

SCHOLLS WELLNESS COMPANY LLC INGROWN TOENAIL- sodium sulfide gel
Scholls Wellness Company LLC

90000062 Ingrown Toenail

Drug Facts

Active Ingredient Purpose

Sodium sulfide 1%.....Ingrown toenail reliever

Purpose

Ingrown toenail reliever

Use

for temporary relief of pain and discomfort from ingrown toenails

Warnings

For external use only

Do not use

- on open sores

Ask a doctor before use if you have

- diabetes
- poor blood circulation
- gout

When using this product

- **use with a retainer ring**
- **avoid contact with eyes. If product gets in eyes, flush with water for 15 minutes and get medical help right away.**

Stop use and ask a doctor if

- redness or swelling of your toe increases
- discharge is present around the nail
- symptoms last more than 7 days or clear up and occur again within a few days

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Directions

adults and children 12 years and over:

wash affected area and dry thoroughly

place retainer ring on toe with slot over the area where the ingrown nail and the skin meet. Smooth ring down firmly.

cut open tip of tube on score mark. Apply enough gel product to fill the slot in the ring. Immediately replace cap on tube.

place round center section of bandage directly over the gel-filled ring to seal the gel in place. Smooth ends of bandage strip around toe.

repeat twice daily (morning and night) for up to 7 days until pain and discomfort is relieved or until the nail can be lifted out of the nail groove and easily trimmed.

children under 12 years: ask a doctor

Other information

- save carton for full labeling
- keep tube tightly closed when not in use
- store between 20° to 25°C (68° to 77°F)

Inactive ingredients

edetate disodium, hydroxyethyl cellulose, potassium acetate, water

Questions

Questions? 1-866-360-3226



SCHOLLS WELLNESS COMPANY LLC INGROWN TOENAIL

sodium sulfide gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:73469-0062
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SODIUM SULFIDE (UNII: YGR27ZW0Y7) (SULFIDE ION - UNII:G15I91XETI)	SODIUM SULFIDE	1 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
POTASSIUM ACETATE (UNII: M911911U02)	
WATER (UNII: 059QF0KO0R)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
HYDROXYETHYL CELLULOSE, UNSPECIFIED (UNII: T4V6TWG28D)	

Product Characteristics

Color	green	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73469-0062-1	1 in 1 CARTON	01/01/2021	
1		1 g in 1 TUBE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M029	01/01/2021	

Labeler - Scholls Wellness Company LLC (117174744)

Registrant - Scapa Tapes North America, LLC (079995435)

Revised: 10/2023

Scholls Wellness Company LLC