ALCOHOL WET WIPES- alcohol cloth Robert Gordon Ind. Ltd.

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.

Alcohol Wet Wipes

Drug Facts

Active ingredient

Ethyl alcohol 75% (v/v)

Purpose

Antiseptic

Uses

- to decrease bacteria on the skin that could cause disease
- recommended for repeated use

Warnings

- For external use only
- Flammable
- Keep away from fire or flame

Do not use

- on children less than 2 months old
- on open skin wounds

When using this product

- keep out of eyes, ears, In case of contact with eyes, flush thoroughly with water.
- avoid contact with broken skin
- do not inhale or ingest

Stop use and ask a doctor if

- irritation or redness develops
- 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

- Place enough product on hands to cover all surfaces.
- Rub hands together until dry
- Supervise children under 6 years of age when using this product to avoid swallowing.

Other information

- Do not store above 105°F
- May discolor some fabrics
- Harmful to wood finishes and plastics

Inactive ingredients

aminomethyl propanol, carbomer homopolymer, glycerin, propylene glycol, purified water USP

Package Labeling:50count



Package Labeling:100count

ALCOHOL WET WIPES





100 sheets













ALCOHOL WET WIPES

alcohol cloth

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:74979-005

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name Basis of Strength Strength ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M) ALCOHOL 0.75 mL in 1 mL

Inactive Ingredients

Ingredient Name	Strength
AMINO METHYLPRO PANOL (UNII: LU49 E6626Q)	
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0 A5MM307FC)	
GLYCERIN (UNII: PDC6A3C0OX)	

WATER (UNII: 059QF0KO0R)

PROPYLENE GLYCOL (UNII: 6DC9Q167V3)

Packaging

# Item Code	Marketing Start		Marketing End Date
1 NDC:74979- 005-01	50 in 1 CANISTER	08/20/2020	Dute

1		3.56 mL in 1 PATCH; Type 2: Prefilled Drug Delivery Device/System (syringe, patch, etc.)		
2	NDC:74979- 005-02	100 in 1 CANISTER	08/20/2020	
2		3.56 mL in 1 PATCH; Type 2: Prefilled Drug Delivery Device/System (syringe, patch, etc.)		

Marketing Information					
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
OTC monograph not final	part333E	08/20/2020			

Labeler - Robert Gordon Ind. Ltd. (070064709)

Revised: 9/2020 Robert Gordon Ind. Ltd.