OVATION INSTANT HAND SANITIZER- benzethonium chloride liquid
QuestVapco Corporation

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.

Drug Facts

Active Ingredient

Benzenethonium chloride USP 0.2%

Purpose

Sanitizer

Uses

Handwash to help decrease bacteria on the skin.

Recommended for repeated use.

Warnings

For external use only

Do Not Use

In the eyes.

When using this product

Do not get in eyes. If contact occurs, flush eyes with water and contact a doctor immediately. If swallowed, seek medical attention.

Stop use and ask a doctor if

Skin irritation develops.

Keep out of reach of children.
Children under 6 years old should be supervised by an adult when using this product.

Inactive ingredients
Water, glycerin, dimethicone, DMDM Hydantoin, isdopropynyl butylcarbamate, fragrance, hydroxyethyl cellulose

Directions

Press dispenser twice to deliver two squirts (about a quarter size) of foaming product onto the palm of your hand. Rub hands together until hands are dry. Wash with soap and water at earliest opportunity.

OVATION INSTANT HAND SANITIZER

benzethonium chloride liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:67858-001
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
Benzethonium chloride (UNII: PH41D05744) (Benzethonium - UNII:1VU15B70BP)	Benzethonium chloride	1 mg in 50 mL

Inactive Ingredients	
Ingredient Name	**Strength**
Water (UNII: 059QF0KO0R)	

Hydroxypropyl cellulose (UNII: RFW2ET671P)	
Glycerin (UNII: PDC6A3C0OX)	
Dimethicone (UNII: 92RU3N3Y1O)	
dmdm hydantoin (UNII: BYR0546TOW)	
Sodium Hydroxide (UNII: 55X04QC32I)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:67858-001-50	50 mL in 1 BOTTLE, PUMP		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	04/01/2011	

Labeler - QuestVapco Corporation (103840377)

Registrant - QuestVapco Corporation (103840377)

Establishment

Name	Address	ID/FEI	Business Operations
QuestVapco Corporation		103840377	manufacture

Revised: 6/2011 QuestVapco Corporation