

**ANTISEPTIC WIPES- alcohol, benzalkonium chloride cloth
D&A USA, 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.

Antiseptic Wipes

Active Ingredients

Ethyl alcohol 15%. v/v

Benzalkonium Chloride 0.1%. v/v

Purpose

Antiseptic

Warnings

- For external use only.

Do not use

on children less than three years old and on skin wounds.

When using this product

Keep out of eyes. In case of contact, rinse eyes thoroughly with water. Avoid contact with broken skin.

Stop use and ask a doctor if

irritation or rash appears and lasts.

Keep out of reach of children

If swallowed, get medical help or contact a Poison Control Center right away.

Directions

- Open lid, gently pull back resealable label, remove and use wipe as required.
- Reseal back after use to avoid evaporation of alcohol.
- Discard after single use Required dispose of used wipe in trash receptable.
- Do not flush.

Uses

- For hand washing to decrease bacteria on the skin.
- Apply topically to the skin to help prevent cross contamination.

Other information

Store at room temperature, avoid sunlight irradiating.

Inactive ingredients

Water, Cetylpyridinium chloride, Chlorphenesin, Biguanide, Propylene glycol

Package Label - Principal Display Panel

Drug Facts

Active ingredients

- Ethyl alcohol 15% v/v Antiseptic
- Benzalkonium Chloride 0.1% v/v Antiseptic

Purpose

- Antiseptic
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D&A USA™
EASY STYLISH SOLUTION



CONTAINS
75% ALCOHOL  STOSTE
ANTISEPTIC WIPES
for hand cleaning

72 Wipes
7.87 x 5.51in (20 x 14cm)

contains
**75%
ALCOHOL**



D&A USA

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Product Name: D&A USA Antiseptic Wipes
Sanitary Certification: CN-2016-02-0008
Sanitary Standard: WS575-2017
Executive Standard: GB/T27728
Application: Cleaning Skin / Hand wipes
MFG: Printing On Bag/Bottle

Developed in USA
Made in China

#20D4008



THIS IS
NOT A
BABY WIPE



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ANTISEPTIC WIPES

alcohol, benzalkonium chloride cloth

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:80956-001
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	15 mL in 100 g
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.1 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
BIGUANIDE (UNII: FB4Q52I9K2)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
CETYLPIRIDINIUM CHLORIDE (UNII: D9OM4SK49P)	
CHLORPHENESIN (UNII: I670DAL4SZ)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:80956-001-01	72 in 1 BAG	10/28/2020	
1		5.6 g in 1 PATCH; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	10/28/2020	

Labeler - D&A USA, LLC (117661331)