JANITEX 75% ALCOHOL WET WIPES- 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 75% ALCOHOL WIPES

Drug Facts

Active Ingredient

Ethyl Alcohol 75%

Purpose

Antiseptic

Uses

- For hand sanitizing to decrease bacteria and germs on skin.
- Apply topically to the skin to help prevent cross contamination.
- Dries in seconds.

Warnings

- For external use only. Harmful if swallowed.
- Do not use as baby wipes.
- Do not flush down the toilet.
- Do not use in or contact the eyes.

Do Not Use

- Highly flammable. Keep away from fire or flame.
- Do not eat or drink when handling
- Avoid contact with open or exposed wounds.

When Using

Keep out of eyes, ears, and mouth.

In case of contact with eyes, rinse eyes thoroughly with water

Stop Use

Stop use if too much skin irritation and sensitivity develops or increases

KEEP OUT OF REACH OF CHILDREN

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

Directions

- Lift the tab on the top of the lid to open.
- Open moisture seal film package before use.
- Pull out first wipe through the dispensing center of the canister.

Next sheets pops up automatically

- Close the lid to retain moisture.
- Wipe clean with this product. Let air dry

Other Informations

- Store at room temperature 15-30 (59-86°F).
- Lot No. Manufacture date and expiration date can be found on package.

Inactive Ingredients Aqua, Glycerin, Parfum

ANTIVIRUS

ANTI - FUNGAL

Kills 99.9% Bacteria & Germs

FDA

75% Alcohol Concentration

99.9% Efficient Sterilization

Non-Woven Skin

Moisturizes and Doet Not Dry

BLEACH FREE

Made in Turkey

Distributed by Pacific Health Systems Inc.

19 Commerce Road Unit E Fairfield, NJ 07004

info@janitexus.com www.janitexus.com

Packaging















Drug Facts













DRUG FACTS LABEL

Drug Facts

Active Ingredient

Ethyl Alcohol 75%

Purpose

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Drug Facts (continued)

When Using

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Drug Facts (continued)

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Tipo de Usuario: Usuario General
Tipo de Producto: Toallitas húmedas con alcohol
Tipo de Formulación: Toallitas Impregnadas en Solución
LEA LA ETIQUETA O EL PROSPECTO ANTES DE USAR EL PRODUCTO.

- Los niños deben usarlo bajo la supervisión de un adulto.
 Para evitar riesgos de salud humana y ambiental, siga las
- instrucciones de uso.

- Para limpiar su piel, tome tantas toallitas húmedas como necesite y limpiela suavemente.
 Este es un producto desarrollado para la higiene personal.
- Puede usar el producto con la frecuencia que desee. La vida útil es de 2 años cuando se almacena a temperatura ambiente y lejos de la luz solar.

La vida útil es de 3 meses después de abrir el paquete.

Sintomas de Envenenamiento: Sin síntomas. ALCOHOL DESNATURALIZADO (DENAT) CAS No: 64-17-5 LD50 (oral, rata): 7060 mg/kg ALCOHOL ISOPROPILICO CAS No: 67-63-0 LD50

(oral, rata): 5280 mg/kg

Medidas de Primeros Auxillos: Evite el contacto visual directo. En caso
de contacto con los ojos, las lentes de contacto, si las hay, deben retirarse
antes del lavado. Abre los párpados y lave los ojos inmediatamente con
abundante agua. Si continúa cualquier molestia, consulte a un médico.
Antidoto e Información Requerida: No se conoce ningún antidoto

Área y Forma de Uso: Áreas públicas y personales, consumidores

Dosis de Uso: Se puede usar en la cantidad requerida para la limpieza e

ngiene de la piel. Freeuencia de Uso: Adecuado para uso frecuente. Sustancias Activas (%): ALCOHOL DESNATURALIZADO (DENAT) CAS No: 64-17-5 (75,00%) (en volumen), ALCOHOL ISOPROPÍLICO CAS No:

67-63-0 (3,00%) (en volumen) Sustancias Auxiliares: AQUA, GLICERINA, PARFUM

JANITEX 75% ALCOHOL WET WIPES

ethyl alcohol cloth

Product Information Product Type HUMAN OTC DRUG Item Code (Source) NDC:79519-020 **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)					
GLYCERIN (UNII: PDC6A3C0OX)					

P	Packaging						
#	Item Code	Package Description	Marketing Start Date	Marketing End Date			
1	NDC:79519-020- 03	40 in 1 PACKET	06/30/2020				
1		4.16 mL in 1 PACKAGE; Type 0: Not a Combination Product					
2	NDC:79519-020- 02	60 in 1 BOX	06/30/2020				
2		4.16 mL in 1 PACKAGE; Type 0: Not a Combination Product					
3	NDC:79519-020-01	80 in 1 CANISTER	06/30/2020				
3		4.16 mL in 1 PACKAGE; Type 0: Not a Combination Product					
4	NDC:79519-020- 04	80 in 1 CANISTER	09/01/2020				
4		4.16 mL 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	06/30/2020			

Labeler - Pacific Health Systems Inc. (080638960)

Revised: 9/2020 Pacific Health Systems Inc.