DEFENSE ZONE HAND SANITIZER- ethyl alcohol spray
Prime Packaging

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.

Defense Zone Sanitizer Spray

Active Ingredients

Ethyl Alcohol 70%

Purpose

Antiseptic

Uses

to decrease bacteria on the skin that could cause disease. recommrnded for repeated use

Warning

For external use only

Dot not use

Do not use on damaged or broken skin

When using this product

Keep away from face to avoid breathing it. Keep out of eyes. rinse with water to remove.

Stop use and ask a doctor

If irritation and redness occurs if condition persistes for more than 72 hours.

Keep out of reach of children

If product is swallowed, get medical help or contact a Posion Control Center right away.

Flammable

Avoid fire, flame, heat and smoking. Contents under pressre. do not punture or incinerate. Store at temperatures below 120°F(50°C).

Directions

wet hands thoroughly with product and allow to dry without wiping. for children under 6, use only under adult supervision. not recommended for infants.

Other Information

do not store above 104°F. may discolor some fabrics. harmful to wood finishes and plastics.

Inactive ingredients

Aloe Barbadensis Leaf Juice,Glycerin,Isopropyl Myristate, Tocopherol,Water.

Questions

Call 1 855-548-4653

Principal display label

DEFENSE ZONE HAND SANITIZER
ethyl alcohol spray

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:13630-0161
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	0.7 mL in 1 mL

Inactive Ingredients

Ingredient Name	Strength
TOCOPHEROL (UNII: R0ZB2556P8)	
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	

Product Characteristics

Color	white (Clear)	**Score**	
Shape		**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:13630-0161-4	177 mL in 1 CAN; Type 0: Not a Combination Product	09/21/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	09/21/2020	

Labeler - Prime Packaging (805987059)

Registrant - Prime Packaging (805987059)

Establishment

Name	Address	ID/FEI	Business Operations
Prime Enterprises		101946028	analysis(13630-0161) , manufacture(13630-0161)

Establishment

Name	Address	ID/FEI	Business Operations

Prime Packaging		805987059	label(13630-0161) , pack(13630-0161)