DR.CLEANEW HAND SANITIZER- alcohol liquid NR 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.

SANITIZER

PURPOSE

Antimicrobial

Active Ingredients

ALCOHOL

Use

Hand sanitizer to help reduce bacteria that potentially can cause disease. Recommended for external use only.

Warnings

For external use only

Keep away from fire of flame

Do not spray on clothes because it can stain.

Be caeful not to get into your eyes, and if so wash thoroughly with water

Keep out of reach of children

Keep out of reach of children.

Stop use and ask a doctor

Take care not to get into the eye.

If a children has swallowed it go to the hospital immediately and see a doctor

Other information

beause replacing in other containers can cause accidents or deteriorate quality, put it in the original container and keep

it tightly closed.

store away from high temperature and fire.

indication

sprinkle evenly on your hands

do not use

do not use on intraoral, mucosal and damaged skin carefully use for the people with allergic reactions, immediately discontinue use and consult a dermatologist if hypersensitivity symptoms such as rash itching etc occur skin irritation symptoms occur

inactive ingredient

water, citric acid

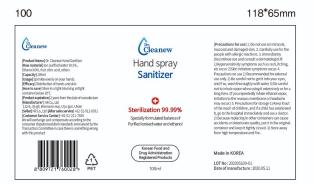
Direction

disinfection of hands and skin

Spray on hands or spray where disinfection is needed

label

79191-301-01



500 185*150mm





[Product Name] Dr. Cleanew Hand Sanitizer [Raw material] Ion purified water 39.5%,

Ethanol 60%, fruit citric acid, others [Capacity] 100ml / 500ml [Usage] Sprinkle evenly on your hands. [Efficacy] Disinfection of hands and skin [How to save] Store in a light-blocking airtight container below 30°C [Product expiration] 2 years from the date of manufacture

[Manufacturer] NR Co., Ltd Corporation 112-8, Jiji-gil, Beomseo-eup, Ulju-gun, Ulsan [Seller] NR Co., Ltd

[After sales service] +82-52-912-0951 [Customer Service Center] +82-52-221-7650 We will exchange and compensate according to the consumer dispute resolution standards announced by the Transaction Committee in case there is something wrong with this product





99.99%

Specially formulated balance of purified ionised water and ethanol

Korean Food and **Drug Administration** Registered Products

500ml

[Precautions for use] 1. Do not use on intraoral, mucosal and damaged skin. 2. Carefully use for the people with allergic reactions.

3. Immediately discontinue use and consult a dermatologist If: 1)Hypersensitivity symptoms such as rash, itching, etc occur 2)Skin irritation symptoms occur. 4. Precautions on use 1) Recommended for external use only.
2) Be careful not to get it into your eyes, and if so, wash thoroughly with water 3) Be careful not to inhale vapor when using it extensively or for a long time. (If you repeatedly inhale ethanol vapor, irritation to the mucous membrane or headache may occur.)
5. Precautions for storage 1) Keep it out of the reach of children, and if a child has swallowed it, go to the hospital immediately and see a doctor. 2) Because replacing in other containers can cause accidents or deteriorate quality, put it in the original container and keep it tightly closed. 3) Store away from high temperature and fire.

Made in KOREA

Date of use: 2 years after opening LOT No: 202005100-0 Date of manufacture: 2020.05

DR.CLEANEW HAND SANITIZER

alcohol liquid

Product Information

NDC:79191-301 HUMAN OTC DRUG Product Type Item Code (Source)

TOPICAL Route of Administration

Active Ingredient/Active Moiety

Ingredient Name Basis of Strength Strength ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M) ALCOHOL 60 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength			
CITRIC ACID MONO HYDRATE (UNII: 2968 PHW8 QP)	0.5 mL in 100 mL			
WATER (UNII: 059QF0KO0R)	39.5 mL in 100 mL			

Packaging

# Item Code	Package Description	Marketing Start Date	Marketing End Date
1 NDC:79 19 1-30 1- 0 1	100 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	06/25/2020	
2 NDC:79 19 1-30 1- 0 2	500 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	06/25/2020	

Marketing Information						
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date			
OTC monograph not final	part333A	06/25/2020				

Labeler - NR CO.,Ltd. (690430136)

Registrant - NR CO.,Ltd. (690430136)

Establishment					
Name	Address	ID/FEI	Business Operations		
NR CO.,Ltd.		690430136	manufacture (79 19 1-30 1)		

Revised: 6/2020 NR CO.,Ltd.