

Division: Pharmacy Policy	Subject: Prior Authorization Criteria
Original Development Date:	August 4, 2021
Original Effective Date: Revision Date:	
Revision Date.	

${\bf ZOKINVY}^{TM} \ (lona farnib)$

LENGTH OF AUTHORIZATION: Up to one year

INITIAL REVIEW CRITERIA:

- Patient must be 12 months of age and documented BSA \geq 0.39 m².
- Diagnosis of one of the following as confirmed by genetic testing:
 - O Hutchinson-Gilford Progeria Syndrome (HGPS)
 - o Processing-deficient Progeroid Laminopathy with either
 - Heterozygous *LMNA* mutation with progerin-like protein accumulation

or

- Homozygous or compound heterozygous *ZMPSTE24* mutations
- Prescribed by, or in consultation, with a specialist, document specialty type.

CONTINUATION OF THERAPY:

- Patient met initial review criteria.
- Documentation of positive clinical response.
- Dosing is appropriate as per labeling or is supported by compendia.

DOSING AND ADMINISTRATION:

- Refer to product labeling at https://www.accessdata.fda.gov/scripts/cder/daf/
- Dosage Forms: 50 mg and 75 mg capsules

