# TOTAL SOLUTIONS ALCOHOL-FREE HAND SANITIZING FOAMbenzalkonium liquid Athea Laboratories, Inc.

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## 0282-Total Solutions Alcohol-Free Hand Sanitizing Foam

## Active ingredient

Benzalkonium Chloride 0.12%

### **Purpose**

**Antiseptic** 

#### Uses

- For handwashing to decrease bacteria on the skin.
- After changing diapers.
- After assisting ill persons.
- Before contact with a person under medical care or treatment.
- Recommended for repeat use.

## Warnings

For external use only.

#### Do not use

in the eyes.

# When using this product

do not get into eyes. If contact occurs, rinse thoroughly with water.

# Stop use and ask a doctor if

irritation and redness develop. If condition persists for more that 72 hours consult a doctor.

# 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.
- Children under six years of age should be supervised whtn using this product.

## Inactive ingredients

Water, Aloe Barbadensis Leaf Juice, Polysorbate 20, Glycerin, Potassium Sorbate, Fragrance, Citric Acid, Disodium EDTA, Tocopheryl Acetate (Vitamin E).

### 1 gallon



# TOTAL SOLUTIONS ALCOHOL-FREE HAND SANITIZING FOAM

benzalkonium liquid

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:62819-282
Route of Administration	TOPICAL		

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

Inactive Ingredients			
Ingredient Name	Strength		
ALOE BARBADENSIS LEAF (UNII: ZY81Z83H0X)			
GLYCERIN (UNII: PDC6A3C0OX)			
ALPHA-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)			
POLYSORBATE 20 (UNII: 7T1F30V5YH)			
WATER (UNII: 059QF0KO0R)			
EDETATE DISODIUM (UNII: 7FLD91C86K)			
CITRIC ACID (UNII: 2968PHW8QP)			
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)			

Purpose

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:62819- 282-00	3785 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/03/2011	

Marketing Information				
Marketing Application Number or Monograp Category Citation		Marketing Start Date	Marketing End Date	
OTC Monograph Drug	505G(a)(3)	01/03/2011		

# Labeler - Athea Laboratories, Inc. (078281547)

Establishment				
Name	Address	ID/FEI	Business Operations	
Athea Laboratories, Inc.		006117816	label(62819-282), manufacture(62819-282)	

Revised: 1/2025 Athea Laboratories, Inc.