SUGAR COOKIE SANITIZER 01- alcohol spray
Shenzhen Lantern Scicence Co.,Ltd.

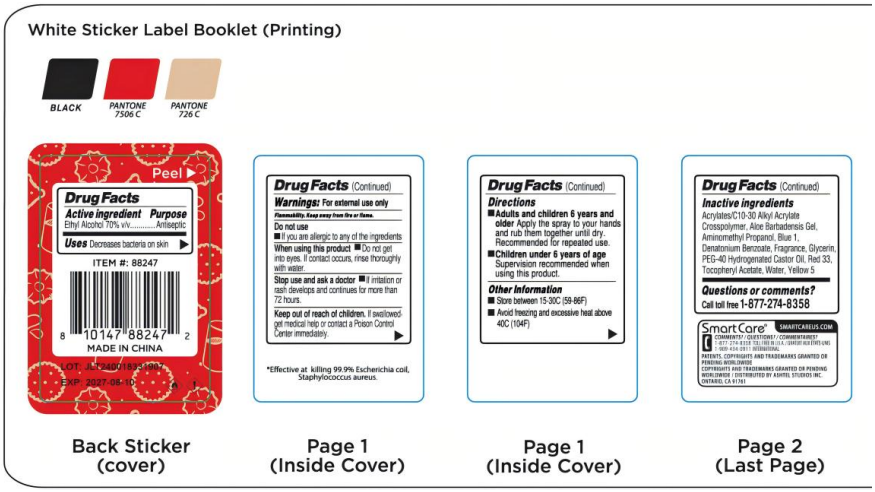
Sugar Cookie Sanitizer

Drug Facts

drug facts

Front View | Back View

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



Back Sticker (cover) | Page 1 (Inside Cover) | Page 1 (Inside Cover) | Page 2 (Last Page)

Active Ingredient

Active ingredient Purpose

Ethyl Alcohol 70% v/v Antiseptic

Uses

Decreases bacieria on skin

Warning

For external use only

Do not use

If you are allergic to any of the ingredients

When using this product

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

Stop use and ask a doctor

If irritation or rash develops and continues for more than72 hours.

keep out of reach of children

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

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.

Directions

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

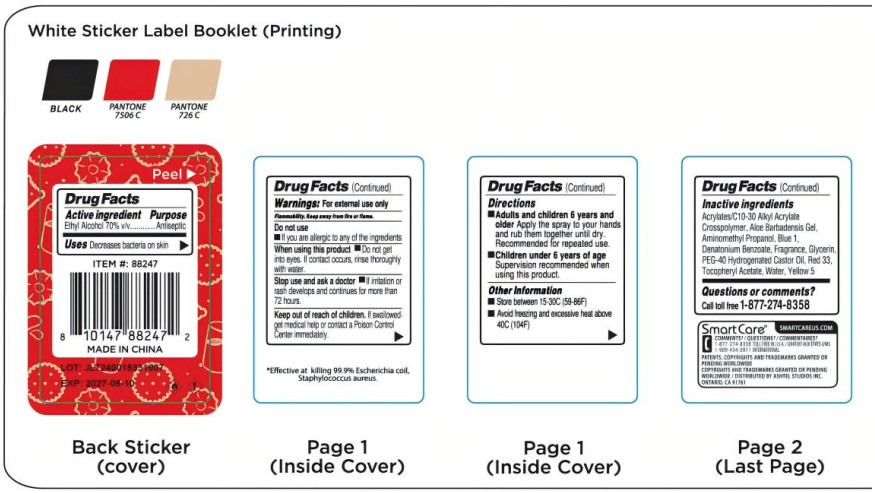
other Information

Storebetween15-30C(59-86F)
Avoid freezing and excessive heat above 40C (104F)

Spray 3-4 times each time.
Used in the skin area of the hands.

packing

Packaging



SUGAR COOKIE SANITIZER 01

alcohol spray

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:54860-426
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
POLYOXYL 40 HYDROGENATED CASTOR OIL (UNII: 7YC686GQ8F)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
ACRYLATES/C10-30 ALKYL ACRYLATE CROSSPOLYMER (60000 MPA.S) (UNII: 8Z5ZAL5H3V)	
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
GLYCERIN (UNII: PDC6A3C0OX)	
WATER (UNII: 059QF0KO0R)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
DENATONIUM BENZOATE (UNII: 4YK5Z54AT2)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:54860-426-01	40 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	08/06/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M003	08/06/2024	

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

Establishment

Name	Address	ID/FEI	Business Operations
Shenzhen Lantern Science Co.,Ltd.		421222423	manufacture(54860-426)

Revised: 8/2024 Shenzhen Lantern Scicence Co.,Ltd.