

WIPESPLUS HAND SANITIZING WIPES- ethyl alcohol cloth
Progressive Products, 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.

Drug Facts

Active Ingredient

Ethyl Alcohol 70%

Purpose

Antiseptic

Uses

- For hand sanitizing to decrease bacteria on the skin.
- Apply topically to the skin to help prevent cross contamination
- Not recommended for repeated use.
- Dries in seconds.

Warnings

For external use only.

Flammable, keep away from fire or flame.

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Do not use

- in or near the eyes
- on open skin wounds or burns.

Stop use and ask a doctor

- if irritation and redness develop
- condition persists for more than 72 hours.

Keep out of reach of children

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

Directions

- Remove lid.
- Pull wipe from center of roll and thread through opening in lid. Do not push finger through opening.
- Replace lid, pull wipe up, and then at 45° angle. The next wipe dispenses automatically.
- Close lid to retain moisture.

Other Information

- Store at room temperature.
- Dispose of used wipe in trash, do not flush into toilet.

Inactive Ingredients

Aloe Barbadensis Extract, Benzalkonium Chloride, Citric Acid, Fragrance, Glycerin, Propylene Glycol, Tocopheryl Acetate, Water

Principal Display Panel

wipesplus

HAND SANITIZING WIPES

- Kills 99.9% of germs

240 CT

DRUG FACTS (CONTINUED)

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INACTIVE INGREDIENTS

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OTHER INFORMATION: Lot No. and Expiration Date can be found on canister.

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Distributed by:

PROGRESSIVE PRODUCTS
211 S. Ridge Street, Dye Brook, NY 10573
Call 1-800-477-6022
Re-Order / Item No. 33984
www.wipesplus.com

Do Not Flush 240 CT PRE-MOISTENED TOWELETTES 6" x 6.5" (16.2 cm x 17.1 cm)

- Kills 99.9% of germs

WIPESPLUS HAND SANITIZING WIPES

ethyl alcohol cloth

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:67151-737
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	0.7 mL

Inactive Ingredients

Ingredient Name	Strength
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
WATER (UNII: 059QF0KO0R)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
GLYCERIN (UNII: PDC6A3C0OX)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:67151-737-01	240 in 1 PACKAGE; Type 0: Not a Combination Product	07/13/2022	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	07/13/2022	

Labeler - Progressive Products, LLC (127111792)

Revised: 10/2022

Progressive Products, LLC