ANTIBACTERIAL SANITIZING AND MOISTURIZING HAND- benzalkonium chloride cream QBLUE TECHNOLOGIES, 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.

QBlue Hand Sanitizer Solution 50mL (83388-003-50)

Drug Facts - Active Ingredient

Benzalkonium Chloride 0.09%

Purpose

Antiseptic

Uses

Hand sanitizer to help reduce bacteria that potentially can cause disease.

Warnings

For external use only

Avoid contact with eyes.

In case of contact, rinse eyes thoroughly with water.

Do not Use

on open skin wounds.

If irritation develops

Discontinue use and consult a doctor.

Keep out of reach of children.

If swallowed, get medical help or contact poison control center right away.

Directions

Apply product thoroughly to hands. Rub hands together until absorbed. Do not wipe off or rinse. Apply to hands as needed.

Other Information

Avoid direct sunlight. Store below 50°C. Do not refrigerate.

Inactive Ingredients

Purified Water, Mineral Oil, Glyceryl Stearate, Stearic Acid, Fragrance, Cetearyl Alcohol, Acrylates /C10-30 Alkyl Acrylate, Glycerin, Sodium Hydroxide, Crosspolymer, Vitamin B, Vitamin E, Titanium Dioxide.

Questions or comment?

1-833-387-9949

PDP

NDC: 83388-001-15



Drug Facts (continued)

Warnings For external use only

Do not use on open skin wounds. **Avoid contact with eyes,** in case of contact, rinse eyes thoroughly with water.

If irritation develops, discontinue use and consult a doctor.

Keep out of reach of children

If swallowed, get medical help or contact a poison control center right away.

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ANTIBACTERIAL SANITIZING AND MOISTURIZING HAND

benzalkonium chloride cream

Prod	uct	Inform	ation

Product Type HUMAN OTC DRUG Item Code (Source) NDC:83388-003

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.9 mg in 1 mL

Inactive Ingredients			
Ingredient Name	Strength		
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)			
GLYCERIN (UNII: PDC6A3C0OX)			
SODIUM HYDROXIDE (UNII: 55X04QC32I)			
.ALPHATOCOPHEROL (UNII: H4N855PNZ1)			
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)			
GLYCERYL STEARATE SE (UNII: FCZ5MH785I)			
STEARIC ACID (UNII: 4ELV7Z65AP)			
MINERAL OIL (UNII: T5L8T28FGP)			
WATER (UNII: 059QF0KO0R)			
CARBOMER COPOLYMER TYPE A (UNII: 71DD5V995L)			

F	Packaging						
#	Item Code	Package Description	Marketing Start Date	Marketing End Date			
1	NDC:83388- 003-50	1 in 1 BOX	04/10/2023				
1		50 mL in 1 CONTAINER; Type 0: Not a Combination Product					

Marketing Information					
Application Number or Monograph Citation	Marketing Start Date	Marketing End Date			
part333A	04/10/2023				
	Application Number or Monograph Citation	Application Number or Monograph Marketing Start Citation Date			

Labeler - QBLUE TECHNOLOGIES, INC. (118928041)

Revised: 4/2023 QBLUE TECHNOLOGIES, INC.