CHARCOAL FILTERED GEL HAND SANITIZER- alcohol gel Solvents and Petroleum Service 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.

Charcoal Filtered Clean All Gel Hand Sanitizer

Active Ingredient(s)

Drug Facts Active ingredient 65% Ethyl Alcohol*Alcohol 65%

Purpose

Purpose Antiseptic Antisepti

Antiseptic, Hand Sanitizer

Use

to reduce bacteria on hands, • when soap and water are not available

For use when soap and

Warnings

Warnings

for external use only

water are not available.

- flammable, keep away from heat and flame
- do not use on open wounds
 avoid eyes, ears and mouth. In case of contact, rinse thoroughly

with water For external use only. Flammable. Keep away from heat or flame

Stop use and ask a doctor if irritation or rash occurs. you may report side effects to 1-800-315-4467 **Stop and ask a doctor if**

irritation or rash occurs

you may report side effects to 1-800-315-4467

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

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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.
- do not use on children less than 2 months old

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Other information

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- Store between 59° 86°F (15 30°C).
 Avoid freezing 32°F (0°C)
- Avoid excessive heat above 104°F (40°C).
- Store between 15-30C (59-86F)
- Avoid freezing and excessive heat above 40C (104F) •

Inactive ingredients

acrylate polymer, bittering agent, glycerin, hydrogen peroxide, polyethylene glycol, water

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Questions

1-800-315-4467 moday through friday 8am-5pm est

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Package Label - Principal Display Panel



CHARCOAL FILTERED GEL HAND SANITIZER

alcohol gel

Product Information
Product Type

HUMAN OTC DRUG

Item Code (Source)

NDC:75660-154

Route of Administration

TOPICAL

Active Ingredient/Active Moiety						
Ingredient Name	Basis of Strength	Strength				
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	65 mL in $100 mL$				

Inactive Ingredients Ingredient Name Strength ACRYLATES/VINYL ISODECANO ATE CROSSPOLYMER (10000 MPA.S NEUTRALIZED AT 0.5%) (UNII: 0.5 mL in 100 mL 2N8MDB79NA) POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A) 2 mL in 100 mL GLYCERIN (UNII: PDC6A3C0OX) 1.35 mL in 100 mL 0.18 mL HYDROGEN PERO XIDE (UNII: BBX060AN9V) in 100 mL 30.96 mL WATER (UNII: 059QF0KO0R) in 100 mL 0.01 mL **DENATO NIUM BENZO ATE** (UNII: 4YK5Z54AT2) in 100 mL

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date		
1	NDC:75660-154- 01	3785.41 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/14/2020			
2	NDC:75660-154- 06	6 in 1 BOX	10/14/2020			
2	NDC:75660-154- 01	3785.41 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product				
3	NDC:75660-154- 04	4 in 1 BOX	10/14/2020			
3	NDC:75660-154- 01	3785.41 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product				
4	NDC:75660-154- 05	18927.05 mL in 1 CONTAINER; Type 0: Not a Combination Product	10/14/2020			
5	NDC:75660-154- 44	118.29 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/14/2020			
6	NDC:75660-154- 08	236.59 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/14/2020			
7	NDC:75660-154- 16	473.17 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/14/2020			
8	NDC:75660-154- 25	9463.52 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/14/2020			
Marketing Information						
Marketing Category Application Number or Monograph Citation Marketing Start Date Marketing End Date						

 $10/14/20\,20$

OTC monograph not final part333E

Labeler - Solvents and Petroleum Service Inc (013277454)

Establishment								
Name	Address	ID/FEI	Business Operations					
Clean All		117505240	manufacture(75660-154)					

Revised: 10/2020

Solvents and Petroleum Service Inc