RAINBOW HAND SANITIZER YELLOW- 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.

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 YELLOW NO. 5

Product label


rainbow
Yellow Sanitizer Gel

Drug Facts

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Batch No./Exp.date : BS21CH10 / Separate display
Customer Service Center
Call us +82-70-5222-2741 or Email us belleson@belleson.kr
Distributed by BELLESON INC.
30 Yeomgokro-14-beongil, Seogu
Incheon, 22839 South Korea

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RAINBOW HAND SANITIZER YELLOW

ethyl alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:75063-0016
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

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)	
POLYSORBATE 20 (UNII: 7T1F30V5YH)	
TROLAMINE (UNII: 9O3K93S3TK)	
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	
CAMELLIA SINENSIS ROOT (UNII: 8H54O0V2K3)	
LICORICE (UNII: 61ZBX54883)	
ROSEMARY (UNII: IJ67X351P9)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
ARTEMISIA PRINCEPS LEAF (UNII: SY077EW02G)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:75063-0016-1	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)

Revised: 10/2021 Belleson Inc