MPULSE ANTIMICROBIAL HAND SANITIZER- benzalkonium chloride liquid ANTIMICROBIAL SOLUTIONS, LLC

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 0.13%

Uses

- Helps reduce bacteria on the skin.
- Recommended for repeated use.

Warnings

- For external use only.
- Avoid contact with eyes.
- In case of contact, flush thoroughly with water.

Keep out of reach of children.

Directions

Apply product to hands and rub until dry.

Other information

Store in cool dry place between 35° and 103°F.

Inactive ingredients

Deionized Water, Propylene Glycol, Triethylene Glycol, Phospholipids Complex, Organo Silane Quaternaries.

Product Label



Moisturizes and Extends Protection Germ Killing Control Increases with Repeated Use Alcohol Free • Non-Toxic • Non-Flammable

MEDI DEFENSE mPulse HAND SANITIZER

DRUG FACTS

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KILLS 99.99% OF GERMS ON CONTACT

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301 Clanton Road, Suite C-104 Charlotte, NC 28217 1-855-877-5262 www.thinkclearstream.com MADE IN USA

NDC:

MPULSE ANTIMICROBIAL HAND SANITIZER

benzalkonium chloride liquid

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:78941-001

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)

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Inactive Ingredients Ingredient Name Strength WATER (UNII: 059QF0KO0R) PROPYLENE GLYCOL (UNII: 6DC9Q167V3) TRIETHYLENE GLYCOL (UNII: 3P5SU53360)

P	Packaging						
#	Item Code	Package Description	Marketing Start Date	Marketing End Date			
1	NDC:78941-001- 01	0.059 L in 1 BOTTLE; Type 0: Not a Combination Product	06/03/2020				
2	NDC:78941-001- 02	0.236 L in 1 BOTTLE; Type 0: Not a Combination Product	06/03/2020				
3	NDC:78941-001- 03	0.355 L in 1 BOTTLE; Type 0: Not a Combination Product	06/03/2020				
1	NDC:78941-001-	0 473 L in 1 ROTTI F. Type 0. Not a Combination Product	UE/U3/2U2U				

4	04	0.4/3 L III 1 DO 11 LE, 1 ype 0. IVOL a COMOMANON FIOUNCE	00/03/2020	
5	NDC:78941-001- 05	0.946 L in 1 BOTTLE; Type 0: Not a Combination Product	06/03/2020	
6	NDC:78941-001- 06	3.785 L in 1 JUG; Type 0: Not a Combination Product	06/03/2020	
7	NDC:78941-001- 07	18.927 L in 1 JUG; Type 0: Not a Combination Product	06/03/2020	
8	NDC:78941-001- 08	208.198 L in 1 DRUM; Type 0: Not a Combination Product	06/03/2020	
9	NDC:78941-001- 09	1040.990 L in 1 CONTAINER; Type 0: Not a Combination Product	06/03/2020	

Marketing Information					
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
OTC monograph not final	part333A	06/03/2020			

Labeler - Antimicrobial solutions, LLC (130702022)

Revised: 6/2020 ANTIMICROBIAL SOLUTIONS, LLC