SOFT FOAM HAND SANITIZER- benzalkonium chloride liquid Freedom Technologies 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.

Soft Foam Hand Sanitizer

Benzalkonium Chloride

Antibacterial

Antibacertial foaming hand cleaner. Use in daycare, nursing homes, restaurants, doctors offices, dental offices, schools and clinics

For external use only. Avoid contact with eyes. If contact occurs, rinse with water. Discontinue use if irritation or redness develops. If irritation persist for more than 72 hours, consult a physician.

KEEP OUT OF REACH OF CHILDREN. If swallowed, get medical help or contact a Poison control Center right away.

Ready to use without dilution. Dispense 1 pump of product onto plalm of hand and rub thoroughly over all surfaces of hands unitl dry.

Water, Decyl Glucoside, Glycerin, DMDM Hydantoin.



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SOFT FOAM HAND SANITIZER

Drug Facts			
Active ingredient[s] Benzalkonium Chloride 0.13%	Purpose Antiseptic		
Use as part of your daily routine when soap and water are not available, or s			
Warnings For external use only. • Keep out of reach of children. • Avoid eye contact. Should contact occur, rinse thoroughly with water. • If swallowed or irritation develops, discontinue use and consult a health ca	re professional.		
Directions • Place in palm full (5 grams) in one hand. Spread onto both hands to wrist a	nd rub into skin until dry.		HEALTH (SALUD)
Other information Store between 15-30C (59-86F) Avoid freezing and excessive heat above 40C (104F) 			FLAMMABILITY
Inactive ingredients Water (aqua), Decyl Glucoside, Glycerin, DMDM H	ydantoin		(INFLAMMABLE IDAD)
Manufactured by Freedom Technologies, LLC Paramount, CA 90723 Made in the USA			REACTIVITY (REACTIVATED)
			PERSONAL PROTECTION
			THESE HMIS RATINGS ARE SUPPLIED AT BUYERS REQUES SELLER ASSUMES NO OBLIGATION OR LIABILITY FOR THE ADVICE OR ASSISTANCE GIVEN OR RESULTS OBTAINED
FT082020		5 GALLC	SELLER ASSUMES NO OBLIGATION OR LIABILITY FOR THE ADVICE OR ASSISTANCE GIVEN OR RESULTS OBTAINED.
ot #:	Net Weight:		

SOFT FOAM HAND S	SANITIZER				
benzalkonium chloride liquid					
Product Information					
Product Type	HUMAN OTC DRUG	Item Code (Source)		NDC:80891-234	
Route of Administration	TOPICAL				
Active Ingredient/Active Moiety					
Ingredient Name Basis of Strength				Strength	
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)		BENZ ALKONIUM CHLORIDE	0.0013 mg in 0.0013 mg		
Inactive Ingredients					
Ingredient Name				Strength	
DECYL GLUCOSIDE (UNII: Z17H9	7EA6Y)				
GLYCERIN (UNII: PDC6A3C0OX)					

DN	MDM HYDANTOI	(UNII: BYR0546TOW)				
WATER (UNII: 059QF0KO0R)						
Pa	ackaging					
#	Item Code	Package Description	Marketing Start Date	Marketing End Date		
1	NDC:80891- 234-02	1 in 1 BOX	10/01/2020			
1		3780 mg in 1 BOTTLE; Type 0: Not a Combination Product				
Marketing Information						
	Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
OT fin	C monograph no al	part333A	10/01/2020			

Labeler - Freedom Technologies LLC (057382260)

Establishment					
Name	Address	ID/FEI	Business Operations		
Freedom Technologies LLC		057382260	manufacture(80891-234)		

Revised: 1/2022

Freedom Technologies LLC