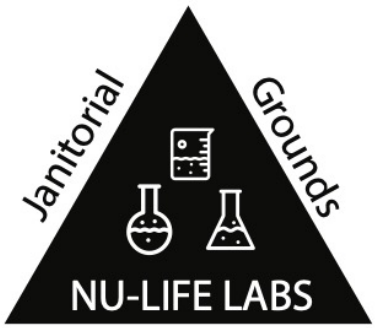
NL HAND SANI WIPES- antimicrobial hand wipe cloth
Nu-Life Labs

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.



Maintenance

NL HAND SANI-WIPES

Anti-Microbial Hand Wipes
With Ethyl Alcohol and Benzalkonium Chloride
One-Step Cleaning
Helps Wipe Away Germs
Pleasing Citrus Fragrance

Nu-Life Laboratories
P.O. Box 15793
Lenexa, KS 66285
Ph: 913-649-2625

Drug Facts

Active ingredient[s]	**Purpose**
Benzalkonium Chloride USP 0.14%	Antiseptic

Use[s]
Hand Sanitizer to help reduce bacteria that potentially can cause disease.

Warnings
For external use only.
Do not use
- In children less than 2 months of age
- On open skin wounds

When using this product Keep out of eyes, ears and mouth. In case of contact with eyes, rinse eyes thoroughly with water.
Stop use and ask a doctor if irritation or rash occurs. These may be signs of a serious condition.
Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Directions
- Wet hand thoroughly with product and allow to dry.
- Discard wipe in trash receptical after use. Do not flush.
- Supervise children under 6 years of age when using this product to avoid swallowing.

Other information
- Store between 15°C – 30° C (59°F - 86°F)
- Avoid from freezing and excessive heat above 40°C (104°F)

Inactive ingredients: water, ethyl alcohol, glycerine, alcohol ethoxylate, lauramine oxide, citric acid, fragrance

NDC 79257-594-01

50 Wipes
200g

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NDC 79257-594-06

6 Tubs
50 Wipes/Tub

NL HAND SANI WIPES

antimicrobial hand wipe cloth

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:79257-594
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.14 g in 100 g

Inactive Ingredients	
Ingredient Name	**Strength**
ALCOHOL (UNII: 3K9958V90M)	
GLYCERIN (UNII: PDC6A3C0OX)	
LAURAMINE OXIDE (UNII: 4F6FC4MI8W)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	

LEMON OIL (UNII: I9GRO824LL)	
C9-11 PARETH-6 (UNII: KCE0V8JT7W)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:79257-594-06	1 in 1 BOX	09/10/2020	
1	NDC:79257-594-01	200 g in 1 JAR; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	09/10/2020	

Labeler - Nu-Life Labs (065349581)

Establishment

Name	Address	ID/FEI	Business Operations
R. N. Eaton & Company, Inc.		056388804	manufacture(79257-594)

Revised: 3/2021 Nu-Life Labs