

ANGEL OF MINE FOAMING HAND SANITIZER- benzalkonium chloride liquid
Greenbrier International, Inc

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.

angel of mine Foaming Hand Sanitizer

Active ingredient

Benzalkonium chloride 0.13%

Purpose

Antiseptic

Use

to decrease bacteria on the skin

Warnings

For external use only, hands only

When using this product

■ avoid contact with the eyes. If contact occurs, rinse eyes thoroughly with water ■ avoid contact with broken skin

Stop use and ask a doctor if

■ irritation or redness develops ■ 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 hands thoroughly with product and allow to dry without wiping ■ for children under 6, use only under adult supervision

Inactive ingredients

water, cetrimonium chloride, disodium cocoamphodiacetate, PEG-12 dimethicone, citrus reticulata X C sinensis peel extract, tocopheryl acetate, methylchloroisithiazolinone, methylisothiazolinone

Package Label



ANGEL OF MINE FOAMING HAND SANITIZER

benzalkonium chloride liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:33992-1450
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.13 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
CETRIMONIUM CHLORIDE (UNII: UC9PE95IBP)	
WATER (UNII: 059QF0KO0R)	
DISODIUM CO CO AMPHODIACETATE (UNII: 18L9G3U51M)	
.ALPHA.-TO COPHEROL ACETATE (UNII: 9E8X80D2L0)	
PEG-12 DIMETHICONE (UNII: ZEL54N6W95)	
TANGERINE PEEL (UNII: JU3D414057)	

METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)

METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:33992-1450-8	236 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	04/17/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	04/17/2020	

Labeler - Greenbrier International, Inc (610322518)

Revised: 4/2020

Greenbrier International, Inc