SUPPRESS HAND SANITIZING WIPES- alcohol cloth
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 Sanitizing Wipes

Active Ingredient(s)

Ethyl Alcohol 75% v/v

Purpose

Antiseptic

Use

- Hand Sanitizer to help decrease bacteria on the skin.
- Recommended for repeated use.

Warnings

For external use only. Flammable. Keep away from heat or flame

Do not use

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

When using this product keep out of eyes, ears, and mouth. In case of contact with eyes, rinse eyes thoroughly with water.

Stop use and ask a doctor if irritation or rash occurs. These may be signs of a serious condition.

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

Directions

- Thoroughly wipe hands, fingers and wrists. Be sure to use the entire wipe.
- Rub hands together until dry.
- For dirty hands, use first wipe to clean hands, then discard wipe. Sanitize with a second wipe.
- Discard after single use.
- Supervise children under 6 years of age when using this product to avoid swallowing.

Other information

- Store between 15-30C (59-86F)
- Avoid freezing and excessive heat above 40C (104F)
- Dispose of wipe in the proper container after use.
- Do not flush down the toilet.
- Lot No. and Expiration Date can be found on canister.

Inactive ingredients

purified water USP

Distributed By

Distributed by:

HD Group LLC

335 N Edgewood Ln, Suite 130

Eagle, ID 83616

SUPPRESS Hand Sanitizing Wipes- 150 Wipes Label

150 Wipes

NDC: 77343-101-01

SUPPRESS Hand Sanitizing Wipes- 75 Wipes Label

75 Wipes

NDC: 77343-101-02

SUPPRESS HAND SANITIZING WIPES

alcohol cloth

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:77343-101
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**
WATER (UNII: 059QF0KO0R)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:77343-101-01	612 mL in 1 CANISTER; Type 0: Not a Combination Product	06/01/2020	
2	NDC:77343-101-02	364 mL in 1 CANISTER; Type 0: Not a Combination Product	06/01/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	06/01/2020	

Labeler - HD Group LLC (833742898)

Revised: 1/2022 HD Group LLC