

DAWNMIST ALCOHOL HAND SANITIZER- alcohol gel
Dukal LLC

Drug Facts

Active ingredient

Ethyl alcohol 63%

Purpose

Antiseptic

Uses

- to decrease bacteria on the skin that could cause disease
- recommended for repeated use

Warnings

For external use only: hands

Flammable, keep away from fire or flame.

When using this product

- keep out of eyes. In case of contact with eyes, flush thoroughly with water.
- avoid contact with broken skin
- do not inhale or ingest

Stop use and ask a doctor if

- irritation or redness develops
- condition persists for more than 72 hours

Keep out of reach of children.

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

Directions

- wet hands thoroughly with product and allow to dry without wiping
- for children under 6, use only under adult supervision
- not recommended for infants

Other information

- do not store above 105°F
- may discolor some fabrics
- harmful to wood finishes and plastics

Inactive ingredients

acrylates/C10-30 alkyl acrylate crosspolymer, benzophenone-4, fragrance, glycerin, tocopherol, water

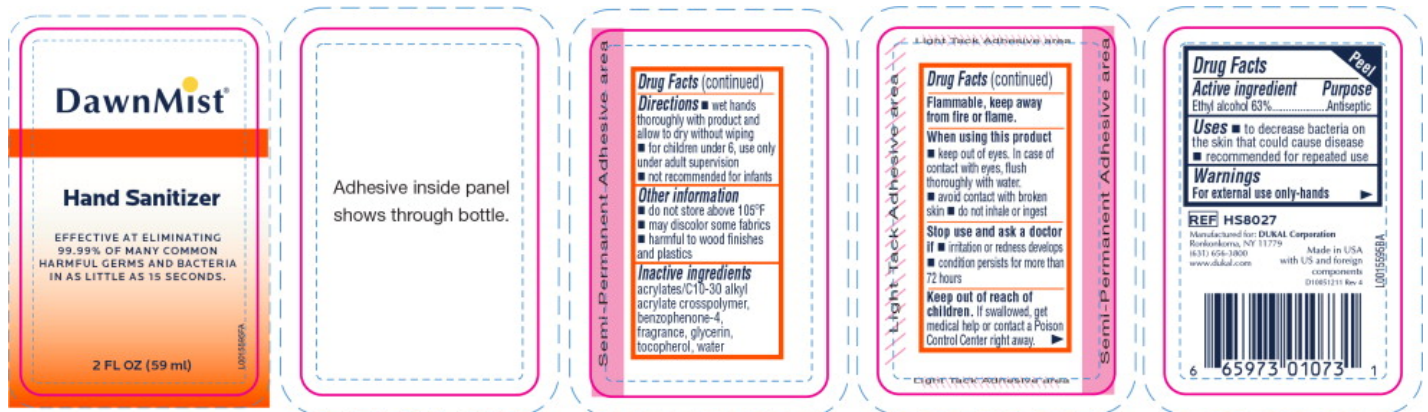
Principal Display Panel - DawnMist Alcohol Hand Sanitizer 59 mL Bottle Label

DawnMist®

Hand Sanitizer

EFFECTIVE AT ELIMINATING
99.99% OF MANY COMMON
HARMFUL GERMS AND BACTERIA
IN AS LITTLE AS 15 SECONDS.

2 FL. OZ. (59 ml)



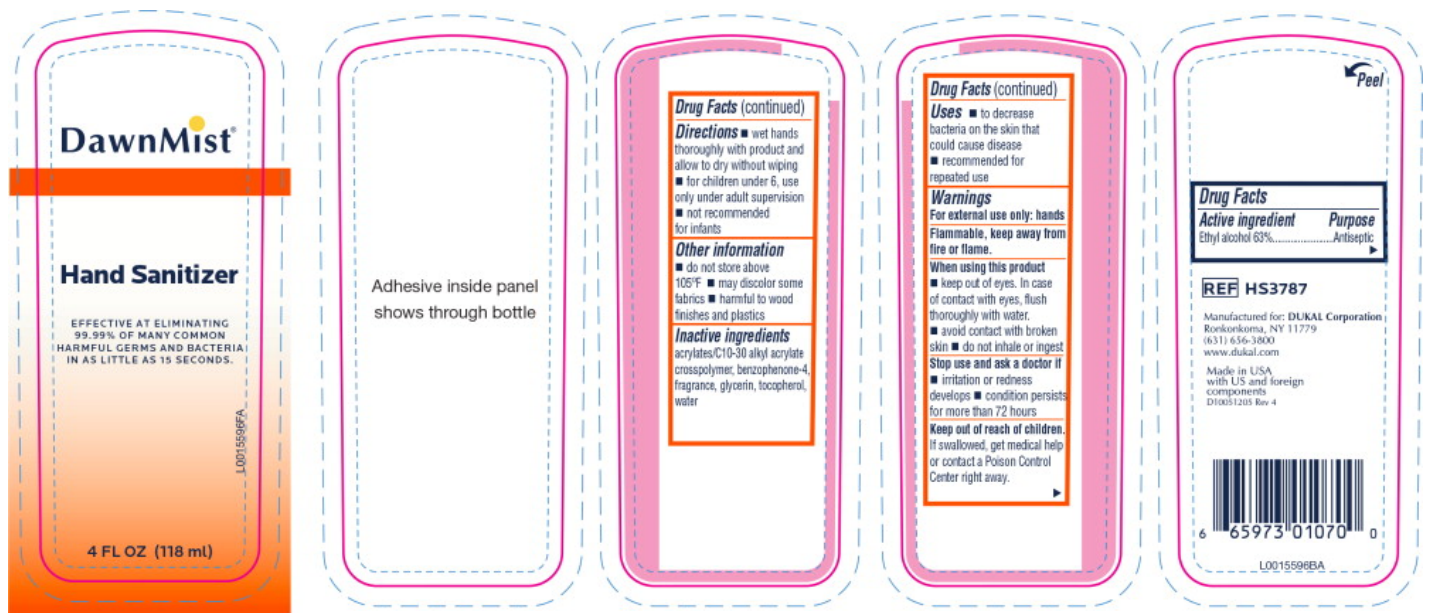
Principal Display Panel - DawnMist Alcohol Hand Sanitizer 118 mL Bottle Label

DawnMist®

Hand Sanitizer

EFFECTIVE AT ELIMINATING
99.99% OF MANY COMMON
HARMFUL GERMS AND BACTERIA
IN AS LITTLE AS 15 SECONDS.

4 FL. OZ. (118 ml)



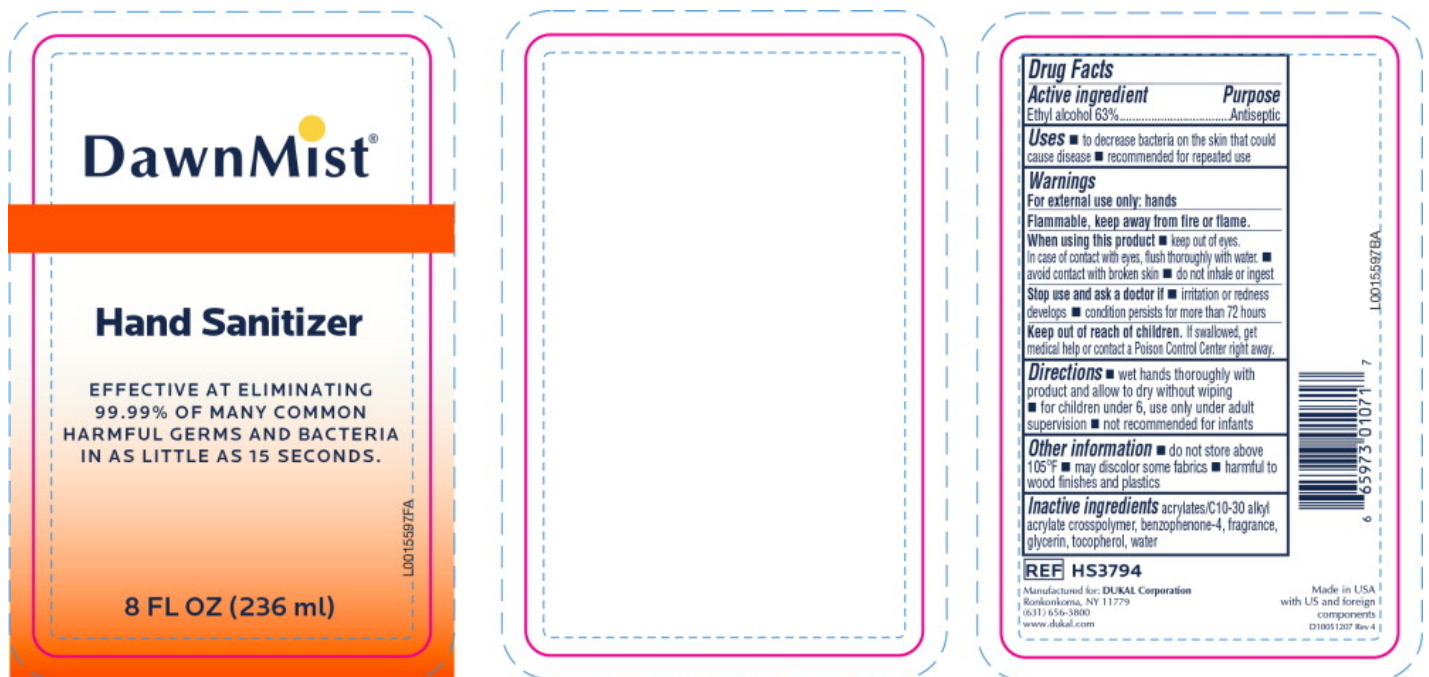
Principal Display Panel - DawnMist Alcohol Hand Sanitizer 236 mL Bottle Label

DawnMist®

Hand Sanitizer

EFFECTIVE AT ELIMINATING
99.99% OF MANY COMMON
HARMFUL GERMS AND BACTERIA
IN AS LITTLE AS 15 SECONDS.

8 FL. OZ. (236 ml)



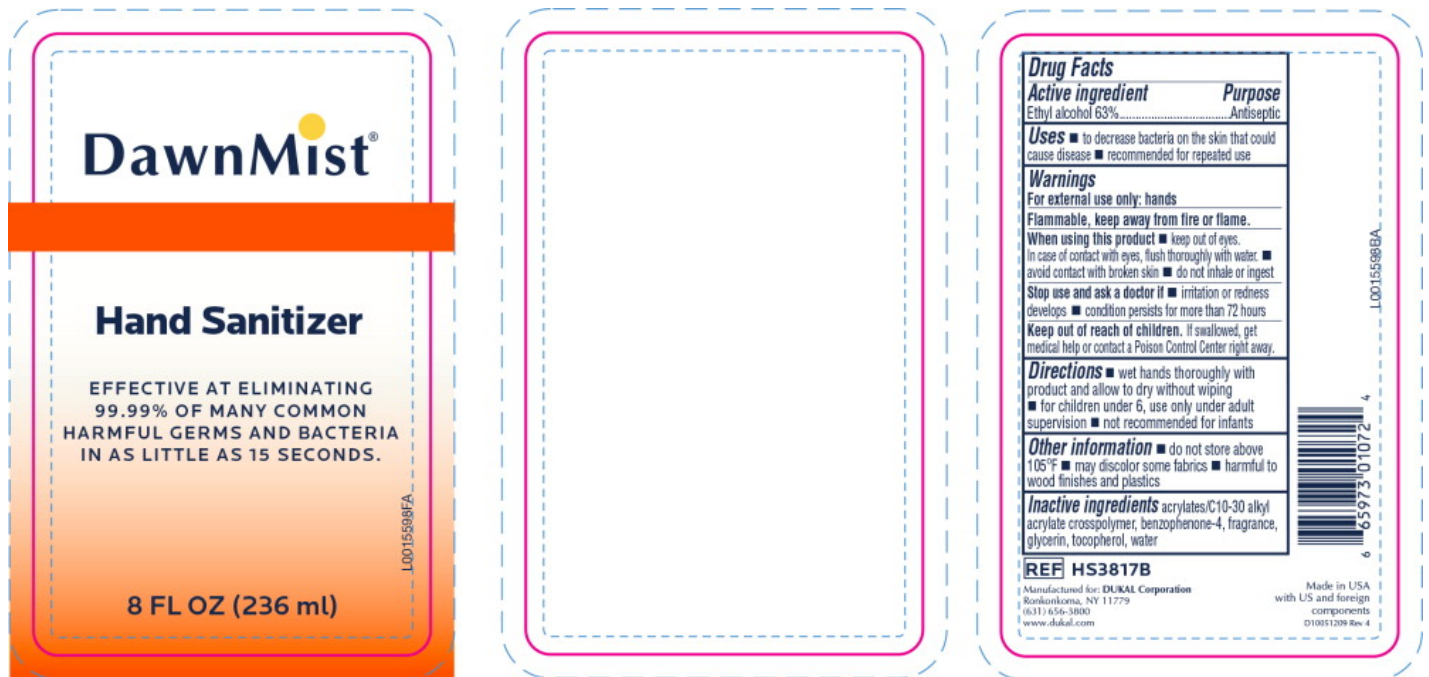
Principal Display Panel - DawnMist Alcohol Hand Sanitizer 236 mL Bottle Label

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8 FL. OZ. (236 ml)



DAWNMIST ALCOHOL HAND SANITIZER			
alcohol gel			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:65517-1020
Route of Administration	TOPICAL		
Active Ingredient/Active Moiety			
Ingredient Name		Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)		ALCOHOL	0.63 mL in 1 mL
Inactive Ingredients			
Ingredient Name			Strength
CARBOMER INTERPOLYMER TYPE A (ALLYL SUCROSE CROSSLINKED) (UNII: 59TL3WG5CO)			
SULISOBENZONE (UNII: 1W6L629B4K)			
GLYCERIN (UNII: PDC6A3C0OX)			

TOCOPHEROL (UNII: R0ZB2556P8)

WATER (UNII: 059QF0KO0R)

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:65517-1020-1	59 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/14/2013	
2	NDC:65517-1020-2	118 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/14/2013	
3	NDC:65517-1020-3	236 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/14/2013	
4	NDC:65517-1020-4	236 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	01/14/2013	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M003	01/14/2013	

Labeler - Dukal LLC (791014871)

Revised: 10/2023

Dukal LLC