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.

Zink Brntag Crayola Grn Sanitizer Gel Alc75

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

FD&C Blue 1, FD&C Yellow 5, Glycerin, Hydroxypropyl Cellulose, Water

Inner Package Label



2 FL OZ / 60 mL



Drug Facts

Active ingredient Purpose

Alcohol 75 % v/v Antiseptic

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Directions Place enough product on your hands to cover all surfaces. Rub hands together until dry.

Drug Facts (continued)

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ABOUT US

Crayola A Hallmark Company

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Official Licensed Product

QUALITY GUARANTEE

Skymall Holdings LLC, guarantees the quality of this Crayola product. If this product does not properly, please contact us at (1-848-244-2343)



Distributed by Skymall Holdings LLC Edison, NJ 08837



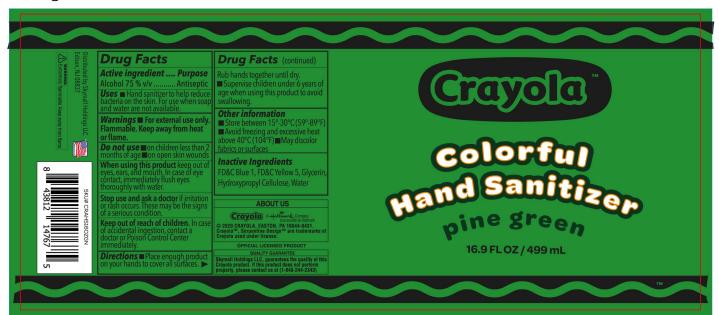
Outer Package Label



Package Label



Package Label



HAND SANITIZER

alcohol gel

Product Information							
Product Type	HUMAN OTC DRUG	Item Code	(Source)	NDC:79332-004			
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		
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)			
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)			
WATER (UNII: 059QF0KO0R)			
HYDROXYPROPYL CELLULOSE, UNSPECIFIED (UNII: 9XZ8H6N6OH)			
GLYCERIN (UNII: PDC6A3C0OX)			

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:79332-004- 08	8 in 1 BOX	08/21/2020	
1	NDC:79332-004- 01	$60\ mL$ in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
2	NDC:79332-004- 04	4 in 1 BOX	08/21/2020	
2	NDC:79332-004- 01	60 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
3	NDC:79332-004- 02	1 in 1 BOX	08/21/2020	
3	NDC:79332-004- 01	60 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
4	NDC:79332-004- 80	236 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	09/10/2020	
5	NDC:79332-004- 16	499 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	09/10/2020	

Marketing Information						
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date			
OTC monograph not final	part333A	08/21/2020				

Labeler - Zink Holdings LLC (117561710)

Revised: 9/2020 Zink Holdings LLC