PL HAND SANITIZER WIPE- alcohol cloth
Newport MFG LLC

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.

PL Hand Sanitizer Wipe

Active Ingredient(s)

Ethyl Alcohol 75% v/v. Purpose: Antimicrobial

Purpose

Antimicrobial

Use

- To decrease bacteria on skin
- Recommended for repeated use

Warnings

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

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.

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Directions

- Wet hands thouroughly with product and allow to dry without wiping.
- for children under 5, use only with adult supervision.
- not recommended for use on infants

Other information

- Store between 20-25C (67-77F)

Inactive ingredients

Aloe Barbadensis Leaf Extract, Citrus Limon Peel Oil, D-Limonene, Glycerin, Tocopherol Acetate

(Vitamin E), Water

Package Label - Principal Display Panel

3 mL NDC: 80479-275-69

PL HAND SANITIZER WIPE

alcohol cloth

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:80479-275
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	75 mL in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
LEMON OIL (UNII: I9GRO824LL)	
LIMONENE, (+)- (UNII: GFD7C86Q1W)	
GLYCERIN (UNII: PDC6A3C0OX)	
WATER (UNII: 059QF0KO0R)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:80479-275-69	3 mL in 1 POUCH; Type 0: Not a Combination Product	03/30/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	03/30/2020	

Labeler - Newport MFG LLC (117654225)

Establishment

Name	Address	ID/FEI	Business Operations
Newport MFG LLC		117654225	manufacture(80479-275)

Revised: 10/2020 Newport MFG LLC