

SUMMIT 377ALCOHOL FREE FOAM HAND SANITIZER- benzalkonium chloride solution

R.L. Williams Company

Summit 377 Alcohol Free Foam Hand Sanitizer

Active Ingredient

Benzalkonium Chloride 0.1%

Purpose

Antibacterial

Use

For hand washing to decrease bacteria on the skin

Warnings

For external use only.

When using this product avoid contact with eyes. In case of eye contact, flush eyes with water.

Stop use and ask a doctor if irritation or redness develops, or if 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

- Pump a small amount of foam into palm of the hand.
- Rub thoroughly over all surfaces of both hands.
- Rub hands briskly until dry.

Inactive Ingredients

Water, Glycerin, Propylene Glycol, Phenoxyethanol, Lauramine Oxide, Panthenol, Citric Acid



SUMMIT 377

ALCOHOL FREE FOAM HAND SANITIZER

HELPS
MINIMIZE
BACTERIA

Benzalkonium Chloride 0.1%



PLEASE RECYCLE

28.7 fl. oz. (850 mL)

LBL2032-1.0

764-539421-12 260323-0950

Drug Facts

Active Ingredient

Benzalkonium Chloride 0.1% Antibacterial

Purpose

Use For hand washing to decrease bacteria on the skin

Warnings

For external use only.

When using this product avoid contact with eyes. In case of eye contact, flush eyes with water.

Stop use and ask a doctor if irritation or redness develops, or if 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

- Pump a small amount of foam into palm of the hand.
- Rub thoroughly over all surfaces of both hands.
- Rub hands together briskly until dry.

Inactive Ingredients

Water, Glycerin, Propylene Glycol, Phenoxyethanol, Lauramine Oxide, Panthenol, Citric Acid



Manufactured for: **R.L. Williams Company**
409 Thornburg Drive SE • Conover, NC 28613

LBL2033-1.0 764-716390-12 260323-0950

SUMMIT 377ALCOHOL FREE FOAM HAND SANITIZER

benzalkonium chloride solution

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:23590-600
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	1 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
PANTHENOL (UNII: WW9CM0067Z)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	

GLYCERIN (UNII: PDC6A3C0OX)	
LAURAMINE OXIDE (UNII: 4F6FC4MI8W)	
WATER (UNII: 059QF0KO0R)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	

Product Characteristics

Color	white (water white - colorless, dispensed as a white foam)	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:23590-600-10	850 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	04/30/2026	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	04/30/2026	

Labeler - R.L. Williams Company (099976362)

Revised: 4/2026

R.L. Williams Company