EAGLE PROMOTIONS- instant hand sanitizer gel gel
Nutrix International, 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.

Active Ingredient[s]

Alcohol 70% v/v

Use[s]

Hand Sanitizer to help reduce bacteria that potentially can cause disease. For use when soap and water are not available.

Warnings

For external use only. Flammable. Keep away from heat or flame.

Do not use

- On children less than 2 months of age.
- On open skin wounds.

When using

Keep out of eyes, ears, and mouth. In case of contact with eyes, rinse eyes thoroughly with water.

Stop use

If irritation or rash occurs. These may be signs of a serious condition.

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 avoid swallowing.

Other Information

- Store between 15-30°C (59-86°F)
- Avoid freezing and excessive heat above 40°C (104°F)

Inactive Ingredients

Water (Aqua), Propylene Glycol, Glycerin, Methyl Hydroxypropyl Cellulose, Melaleuca Alternifolia (Tea Tree) Leaf Oil.

Purpose

Antiseptic

Package

Full Package label

EAGLE PROMOTIONS

instant hand sanitizer gel gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:73333-284
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	0.74 g in 284 g

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	0.233 g in 284 g
TEA TREE OIL (UNII: VIF565UC2G)	0.002 g in 284 g
GLYCERIN (UNII: PDC6A3C0OX)	0.008 g in 284 g
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	0.01 g in 284 g
HYPROMELLOSES (UNII: 3NXW29V3WO)	0.007 g in 284 g

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73333-284-01	284 g in 1 BOTTLE; Type 0: Not a Combination Product	05/21/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	05/21/2020	

Labeler - Nutrix International, LLC. (117341868)

Establishment

Name	Address	ID/FEI	Business Operations
Nutrix International, LLC.		117341868	manufacture(73333-284) , pack(73333-284) , label(73333-284)

Revised: 5/2020 Nutrix International, LLC.