WISH ULTRA ALCOHOL SANITIZING SPRING WATERFALL- ethyl alcohol spray Click 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.

WISH ULTRA ALCOHOL SANITIZING SPRAY: SPRING WATERFALL

Active Ingredients

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

- -Flammable.Keep away from heat and flame/fire
- -Keep away from heat, hot surfaces, open flames and ignition sources
- -Pressurized container. Do not pierce or burn even after use
- -Protect from sunlight
- -For external use only
- -On children less than 2 months of age
- -On open skin wounds

-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
- -irritation persist
- -if product is swallowed

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 the 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.

-store between 15-30 degree Celcius

-Do not expose to temperature exceeding 50 degree celcius

Ethanol, Butane, Propane, Isobutane, Parfum.

Label



WISH ULTRA ALCOHOL SANITIZING SPRING WATERFALL

ethyl alcohol spray

Product Information					
Product T ype	HUMAN OTC DRUG	Item Code (Source)	NDC:71611-022		
Route of Administration	TOPICAL				

	Active Moiety			
Ingredient Name		Basis of Strength		Strength
ALCOHOL (UNII: 3K99	58 V90M) (ALCOHOL - UNII:3K9958 V90M)	ALCOHOL		75 mL in 100 mL
Inactive Ingredien	ts			
			Strength	
ISOBUTANE (UNII: BXR	249 TP6 11)			
PROPANE (UNII: T75W9	911L6)			
BUTANE (UNII: 6LV4FO	D/3D)			
	N45K)			
Packaging # Item Code	Package Description	Marketing Star	rt Date	Marketing End Da
# Item Code			rt Date	Marketing End Da
# Item Code 1 NDC:71611-022-55 55	Package Description 50 mL in 1 BOTTLE; Type 0: Not a Combination Product		't Date	Marketing End Da
 # Item Code 1 NDC:71611-022-55 55 Marketing Info 	Package Description 50 mL in 1 BOTTLE; Type 0: Not a Combination Product rmation	09/26/2020		
# Item Code 1 NDC:71611-022-55 55	Package Description 50 mL in 1 BOTTLE; Type 0: Not a Combination Product rmation Ymation Application Number or Monograph Citation			Marketing End Da Marketing End Da

Labeler - Click Products LLC (080766174)

Registrant - Click Products LLC (080766174)

Establishment					
Name	Address	ID/FEI	Business Operations		
ENDEKS KIMYA SANAYI VE TICARET ANONIM SIRKETI		565678914	manufacture(71611-022)		

Establishment

Name	Address	ID/FEI	Business Operations
ATAK FARMA KOZMETIK VE KIMYA SANAYI TICARET ANONIM SIRKETI		566218248	manufacture(71611-022)

Revised: 9/2020

Click Products LLC