CLINICARE PRO SURFACE SANITIZER- benzalkonium chloride, didecyl dimethyl ammonium chloride liquid Meroven 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.

Clinicare Pro Surface Sanitizer

Drug Facts

Active ingredient

- Benzalkonium Chloride 0.13%
- Didecyl dimethyl ammonium chloride 0.15%

Purpose

Antiseptic

Uses ■ surface sanitizer to decrease germs

- recommended for repeated use
- for use when soap and water are not available

Warnings

For surface use only

Do not use ■ on open skin wounds

When using this product • do not get into eyes. In case of contact, rinse eyes thoroughly with water **Stop use and ask a doctor if** • irritation and redness develop • 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 • wet surface thoroughly with product and allow to dry without wiping

Other information ■ store between 15-30°C (59-86°F)

avoid freezing and excessive heat above 40°C (104°F)

Inactive ingredients • Propylene glycol (plasticizer) • Laureth-4 (Surfactant) • Water • Glycolic acid • 1-Octadecanaminium NN dimethyl (3-trihydroxysily) propyl chloride • Dimethyloctadecyl (3-trimethoxysilyl) Propyl Ammonium Chloride

Questions? +1-312-803-1738

You may also report serious side effects to this phone number.

EXTEN-DAY

24 HOUR

Up to 24 hours continuous protection

Kills 99.99% of germs*

Bonds to most surfaces

Fast Acting 15 Second Formula

*Effective at eliminating more than 99.99% of many common harmful germs and bacteria in as little as 15 seconds; it forms a non-toxic barrier with lasting effects for up to 24 hours and does not produce the harmful, dermatological effects associated with traditional, high alcohol based hand sanitizers Manufactured for: Meroven LLC - 7901 4th St. N STE 300,St.

Petersburg, FL., 33702 Country of origin: Spain

Packaging



Drug Facts Active ingredient Purpose Benzalkonium Chloride......0.13 % Antiseptic Didecyl dimethyl ammonium chloride... 0.15% Antiseptic Uses • surface sanitizer to decrease germs recommended for repeated use · for use when soap and water are not available Warnings For surface use only Do not use • on open skin wounds. When using this product . do not get into eyes. In case of contact, rinse eyes thoroughly with water Stop use and ask a doctor if . irritation and redness develop • 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 • wet surface thoroughly with product and allow to dry without wiping Other information • store between 15-30°C (59-86°F avoid freezing and excessive heat above 40°C (104°F) Inactive ingredients - Propylene glycol (plasticizer) · Laureth-4 (Surfactant) · Water · Glycolic acid 1-Octadecanaminium NN dimethyl (3-trihydroxysily) propyl chloride . Dimethyloctadecyl (3-trimethoxysilyl) Propyl Ammonium Chloride Questions?+1-312-803-1738 You may also report serious side effects to this phone number. Mon-Fri 9:00 AM - 5:00 PM *Effective at eliminating more than 99.99% of many common harmful germs and bacteria in as little as 15 seconds; it forms a

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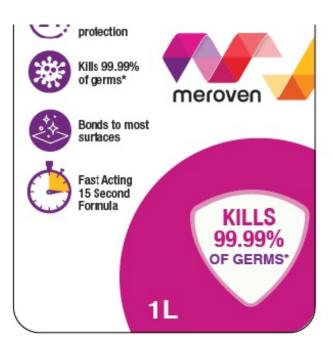
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benzalkonium chloride, didecyl dimethyl ammonium chloride liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:74653-104
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Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength	
BENZALKO NIUM CHLO RIDE (UNII: F5UM2KM3W7) (BENZALKO NIUM - UNII:7N6 JUD5X6 Y)	BENZALKONIUM CHLORIDE	1.3 mg in 1 mL	
DIDECYLDIMO NIUM CHLO RIDE (UNII: JXN40 O 9 Y 9 B) (DIDECYLDIMO NIUM - UNII: Z7F472 XQPA)	DIDECYLDIMONIUM CHLORIDE	1.5 mg in 1 mL	

Inactive Ingredients

Ingredient Name	Strength
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
LAURETH-4 (UNII: 6 HQ855798J)	
WATER (UNII: 059QF0KO0R)	
GLYCOLIC ACID (UNII: 0 WT12SX38S)	
O CTADECYLDIMETHYL(3-TRIHYDRO XYSILYLPRO PYL) AMMO NIUM CHLO RIDE (UNII: GLJ50 K8 6 6 T)	
DIMETHYLO CTADECYL(3-(TRIMETHO XYSILYL)PRO PYL)AMMO NIUM CHLO RIDE (UNII: IQ36 O 85 WQ4)	

F	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:74653-104-21	80 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/23/2020		
2	NDC:74653-104-22	500 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/23/2020		
3	NDC:74653-104-23	1000 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/23/2020		
4	NDC:74653-104-24	3000 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/23/2020		
5	NDC:74653-104-25	5000 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/23/2020		

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part333A	06/23/2020		

Labeler - Meroven LLC (117473235)

Revised: 6/2020 Meroven LLC