DISINFECTION WASH FREE HAND SANITIZER- sodium hypochlorite liquid Davemed Healthcare Co., Ltd.

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.

Disinfection Wash Free Hand Sanitizer

Active ingredient

Sodium hypochlorite

Purpose

Antiseptic

Uses

Hand sanitizer to help reduce bacteria on the skin recommended for repeated use.

Warnings

- Flammable
- Keep away from fire or flame
- For external use only

When using this product

- avoid contact with eyes in case of eye contact immediately flush eyes with water, call a doctor
- avoid contact with broken skin
- Discontinue use if irritation or redness develops. If condition persists from more than 72 hours, consult a doctor

Keep out of reach of children. Children should only use this product unde adult supervision. Do not drink. Not edible. In case of accidental ingestion, seek professional assistance or contact a Poison Control Center immediately.

Other information

- Do not store above 150F.
- May discolor some fabrics
- Harmful to wood finishes and plastics

Directions

• Place enough product in your palm to thoroughly spread on both hands and rub into the skin until dry. Recommended for repeated use.

Inactive ingredients

aloe barbadensis leaf extract, caprylyl glycol, carbomer, glycerin siopropyl alcohol, isopropyl alcohol myristate, Hyrochloric acid, sodium chloride

NDC: 70897-010



Disinfection
Wash Free
Hand Sanitizer

Contains:

sodium hypochlorite 120~160mg/L

KILLS 99.99% GERMS
WITHOUT WATER

04032020-5

8.5 FL OZ (250 ml)

Drug Facts

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Manufacturer in

Davemed Healthcare Co., Ltd

No.150 Guomao Street, Jinhua, Zhejiang, China www.davemedical.com Made in China

DISINFECTION WASH FREE HAND SANITIZER

sodium hypochlorite liquid

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:70897-010

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SODIUM HYPOCHLORITE (UNII: DY38 VHM5OD) (HYPOCHLORITE ION - UNII:T5UM7HB19N)	SODIUM HYPOCHLORITE	120 mg in 1000 mL

Inactive Ingredients	
Ingredient Name	Strength
CARBO MER HO MO PO LYMER, UNSPECIFIED TYPE (UNII: 0 A5MM307FC)	
ISO PRO PYL MYRISTATE (UNII: 0 RE8K4LNIS)	

CAPRYLYL GLYCOL (UNII: 00 YIU5438 U)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
GLYCERIN (UNII: PDC6A3C0OX)	
ALOE VERA LEAF (UNII: ZY8 1Z8 3H0 X)	
HYDRO CHLO RIC ACID (UNII: QTT17582CB)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	

]	Packaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70897-010-01	250 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/02/2020	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		04/02/2020	

Labeler - Davemed Healthcare Co., Ltd. (529128716)

Establishment				
Name	Address	ID/FEI	Business Operations	
Davemed Healthcare Co., Ltd		529128716	manufacture(70897-010)	

Revised: 4/2020 Davemed Healthcare Co., Ltd.