

MYMULIKE CORN REMOVAL PADS- corn removal pads patch
Zhengzhou Miaoqi Pharmaceutical Technology Co., Ltd

Active Ingredient

Salicylic acid 40%

Purpose

Corn and callus remover

Use

for the removal of corns from foot and hand.

Warnings

for external use only

Do not use

on irritated skin or any area that is infected or reddened.

When Using

if product gets in eyes,flush with water for 15 minutes.
Ask a doctor before use if you have

- diabetes
- poor blood circulation
- under 14 years old
- pregnant and lactating women

Stop Use

discomfort persists.

Ask Doctor

discomfort persists.

Keep Out Of Reach Of Children

If swallowed, get medical help or contact a Poison Control Center right away.

Directions

- Wash affected corn and dry thoroughly.
- Please cut the medicated patch as a suitable size to cover your corn.
- Place firmly the medicated round part over the center of corn. Adhesive next to skin.
- Repeat the treatment as needed until corn is removed. (The result may vary from person.)
- May soak corn in warm water for 5 minutes to assist in the removal.

Other information

store between 20° to 25°C (68° to 77°F)

Inactive ingredients

hard paraffin, lanolin, phenol

Questions

MYMULIKE@gmail.com

PRINCIPAL DISPLAY PANEL

MYMU大小鸡眼贴-美版
115*90*18mm



尊敬的客户：此图印刷依据，请务必严谨审核尺寸、文字、盒型、图案等内容，确认无误后签字回传。因电脑和手机电子屏显色各有不同，以及印刷油墨、合拼版等因素影响，不同印刷批次会有少许色差，最终颜色以实际印刷为准。本公司会尽量调整减少色差情况，敬请谅解！（如要求非常严格请做特殊交代）

	产品类型	盒子	成品尺寸	见标注	材质/工艺/数量	
	设计类别	换批件	设计师	郑鹏举 19138056898	日期	2023.10.24
	客户签字	生产厂家:	公司名称:	姓名/日期:	投诉及建议电话	13083692526

MYMULIKE CORN REMOVAL PADS

corn removal pads patch

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:83781-003
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SALICYLIC ACID (UNII: O414PZ4LPZ) (SALICYLIC ACID - UNII:O414PZ4LPZ)	SALICYLIC ACID	40 g in 100

Inactive Ingredients

Ingredient Name	Strength

PARAFFIN (UNII: I9O0E3H2ZE)				
PHENOL (UNII: 339NCG44TV)				
LANOLIN (UNII: 7EV65EAW6H)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:83781-003-01	24 in 1 BOX; Type 0: Not a Combination Product	11/05/2023	
Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC Monograph Drug	M030	11/05/2023		

Labeler - Zhengzhou Miaoqi Pharmaceutical Technology Co., Ltd (701762807)

Establishment				
Name	Address	ID/FEI	Business Operations	
Zhengzhou Miaoqi Pharmaceutical Technology Co., Ltd		701762807	label(83781-003) , manufacture(83781-003)	

Revised: 11/2023

Zhengzhou Miaoqi Pharmaceutical Technology Co., Ltd