

SMART CARE HAND SANITIZER VANILLA SUGAR 01- alcohol spray Shenzhen Lantern Science Co.,Ltd.

Smart Care Hand Sanitizer Vanilla Sugar

Drug Facts

drug facts



Front View

Back View

Product dimensions: (W)58mm x (H)104mm x (D)17mm

White Sticker Label Booklet (Printing)



Back Sticker
(cover)



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Active Ingredient

Active ingredient Purpose

Ethyl Alcohol 70% v/v Antiseptic

Uses

Decreases bacteria on skin

Warning

For external use only

When using this product

Do not get into eyes. If contact occurs, rinse thoroughly with water.

Do not use

If you are allergic to any of the ingredients

Stop use and ask a doctor

If irritation or rash develops and continues for more than 72 hours.

keep out of reach of children

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

Directions

Adults and children 6 years and older
Spray onto hands and rub together until dry. Recommended for repeat use.
Children under 6 years of age
should be supervised when using this product.

Inactive ingredients

Acrylates/C10-30 Alkyl Acrylate Crosspolymer, Aloe Barbadensis Gel, Aminomethyl Propanol, Blue 1, Denatonium Benzoate, Fragrance, Glycerin, Peg-40 Hydrogenated Castor Oil, Red 33, Tocopheryl Acetate, water, Yellow 5

other Information

Store between 15-30°C (59-86°F)
Avoid freezing and excessive heat above 40°C (104°F)
Spray 3-4 times each time.
Used in the skin area of the hands.

packing

Packaging



Front View

Back View

Product dimensions: (W)58mm x (H)104mm x (D)17mm

White Sticker Label Booklet (Printing)

BLACK PANTONE 6028 C

Peel ▶

Drug Facts
Active ingredient Purpose
Ethyl Alcohol 70% v/v Antiseptic
Uses Decreases bacteria on skin ▶

ITEM # 88354

Drug Facts (Continued)
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DISTRIBUTED BY:
ASHTEL STUDIOS, INC.
ONTARIO, CALIFORNIA 91761

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Questions or comments?
Call toll free 1-877-274-8358

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SMART CARE HAND SANITIZER VANILLA SUGAR 01

alcohol spray

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:54860-430
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	70 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
GLYCERIN (UNII: PDC6A3C0OX)	
WATER (UNII: 059QF0KO0R)	
ACRYLATES/C10-30 ALKYL ACRYLATE CROSSPOLYMER (60000 MPA.S) (UNII: 8Z5ZAL5H3V)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
DENATONIUM BENZOATE (UNII: 4YK5Z54AT2)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	
PEG-40 HYDROGENATED CASTOR OIL (UNII: 7YC686GQ8F)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:54860-430-01	30 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	12/04/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M003	12/04/2024	

Labeler - Shenzhen Lantern Science Co.,Ltd. (421222423)

Establishment

Name	Address	ID/FEI	Business Operations
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Revised: 12/2024

Shenzhen Lantern Science Co.,Ltd.