

**NORO-X FOAMING HANDSOAP- o-cymen-5-ol soap
BIO3S 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.

77935-507 NORO-X Foaming Hand Soap 500mL

Active ingredients

o-Cymen-5-ol 0.095%

Purposes

o-Cymen-5-ol 0.095%..... Antibacterial

Uses

Uses for hand washing to decrease bacteria on the skin

Warnings

Warnings For external use only.

Warnings

When using this product, avoid contact with eyes. In case of contact with eyes, rinse with water.

Warnings

Stop use and ask a doctor if irritation or redness develops.

Warnings

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

Directions

- pump into hands
- lather with soap for at least 30 seconds
- rinse hands with water

Inactive ingredients

Water, Disodium Laureth Sulfosuccinate, Glycerin, Cocamidopropyl Betaine, Disodium Cocoamphodiacetate, Lauryl Glucoside, Sodium Chloride, Capiylii Glycol, Ethihexylglycerin, Menthol, Fragrance, Allantoin, Sodium Benzoate, Citric Acid, Bacillus/Canavalia Ensiformis Seed Ferment Extract, Disodium EDTA, Hexylene Glycol, Butylene Glycol, Chrysanthellum Indicum Extract, RhusSemialata Gall Extract, Propolis Extract, Pinus Densiflora Leaf Extract, Scutellaria Baicalensis Root Extract

Display Panel

노로X 손세정제 500ml

규격 : 252 X 82



NORO-X FOAMING HANDSOAP

o-cymen-5-ol soap

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:77935-507
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
O-CYMEN-5-OL (UNII: H41B6Q1I9L) (O-CYMEN-5-OL - UNII:H41B6Q1I9L)	O-CYMEN-5-OL	0.095 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
DISODIUM LAURETH SULFOSUCCINATE (UNII: D6DH1DTN7E)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
LAURYL GLUCOSIDE (UNII: 76LN7P7UCU)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
EDETATE DISODIUM ANHYDROUS (UNII: 8NLQ36F6MM)	
CHRYSANTHELLUM INDICUM TOP (UNII: STJ856D1Z0)	
MENTHOL (UNII: L7T10EIP3A)	
ALLANTOIN (UNII: 344S277G0Z)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
CAPRYLYL GLYCOL (UNII: 00YIU5438U)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
PINUS DENSIFLORA LEAF (UNII: Q1Q9P50WY)	
HEXYLENE GLYCOL (UNII: KEH0A3F75J)	
FRAGRANCE CLEAN ORC0600327 (UNII: 329LCV5BTF)	
GLYCERIN (UNII: PDC6A3C0OX)	
WATER (UNII: 059QF0KO0R)	
DISODIUM COCOAMPHODIACETATE (UNII: 18L9G3U51M)	
PROPOLIS WAX (UNII: 6Y8XYV2NOF)	
SCUTELLARIA BAICALENSIS ROOT (UNII: 7J95K7ID2S)	
RHUS CHINENSIS GALL (UNII: 4W3Y2V7J3R)	
CANAVALIA ENSIFORMIS WHOLE (UNII: U485ST9OUN)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:77935-507-01	500 mL in 1 BOTTLE; Type 0: Not a Combination Product	12/27/2022	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		12/27/2022	

Labeler - BIO3S Co.,Ltd. (694813103)

Registrant - BIO3S Co.,Ltd. (694813103)

Establishment

Name	Address	ID/FEI	Business Operations
BIO3S Co.,Ltd.		694813103	manufacture(77935-507)

Revised: 2/2023

BIO3S Co.,Ltd.