

Ceredase® (alglucerase) & Cerezyme® (imiglucerase) PreDetermination of Medical Benefits

Complete form in its entirety and fax to UM Call Center at (404) 848-2448

Click on grey boxes to type

Request Date: _____ / _____ / _____

- Initial Authorization Request Re-Authorization Request; List Prior Auth Ref #: _____
- Medication(s) is to be dispensed, delivered, and managed by PrecisionRx Specialty Solutions (800.870.6419) Yes No

1. PATIENT INFORMATION			
Patient Last Name	Patient First Name	Patient ID Number	Patient DOB / /
Address	City	State / Zip Code /	Contact Phone Number () -
Date of Diagnosis / /	Primary Diagnosis	ICD-9 Code(s)	Patient's Current Weight

2. PHYSICIAN INFORMATION			
Physician Last Name	Physician First Name	Physician DEA or NPI Number	Physician Tax ID
Address	City	State	Zip Code
Office Phone Number () -	Office Fax Number () -	Office Contact Name	Physician Specialty

3. MEDICATION INFORMATION – This section serves as the active prescription – signature required.			
Drug Name <input type="checkbox"/> Ceredase <input type="checkbox"/> Cerezyme	HCPCS or CPT Code(s) <input type="checkbox"/> J0205 <input type="checkbox"/> J1785	Strength / Dose	
Direction for Use (SIG)			
Date patient is scheduled to be treated (need by date) / /	Service From Date / /	Service Thru Date / /	Number of Refills
Place of Service <input type="checkbox"/> MD Office <input type="checkbox"/> Pt's Home <input type="checkbox"/> Other: (please specify)			
Prescriber Signature			Date / /

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4. APPROVAL CRITERIA: CHECK ALL BOXES THAT APPLY

NOTE: **To avoid delays**, please complete this form in its entirety. Incomplete forms that are missing pertinent information will be pended.
 If indicated, please provide **ALL** supporting lab results, progress notes, etc.

(1) Gaucher Disease Type I (adult)

- Yes No Is the pt confirmed to have type 1 Gaucher disease by one of the following methods?
 Glucocerebrosidase activity in the WBCs or skin fibroblasts is \leq 30% of normal activity, or
 Genotype testing indicates mutation of 2 alleles of the glucocerebrosidase genome.
- Yes No Does the pt have significant clinical manifestations of skeletal disease with one or more of the following?
 Avascular necrosis Osteopenia
 Erlenmeyer flask deformity Osteosclerosis
 Lytic disease Pathological fracture
 Marrow infiltration Radiological evidence of joint deterioration
- Yes No Does pt present with at least 2 of the following?
 Clinically significant hepatomegaly Hgb \leq 11.5 g/dL for females, or
 Clinically significant splenomegaly Hgb \leq 12.5 g/dL for males, or
 Platelet count \leq 120,000 mm³ Hgb is 1.0 g/dL below lower limit of normal for age and sex

(2) Gaucher Disease Type I (children)

- Yes No Is the pt confirmed to have type 1 Gaucher disease by one of the following methods?
 Glucocerebrosidase activity in the WBCs or skin fibroblasts is \leq 30% of normal activity, or
 Genotype testing indicates mutation of 2 alleles of the glucocerebrosidase genome.
- Yes No Does the pt have significant clinical manifestations of Gaucher Disease with one or more of the following?
 Abdominal or bone pain Documented growth failure not associated with other conditions
 Cachexia Evidence of skeletal involvement, not limited to Erlenmeyer flask deformity
 Exertional limitations Anemia with Hgb < 2.0 g/L below limit of normal for age and sex
 Fatigue Platelet count < 60,000 mm³ and/or documented abnormal bleeding episodes

(3) Gaucher Disease Type 3 (adult)

- Yes No Is pt confirmed to have type 3 Gaucher disease by genotype testing indicating 2 or more alleles for neuropathic Gaucher disease?
- Yes No Does pt have significant clinical manifestations of skeletal disease with one or more of the following?
 Avascular necrosis Osteopenia
 Erlenmeyer flask deformity Osteosclerosis
 Lytic disease Pathological fracture
 Marrow infiltration Radiological evidence of joint deterioration
- Yes No Does the pt present with at least **two (2)** of the following?
 Clinically significant hepatomegaly Hgb is 1.0 g/dL below lower limit of normal for age and sex
 Clinically significant splenomegaly Platelet count \leq 120,000 mm³
- Yes No Are the neurological findings consistent with the presence of type 3 Gaucher disease, confirmed by the following tests?
 Neurological exam by a neurologist Measurement of peripheral hearing
 Eye movement exam by an ophthalmologist Brain imaging
 Neuro-ophthalmological investigation w/direct ophthalmoscopy Electroencephalography (EEG)
 Intelligence quotient testing (when appropriate & reasonable) Diagnostic brain stem evoked responses

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APPROVAL CRITERIA: CHECK ALL BOXES THAT APPLY (continue)

(4) Gaucher Disease Type 3 (children)

- Yes No Is the pt confirmed to have type 3 Gaucher disease by genotype testing indicating 2 or more alleles for neuropathic Gaucher disease?
- Yes No Does the pt have significant clinical disease manifestations with one or more of the following?
- | | |
|---|---|
| <input type="checkbox"/> Abdominal or bone pain | <input type="checkbox"/> Documented growth failure not associated with other conditions |
| <input type="checkbox"/> Cachexia | <input type="checkbox"/> Evidence of skeletal involvement, not limited to Erlenmeyer flask deformity |
| <input type="checkbox"/> Exertional limitations | <input type="checkbox"/> Anemia with Hgb < 2.0 g/L below limit of normal for age and sex |
| <input type="checkbox"/> Fatigue | <input type="checkbox"/> Platelet count < 60,000 mm ³ and/or documented abnormal bleeding episodes |
- Yes No Are the neurological findings consistent with the presence of type 3 Gaucher disease, confirmed by the following tests?
- | | |
|--|---|
| <input type="checkbox"/> Neurological exam by a neurologist | <input type="checkbox"/> Measurement of peripheral hearing |
| <input type="checkbox"/> Eye movement exam by an ophthalmologist | <input type="checkbox"/> Brain imaging |
| <input type="checkbox"/> Neuro-ophthalmological investigation w/direct ophthalmoscopy | <input type="checkbox"/> Electroencephalography (EEG) |
| <input type="checkbox"/> Intelligence quotient testing (when appropriate & reasonable) | <input type="checkbox"/> Diagnostic brain stem evoked responses |

(5) Other Use(s) (This will not be reviewed unless all supporting evidence/documentation, labs, etc., are attached.)

5. PHYSICIAN SIGNATURE

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Prescriber Signature	Date
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Prior Authorization is not the practice of medicine or the substitute for the independent medical judgment of a treating physician. Only a treating physician can determine what medications are appropriate for a patient. Please refer to the applicable plan for the detailed information regarding benefits, conditions, limitations, and exclusions.

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Medical Policy Reference can be found at: www.bcbsga.com

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