

DERMARITE ANTIBACTERIAL WET WIPES- alcohol cloth
Shaoxing Hengsheng New Material Technology Development 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.

79770-001 DermaRite Antibacterial Wet Wipes (75% Alcohol)

Ethyl Alcohol 75%

antiseptic

USE

For hand sanitizing to decrease bacteria on the skin

Warning

For external use only

Do not use in or contact the eyes

This wipes are meant for general cleaning

This wipes do not replace wipes used for medical procedures

keep this out of reach of children.Store below 95 F(35 C)

If swallowed,get medical help or contract a Poison Control Center immediately

Keep in a dry place away from fire

Discontinue use if irritation and redness develop.If the condition persists for more than 72 hours,consult a physician.

Please close the lid after dispensing to maintain purity Alcohol by nature evaporates quickly.

Keep this out of reach of children

Directions

. Wet hands thoroughly with product and allow to dry without wiping.

Inactive ingredients

Purified Water,Glycerin,Aloe Barbadensis extract,Phenoxyethanol,Ethylhexylglycerin, Propylene Glycol.

Drug Facts

ACTIVE INGREDIENTS

Purpose

Ethyl Alcohol 75%Antiseptic

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DERMARITE ANTIBACTERIAL WET WIPES

alcohol cloth

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:79770-001
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	75 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
GLYCERIN (UNII: PDC6A3C0OX)	
WATER (UNII: 059QF0KO0R)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:79770-001-01	10 in 1 BAG	07/20/2020	
1		4.4 g in 1 NOT APPLICABLE; Type 0: Not a Combination Product		
2	NDC:79770-001-02	20 in 1 BAG	07/20/2020	
2		4.4 g in 1 NOT APPLICABLE; Type 0: Not a Combination Product		
3	NDC:79770-001-03	30 in 1 BAG	07/20/2020	
3		4.4 g in 1 NOT APPLICABLE; Type 0: Not a Combination Product		
4	NDC:79770-001-04	50 in 1 BAG	07/20/2020	
4		4.4 g in 1 NOT APPLICABLE; Type 0: Not a Combination Product		
5	NDC:79770-001-05	80 in 1 BAG	07/20/2020	
5		4.4 g in 1 NOT APPLICABLE; Type 0: Not a Combination Product		
6	NDC:79770-001-06	100 in 1 BAG	07/20/2020	
6		4.4 g in 1 NOT APPLICABLE; Type 0: Not a Combination Product		
7	NDC:79770-001-07	110 in 1 PAIL	07/20/2020	

7		4.4 g in 1 NOT APPLICABLE; Type 0: Not a Combination Product		
8	NDC:79770-001-08	160 in 1 PAIL	07/20/2020	
8		4.4 g in 1 NOT APPLICABLE; Type 0: Not a Combination Product		
9	NDC:79770-001-09	400 in 1 PAIL	07/20/2020	
9		4.4 g in 1 NOT APPLICABLE; Type 0: Not a Combination Product		
10	NDC:79770-001-10	800 in 1 PAIL	07/20/2020	
10		4.4 g in 1 NOT APPLICABLE; 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	07/20/2020	

Labeler - Shaoxing Hengsheng New Material Technology Development Co., Ltd. (526871362)

Establishment

Name	Address	ID/FEI	Business Operations
Shaoxing Hengsheng New Material Technology Development Co., Ltd.		526871362	manufacture(79770-001)

Revised: 7/2020

Shaoxing Hengsheng New Material Technology Development Co., Ltd.