



Zokinvy

Prior Authorization Request

CVS Caremark administers the prescription benefit plan for the patient identified. This patient's benefit plan requires prior authorization for certain medications in order for the drug to be covered. To make an appropriate determination, providing the most accurate diagnosis for the use of the prescribed medication is necessary. **Please respond below and fax this form to CVS Caremark toll-free at 1-866-249-6155.** If you have questions regarding the prior authorization, please contact CVS Caremark at **1-866-814-5506**. For inquiries or questions related to the patient's eligibility, drug copay or medication delivery; please contact the Specialty Customer Care Team: CaremarkConnect® 1-800-237-2767.

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Patient's Name: _____ **Date:** _____
Patient's ID: _____ **Patient's Date of Birth:** _____
Physician's Name: _____ **NPI#:** _____
Specialty: _____ **Physician Office Fax:** _____
Physician Office Telephone: _____
Request Initiated For: _____

- What is the diagnosis?
 - Hutchinson-Gilford Progeria Syndrome
 - Processing Deficient Progeroid Laminopathy with Progerin-Like Protein Accumulation
 - Processing Deficient Progeroid Laminopathy without Progerin-Like Protein Accumulation
 - Other _____
- What is the ICD-10 code? _____
- What is the patient's body surface area (BSA)? _____ meters squared
- Is the patient currently receiving treatment with the requested medication?
 - Yes No *If No, skip to diagnosis section*
- Has the patient experienced a benefit from therapy? Yes No *No further questions*

Complete the following section based on the patient's diagnosis, if applicable.

Section A: Hutchinson-Gilford Progeria Syndrome

- Has the diagnosis of Hutchinson-Gilford Progeria Syndrome been confirmed with genetic testing indicating the patient has an *LMNA* mutation? **ACTION REQUIRED: If Yes, attach supporting chart note(s).**
 - Yes
 - No
 - Unknown

Section B: Processing Deficient Progeroid Laminopathy with Progerin-Like Protein Accumulation

- Has the diagnosis of Processing Deficient Progeroid Laminopathy with Progerin-Like Protein Accumulation been confirmed with genetic testing indicating the patient has a heterozygous *LMNA* mutation? **ACTION REQUIRED: If Yes, attach supporting chart note(s).**
 - Yes
 - No
 - Unknown

Section C: Processing Deficient Progeroid Laminopathy without Progerin-Like Protein Accumulation

Send completed form to: Case Review Unit CVS Caremark Prior Authorization Fax: 1-866-249-6155

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8. Has the diagnosis of Processing Deficient Progeroid Laminopathy without Progerin-Like Protein Accumulation been confirmed with genetic testing indicating the patient has a homozygous or compound heterozygous *ZMPSTE24* mutation? ***ACTION REQUIRED: If Yes, attach supporting chart note(s).***
- Yes
 - No
 - Unknown

I attest that this information is accurate and true, and that documentation supporting this information is available for review if requested by CVS Caremark or the benefit plan sponsor.

X _____

Prescriber or Authorized Signature

Date (mm/dd/yy)

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