SCHOLLS WELLNESS COMPANY LLC INGROWN TOENAIL- sodium sulfide gel
Scapa Tapes North America, 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.

Ingrown Toenail (Scapa)

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 Poisen 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 20º to 25ºC (68º to 77ºF)

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:66843-721
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 10 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:66843-721-41	1 in 1 CARTON	01/01/2021	08/31/2023
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 final	part358D	01/01/2021	

Labeler - Scapa Tapes North America, LLC (079995435)

Registrant - Scapa Tapes North America, LLC (079995435)

Revised: 8/2023 Scapa Tapes North America, LLC