ALTRA NON-ALCOHOL FOAMING HAND SANITIZER WITH ALLANTOIN AND ALOE-
benzalkonium chloride solution
Buckeye International, 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.

Active ingredient

Benzalkonium Chloride 0.12%

Purpose

Antibacterial

Uses

- Hand sanitizer to help reduce bacteria on the skin that could cause disease
- Recommended for repeated use

Warnings

For external use only.

When using this product do not use in or near eyes.

If in eyes, flush thoroughly with water.

If irritation or rash appears and persists, stop use and see a physician.

Keep out of reach of children.

If swallowed, call a physician or Poison Control Center immediately.

Directions

- Dispense an adequate amount of hand sanitizer
- Rub hads together until completely dry

Inactive ingredients

Water (Aqua), Cocamidopropyl Betaine, Lauramine Oxide, Tetrasodium EDTA, Allantoin, Fragrance (Parfum), Aloe Barbadensis Leaf Juice, Methylchloroisothiazolinone, Methylisothiazolinone

Questions?

Call Buckeye International, Inc. 314-291-1900

Monday through Friday 8:00 a.m. to 5:00 p.m. CST

ALTRA

Non-Alcohol

Foaming hand

Sanitizer

with Allantoin and Aloe

Hand Hygiene

Manufactured by

Buckeye International

2700 Wagner Place Maryland Heights

MO 63043 USA (314) 291-1900

Product # 71151120

Net Contents 1250 mL (42.2 fl oz)

ALTRA NON-ALCOHOL FOAMING HAND SANITIZER WITH ALLANTOIN AND ALOE

benzalkonium chloride solution

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:30805-015
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.12 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
LAURAMINE OXIDE (UNII: 4F6FC4MI8W)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
ALLANTOIN (UNII: 344S277G0Z)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
WATER (UNII: 059QF0KO0R)	
EDETATE SODIUM (UNII: MP1J8420LU)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:30805-015-05	50 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/15/2020	
2	NDC:30805-015-07	550 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/15/2020	
3	NDC:30805-015-09	1000 mL in 1 BAG; Type 0: Not a Combination Product	04/15/2020	
4	NDC:30805-015-02	1250 mL in 1 BAG; Type 0: Not a Combination Product	04/15/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	04/15/2020	

Labeler - Buckeye International, Inc. (077132280)

Establishment

Name	Address	ID/FEI	Business Operations
Buckeye International, Inc.		077132280	manufacture(30805-015)

Revised: 4/2020 Buckeye International, Inc.