#### HAND SANITIZER- benzalkonium chloride liquid HAND SANITIZER- benzalkonium chloride spray Northmed SIA

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.

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Benzalkonium Chloride (.20%). Purpose: Antiseptic

Antiseptic. Hand Sanitizer.

Sanitizer to help reduce bacteria on hands. Recommended for repeated use.

For external use only.

Should not be sotred below 41F/5C or above 104F/40C.

Do not use in or near eyes. In case of contact with eyes, flush thoroughly with water. Avoid contact with broken skin.

Stop use and ask a doctor if irritation and rendess develop and persists for more than 72 hours.

Keep out of reach of children. If swallowed, get medical help or contact a Poison control center immediately.

Apply to hands and rub into the surface of the hands until completely absorbed.

Water, Glycerin, Panthenol, Aloe Vera, Tetrasodium Glutamate Diacetate.

1000 mL (33.8 fl. oz.) NDC: 78522-104-00



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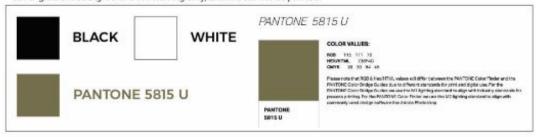
YOUR ECO-FRIENDLY PERSONAL PROTECTION 33.8 fl. oz. (1,000 ml)

Distributed by: Point One International, Ltd. Olmsted Falls, OH 44138 Made in Latvia

2 fl. oz. 59ml. NDC: 78522-100-00



\*Silver gradient background is for viewing only, and should not be printed.



### **HAND SANITIZER**

benzalkonium chloride liquid

#### **Product Information**

Product Type HUMAN OTC DRUG Item Code (Source) NDC:78522-104

**Route of Administration** TOPICAL

## **Active Ingredient/Active Moiety**

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Ingredient Name	<b>Basis of Strength</b>	Strength		
<b>BENZALKONIUM CHLORIDE</b> (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII: 7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	2 mg in 59 mL		

Inactive Ingredients		
Ingredient Name	Strength	
TETRASODIUM GLUTAMATE DIACETATE (UNII: 5EHL50I4MY)		
ALOE VERA LEAF (UNII: ZY81Z83H0X)		
PANTHENOL (UNII: WV9CM0O67Z)		
GLYCERIN (UNII: PDC6A3C0OX)		
WATER (UNII: 059QF0KO0R)		

#### **Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:78522- 104-00	1000 mL in 1 BOTTLE; Type 0: Not a Combination Product	10/27/2021	

# **Marketing Information**

Marketing	Application Number or Monograph	Marketing Start	Marketing End
Category	Citation	Date	Date
OTC monograph not final	part333A	10/27/2021	

## **HAND SANITIZER**

benzalkonium chloride spray

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:78522-100
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**Route of Administration** TOPICAL

Active Ingredient/Active Moiety		
Ingredient Name	<b>Basis of Strength</b>	Strength
<b>BENZALKONIUM CHLORIDE</b> (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII: 7N6JUD5X6Y)	BENZ ALKONIUM CHLORIDE	2 mg in 59 mL

Inactive Ingredients			
Ingredient Name	Strength		
WATER (UNII: 059QF0KO0R)			
ALOE VERA LEAF (UNII: ZY81Z83H0X)			
PANTHENOL (UNII: WW9CM0067Z)			
GLYCERIN (UNII: PDC6A3C0OX)			
TETRASODIUM GLUTAMATE DIACETATE (UNII: 5EHL50I4MY)			

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:78522- 100-00	59 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	01/05/2021	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	01/05/2021	
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# Labeler - Northmed SIA (662588132)

Establishment			
Name	Address	ID/FEI	Business Operations
Northmed SIA		662588132	manufacture(78522-100, 78522-104) , pack(78522-104) , label(78522-104)

Revised: 10/2021 Northmed SIA