M MARVEL SANI CARE MULTI FUNCTION DISINFECTANT LIQUID NO RINSE-benzalkonium chloride spray
GUANGZHOU TECHIN DEVELOPMENT CO., 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.

80107-003

Active Ingredient

Benzalkonium Chloride 0.02%

Purpose

Antiseptic

Use

For Daily and Surgical hand disinfection to reduce bacteria on hands

Warnings

For external use only.
Keep away from fires and high temperatures.

Do Not Use: On open skin wounds.

When using this product: Avoid eyes.
In case of contact with eyes, rinse eyes thoroughly with water.

Stop use If irritation or rash occurs. If condition persists, contact a health care practitioner.

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Directions

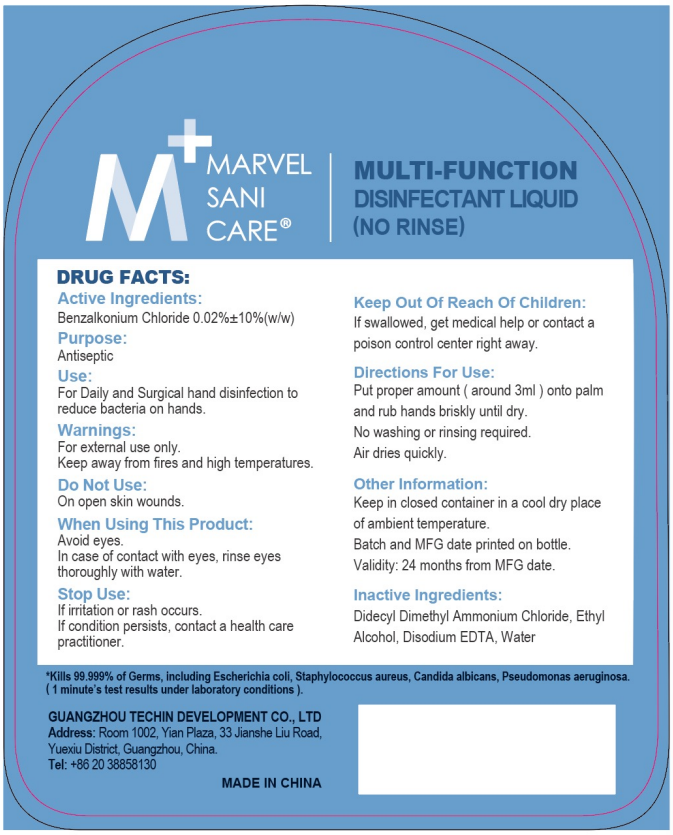
- Put proper amount (around 3ml) onto palm and rub hands briskly until dry.
- No washing or rinsing required. Air dries quickly.

Other information

Keep in closed container in a cool dry place of ambient temperature.
Batch and MFG date printed on bottle.
Validity: 24 months from MFG date.
MADE IN CHINA

Inactive ingredients

Didecyl Dimethyl Ammonium Chloride, Ethyl Alcohol, Disodium EDTA, Water



M MARVEL SANI CARE MULTI FUNCTION DISINFECTANT LIQUID NO RINSE

benzalkonium chloride spray

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:80107-003
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.02 g in 100 mL

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

ALCOHOL (UNII: 3K9958V90M)	
DISODIUM HEDTA (UNII: KME849MC7A)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:80107-003-01	60 mL in 1 BOTTLE; Type 0: Not a Combination Product	12/21/2020	
2	NDC:80107-003-02	250 mL in 1 BOTTLE; Type 0: Not a Combination Product	12/21/2020	
3	NDC:80107-003-03	500 mL in 1 BOTTLE; Type 0: Not a Combination Product	12/21/2020	
4	NDC:80107-003-04	750 mL in 1 BOTTLE; Type 0: Not a Combination Product	12/21/2020	
5	NDC:80107-003-05	1000 mL in 1 BOTTLE; Type 0: Not a Combination Product	12/21/2020	
6	NDC:80107-003-06	3785 mL in 1 BOTTLE; Type 0: Not a Combination Product	12/21/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	12/21/2020	

Labeler - GUANGZHOU TECHIN DEVELOPMENT CO., LTD (527219392)

Establishment

Name	Address	ID/FEI	Business Operations
GUANGZHOU TECHIN DEVELOPMENT CO., LTD		527219392	manufacture(80107-003)

Revised: 2/2022 GUANGZHOU TECHIN DEVELOPMENT CO., LTD