SUPPRESS HAND SANITIZER- alcohol gel
HD Group 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.

SUPPRESS Hand Sanitizer

Active Ingredient(s)

Alcohol 75% v/v. Purpose: Antiseptic

Purpose

Antiseptic, Hand Sanitizer

Use

to decrease bacteria on the skin.

Warnings

For external use only. Do not inhale or ingest. Flammable. Keep away from heat or flame when using this product.

Do not use

- in children less than 2 months of age
- on open skin wounds

When using this product keep away from eyes. In case of contact with eyes, flush thoroughly with water. Avoid contact with broken skin.

Stop use and ask a physician if skin irritation occurs.

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

Directions

- Apply to hands thoroughly and rub vigorously until dry.
- For children under 6 years, use only with adult supervision.
- Not recommended for infants.

Other information

- Do not store above 104F (40C)

- May discolor some fabrics
- harmful to some wood finishes

Inactive ingredients

purified water USP

PEG-40 hydrogenated castor oil

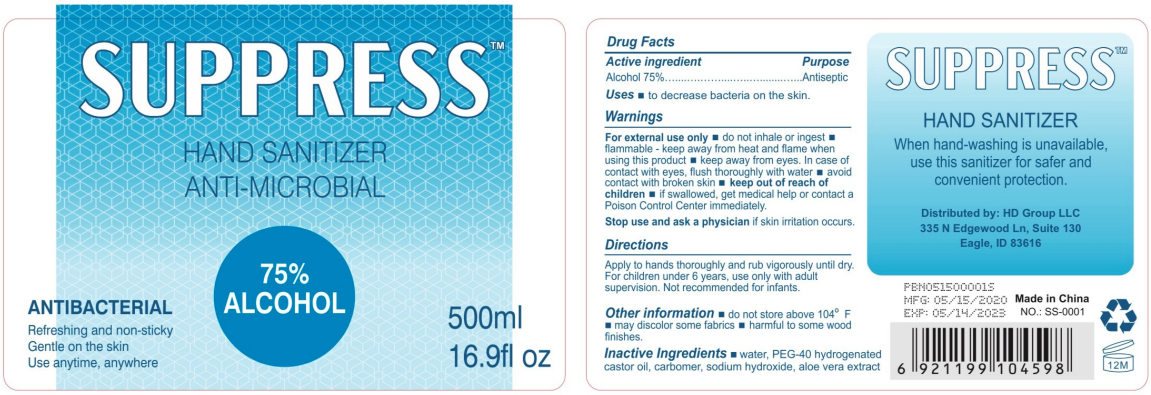
carbomer

sodium hydroxide

aloe vera extract

Package Label - Principal Display Panel

500 mL NDC: 77343-111-01



Distributed By:

Distributed by: HD Group LLC 335 N Edgewood Ln, Suite 130 Eagle, ID 83616

SUPPRESS HAND SANITIZER

alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:77343-111
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	75 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
PEG-40 CASTOR OIL (UNII: 4ERD2076EF)	
CARBOMER 940 (UNII: 4Q93RCW27E)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:77343-111-01	500 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/30/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	03/30/2020	

Labeler - HD Group LLC (833742898)

Revised: 1/2022 HD Group LLC