## HAND SANITIZER- alcohol gel Zink Holdings 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.

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## Zink Brntg Rhnstne Sanitizer Gel 75

### **Active Ingredient**

Alcohol 75%v/v

#### **Purpose**

Antiseptic

#### Uses

■ Hand Sanitizer to help reduce bacteria on the skin. 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 this product

keep out of eyes, ears, ears and mouth. In case of eye contact immediately flush eyes thoroughly with water

#### Stop use and ask a doctor

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

#### Keep out of reach of children

In case of accidentail ingestion, contact a doctor or Poison Control Center immediately.

#### 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

■ Store between 15°-30°C (59°-89°F) ■ Avoid freezing and excessive heat above 40°C (104°F) ■ May discolor fabrics or surfaces.

## **Inactive ingredients**

glycerin, hydroxypropyl cellulose, water

# **Package Label**



#### HAND SANITIZER

alcohol gel

Product Information	oduct Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:79332-001	
Route of Administration	TOPICAL			

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

Inactive Ingredients		
Ingredient Name	Strength	
WATER (UNII: 059QF0KO0R)		
HYDRO XYPRO PYL CELLULO SE, UNSPECIFIED (UNII: 9 XZ8 H6 N6 O H)		
GLYCERIN (UNII: PDC6A3C0OX)		

I	Packaging			
#	t Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:79332-001- 08	237 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/22/2020	
	NIDC -70222 001	EAR mi in 1 DOTTLE DI ACTIC. Type O. Not a Combination		

2	NDC:/9332-001- 16	Product	06/22/2020	
3	NDC:79332-001- 28	3790 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/22/2020	
4	NDC:79332-001- 32	946 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	09/14/2020	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	06/22/2020	

# Labeler - Zink Holdings LLC (117561710)

Revised: 9/2020 Zink Holdings LLC