

DAWNMIST ALCOHOL HAND SANITIZER- alcohol gel

Dukal Corporation

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.

DawnMist Hand Sanitizer

DawnMist Alcohol Hand Sanitizer

Active Ingredient

Ethyl Alcohol 62%

Purpose

Antiseptic

Use

To decrease bacteria on the skin that could cause disease

Warning

For external use only

Flammable, keep away from fire or flame.

When using this product

Keep out of eyes, ears and mouth. In case of contact with eyes, flush thoroughly with water.

Stop use and ask a doctor if

irritation or redness develops

Keep out of reach of children

If swallowed, get medical help or contact Poison Control Center right away.

Directions

wet hands thoroughly with products and allow to dry without wiping.

for children under 6, use only under adult supervision

Other Information

Avoid freezing and excessive heat above 104°F

Inactive Ingredients

Acrylates/C10-30Alkyl Acrylate Cross Polymer, Aloe Barbadensis Leaf Juice, Fragrance, Glycerin, Maltodextrin, Propylene Glycol, Triethanolamine, Water

Principle Display Panel

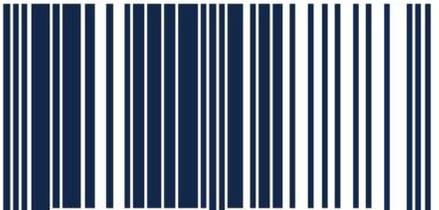
NDC 65517-1023-1

DawnMist[®]

Hand Sanitizer

Drug Facts	
Active ingredient	Purpose
Ethyl alcohol 62%.....	Antiseptic

REF HS8004



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4 FL OZ (118 ml)

Drug Facts (continued)

Uses ■ to decrease bacteria on the skin that could cause disease

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Questions? 1-800-243-0741

Manufactured for:
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Made in China
D04082003

NDC 65517-1023-1

DawnMist[®]

Hand Sanitizer

Drug Facts

Active Ingredient Purpose

Ethyl alcohol 62%.....Antiseptic

REF HS8004

4 FL. OZ. (118 ml)

DAWNMIST ALCOHOL HAND SANITIZER

alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:65517-1023
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	62 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
TROLAMINE (UNII: 9O3K93S3TK)	
WATER (UNII: 059QF0K00R)	
CARBOMER INTERPOLYMER TYPE A (55000 CPS) (UNII: 59TL3WG5CO)	
MALTODEXTRIN (UNII: 7CVR7L4A2D)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
GLYCERIN (UNII: PDC6A3C0OX)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:65517-1023-1	118 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/09/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	04/09/2020	

Labeler - Dukal Corporation (791014871)

Revised: 4/2020

Dukal Corporation