VIROCLEAR- benzalkonium chloride liquid
Bioinnovate Pty Ltd

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 0.12% w/w.

Purpose

Antiseptic, Hand Sanitizer

Uses

- Hand sanitizer to decrease bacteria on the skin.
- For use when soap and water are not available.

Warnings

For external use only.

When using this product

When using this product, keep out of eyes, ears and mouth. In case ofcontact with eyes, rinse eyes thoroughly with water.

Stop use and ask a doctor

Discontinue use if irritation and redness develops.

Keep out of reach of children

In case of accidental ingestion, seek medical attention or contact a poison control centre immediately.

Directions

Apply 2-3 sprays onto dry hands and rub hands together vigorously.

Other information

Store in a cool dry place below 104º F.

Inactive Ingredients

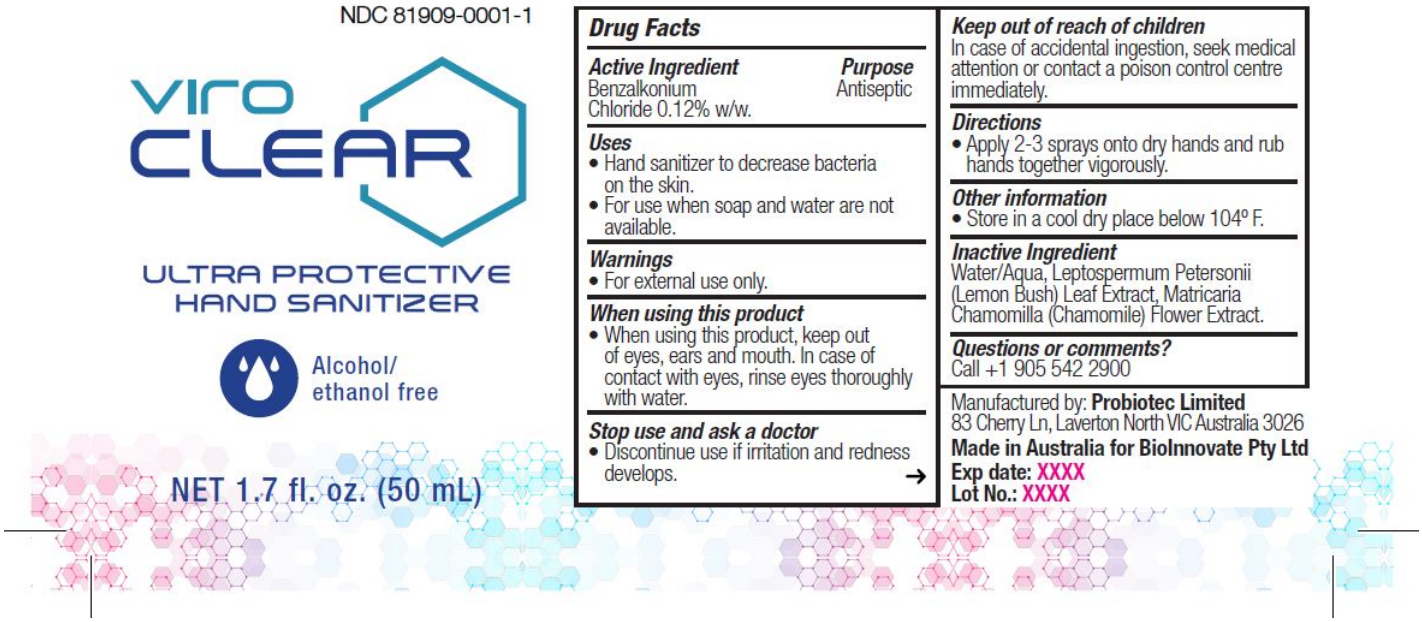
Water/Aqua, Leptospermum Petersonii (Lemon Bush) Leaf Extract, Matricaria Chamomilla (Chamomile) Flower Extract.

Questions or comments

Call +1 905 542 2900

Package Label - Principal Display Panel

50 mL; NDC 81909-0001-1



VIROCLEAR

benzalkonium chloride liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:81909-0001
Route of Administration	TOPICAL		

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

Inactive Ingredients	
Ingredient Name	**Strength**
LEPTOSPERMUM PETERSONII LEAF OIL (UNII: N37UWG52T3)	

CHAMOMILE (UNII: FGL3685T2X)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:81909-0001-1	50 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	06/07/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	06/07/2021	

Labeler - Bioinnovate Pty Ltd (747092416)

Revised: 6/2021

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