JANITEX DISINFECTING WIPES 75% ALCOHOL- ethyl alcohol cloth
Pacific Health Systems Inc.

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.

JANITEX DISINFECTING ALCOHOL WIPES

Drug Facts

Active Ingredients

Ethyl Alcohol 75 %

Purpose

Antiseptic

Uses

- To decrease bacteria on the skin that could cause disease

Warnings

- **For external use only**, harmful if swallowed.
- **Do not use** as baby wipe.
- **Do not flush.**
- **Do not use** on open skin wounds.
- **Do not use** in or near the eyes. In case of contact, rinse eyes thoroughly with water.
- **Stop use and ask a doctor** if irritation or allergic reaction occurs.
- **Flammable - Keep away from fire or flame**

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

Directions

• Open container plastic cover. Tear off moisture seal film. Pull out wipe corner from the center of wipe roll; twist wipe corner through the dispensing center of the canister. Next sheets pop up automatically. Keep center cap closed to prevent moisture loss. • No need to rinse after usage. Allow skin to dry without wiping. For adults and children 2 years and over. • Discard properly after use. • Supervise children under 6 years of age when using this product to avoid swallowing.

Other Information

- Store in dry and cool place and away from oxidizers.

Inactive ingredients

Aqua, Glycerin, Propylene Glycol, Parfum, Tocopheryl Acetate, Citric Acid, Aloe Barbadensis Extract

75% Alcohol

75% Alcohol Concentration

E Vitamin

+ moisturizer

non-woven fabric

Bleach Free

Ultra Protection

DISINFECTING WIPES

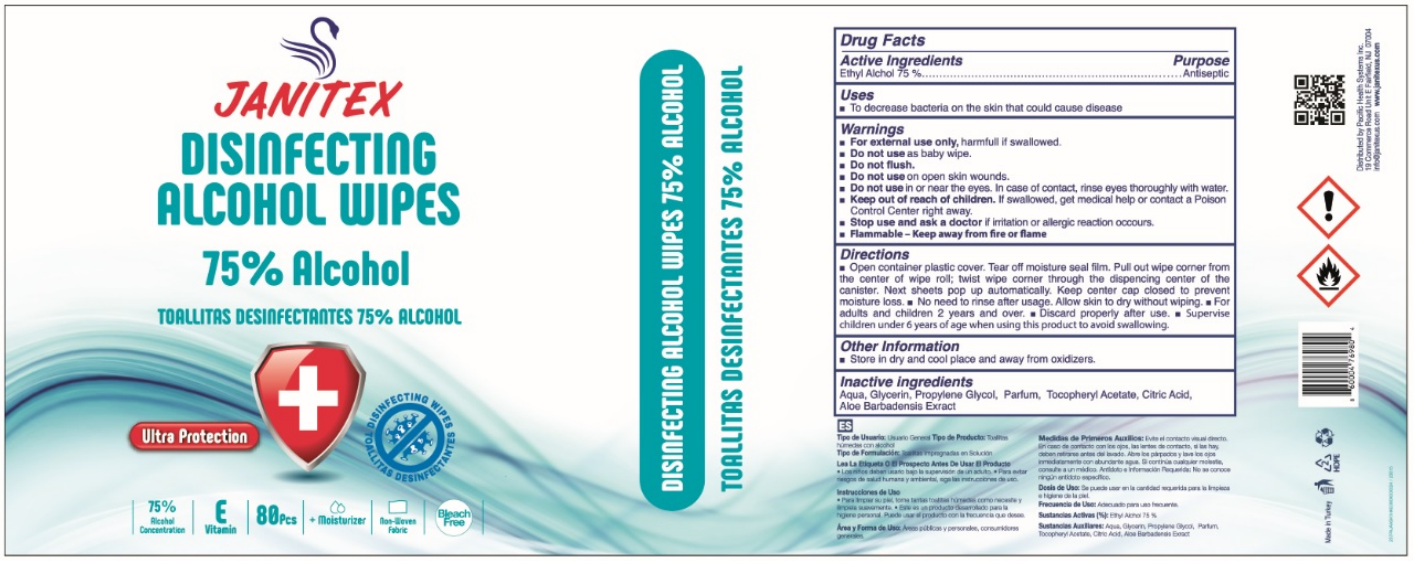
Distributed by Pacific Health Systems Inc.

19 Commerce Road Unit E Fairfield, NJ 07004

info@janitexus.com **www.janitexus.com**

Made in Turkey

Packaging



JANITEX DISINFECTING WIPES 75% ALCOHOL

ethyl alcohol cloth

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:79519-060
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	75 g in 100 g

Inactive Ingredients	
Ingredient Name	**Strength**

WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:79519-060-01	80 in 1 CANISTER	08/11/2020	
1		1.83 g in 1 PACKAGE; 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	08/11/2020	

Labeler - Pacific Health Systems Inc. (080638960)

Revised: 8/2020 Pacific Health Systems Inc.