

1438 DYNASOAP- antibacterial hand soap soap
Dynarex Corporation

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.

1438 Dyna Soap - Antibacterial Hand Soap 67777-319-01

Active Ingredient

Benzalkonium Chloride 0.13%.....Antiseptic handwash

Purpose

Antibacterial Hand Soap

Use

Cleaning and antiseptic cleansing of hands and skin.

Warnings

For external use only.

When using this product

Avoid contact with eyes. In case of contact, flush with water.

Stop use and ask a doctor if.

Irritation and redness develop.

Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center right away.

Directions.


Wet hands and forearms. Apply 5 milliliters (teaspoon) or palmful to hands and forearms. Scrub thoroughly for 1 minute. Rinse and repeat.


Inactive Ingredients

Benzyl Alcohol, Citric Acid Monohydrate, Cocamide DEA, Cocamidopropyl Betaine, Glycerin, Lauryl Glucoside, Methylchloroisothiazolinone, Methylisothiazolinone, Parfum, Sodium Chloride, Sodium Laureth Sulfate, Sulisobenzone, Water.


Label

Drug Facts	
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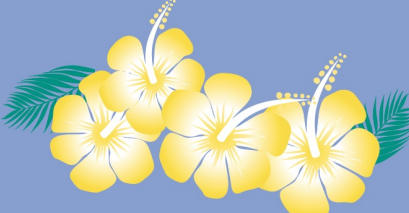
Reorder No. 1438  Recyclable Bottle



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**TROPICAL BEACH
ANTIBACTERIAL HAND SOAP
GENTLY CLEANS & PROTECTS**



7.5 FL. OZ. (221.8 mL)

Manufactured for: Dynarex Corporation
10 Glenshaw Street • Orangeburg, NY 10962
USA • www.dynarex.com
Made in Turkey

1438 DynaSoap Label

1438 DYNASOAP

antibacterial hand soap soap

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:67777-319
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII: 7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.13 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
SULISOBENZONE (UNII: 1W6L629B4K)	
SODIUM LAURETH SULFATE (UNII: BPV390UAP0)	
LAURYL GLUCOSIDE (UNII: 76LN7P7UCU)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
GLYCERIN (UNII: PDC6A3C0OX)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	
COCO DIETHANOLAMIDE (UNII: 92005F972D)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
WATER (UNII: 059QF0KO0R)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	

Product Characteristics

Color	blue	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:67777-319-01	24 in 1 CASE	08/01/2020	
1	NDC:67777-319-02	0.13 g in 1 BOTTLE, PUMP; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	08/01/2020	

Labeler - Dynarex Corporation (008124539)

Registrant - Dynarex Corporation (008124539)

Establishment

Name	Address	ID/FEI	Business Operations
Hobi Kozmetik Imalat Sanayi Ve Ticaret Anonim Sirketi		533139227	manufacture(67777-319)

