

NAIL FUNGUS TREATMENT- tomumia nail fungus liquid
Consilii LLC

83299-030

Active Ingredient

Chlorhexidine acetate (3%)

Purpose

for Toenails & Fingernails /Restores Fungus Damaged Nails

Use

(Usage)

Clean the affected area with soap and warm water and dry thoroughly.
Gently file thickened part of the fungal nail with a nail file until it's flat.
Using the brush, apply a thin layer of liquid onto the affected area.
Repeat this process 2 times daily for
1-2 weeks (or) until the infection is gone

Warnings

For external use only

Keep out of the reach of children.

Do not use

Avoid contact with eyes and mouth. Do not use it on the irritated skin. Not suitable for pregnant and breastfeeding.

When Using

Using the brush, apply a thin layer of liquid onto the affected area

Stop Use

If swallowed, get medical assistance or contact the poison control center.

Ask Doctor

If swallowed, get medical assistance or contact the poison control center.

Keep Out Of Reach Of Children

Keep out of the reach of children.

Directions

1. For thickened and hard nails, it must be thinned before use.
2. If there is a slight deposit, it is the Chinese medicinal material, feel free to use it.
3. After healthy nails grow, take care and do not file them. Always keep the hands and feet clean. After the nails are normal, continue to use for 30 days.

Other information

Store at room temperature
Avoid excessive heat 37°C (99°F)

Inactive ingredients

Impatiens
Sophora flavescens
Tougu grass
Stemona
Angelica dahurica dahurica
Vitex bark
Wormwood leaf
chlorhexidine
acetate
water

Questions

1-9146081258

PRINCIPAL DISPLAY PANEL

Brand / Pulchra

Size / Package size: 9 cm x 4 cm x 4 cm
Bottle label size: 10.5 cm x 4.5 cm

Color / ■ PANTONE Reflex Blue C
RGB 0 20 137 / CMYK 100 87 0 20

■ PANTONE P 103-16 C
RGB 46 43 87 / CMYK 97 81 0 51

■ PANTONE Yellow C
RGB 254 221 0 / CMYK 0 1 100 0

Gradient



Package design



Label design



NAIL FUNGUS TREATMENT

tomumia nail fungus liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:83299-030
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CHLORHEXIDINE ACETATE (UNII: 5908Z UF22Y) (CHLORHEXIDINE - UNII:R4KO0DY52L)	CHLORHEXIDINE ACETATE	0.3 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
ANGELICA DAHURICA SEED (UNII: 9M69B50875)	
CHLORHEXIDINE (UNII: R4KO0DY52L)	
ILEX CORNUTA WHOLE (UNII: 9WEY0RQ9HO)	
VITEXIN (UNII: 9VP70K75OK)	
SOPHORA FLAVESCENS WHOLE (UNII: X8KX602M5L)	
IMPATIENS WALLERIANA WHOLE (UNII: 8W4759RC84)	
ACETATE ION (UNII: 569DQM74SC)	
STEMONA TUBEROSA WHOLE (UNII: 9373Z OT316)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:83299-030-01	30 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/13/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M005	06/13/2024	

Labeler - Consilii LLC (118891890)

Establishment

Name	Address	ID/FEI	Business Operations
Consilii LLC		118891890	label(83299-030) , manufacture(83299-030)

Revised: 6/2024

Consilii LLC