

5.90.16

Section:	Prescription Drugs	Effective Date:	April 1, 2021
Subsection:	Topical Products	Original Policy Date:	July 31, 2015
Subject:	Solaraze	Page:	1 of 3

Last Review Date: March 12, 2021

Solaraze

Description

Solaraze (diclofenac sodium)

Background

Solaraze (diclofenac sodium) gel is a prescription medicine used on the skin for actinic keratoses. Actinic keratosis (AK), also called solar keratosis, which is a chronic (long-term) condition of the skin caused by a chemical reaction to ultraviolet (UV) rays. Actinic keratosis can be linked to the development of skin cancer (1).

Regulatory Status

FDA-approved indications: Solaraze gel is indicated for the topical treatment of actinic keratoses (AK). Sun avoidance is indicated during therapy (1).

Safety and effectiveness of Solaraze in pediatric patients less than 18 years of age have not been established (1).

Related policies

Aldara, Klysryi, Zyclara

Policy

This policy statement applies to clinical review performed for pre-service (Prior Approval, Precertification, Advanced Benefit Determination, etc.) and/or post-service claims.

Solaraze may be considered **medically necessary** in patients 18 years of age or older with actinic keratosis (AK) and if the conditions indicated below are met.

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Solaraze may be considered **investigational** in patients less than 18 years of age and for all other indications.

Prior-Approval Requirements

Age 18 years of age or older

Diagnosis

Patient must have the following:

Actinic keratosis (AK)

AND ALL of the following:

1. Inadequate treatment response, intolerance, or contraindication to a topical purine analog and topical antineoplastic

Prior – Approval *Renewal* Requirements

Age 18 years of age or older

Diagnosis

Patient must have the following:

Actinic keratosis (AK)

AND ALL of the following:

1. Re-evaluation of lesion(s) for improvement

Policy Guidelines

Pre - PA Allowance

None

Prior - Approval Limits

Duration 3 months

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Prior – Approval *Renewal* Limits

Duration 3 months (One renewal only)

Rationale

Summary

Solaraze (diclofenac sodium) gel is a prescription medicine used on the skin for actinic keratoses. Actinic keratosis (AK), also called solar keratosis, which is a chronic (long-term) condition of the skin. It is caused by a chemical reaction to ultraviolet (UV) rays. AK can be linked to the development of skin cancer. Safety and effectiveness of Solaraze in pediatric patients less than 18 years of age have not been established (1).

Prior approval is required to ensure the safe, clinically appropriate and cost-effective use of Solaraze while maintaining optimal therapeutic outcomes.

References

1. Solaraze [package Insert]. Melville, NY: PharmaDerm.; May 2016.

Policy History

Date	Action
July 2015	Addition to PA
September 2015	Annual review
December 2016	Annual review and reference update Addition of age requirement to renewal section. Policy number change from 5.14.16 to 5.90.16
September 2017	Annual editorial review
September 2018	Annual review
September 2019	Annual review
September 2020	Annual review
March 2021	Annual editorial review

Keywords

This policy was approved by the FEP® Pharmacy and Medical Policy Committee on March 12, 2021 and is effective on April 1, 2021.