

Field Name	Field Description
Prior Authorization Group Description	Emergency Use Authorization (EUA) Drugs/Products for COVID-19
Drugs	Any drug/product approved by EUA for COVID-19
Covered Uses	Medically accepted indications are defined using the following sources: the Food and Drug Administration (FDA), Micromedex, American Hospital Formulary Service (AHFS), United States Pharmacopeia Drug Information for the Healthcare Professional (USP DI), the Emergency Use Authorization for the drug/product in question, and the Drug Package Insert (PPI).
Exclusion Criteria	See “Other Criteria”
Required Medical Information	See “Other Criteria”
Age Restrictions	As outlined within current FDA Emergency Use Authorization (EUA) guidelines
Prescriber Restrictions	N/A
Coverage Duration	As outlined within current FDA Emergency Use Authorization (EUA) guidelines
Other Criteria	<p>Emergency Use Authorization for COVID-19 related drugs/products (all must apply):</p> <ul style="list-style-type: none"> <li>• The requested drug/product has a currently active Emergency Use Authorization as issued by the U.S. Food and Drug Administration.</li> <li>• Use of the requested drug/product is consistent with the current terms and conditions of the emergency use authorization (such as appropriate age/weight, <u>formulation</u>, disease severity, concurrent use with other medications or medical interventions, etc.).</li> <li>• Attestation that the provider is not requesting reimbursement for ingredient cost of drug when drug is provided by U.S. government at no charge</li> </ul>
Revision/Review Date 1/2023	<b>Medical Director/clinical reviewer must override criteria when, in his/her professional judgement, the requested item is medically necessary.</b>