

MEDICARE HAND SANITIZER (500ML)- ethyl alcohol gel
B&D LIFE HEALTH CO., LTD.

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.

Medicare Hand Sanitizer (500ml)

Active ingredient

Ethanol (ethyl alcohol) (62%)

Purpose

Antibacterial

Uses

Hand sanitizer for decreasing bacteria on skin

Warnings

For external use only

Do not use if you are allergic to any of the ingredients

When using this product avoid contact with eyes. If contact occurs, rinse eyes thoroughly with water. avoid contact with broken, irritated, or itching skin. Do not puncture or incinerate.

Stop use and ask a doctor if irritation or redness develops and condition persists for more than 72 hours.

If pregnant or breast-feeding, ask a health professional before use

Keep out of reach of children

If swallowed, get medical help or contact a Poison Control Center (1-800-222-1222) right away.

Directions

Apply the adequate amount on hands

Inactive ingredients

Purified water, Glycerin, L-menthol, Carbomer, Aminomethyl Propanol, Aloe extract

Medicare Hand Sanitizer (500ml)



의약품

메디케어 새니타이저겔

Medicare sanitizer
hygienic life care solution

손소독제



500ml / 16.9 fl oz

Product Name: Medicare Hand Sanitizer

Drug Facts	
Active Ingredients	Purpose
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Uses Hand sanitizer for decreasing bacteria on skin	
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Distributed by: B&D AMERICA INC.
5825 LINCOLN AVE. SUITE 402, BUENA PARK, CA 90620
Made in Korea

MEDICARE HAND SANITIZER (500ML)

ethyl alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:73786-101
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	310 g in 500 mL

Inactive Ingredients

Ingredient Name	Strength
GLYCERIN (UNII: PDC6A3C0OX)	
LEVOMENTHOL (UNII: BZ1R15MTK7)	
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	
ALOE (UNII: V5VD430YW9)	
WATER (UNII: 059QF0K00R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73786-101-01	500 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/25/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	03/25/2020	

Labeler - B&D LIFE HEALTH CO., LTD. (689849886)

Registrant - B&D LIFE HEALTH CO., LTD. (689849886)

Establishment

Name	Address	ID/FEI	Business Operations
HANKOOK MEDICARE CO., LTD.		694765274	manufacture(73786-101)

Revised: 3/2020

B&D LIFE HEALTH CO., LTD.