HAND SANITIZER- ethyl alcohol gel
Skaffles Group Limited Liability Company

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.

77720-012 HAND SANITIZER 71% Ethyl Alcohol

Active Ingredient

Ethyl Alcohol 71%

Purpose

Antiseptic

USE

Hand sanitizing to help reduce bacteria on the skin. Recommended for repeated use.

Warning

For external use only. Flammable. Keep away from fire or flame. When using this product do not use in or near the eyes. In case of contact, rinse eyes thoroughly with water.
Discontinue use if irritation and redness develop. lf condition persists for more than 72 hours consult a physician.

Keep out of reach of children. If swallowed, get medical
help or contact a Poison Control Center right away.

Directions

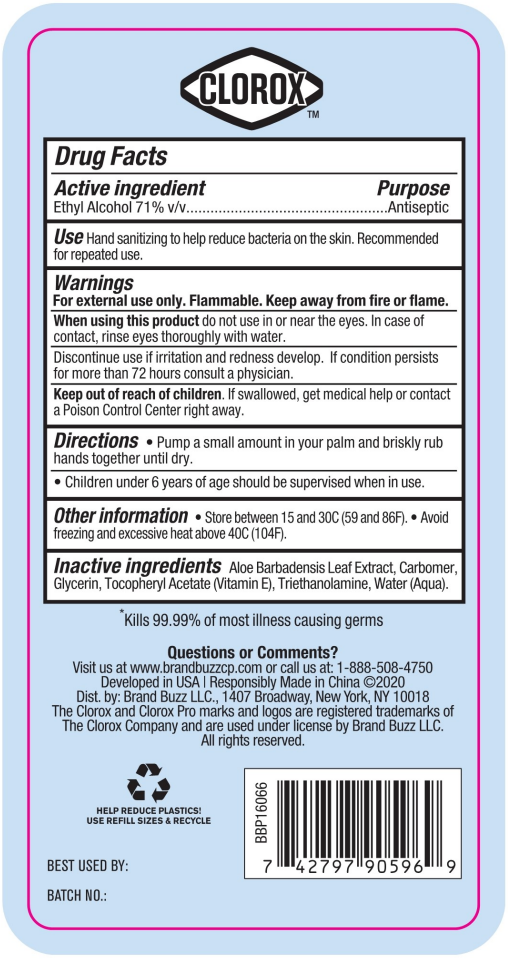
Directions . Pump a small amount in your palm and briskly rub hands together until dry.
Children under 6 years of age should be supervised when in use. Other information . Store between 15 and 30C(59 and 86F).· Avoid freezing and excessive heat above 40C(104F).

Inactive ingredients

Aloe Barbadensis Leaf Extract, Carbomer,Glycerin,Tocopheryl Acetate (Vitamin E), Triethanolamine, Water(Aqua).

4.72 in.

Front



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CLOROX Hand Sanitizer

Drug Facts

Active ingredient — **Purpose**

Ethyl Alcohol 71% v/v ..... Antiseptic

Use Hand sanitizing to help reduce bacteria on the skin. Recommended for repeated use.

Warnings

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

When using this product do not use in or near the eyes. In case of contact, rinse eyes thoroughly with water.

Discontinue use if irritation and redness develop. If condition persists for more than 72 hours consult a physician.

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Directions • Dispense a small amount in your palm and briskly rub hands together until dry.

• Children under 6 years of age should be supervised when in use.

Other information • Other information Store between 15 and 30C (59 and 86F).
• Avoid freezing and excessive heat above 40C (104F).

Inactive ingredients Aloe Barbadensis Leaf Extract, Carbomer, Glycerin, Tocopheryl Acetate (Vitamin E), Triethanolamine, Water (Aqua).

USE REFILL WITH CLOROX DISPENSER: model # 1407_0120

33.8 fl oz (1 qt 1.81 oz) / 1 L

Questions or Comments?
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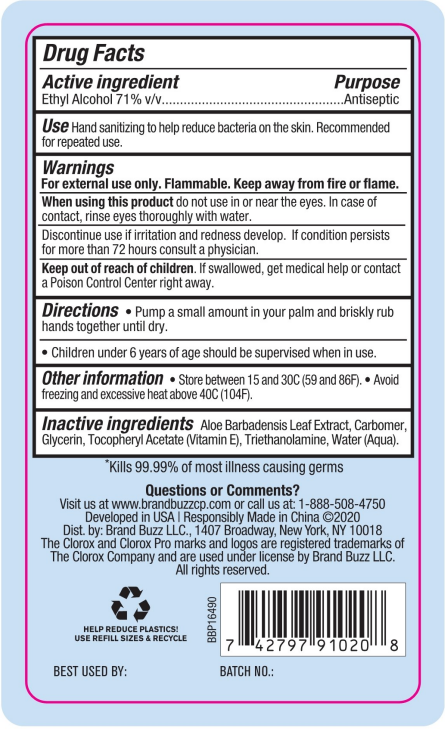
BEST USED BY:

BATCH NO.:

7 42797 90595 2

BBP16065

front



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HAND SANITIZER

ethyl alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:77720-012
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	71 mL in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
GLYCERIN (UNII: PDC6A3C0OX)	
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
WATER (UNII: 059QF0KO0R)	
TROLAMINE (UNII: 9O3K93S3TK)	
ALPHA-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:77720-012-01	60 mL in 1 BOTTLE; Type 0: Not a Combination Product	11/15/2020	
2	NDC:77720-012-08	340 mL in 1 BOTTLE; Type 0: Not a Combination Product	11/15/2020	
3	NDC:77720-012-04	1000 mL in 1 BOTTLE; Type 0: Not a Combination Product	11/15/2020	
4	NDC:77720-012-03	500 mL in 1 BOTTLE; Type 0: Not a Combination Product	11/15/2020	
5	NDC:77720-012-05	30 mL in 1 BOTTLE; Type 0: Not a Combination Product	11/15/2020	
6	NDC:77720-012-06	250 mL in 1 BOTTLE; Type 0: Not a Combination Product	11/15/2020	
7	NDC:77720-012-07	75 mL in 1 BOTTLE; Type 0: Not a Combination Product	11/15/2020	
8	NDC:77720-012-02	237 mL in 1 BOTTLE; Type 0: Not a Combination Product	11/15/2020	
9	NDC:77720-012-09	170 mL in 1 BOTTLE; Type 0: Not a Combination Product	11/15/2020	
10	NDC:77720-012-10	1000 mL in 1 BOTTLE; Type 0: Not a Combination Product	11/15/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	11/15/2020	

Labeler - Skaffles Group Limited Liability Company (831115642)

Establishment

Name	Address	ID/FEI	Business Operations
Zhejiang Guoyao Jingyue Aerosol Co., Ltd.		554529812	manufacture(77720-012)

Revised: 11/2020 Skaffles Group Limited Liability Company