

SENTARA HEALTH PLANS

PHARMACY PRIOR AUTHORIZATION/STEP-EDIT REQUEST*

Directions: The prescribing physician must sign and clearly print name (preprinted stamps not valid) on this request. All other information may be filled in by office staff; **(Pharmacy) 1-800-750-9692.** No additional phone calls will be necessary if all information (including phone and fax #s) on this form is correct. If the information provided is not complete, correct, or legible, the authorization process can be delayed.

Drug Requested: Revcovi™ (elapegademase-lvlr) **(Pharmacy)**

MEMBER & PRESCRIBER INFORMATION: Authorization may be delayed if incomplete.

Member Name: _____

Member Sentara #: _____ Date of Birth: _____

Prescriber Name: _____

Prescriber Signature: _____ Date: _____

Office Contact Name: _____

Phone Number: _____ Fax Number: _____

DEA OR NPI #: _____

DRUG INFORMATION: Complete information below or authorization will be delayed if incomplete.

Drug Form/Strength: _____

Dosing Schedule: _____ Length of Therapy: _____

Diagnosis: _____ ICD Code, if applicable: _____

Dosing Limit: (see below)

A. Quantity Limit (max daily dose): Pharmacy Benefit - 2.4 mg/1.5 mL single dose vial: 20 vials per 7 days

B. Max Units (per dose and over time): Medical Benefit (J3590) - 23 mg twice weekly

CLINICAL CRITERIA: Check below **ALL** that apply. **ALL** criteria **must** be met for approval. **ALL** documentation including labs or chart notes **must** be submitted or request will be denied.

Initial Authorization Approval – 12 months

• **Coverage is provided in the following conditions:**

- Must not be used in combination with pegademase-bovine

AND

- Member has severe combined immunodeficiency disease (SCID) with a definitive diagnosis of adenosine deaminase deficiency as determined by one of the following:
- Deficient ADA catalytic activity (<1% of normal) in hemolysates (in untransfused individuals) or in extracts of other cells (e.g., blood mononuclear cells, fibroblasts)

(Continued on next page)

OR

- Detection of pathogenic mutations in the ADA gene by molecular genetic testing;

AND

- Member has a marked elevation of the metabolite dATP or total dAdo nucleotides (the sum of dAMP, dADP and dATP) in erythrocytes;

AND

- Member is not a candidate for or has failed hematopoietic cell transplantation (HCT);

AND

- Member does not have severe thrombocytopenia (<50,000/microL);

AND

- Baseline lab values for plasma ADA activity, red blood cell deoxyadenosine triphosphate (dATP), trough deoxyadenosine nucleotide (dAXP) levels and/or total lymphocyte counts must be submitted with this request
- Is the member transitioning from Adagen® to Revcovi™? Yes No
- Member's current height and weight must be noted and dosing will be based on ideal body weight
Height: _____ Weight: _____

Reauthorization of Therapy: Yearly reauthorization is required for continuation of therapy. (12 month approval)

CLINICAL CRITERIA: Check below **ALL** that apply. **ALL** criteria **must** be met for approval. **ALL** documentation including labs or chart notes **must** be submitted or request will be denied.

• Authorizations can be renewed based on the following criteria:

- Member continues to meet the criteria identified in Initial Approval Section.

AND

- Absence of unacceptable toxicity from the drug. Examples of unacceptable toxicity include severe injection site reactions (e.g., bleeding), severe thrombocytopenia, etc.;

AND

- Adequate documentation of disease stability and/or improvement as indicated by one or more of the following must be submitted:
 - Increase in plasma ADA activity (target trough level ≥ 15 mmol/hr/L)
 - Red blood cell dATP level decreased (target ≤ 0.005 to 0.015 mmol/L)
 - Improvement in immune function with diminished frequency/complications of infection as evidenced in improvement in the ability to produce antibodies

(Continued on next page)

- Improvement in red blood cell dAXP levels (target trough level ≤ 0.02 mmol/L)
- Member's current height and weight must be noted and dosing will be based on ideal body weight

Height: _____ Weight: _____

Medication being provided by Specialty Pharmacy - PropriumRx

****Use of samples to initiate therapy *does not* meet step edit/ preauthorization criteria.****

Previous therapies will be verified through pharmacy paid claims or submitted chart notes.