

MARY KAY SATIN HANDS SHEA HAND SANITIZER- alcohol solution
Mary Kay Inc.

Mary Kay Satin Hands Shea Hand Sanitizer
Drug Facts

Active ingredient(s)

Alcohol 75% v/v

Purpose

Antiseptic

Uses

Hand sanitizer to help reduce bacteria on the skin. For use when soap and water are not available.

Warnings

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

Do not use

- on 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 product is swallowed, get medical help or contact a Poison Control Center right away.

Directions

- Spray 3+ pumps on hands, insuring both are thoroughly covered. Rub hands

together briskly until dry.

- Supervise children under 6 years of age when using this product.

Other information

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

Inactive ingredients

aloe barbadensis leaf extract, betaine, butylene glycol, glycerin, peg-50 shea butter, phenoxyethanol, water

Questions?

Call toll-free **1-800-627-9529**

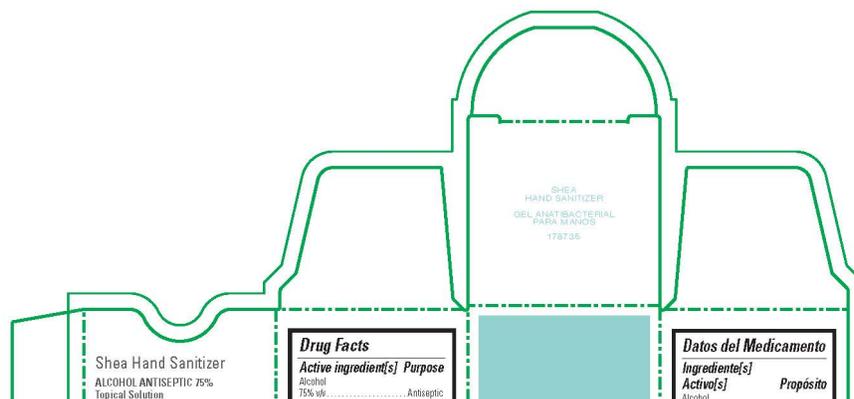
Principal Display Panel - 50 ml carton

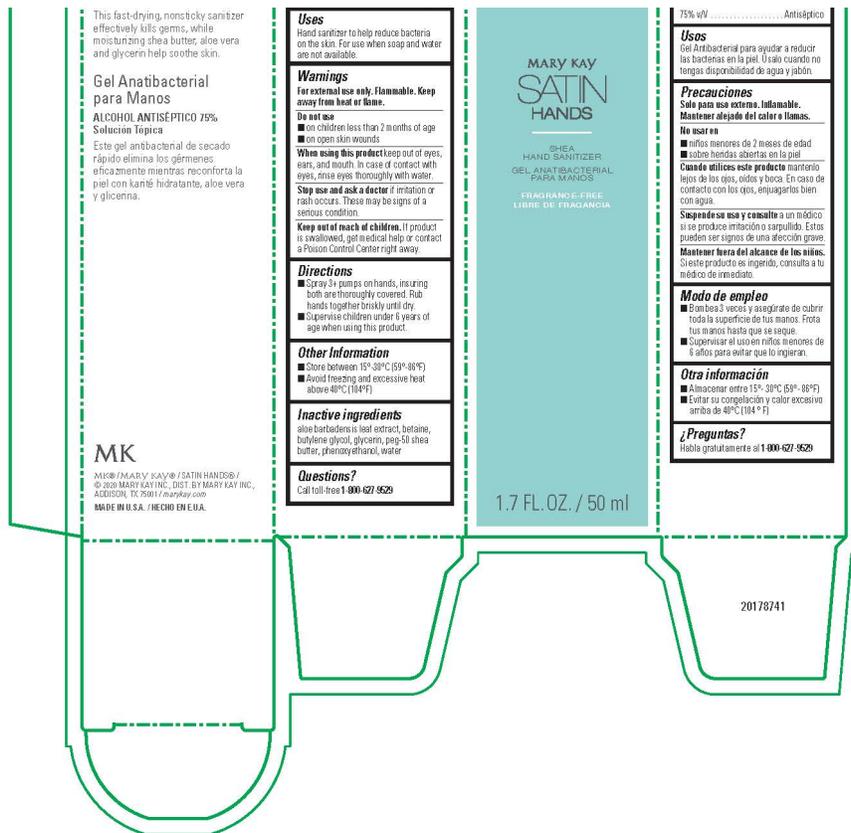
Mary Kay

Satin Hands Shea Hand Sanitizer

Fragrance-Free

1.7 FL. OZ. / 50 ml





MARY KAY SATIN HANDS SHEA HAND SANITIZER

alcohol solution

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:51531-8735
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
BETAINE (UNII: 3SCV180C9W)	
WATER (UNII: 059QF0K00R)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
GLYCERIN (UNII: PDC6A3C0OX)	
PEG-50 SHEA BUTTER (UNII: SF35CE4XLR)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:51531-8735-7	1 in 1 CARTON	11/16/2020	
1		50 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M003	11/16/2020	

Labeler - Mary Kay Inc. (049994452)

Establishment

Name	Address	ID/FEI	Business Operations
Mary Kay Inc.		103978839	manufacture(51531-8735)

Revised: 12/2024

Mary Kay Inc.