

**NAIL TREATMENT LIQUID- salicylic acid, chlorhexidine di(acetate) liquid**  
**Shenzhen Shenyuanye Trading Co., LTD**

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**Nail Treatment Liquid**

SALICYLIC ACID 2%

Chlorhexidine di(acetate) 2%

REMOVES FUNGUS

Uses For all caused by nail fungus

nail discoloration

nail thickening

nail splitting

nail crumbling

For external use on nailfungus(onychomycosis) only, please don't use it on healthy nails.

Don't use it on damaged skin, otherwise, your damaged skin maybe painful.

Don't drink, avoid contact with your eye.

For people with severe skin ulceration, please use it with caution.

Keep out of reach of children

Children use it under the supervision of adults.

Stop use and ask a doctor if

You are allergic to vinegar (such as itching, rash swelling of the lips, eyelids, and shortness of breath)

Direction

Apply it in the day and night (twice a day).

Before using it for the first time, soak the diseased nail with warm water to soften it, use the nail file to polish the nail, make it thin visibly, and scrape off the dirt on the nail (be careful not to hurt the nail bed), then apply the liquid.

After first use, every 2-3 days use the nail file to polish the surface of the diseased nail and then continue to use it.

If swallowed, get medical help or contact a poison control center right away.

Water, Sorbitol, Alcohol, Propylene-Glycol, Angelica Dahurica, Laurocapram, Caramel, Dimethylol.Urea, Sunset Yellow.

For the question, please contact us by [bybaizelu@outlook.com](mailto:bybaizelu@outlook.com)



# NAIL TREATMENT LIQUID

salicylic acid, chlorhexidine di(acetate) liquid

Product Information			
<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:84336-002
<b>Route of Administration</b>	TOPICAL		

Active Ingredient/Active Moiety				
Ingredient Name			Basis of Strength	Strength
CHLORHEXIDINE ACETATE (UNII: 5908ZUF22Y) (CHLORHEXIDINE - UNII:R4KO0DY52L)			CHLORHEXIDINE	2 mg in 100 mL
SALICYLIC ACID (UNII: O414PZ4LPZ) (SALICYLIC ACID - UNII:O414PZ4LPZ)			SALICYLIC ACID	2 mg in 100 mL
Inactive Ingredients				
Ingredient Name				Strength
SORBITOL (UNII: 506T60A25R)				
LAUROCAPRAM (UNII: 1F3X9DRV9X)				
CAMEL (UNII: T9D99G2B1R)				
WATER (UNII: 059QF0KO0R)				
OXYMETHUREA (UNII: N68H97CAWG)				
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)				
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)				
ANGELICA DAHURICA ROOT (UNII: 1V63N2S972)				
ALCOHOL (UNII: 3K9958V90M)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:84336-002-01	30 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/11/2024	
Marketing Information				
Marketing Category	Application Number or Monograph Citation		Marketing Start Date	Marketing End Date
OTC Monograph Drug	M005		06/11/2024	

**Labeler** - Shenzhen Shenyuanye Trading Co., LTD (550376513)

**Registrant** - Shenzhen Shenyuanye Trading Co., LTD (550376513)

### Establishment

Name	Address	ID/FEI	Business Operations
Guangzhou Mizi Biotechnology Co., Ltd.		418001649	manufacture(84336-002)

Revised: 6/2024

Shenzhen Shenyuanye Trading Co., LTD