SPECIALTY GUIDELINE MANAGEMENT

VEKLURY (remdesivir)

POLICY

I. INDICATIONS

The indications below including FDA-approved indications and compendial uses are considered a covered benefit provided that all the approval criteria are met and the member has no exclusions to the prescribed therapy.

FDA-Approved Indication¹

Treatment of adults and pediatric patients (12 years of age and older and weighing at least 40 kg) with coronavirus disease 2019 (COVID-19) requiring hospitalization. Veklury should only be administered in a hospital or in a healthcare setting capable of providing acute care comparable to inpatient hospital care.

All other indications are considered experimental/investigational and not medically necessary.

II. DOCUMENTATION

Submission of the following information is necessary to initiate the prior authorization review: medical records documenting positive COVID-19 infection, estimated glomerular filtration rate (eGFR), hepatic laboratory testing, and prothrombin time.

III. CRITERIA FOR INITIAL APPROVAL

Treatment of COVID-19

Authorization of 30 days may be granted for treatment of COVID-19 when all of the following criteria are met:

- A. Member has a confirmed active infection with COVID-19
- B. Member is 12 years of age or older and weighs at least 40 kg
- C. Member has eGFR ≥ 30 mL/min
- D. Member is sufficiently ill to require hospitalization
- E. Renal function, hepatic function, and prothrombin time have been taken prior to starting Veklury and will be monitored while receiving therapy as clinically appropriate
- F. Member will receive a loading dose of 200mg on Day 1 followed by maintenance dose of 100mg per day starting Day 2
- G. Member will not receive a total duration of therapy of greater than 10 days for an individual infection
- H. Veklury will not be administered in combination with chloroquine or hydroxychloroquine
- Veklury will be administered in a hospital or healthcare setting capable of providing acute care comparable to inpatient hospital care

IV. REFERENCES

1. Veklury [package insert]. Foster City, CA: Gilead Sciences, Inc.; October 2020.

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