GREEN BEAUTY HAND SANITIZER- alcohol gel GTP International Corp

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.

Green Beauty Hand Sanitizer

Drug Facts

Active ingredient

Ethyl alcohol 70%

Purpose

Antiseptic

Uses

to decreases bacteria on the skin that could cause disease

Warnings

For external use only

Flammable, Keep away from heat and flame.

When using this product,

keep out of eyes in case of contact with eyes, flush thoroughly with water, avoid contact with broken skin, do not inhale or ingest

Stop use and ask a doctor if

irritation or redness develops, condition persists for more than 72 hours

Keep out of reach of children.

If swallowed, get medical help or contact Poison Control Center right away.

Directions:

wet hands thoroughly with product and allow to dry without wipping, for children under 6, use only under adult supervision, not recommended for infants

Other information:

do nto store above 105°F may discolor some fabrics, harmful to wood finishes and plastic

Inactive Ingrediets:

water, aloe barbadensis leaf juice, glyceryl caprylate/caprate, glycerin, isopropyl myristate, tocopheryl acetate, acrylates/C10-30 alkyl acrylate crosspolymer, Fragrance. benzophennane-4, blue 1, yellow 5.

Package Labeling:



LEAVES HANDS FEELING SOFT&REFRESHED

Drug Facts

Active ingredient Ethyl alcohol 70% Uses to decreases bacteria on the skin that could cause disease

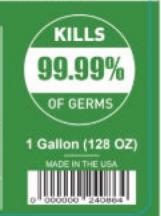
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GREEN BEAUTY HAND SANITIZER

alcohol gel

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Prod	uct	Intori	mation

Product Type HUMAN OTC DRUG Item Code (Source) NDC:79347-000

Route of Administration TOPICAL

Active Ingredient/Active Moiety Ingredient Name Basis of Strength ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M) ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)

Inactive Ingredients			
Ingredient Name	Strength		
WATER (UNII: 059QF0KO0R)			
ALOE VERA LEAF (UNII: ZY81Z83H0X)			
GLYCERYL MONOCAPRYLOCAPRATE (UNII: G7515SW10N)			
GLYCERIN (UNII: PDC6A3C0OX)			
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)			
.ALPHATOCOPHEROL ACETATE (UNII: 9E8X80D2L0)			
CARBOMER INTERPOLYMER TYPE A (ALLYL SUCROSE CROSSLINKED) (UNII: 59TL3WG5CO)			
SULISOBENZONE (UNII: 1W6L629B4K)			
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)			
FD&C YELLOW NO. 5 (UNII: 1753WB2F1M)			

P	Packaging					
#	Item Code	Package Description	Marketing Start Date	Marketing End Date		
1	NDC:79347- 000-01	3800 mL in 1 BOTTLE; Type 0: Not a Combination Product	07/09/2020			

Marketing Information					
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
OTC monograph not final	part333E	07/09/2020			

Labeler - GTP International Corp (118862131)

Revised: 12/2022 GTP International Corp