

HAND SANITIZER- alcohol gel
Skymall 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 Skymall 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

Crayola™

Colorful
Hand Sanitizer
forest green

2 FL OZ / 60 mL



Drug Facts

Active ingredient Purpose

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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

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QUALITY GUARANTEE

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



SKU# CRAHS20ZGN



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:80230-004
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958 V90M) (ALCOHOL - UNII:3K9958 V90M)	ALCOHOL	75 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
FD&C BLUE NO. 1 (UNII: HBR47K3TBD)	
WATER (UNII: 059QF0K00R)	
HYDROXYPROPYL CELLULOSE, UNSPECIFIED (UNII: 9XZ8H6N6OH)	
GLYCERIN (UNII: PDC6A3C0OX)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:80230-004-08	8 in 1 BOX	08/21/2020	
1	NDC:80230-004-01	60 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
2	NDC:80230-004-04	4 in 1 BOX	08/21/2020	
2	NDC:80230-004-01	60 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
3	NDC:80230-004-02	1 in 1 BOX	08/21/2020	
3	NDC:80230-004-01	60 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
4	NDC:80230-004-80	236 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	09/10/2020	
5	NDC:80230-004-16	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 - Skymall Holdings LLC (080000475)

Revised: 9/2020

Skymall Holdings LLC