

Physician Order/Prescription & Statement of Medical Necessity



Please fax completed form to Chiesi Total CareSM staff at 866-272-7079

PATIENT INFORMATION					
Patient Name (Last, First)					
Social Security #	Sex: □ Male	☐ Female Date of	Birth/	/(mm/dd/yyyy)	
Address					
Primary Phone (Required)	Cell Phone		Language: □ Enç	glish Other	
Parent/Guardian (If applicable) Relationship to Patient					
CLINICAL INFORMATION					
Diagnosis: ☐ Adenosine deaminase severe combined immune deficiency (ADA-SCID) ICD-10 code D81.31 (primary)					
□ Secondary ICD-10	• `	, 			
Treatment information					
☐ Initial Rx for ADA-SCID ☐ Continuation on ERT	☐ Restart after Gene T	herapy □ Restart at	fter HSCT		
☐ New patient/returning to therapy ☐ Other		Allergies: None Specify			
Height inches or cm Weig	ht lb or	kg Known Drug Allergies:			
INSURANCE INFORMATION					
Primary Prescription Insurance	Primary Medical Insurance		Secondary Medica	al	
Policy Holder	Policy Holder				
Policy ID #	Policy ID #				
Group #	Group #				
Phone	Phone				
☐ Eligible for Medicare ☐ Eligible for Medicaid ☐ No Insurance					
Please attach copies of patient insurance and prescription cards—front and back.					
REVCOVI (elapegademase-lvir) PRESCRIPTION/ORDER					
Revcovi (elapegademase-lvlr) 2.4 mg/1.5 mL single use vial NDC 10122-502-01					
Instructions: Inject mg intramuscularly (IM) times per week.					
Provide medical supplies, including syringes and needles, to safely administer prescribed medication.					
☐ Skilled nursing visits times for medication administration teaching					
PRESCRIBER/OFFICE INFORMATION					
Prescriber's Name (Print)		Practice/Group Name		· · · · · · · · · · · · · · · · · · ·	
Address			S	uite	
City		State	Zip		
Office Contact Person					
Office Phone					
License # NPI #				····	
By signing below, I certify that I am the prescribing provider mentioned above, that I am part of the Chiesi Total Care Program, that the therapy described above is medically necessary, and that all the medical necessity information is true, accurate, and complete. The patient's records contain supporting documentation that substantiates the utilization and medical necessity of the products marked above. I provide permission to use my personal information and the personal information of the patient provided above to facilitate this request and complete any regulatory or legal requirements associated with this request. I understand that the personal information provided herein may be shared with Chiesi, successors, and their agents and service providers as needed to support this request. I also attest that I have obtained the patient's authorization to release the above information and such other personal information as may be necessary for the Chiesi Total Care Program and/or their agents and service providers. If the patient is 18 years old or younger, I attest that I have obtained permission from the patient's legal guardian. If my patient is eligible for free product, I understand that receiving free product is not contingent on any purchase obligations. I also understand that no free product may be submitted for reimbursement to any payer, including Medicare and Medicaid; nor may I bill any payer for administration of such product. I understand that any falsification, omission, or concealment of material fact may result in criminal liability.					
Prescriber's Signature			D	ato	
Prescriber's Signature Substitution Permitted		Dispense as Written			

INDICATION

Revcovi® (elapegademase-lvlr) is indicated for the treatment of adenosine deaminase severe combined immune deficiency (ADA-SCID) in pediatric and adult patients.

IMPORTANT SAFETY INFORMATION

WARNINGS AND PRECAUTIONS

- Injection site bleeding in patients with thrombocytopenia: Increased risk of local bleeding in patients with thrombocytopenia; should not be used if thrombocytopenia is severe.
- Delay in improvement of immune function: Protect immune deficient patients from infections until improvement in immune function.

ADVERSE REACTIONS

The most commonly reported adverse reactions were cough (50%) and vomiting (33%).

In addition, the following post-marketing reports for the same class of enzyme replacement therapy used in the treatment of ADA-SCID may also be seen with Revcovi treatment:

- Hematologic events: hemolytic anemia, autoimmune hemolytic anemia, thrombocythemia, thrombocytopenia and autoimmune thrombocytopenia
- · Dermatological events: injection site erythema, urticaria
- Lymphomas

IMPORTANT MONITORING INFORMATION

Treatment with Revcovi should be monitored by measuring trough plasma ADA activity and trough dAXP levels for maintenance of therapeutic targets. If a persistent decline in plasma ADA activity occurs, immune function and clinical status should be monitored closely, and precautions should be taken to minimize the risk of infection.

Please see <u>Full Prescribing Information</u> for more information.

