

CORNEX- lactic acid, salicylic acid gel
Pella Pharmaceuticals Co. Ltd

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.

Cornex

Active Ingredients

Each one gram contains: Lactic acid 167mg and salicylic acid 167mg.

Composition

Excipients: Alcohol, Hydroxypropyl Cellulose.

Properties

Cornex is a topical gel for the removal of warts and corns.

Indications

Cornex is used to remove Warts and Corns. A wart is a viral infection of the skin surface layer. The incubation period varies from a few weeks to several months. Warts can be spread by contact of the virus with damaged skin, particularly if warm and moist. This is why one of the most common sources of infection is swimming pools. Corns are hard, thick pads of skin caused by pressure and friction. They usually occur on the feet due to poorly fitting shoes. They can also occur on hands. Cornex Gel is suitable for use by adults, children and the elderly.

Contraindications

Cornex is contraindicated in patients with known hypersensitivity to lactic acid and / or salicylic acid or to any other components of this product.

Do not use Cornex if you are diabetic or suffer from poor blood circulation to your hands or feet.

Precautions

Cornex is formulated for the controlled corrosion of keratin and thus care should be taken to apply the product to the wart only. Avoid applying it to normal surrounding skin. Do not use Cornex on the face, mucous membranes or anogenital regions. Cornex is not recommended for use in infants under 2 years of age. Children under 12 years should only use the product under supervision.

Precaution

As well all the medicines you should keep Cornex out of reach of children.

Drug Interactions

No interaction between Lactic acid and Salicylic acid and other drugs or substances are known or suspected.

Warnings

Remember that warts are infectious. To prevent the infection spreading to others always keep to your own towel. Do not walk about barefoot if warts are on the soles of your feet. Be patient, it can take 6 to 12 weeks to completely remove the wart. Do not use Cornex on moles, birthmarks or unusual skin growths or near eyes or on mucous membranes.

Avoid use on broken or inflamed skin.

Dosage and Administration

Remove any plaster and soak the wart in hot water for five minutes. Dry with your own towel. Rub the surface of the warts carefully with a pumice stone or emery board. Apply Cornex daily to the warts. Use only enough Cornex to cover the warts and avoid applying to the surrounding normal skin.

Allow Cornex to dry and cover with plaster if the wart is large or on foot. Continue using Cornex daily until the wart is completely cleared and the ridge lines of the skin have been restored.

Side Effects

Avoid use on large areas of skin; as this may result in absorption of sufficient quantities to cause adverse effects.

Storage

Store below 30 °C.

How supplied

15 g packs.

Secondary Package



Primary Package



Topical gel for the removal of warts & corns

EACH 1G CONTAINS:

- Lactic Acid (167mg)
- Salicylic Acid (167mg)

DIRECTIONS:

For method of application, read the enclosed leaflet.



Manufactured By:
Pella Pharmaceuticals Co. Ltd. JO
www.dermapharma.com.jo

Store below 30 °C.
Read enclosed leaflet before use.
Keep out of reach of children.

**DO NOT USE ON FACE OR
ANOGENITAL REGIONS
FOR TOPICAL USE ONLY**

CORNEX

lactic acid, salicylic acid gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:82160-238
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LACTIC ACID (UNII: 33X04XA5AT) (LACTIC ACID - UNII:33X04XA5AT)	LACTIC ACID	2505 mg in 15 g
SALICYLIC ACID (UNII: O414PZ4LPZ) (SALICYLIC ACID - UNII:O414PZ4LPZ)	SALICYLIC ACID	2505 mg in 15 g

Inactive Ingredients

Ingredient Name	Strength
ALCOHOL (UNII: 3K9958V90M)	
HYDROXYPROPYL CELLULOSE, UNSPECIFIED (UNII: 9XZ8H6N6OH)	

Packaging

	Marketing Start	Marketing End

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:82160-238-01	1 in 1 CARTON	03/25/2010	
1		15 g in 1 TUBE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		03/25/2010	

Labeler - Pella Pharmaceuticals Co. Ltd (562370925)

Revised: 1/2025

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