WART REMOVER OINTMENT.- you yuping antibacterial ointment ointment Consilii LLC

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

83299-002 Wart Remover Ointment

Active Ingredient

Salicylic acid 1% Chlorhexidine di(acetate) 1%

Purpose

Wart remover

Use

for the removal of common and plantar warts the common wart is easily recognized by the rough "cauliflower-like appearance of the surface.

the plantar wart is recognized by its location only on the bottom of the foot.its tendemess and the interruption of the footprint pattern

Warnings

For external use only. Keep away from fire and flame.

Do not use

if discomfort persists.
if you have diabetes or poor bioodcirculation

WHEN USING SECTION

avoid contact with eyes If the product gets into the eye, flush with water for 15 minutes avoid inhaling vapors cap the tube tightly out of direct sunlight and store at room temperature away from heat

STOP USE section

if discomfort persists.

if you have diabetes or poor blood circulation.

ASK DOCTOR

if discomfort persists. if you have diabetes or poor bioodcirculation

KEEP OUT OF REACH OF CHILDREN

If swallowed, get medical help or contact a Poison Control Center (1-800-222-1222) right away.

Directions

wash the affected area may soak the wart in warm water for 5minutes dry area thoroughly. using the applicator(cotton swab)apply a layer of ointment to sufficiently cover each wart allow it to fully absorb and cover it witha bandage as needed repeat this procedure once or twicedaily as needed (until the wart isremoved) for up to 12 weeks

Other information

Keep away from direct sunlight or heat. Store at room tempetature 15°-30°C (59°-86°F)

Inactive ingredients

Fructus cnidii, radix sophorae flavescentis, herba portulacae, rhizoma atractyiodis macrocephalae, radix angelicae dahuricae.

QUESTIONS

1-914-608-125

PACKAGE LABEL - PRINCIPAL DISPLAY PANEL

83299-002-01



WART REMOVER OINTMENT.

you yuping antibacterial ointment ointment

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:83299-002

Route of Administration TOPICAL

Active Ingredient/Active Moiety Ingredient Name

SALICYLIC ACID (UNII: O414PZ4LPZ) (SALICYLIC ACID - UNII:O414PZ4LPZ)SALICYLIC ACID1 g in 100 gCHLORHEXIDINE (UNII: R4KO0DY52L) (CHLORHEXIDINE - UNII:R4KO0DY52L)CHLORHEXIDINE1 g in 100 g

Basis of Strength Strength

Inactive Ingredients

Ingredient Name	Strength
ANGELICA DAHURICA ROOT (UNII: 1V63N2S972)	
CITRUS RETICULATA WHOLE (UNII: O0OX7CMF92)	
FORSYTHIA SUSPENSA ROOT (UNII: YM9E56VP46)	
PLATYCODON GRANDIFLORUS WHOLE (UNII: WC73QE9274)	
ASARUM SIEBOLDII (UNII: T1134STW1K)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
SOPHORA FLAVESCENS ROOT (UNII: IYR6K8KQ5K)	
PORTULACA OLERACEA WHOLE (UNII: D5J3623SV2)	
ATRACTYLODES MACROCEPHALA ROOT (UNII: 08T3N29QJB)	
ARCTIUM LAPPA WHOLE (UNII: 73070DU1LA)	
BAMBUSA VULGARIS LEAF (UNII: EMY54R518C)	
BORNEOL (UNII: M89NIB437X)	
CNIDIUM MONNIERI FRUIT (UNII: V1IA3S3CUS)	
HONEY (UNII: Y9H1V576FH)	
ALCOHOL (UNII: 3K9958V90M)	

Packaging

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	# Item Code	Package Description	Marketing Start Date	Marketing End Date		
	1 NDC:83299-002-	20 g in 1 BOTTLE; Type 0: Not a Combination Product	04/05/2023			

Marketing Information

Marketing Application Number or Monograph Category Citation		Marketing Start Date	Marketing End Date
OTC monograph final	M028	04/05/2023	

Labeler - Consilii LLC (118891890)

Establishment

Name	Address	ID/FEI	Business Operations
Consilii LLC		118891890	label(83299-002), manufacture(83299-002)

Revised: 4/2023 Consilii LLC