TARGOL SANITIZING- ethyl alcohol aerosol, spray Endeks Kimya Sanayi Ve Ticaret Anonim Sirketi

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.

TARGOL Sanitizing SPRAY EUCALYPTUS

Drug Facts

Active ingredient

Ethyl alcohol 75%

Purpose

Antiseptic

Uses

- hand sanitizer to decrease bacteria on the skin
- recommended for repeated use
- for use when soap and water are not available

Warnings

- Extremely flammable aerosol, keep away from fire / flame.
- Keep away from heat, hot surfaces, open flames and other ignition sources.
- Pressurized container. Do not pierce or burn even after use.
- Protect from the sunlight.
- For external use only.

Do not use

- on children less than 2 months of age
- on open skin wounds

When using this product • Avoid use on/or around eyes, ears, mouth, broken/irritated skin or large areas of body. In case of contact with eyes, rinse thoroughly with water several minutes. • do not inhale or ingest.

Stop and ask doctor if

- irritation persist
- or if product is swallowed

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

Directions

- Shake well before use.
- Hold can upright at 6-7 inches away from surface and spray evenly.
- Allow to air dry for 5 to 10 minutes. Repeat application a necessary
- Supervise children under 6 years of age when using this product to avoid swallowing

Other information

- store between 15-30°C (59-86°F)
- Do not expose to temperatures exceeding 50 C/122 F.

Inactive ingredients

Butane, Propane, Isobutane, Parfum, Glycerin

Kills 99.9% of Most Common Germs*

EUCALYPTUS

ALCOHOL CONTENT 75 %

Sanitizer Spray

*Effective at eliminating 99.9% of many common harmful germs and bacteria in as little as 15 seconds

You may also report any serious side effects to P O Box 328, Wood Ridge, NJ 07075-328

Packaging







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TARGOL SANITIZING

ethyl alcohol aerosol, spray

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:78089-035

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength	
ı	ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	75 mL in 100 mL

Inactive Ingredients	
Ingredient Name	Strength
BUTANE (UNII: 6LV4FOR43R)	
PROPANE (UNII: T75W9911L6)	
ISOBUTANE (UNII: BXR49 TP6 11)	
GLYCERIN (UNII: PDC6A3C0OX)	

P	Packaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
	NDC:78089-035- 01	550 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	09/07/2020	

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part333A	09/07/2020		

Labeler - Endeks Kimya Sanayi Ve Ticaret Anonim Sirketi (565678914)

Establishment			
Name	Address	ID/FEI	Business Operations
Endeks Kimya Sanayi Ve Ticaret Anonim Sirketi		565678914	manufacture(78089-035)

Revised: 8/2020 Endeks Kimya Sanayi Ve Ticaret Anonim Sirketi