# RAINBOW HAND SANITIZER GREEN- ethyl alcohol gel Belleson 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.

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# **Drug Facts**

## **Active ingredient**

Ethyl Alcohol 70% v/v

## **Purpose**

**Antiseptic** 

#### Uses

Hand sanitizer to help reduce bacteria that potentially can cause disease. For use when soap and

water are not available. Warnings For external use only, Flammable. Keep away from heat or flame

# **Warnings**

#### Do not use

- in children less than 2 months of age
- on open skin wounds

**When using this product** keep out of eyes, ears, and mouth. In case of contact with eyes, rinse eyes thoroughly with water.

**Stop use and ask a doctor** if irritation or rash occurs. These may be signs of a serious condition.

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

#### **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 15-30C (59-86F)

Avoid freezing and excessive heat above 40C (104F)

# **Inactive ingredients**

Aqua, Glycerin, Aloe vera leaf, Carbomer, Triethanolamine, Mugwort extract, Calendula officinalis flower, Camellia Sinensis Leaf Extract, Mulberry root extract, Licorice extract, Rosmarinus officinalis (Rosmary) leaf oil, Polysorbate 20, Fragrance, FD&C Blue No. 1, FD&C Yellow No. 5

#### Product label





### **RAINBOW HAND SANITIZER GREEN**

ethyl alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:75063-0012
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	70 mL in 100 mL
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	70 mL in 10

Inactive Ingredients			
Ingredient Name	Strength		
WATER (UNII: 059QF0KO0R)			
GLYCERIN (UNII: PDC6A3C0OX)			
ALOE VERA LEAF (UNII: ZY81Z83H0X)			
CALENDULA OFFICINALIS FLOWER (UNII: P0M7O4Y7YD)			
MORUS ALBA ROOT (UNII: CST1G9BZGD)			
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)			
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)			
POLYSORBATE 20 (UNII: 7T1F30V5YH)			
TROLAMINE (UNII: 903K93S3TK)			
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)			
CAMELLIA SINENSIS ROOT (UNII: 8H54O0V2K3)			
LICORICE (UNII: 61ZBX54883)			
ROSEMARY (UNII: IJ67X351P9)			
ARTEMISIA PRINCEPS LEAF (UNII: SY077EW02G)			

l	P	Packaging			
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
	1		500 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/20/2021	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	10/20/2021	

# Labeler - Belleson Inc (694793004)

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