## SYMMETRY 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.

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## **Drug Facts**

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 hands 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 8:00to 5:00 p.m. CST

For General Hand Sanitizing

Symmetry

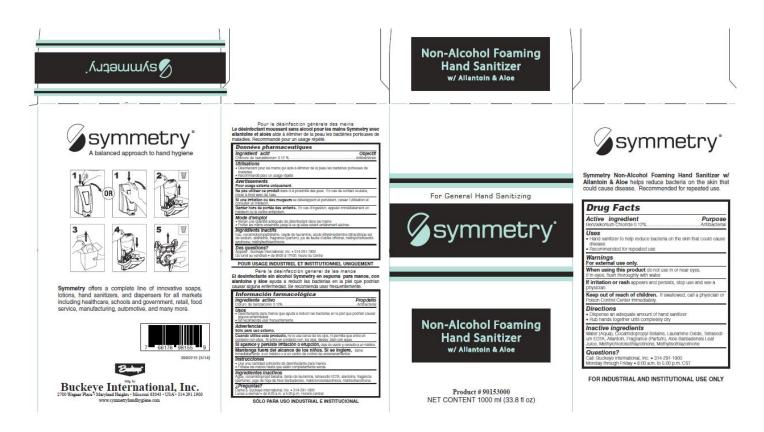
Non-Alcohol Foaming

Hand Sanitizer

w/ Allantoin and Aloe

Product # 90153000

Net Content 1000 ml (33.8 fl oz)



## SYMMETRY NON-ALCOHOL FOAMING HAND SANITIZER WITH ALLANTOIN AND ALOE

benzalkonium chloride solution

Product Information								
Product Type	HUMAN OTC DRUG	Item Code (Source) NDC		NDC:3	2:30805-010			
Route of Administration	TOPICAL							
Active Ingredient/Active Mo	iety							
Ing	Ingredient Name Basis of Strength							
Benzalkonium Chloride (UNII: F5UM	2KM3W7) (Benzalkonium - UNII:7	N6 JUD5 X6 Y)	Benzalkonium Ch	1 lo ride	0.12 g in 100 mL			
Inactive Ingredients								
	Ingredient Name				Strength			
Water (UNII: 059QF0KO0R)								
Cocamidopropyl Betaine (UNII: 50CF3011KX)								
The second se	FJOTIKA)							
LAURAMINE O XIDE (UNII: 4F6FC4M								
	8 W)							
LAURAMINE O XIDE (UNII: 4F6 FC4M	8 W)							
LAURAMINE O XIDE (UNII: 4F6FC4M) EDETATE SO DIUM (UNII: MP1J84201	8 W)							
LAURAMINE O XIDE (UNII: 4F6FC4M) EDETATE SO DIUM (UNII: MP1J84201 Allantoin (UNII: 344S277G0Z)	8W) U)							
LAURAMINE O XIDE (UNII: 4F6FC4M) EDETATE SO DIUM (UNII: MP1J84201 Allantoin (UNII: 344S277G0Z) Aloe Vera Leaf (UNII: ZY81Z83H0X)	8W) U) NE (UNII: DEL7T5QRPN)							
LAURAMINE O XIDE (UNII: 4F6FC4M) EDETATE SO DIUM (UNII: MP1J84201 Allantoin (UNII: 344S277G0Z) Aloe Vera Leaf (UNII: ZY81Z83H0X) METHYLCHLOROISOTHIAZOLING	8W) U) NE (UNII: DEL7T5QRPN)							
LAURAMINE O XIDE (UNII: 4F6FC4M) EDETATE SO DIUM (UNII: MP1J84201 Allantoin (UNII: 344S277G0Z) Aloe Vera Leaf (UNII: ZY81Z83H0X) METHYLCHLOROISOTHIAZOLING	8W) U) NE (UNII: DEL7T5QRPN)							

#	Item Code		Package Description	Marketing Start Date	Marketing End Date			
1	NDC:30805-010- 02	1250	mL in 1 BAG; Type 0: Not a Combination Product	11/0 1/20 11				
2	NDC:30805-010- 05	50 m Prod	L in 1 BOTTLE, PUMP; Type 0: Not a Combination uct	11/0 1/20 11				
3	NDC:30805-010- 07	550 Prod	mL in 1 BOTTLE, PUMP; Type 0: Not a Combination uct	11/0 1/20 11				
4	NDC:30805-010- 09	1000	mL in 1 BAG; Type 0: Not a Combination Product	04/07/2014				
Marketing Information								
Marketing Category		ory	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date			
0	TC monograph not f	inal	part333A	11/0 1/20 11				

## Labeler - Buckeye International, Inc. (077132280)

Establishment									
Name	Address	ID/FEI	<b>Business Operations</b>						
Buckeye International, Inc.		077132280	manufacture(30805-010)						

Revised: 12/2016

Buckeye International, Inc.