

XALIX- salicylic acid solution
KMM Pharmaceuticals, LLC

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, [click here](#).

XALIX

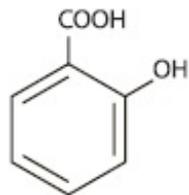
Rx Only

FOR EXTERNAL USE ONLY. NOT FOR OPHTHALMIC USE.

DESCRIPTION:

Each gram contains 280 mg of salicylic acid extended release in a film-forming virucidal solution consisting of acrylates copolymer, isopropyl alcohol, n-Butyl acetate, polyvinyl butyral, and trimethyl pentanyl diisobutyrate.

The pharmacologic activity of this product is generally attributed to the keratolytic activity of salicylic acid, which is incorporated into a polyacrylic, film-forming virucidal solution designed to cover the wart without the need for a bandage. The structural formula of salicylic acid is:



CLINICAL PHARMACOLOGY:

Although the exact mode of action for salicylic acid in the treatment of warts is unknown, its activity appears to be associated with its keratolytic action, which results in mechanical removal of epidermal cells infected with wart viruses.

INDICATIONS AND USAGE:

This product is indicated for the topical treatment and removal of common warts and plantar warts.

CONTRAINDICATIONS:

This product is contraindicated in persons with known or suspected hypersensitivity to any of the ingredients of the product. Patients with diabetes or impaired blood circulation should not use this product. This product also should not be used on moles, birthmarks, and unusual warts with hair growing from them, or warts on the face.

WARNINGS:

KEEP OUT OF REACH OF CHILDREN. This product is flammable. Keep away from fire or flame. Keep bottle tightly closed when not in use.

PRECAUTIONS:

FOR EXTERNAL USE ONLY. NOT FOR OPHTHALMIC USE. If contact with eyes or mucous membranes occurs, immediately flush with water for 15 minutes. This product should not be allowed to

contact normal skin surrounding the wart site, since localized irritation may occur. Treatment should be discontinued if excessive irritation occurs.

ADVERSE REACTIONS:

Localized irritation may occur if this product is applied to normal skin surrounding the wart; however irritation may normally be controlled by temporarily discontinuing use and by applying the medication only to the wart site when treatment is resumed. To report a serious adverse event, call 1-855-899-4237.

DOSAGE AND ADMINISTRATION:

Prior to applying this product, soak wart in warm water for five minutes. Remove any loose tissue by gently rubbing with a washcloth, emery board, or pumice stone. Dry the wart site thoroughly. Using the brush applicator supplied, apply this product twice to the entire wart surface, allowing the first application to dry before applying the second. Continue treatment once or twice a day or as directed by a physician. Be careful not to apply to surrounding skin.

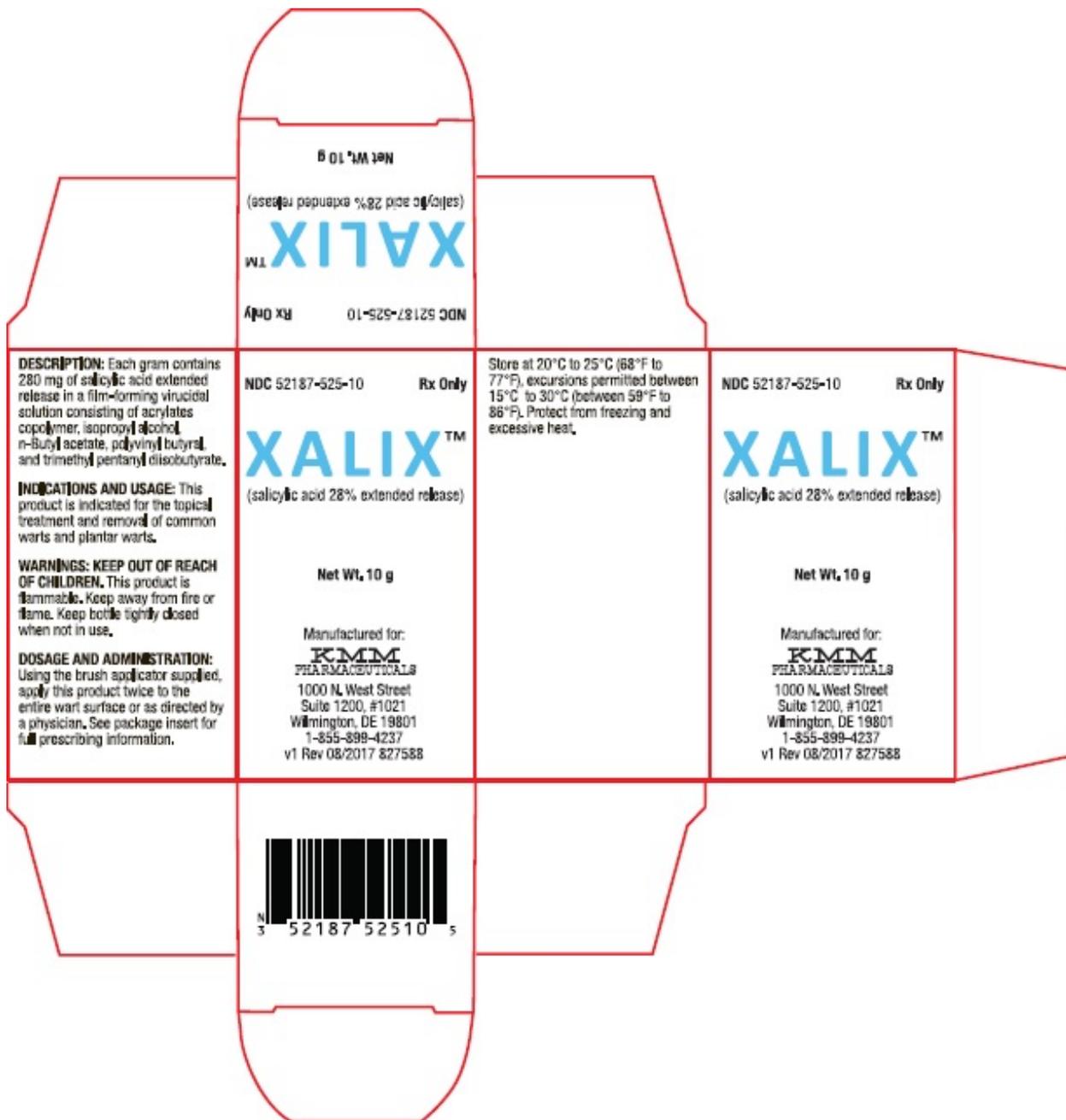
Clinically visible improvement normally occurs during the first or second week of therapy. Resolution may be expected after four to six weeks of this product's use, though some warts may take longer to remove.

HOW SUPPLIED:

10 g bottles, NDC 52187-525-10

Store at 20°C to 25°C (68°F to 77°F), excursions permitted between 15°C to 30°C (between 59°F to 86°F). Brief exposure to temperatures up to 40°C (104°F) may be tolerated provided the mean kinetic temperature does not exceed 25°C (77°F); however, such exposure should be minimized. Protect from freezing and excessive heat. Keep bottle tightly closed.

Manufactured for:
KMM Pharmaceuticals, LLC
1000 N. West Street
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Wilmington, DE 19801



XALIX

salicylic acid solution

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:52187-525
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SALICYLIC ACID (UNII: O414PZ4LPZ) (SALICYLIC ACID - UNII:O414PZ4LPZ)	SALICYLIC ACID	280 mg in 1 g

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:52187-525-10	1 in 1 CARTON	10/25/2017	
1		10 g in 1 BOTTLE, WITH APPLICATOR; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		10/25/2017	

Labeler - KMM Pharmaceuticals, LLC (078521761)

Revised: 10/2017

KMM Pharmaceuticals, LLC