessee Department of Health coordinated a distribution plan of molnupiravir and Paxlovid™ with Walmart pharmacies across the state. This treatment is free, and Tennesseans can visit www.walmart.com/covidmedication to find a participating Walmart pharmacy near them. Initial supply in the state is limited as the first allocation from the federal government was 5,000 courses of molnupiravir and 1,000 courses of Paxlovid™. TDH anticipates additional allocations in the coming weeks as production increases.

While antivirals may help treat COVID-19, vaccination is the best approach to prevent infection. Tennesseans age 5 and above are encouraged to receive the COVID-19 vaccine. Individuals ages 16 and above who received an mRNA vaccine may also be [eligible for a booster shot](#) at six months or more after they complete the initial series. For adults ages 18 and older who received single-shot Johnson & Johnson vaccine, a booster dose is recommended at two or more months after the initial vaccine. More information on vaccine locations, including available vaccine products, is available at vaccines.gov.



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