

REMEVERSE INSTANT FOAM HAND SANITIZER- benzalkonium chloride liquid

RemeVerse

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.

RemeVerse Instant Foam Hand Sanitizer

RemeVerse instant foam hand sanitizer

Water, cetrimonium chloride, cocamidopropyl PG-dimonium chloride phosphate, dihydroxyethyl cocamine oxide, acetamidoethoxyethanol, citric acid, fragrance

RemeVerse Instant Foam Hand Sanitizer

Pump small amount into palm of hand ▪ Rub thoroughly over all surfaces of both hands ▪ Rub hands together briskly until dry ▪ For children under 6, use only under adult supervision

Topical Antimicrobial

Effective at eliminating many common harmful germs in as little as 15 seconds ▪ Recommended for repeated use

Benzalkonium Chloride .1% Active

Benzalkonium Chloride - .1%

Warnings:

For external use only

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

Safety Warning

Stop use and ask a doctor if irritation or redness develops, or if condition persists for more than 72 hours.

Warning - Children

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Purpose - Antimicrobial

Antimicrobial

Label

**CONTAINS
NO ALCOHOL**



- ▶ **Use without water - rub over hands**
- ▶ **Recommended for repeated use**
- ▶ **Moisturizes and conditions to leave skin soft and silky**
- ▶ **Won't stain clothing**
- ▶ **Fast drying; leaves no sticky residue**
- ▶ **Safe and effective**

300 mL / 10.1 FL. OZ.

Made in the USA

Drug Facts

Active Ingredient	Purpose
Benzalkonium Chloride 0.1%	Antimicrobial

Uses

▪ Effective at eliminating many common harmful germs in as little as 15 seconds ▪ Recommended for repeated use

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Directions

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Inactive ingredients

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Remeverse Hand Sanitizer Label 300ml lower resolution.jpg

REMEVERSE INSTANT FOAM HAND SANITIZER

benzalkonium chloride liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:70317-145
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.1 g in 300 mL

Inactive Ingredients

Ingredient Name	Strength
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	0.1 g in 300 mL
FRAGRANCE LAVENDER & CHIA F-153480 (UNII: SXS9CO2TZK)	0.1 g in 300 mL
CETRIMONIUM CHLORIDE (UNII: UC9PE95IBP)	0.1 g in 300 mL
DIHYDROXYETHYL COCAMINE OXIDE (UNII: 8AR51R3BL5)	0.1 g in 300 mL
COCAMIDOPROPYL PG-DIMONIUM CHLORIDE PHOSPHATE (UNII: H2KVQ74JM4)	0.1 g in 300 mL

Product Characteristics

Color		Score	
Shape	ROUND	Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70317-145-04	120 mL in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product	03/31/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	03/31/2020	

Labeler - RemeVerse (050186540)**Registrant** - Lexia LLC (015552120)**Establishment**

Name	Address	ID/FEI	Business Operations
Lexia, LLC		015552120	manufacture(70317-145)

Revised: 4/2020

RemeVerse