ALCOHOL FOAMING HAND SANITIZER- alcohol soap
Lawson Prodcuts, Inc

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.

Ocean Blue Alcohol Foaming Hand Sanitizer

Active Ingredient

Ethyk Alcohol 62%

Warnings

For external use only. Do not ingest.
FLAMMABLE. This product contains ethyl alcohol. Keep away from sources of ignition.
Avoid contact with eyes. In case of eye contact, flush with large quantities of water, seek medical attention if irritation persists.
If skin irritation develops, discontinue use. If irritation persists for more than 72 hours, seek medical attention.
Keep out of reach of children.
If swallowed, get medical help or contact a Poison Control Center right away.

Inactive Ingredients

Water, Propylene Glycol, Acrylic Polymer, Alkyl Phosphate Salts, Fragrance, DC Green #5.

Purpose

Antiseptic

Alcohol Foaming Hand Sanitizer

KEEP OUT OF REACH OF CHILDREN

Directions

Read the entire label before using this product.

Dispense 2 pumps of product onto palm of hand and rub thoroughly over all surfaces of both hands until dry.

Uses

Use hand sanitizer to reduce microorganisms on the skin.
Use this product when soap and water are not available.

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Drug Facts

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- **Keep out of reach of children.** • If swallowed, get medic help or contact a Poison Control Center right away.

Directions

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- Dispense 2 pumps of product onto palm of hand and rub thoroughly over all surfaces of both hands until dry.

Inactive Ingredients

Water, Perfluoroalkyl phosphates diethanolamine salts, PEG-10 Dimethicone, Propylene glycol, Fragrance, D&C Green #5.

WARNING: Cancer - www.P65Warnings.ca.

Sold by:
Lawson Products, Inc.
866-LAWSON4U (866-529-7664)
lawsonproducts.com

8770 W. Bryn Mawr Ave., Suite 900
Chicago, IL 60631-3515
USA

Made in U.S

1573956001
RLB3665A 02765

950 mL (25.4 fl. oz.)

ALCOHOL FOAMING HAND SANITIZER

alcohol soap

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:62428-701
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	0.62 mL in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
D&C GREEN NO. 5 (UNII: 8J6RDU8L9X)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
C20-22 ALKYL PHOSPHATE (UNII: L4VKP0Y7RP)	
WATER (UNII: 059QF0KO0R)	
PEG-10 DIMETHICONE (600 CST) (UNII: 8PR7V1SVM0)	

Product Characteristics			
Color	blue	**Score**	
Shape		**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:62428-701-03	750 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/14/2019	
2	NDC:62428-701-57	550 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/14/2019	
3	NDC:62428-701-53	50 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/14/2019	
4	NDC:62428-701-29	1000 mL in 1 BAG; Type 0: Not a Combination Product	03/14/2019	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	03/14/2019	

Labeler - Lawson Prodcuts, Inc (005438890)

Registrant - Betco Corporation, Ltd (005050158)

Establishment

Name	Address	ID/FEI	Business Operations
Betco Corporation, Ltd		005050158	manufacture(62428-701)

Revised: 3/2019

Lawson Prodcuts, Inc