CORONA PLUS- hand sanitizer liquid
ALTERNATIVE UNIVERSAL MEDICINE INC

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.

Coroa Plus Hand Sanitizer Registration

Product Application / Dosage

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

Keep Out Of Reach Of Children

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away. Do not use in/on children less than 2 month of age. Supervise children under 6 years of age when using this product to avoid swallowing.

Warnings Section

For external use only: hands. Flammable. Keep away from fire or flame. When using the product keep out of eyes, ears, and mouth. In case of contact with eyes, rinse thoroughly with water. Stop use and ask a doctor if irritation or rash occurs. These may be signs of serious condition. Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away. Do not use in/on children less than 2 month of age. Do not use on open wounds.

Inactive Ingredient Section

Inactive ingredients

Glycerin, Hydrogen peroxide, Purified water

Indication And Usage

Uses

Hand Sanitizer to help reduce bacteria on the skin that potentially can cause disease. Recommender for repeated use.

Warnings

For external use only: hands. Flammable. Keep away from fire or flame. When using the product keep out of eyes, ears, and mouth. In case of contact with eyes, rinse thoroughly with water. Stop use and ask a doctor if irritation or rash occurs. These may be signs of serious condition.

Purpose Section

Purpose

Antiseptic

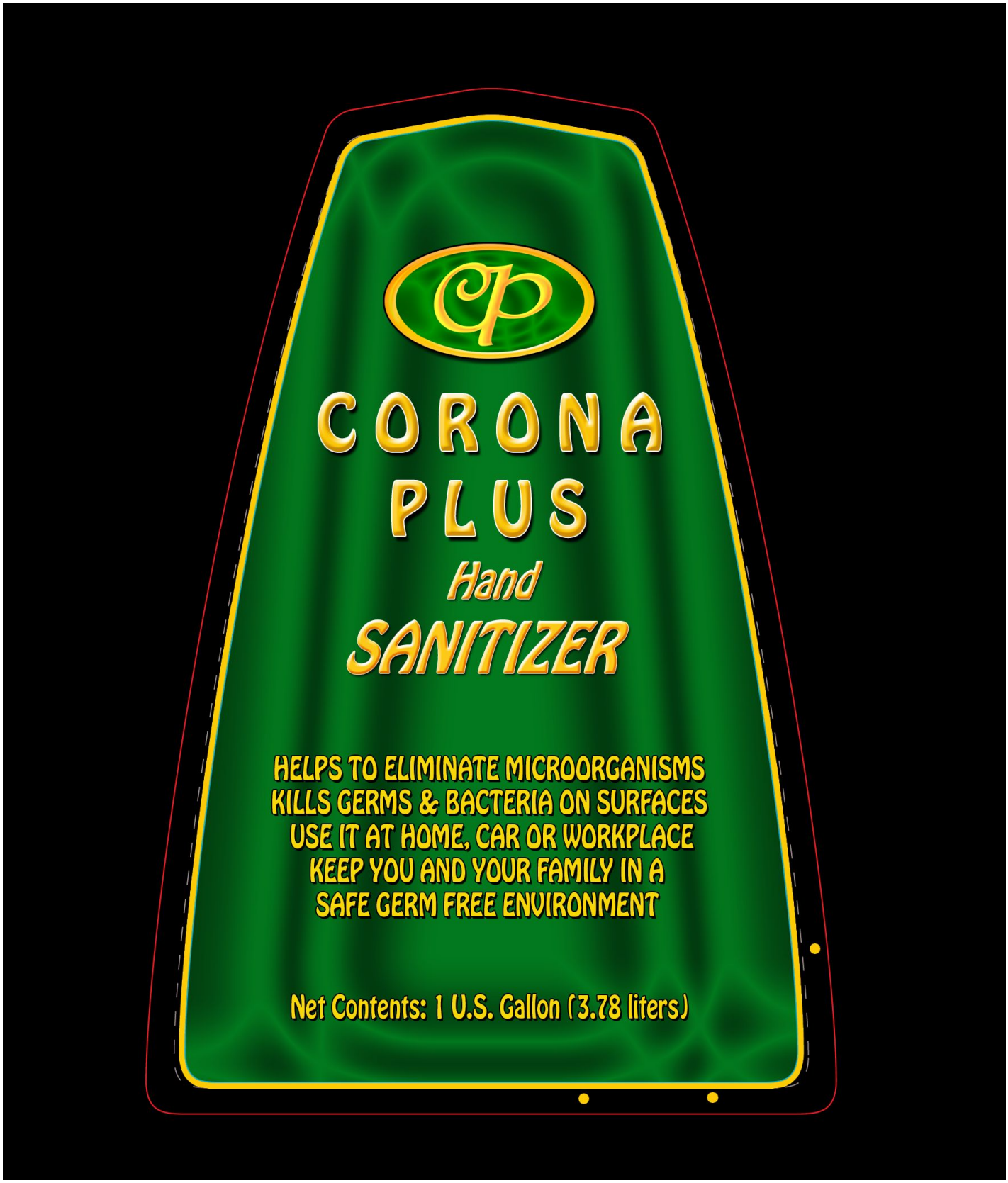
Active Ingredient

Active Ingredients

Ethyl Alcohol 86%vv

Corona Plus Package Label

3.78 ml Corona Plus



Product Lable - NDC - 79441-299-01

Front and Back

CORONA PLUS hand sanitizer liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:79441-299
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	86.18 mL in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
GLYCERIN (UNII: PDC6A3C0OX)	4.17 mL in 100 mL
HYDROGEN PEROXIDE (UNII: BBX060AN9V)	1.5 mL in 100 mL
WATER (UNII: 059QF0KO0R)	8.15 mL in 100 mL

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:79441-299-01	3780 mL in 1 BOTTLE; Type 0: Not a Combination Product	08/15/2020	
2	NDC:79441-299-02	473 mL in 1 BOTTLE; Type 0: Not a Combination Product	10/27/2020	
3	NDC:79441-299-03	946 mL in 1 BOTTLE; Type 0: Not a Combination Product	10/27/2020	
4	NDC:79441-299-04	236.5 mL in 1 BOTTLE; Type 0: Not a Combination Product	10/27/2020	
5	NDC:79441-299-05	118.25 mL in 1 BOTTLE; Type 0: Not a Combination Product	10/27/2020	
6	NDC:79441-299-06	59.125 mL in 1 BOTTLE; Type 0: Not a Combination Product	10/27/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	08/15/2020	

Labeler - ALTERNATIVE UNIVERSAL MEDICINE INC (024341671)

Establishment

Name	Address	ID/FEI	Business Operations
ALTERNATIVE UNIVERSAL MEDICINE INC		024341671	manufacture(79441-299)

Revised: 10/2020 ALTERNATIVE UNIVERSAL MEDICINE INC