

CLEARLY BETTER- foaming hand sanitizer liquid
Intercon Chemical Company

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.

Alcohol Free Foaming Hand Sanitizer
391.000/391AA

claims

Clearly Better

by INTRTCON

ALCOHOL FREE FOAMING HAND SANITIZER

Active ingredient

Benzalkonium chloride 0.13%

Purpose

Antiseptic

Use

- to decrease bacteria on the skin that could cause disease

Warnings

For external use only-hands

When using this product

- avoid contact with eyes. if contact occurs, rinse eyes thoroughly with water.
- avoid contact with broken skin

Stop use and ask a doctor if

- irritation or redness develops
- condition persists for more than 72 hours

Keep out of reach of children.

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

Directions

- wet hands thoroughly with product and allow to dry without wiping
- for children under 6, use only under adult supervision
- not recommended for infants

Inactive ingredients

cetrimonium chloride, diglycerol, disodium cocoamphodiacetate, fragrance, glycerin, hydrochloric acid, methoxy PEG/PPG-7/3 aminopropyl dimethicone, methylchloroisothiazolinone, methylisothiazolinone, tetrasodium EDTA, water

Adverse Reactions Section

DISTRIBUTED BY: INTERCON CHEMICAL CO.

1100 CENTRAL INDUSTRIAL DR.

ST. LOUIS, MO 63110

Principal display panel

ClearlyBetter

BY INTERCON

ALCOHOL FREE

FOAMING

HAND SANITIZER

1150 mL (39 FL OZ)



ClearlyBetter
BY INTERCON®

L0017008FA

**ALCOHOL FREE
FOAMING
HAND SANITIZER**

1150 mL (39 FL OZ)

CLEARLY BETTER

foaming hand sanitizer liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:67502-391
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety				
Ingredient Name			Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII: 7N6JUD5X6Y)			BENZALKONIUM CHLORIDE	1.3 mg in 1 mL
Inactive Ingredients				
Ingredient Name				Strength
CETRIMONIUM CHLORIDE (UNII: UC9PE95IBP)				
DIGLYCERIN (UNII: 3YC120743U)				
DISODIUM COCOAMPHODIACETATE (UNII: 18L9G3U51M)				
glycerin (UNII: PDC6A3C0OX)				
HYDROCHLORIC ACID (UNII: QTT17582CB)				
METHOXY PEG/PPG-7/3 AMINOPROPYL DIMETHICONE (UNII: 4M7P1JZ2V2)				
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)				
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)				
EDETATE SODIUM (UNII: MP1J8420LU)				
water (UNII: 059QF0KOOR)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:67502-391-27	1150 mL in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product	08/28/2017	
Marketing Information				
Marketing Category	Application Number or Monograph Citation		Marketing Start Date	Marketing End Date
OTC monograph not final	part333A		08/28/2017	

Labeler - Intercon Chemical Company (103204970)

Registrant - Vi Jon, LLC (790752542)

Establishment

Name	Address	ID/FEI	Business Operations
Vi-Jon, LLC		790752542	manufacture(67502-391)

Establishment

Name	Address	ID/FEI	Business Operations
Vi-Jon, LLC		088520668	manufacture(67502-391)