7 DAYS NAIL FUNGUS TREATMENT- sodium sulfide liquid
Guangdong Ximonth Technology Co., Ltd.

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.

Active Ingredient(s)

Purpose

Restores damage nail & cuticle

Use

1.File the area affected by the nail to enhance the permeability of the essence.2.Clean the surface with a dry swab /cloth.3.Apply the essence to the entire surface of the infected nail.4.Apply the cream twice a day (morning and evening) for at least 4 weeks.

Warnings

Please keep out of reach of children.Do not swallow.Please clean your hands before use to ensure the best results from the product.Discontinue use if signs of irritation or rash occur.Store in a cool and dry place.

Do not use

Discontinue use if signs of irritation or rash occur.

STOP USE

Discontinue use if signs of irritation or rash occur.

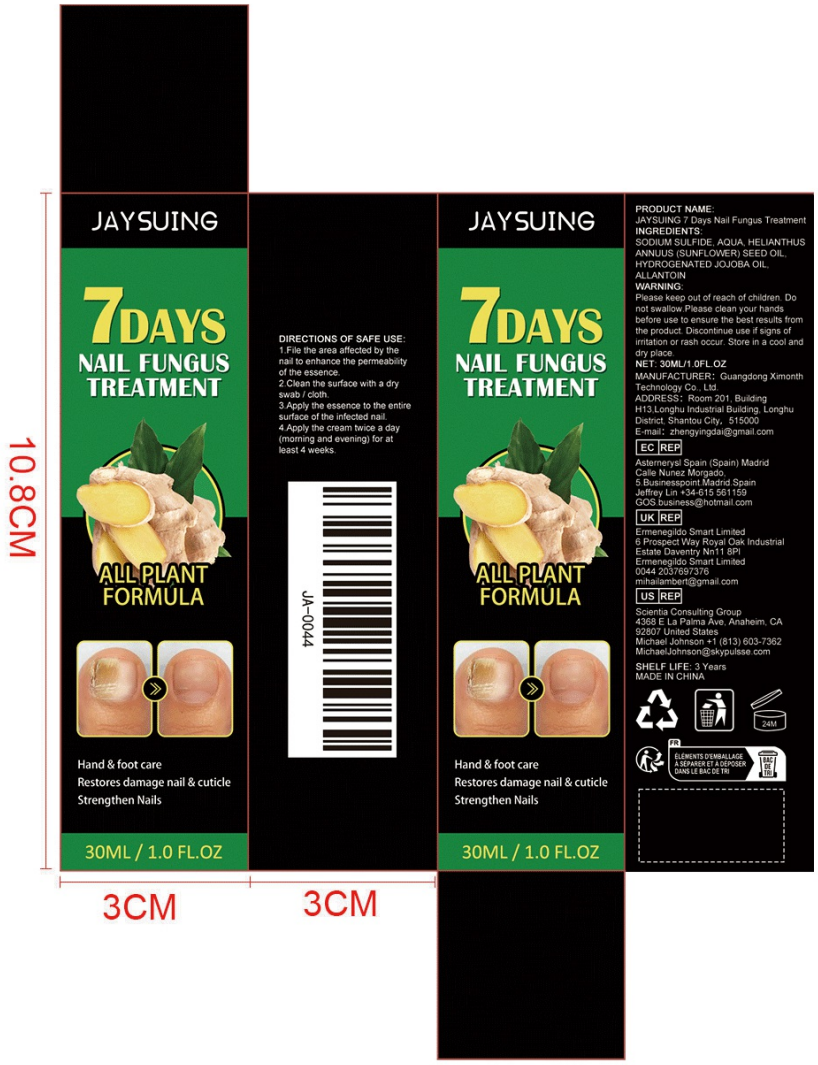
Please keep out of reach of children. Do not swallow.

Avoid freezing and excessive heat above 40C (104F)

Store in a cool and dry place.

AQUA

纸盒
规格尺寸:长3X宽3X高10.8cm



7 DAYS NAIL FUNGUS TREATMENT

sodium sulfide liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:84660-004
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**

SUNFLOWER OIL (UNII: 3W1JG795YI) (SUNFLOWER OIL - UNII:3W1JG795YI)	SUNFLOWER OIL	3 mg in 30 mg
SODIUM SULFIDE (UNII: YGR27ZW0Y7) (SULFIDE ION - UNII:G15I91XETI)	SODIUM SULFIDE	7.5 mg in 30 mg
HYDROGENATED JOJOBA OIL (UNII: 7F674YQ5SO) (HYDROGENATED JOJOBA OIL - UNII:7F674YQ5SO)	HYDROGENATED JOJOBA OIL	6 mg in 30 mg

Inactive Ingredients

Ingredient Name	Strength
ALLANTOIN (UNII: 344S277G0Z)	4.5 mg in 30 mg
WATER (UNII: 059QF0KO0R)	9 mL in 30 mg

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:84660-004-01	30 mg in 1 BOX; Type 0: Not a Combination Product	08/28/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		08/28/2024	

Labeler - Guangdong Ximonth Technology Co., Ltd. (699436264)

Registrant - Guangdong Ximonth Technology Co., Ltd. (699436264)

Establishment

Name	Address	ID/FEI	Business Operations
Guangdong Ximonth Technology Co., Ltd.		699436264	manufacture(84660-004)

Revised: 8/2024 Guangdong Ximonth Technology Co., Ltd.