# 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.

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#### **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**

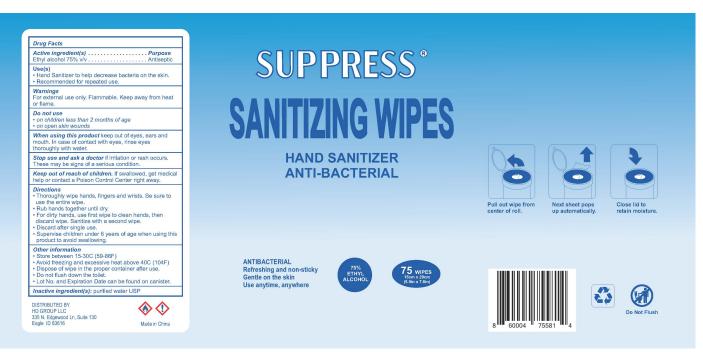
150 Wipes

NDC: 77343-101-01



## **SUPPRESS Hand Sanitizing Wipes-75 Wipes Label**

NDC: 77343-101-02



## **SUPPRESS HAND SANITIZING WIPES**

alcohol cloth

<b>Product Information</b>			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:77343-101
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	<b>Basis of Strength</b>	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	75 mL in 100 mL

<b>Inactive Ingredients</b>		
	Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)		

P	ackaging			
#	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			
ing End te	Mari	Marketing Start Date	Application Number or Monograph Citation	Marketing Category
		06/01/2020	part333A	OTC monograph not final
		06/01/2020	part333A	

## Labeler - HD Group LLC (833742898)

Revised: 1/2022 HD Group LLC