

SKIN ULCER- Imnoop skin ulcer ointment ointment
Hunan Xiong Shi Pharmaceutical Co., Ltd

Disclaimer: This homeopathic product has not been evaluated by the Food and Drug Administration for safety or efficacy. FDA is not aware of scientific evidence to support homeopathy as effective.

Active ingreddient

Calendula oils ---1%----Speed healing.

Inactive ingredients:

Bees Wax, Linoleic Acid, Propolis, Sesame oil, Glycyrrhiza glabra, Panax notoginseng root, Cullen corylifolium seed, Calamus draco willd, Geranium oils.

Ask doctor if

on deep or puncture wounds, animal bites or serious burns.

do not

get into eyes

USES

Promote rapid healing and helps prevent infection of wounds, skin ulcers and sores.

Stop use if

condition worsens, symptoms last more than 7 days or clear up and occur again within a few days.

Warning

For external use only

Other information:

Store at room temperature; may stain fabrics.

Dosage & Administration

Keep out of reach of children

if swallowed, call poison control or seek medical help.

Directions

Wash and dry affected area, apply a thin layer of LMNOOP over affected area, gently massage until fully absorbed, repeat 2-3 times daily.

Lable



Dabetic Ulcer & Sore Treatment

SKIN ULCER

Imnoop skin ulcer ointment ointment

Product Information

Product Type

HUMAN OTC DRUG

Item Code (Source)

NDC:73252-135

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CALENDULA OFFICINALIS FLOWER (UNII: P0M7O4Y7YD) (CALENDULA OFFICINALIS FLOWER - UNII:P0M7O4Y7YD)	CALENDULA OFFICINALIS FLOWER	1 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
PANAX NOTOGINSENG ROOT (UNII: GQX1C1175U)	3 g in 100 g
GLYCYRRHIZA GLABRA (UNII: 2788Z9758H)	3 g in 100 g
CULLEN CORYLIFOLIUM FRUIT (UNII: 78AD6Z52S6)	3 g in 100 g
SESAME OIL (UNII: QX10HYY4QV)	60 g in 100 g
PROPOLIS WAX (UNII: 6Y8XYV2NOF)	5 g in 100 g
YELLOW WAX (UNII: 2ZA36H0S2V)	20 g in 100 g
LINOLEIC ACID (UNII: 9KJL21T0QJ)	5 g in 100 g

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73252-135-01	50 g in 1 BOTTLE; Type 0: Not a Combination Product	10/01/2022	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		10/01/2022	

Labeler - Hunan Xiong Shi Pharmaceutical Co., Ltd (554497403)

Registrant - Hunan Xiong Shi Pharmaceutical Co., Ltd (554497403)

Establishment

Name	Address	ID/FEI	Business Operations
Hunan Xiong Shi Pharmaceutical Co., Ltd		554497403	manufacture(73252-135)

Revised: 8/2022

Hunan Xiong Shi Pharmaceutical Co., Ltd