

## Pharmacy Prior Authorization Form

Fax completed form to: 877.974.4411 toll free, or 616.942.8206

This form applies to: ☐ **Commercial (Traditional)** ☐ **Commercial (Individual/Optimized)**

☒ **Medicaid**

This request is: ☐ **Urgent** (life threatening) ☐ **Non-Urgent** (standard review)

Urgent means the standard review time may seriously jeopardize the life or health of the patient or the patient's ability to regain maximum function.

**Vemlidy<sup>®</sup>** (tenofovir alafenamide)

### Member

Last Name: \_\_\_\_\_

First Name: \_\_\_\_\_

ID #: \_\_\_\_\_

DOB: \_\_\_\_\_ Gender: \_\_\_\_\_

Primary Care Physician: \_\_\_\_\_

Requesting Provider: \_\_\_\_\_

Prov. Phone: \_\_\_\_\_ Prov. Fax: \_\_\_\_\_

Provider Address: \_\_\_\_\_

Provider NPI: \_\_\_\_\_

Contact Name: \_\_\_\_\_

Provider Signature: \_\_\_\_\_

Date: \_\_\_\_\_

### Product Information

☐ New request ☐ Continuation request

Drug product: ☐ Vemlidy 25 mg tablet

Start date (or date of next dose): \_\_\_\_\_

Date of last dose (if applicable): \_\_\_\_\_

Dosing frequency: \_\_\_\_\_

### Precertification Requirements

**Before this drug is covered, the patient must meet all of the following requirements:**

1. Must have a diagnosis of Chronic Hepatitis B infection with compensated liver disease
2. Must be tested for and is HIV negative (if positive must provide further justification for use due to HIV treatment resistance risk)
3. Failure of Entecavir, at up to maximally indicated doses, unless contraindicated or clinically significant adverse effects are experienced
4. Use not recommended in those with CrCl < 15 mL/minute or if Child-Pugh class B or C

**For continuation, patient must have met the following requirements:**

1. HBV DNA testing every three months until undetectable for at least two consecutive visits then decreased to every six months.
2. Aminotransferases testing every three to six months.
3. Creatinine and phosphate testing every 6 months.

**Note:** Authorization for indications, dosing, or a route of administration not approved by the Food and Drug Administration (FDA) or recognized in CMS-accepted compendia (e.g. DrugDex, AHFS, U.S. Pharmacopeia, and also Clinical Pharmacology for oncology indications only) require supporting evidence for coverage. Please provide two published peer-reviewed literature articles supporting the appropriateness of the drug, the dosing of the drug, or the route of administration to be used for the identified indication.

### Additional information

**Note:** Initial approval length 6 months, with requests for continuation required every 6 months thereafter.

**New request**

**Priority Health Precertification Documentation**

**A. What condition is this drug being requested for?**

☐ Chronic Hepatitis B infection with compensated liver disease

☐ Other – the patient's condition is: \_\_\_\_\_

Rationale for use: \_\_\_\_\_

**B. Has the patient been tested for and is HIV negative?**

☐ HIV negative

☐ HIV positive, rationale for use: \_\_\_\_\_

☐ Has not been tested, rationale: \_\_\_\_\_

**C. Has the patient had a failure on Entecavir, up to maximally indicated doses (please provide dose and dates of use)?**

☐ Yes, Dose: \_\_\_\_\_ Dates of use: \_\_\_\_\_

☐ No, rationale: \_\_\_\_\_

**D. What is the patient's creatinine clearance? \_\_\_\_\_**

**E. What is the patients Child-Pugh class? \_\_\_\_\_**

**Request for Continuation**

**Priority Health Precertification Documentation**

**A. What condition is this drug being requested for?**

☐ Chronic Hepatitis B infection with compensated liver disease

☐ Other – the patient's condition is: \_\_\_\_\_

Rationale for use: \_\_\_\_\_

**B. Does patient meet monitoring criteria listed under criteria? Please describe.**

☐ Yes, dates of labs (or attach to request): \_\_\_\_\_

☐ No, rationale: \_\_\_\_\_