HAND SANITIZER- alcohol gel
Landy International

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.

51706-903 hand sanitizer 62% alcohol

Active Ingredient(s)

Ethyl Alcohol 62%

Purpose

Antiseptic

Use

■decreases 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 doctor if■irrtation and redness develop m 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 I not recommended for infants

Inactive ingredients

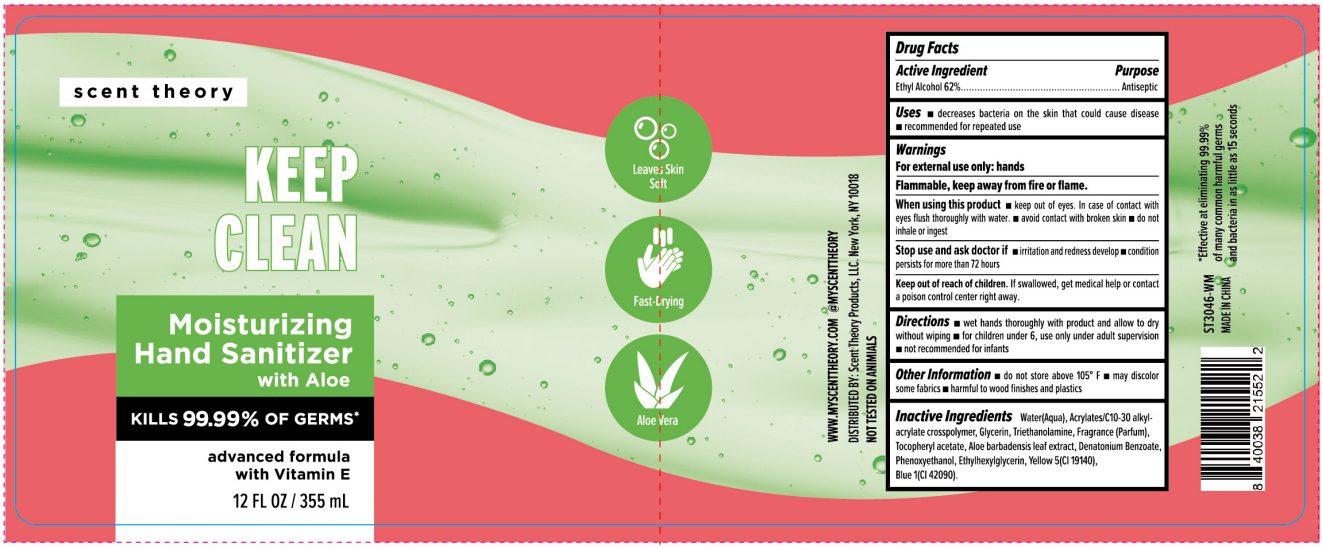
Inactive Ingredients Water(Aqua), AcrlatesC10-30 alky-acrylate crosspolymer, Glycerin, Triethanolamine, Fragrance (Parfum), Tocophery| acetate, Aloe barbadensis leaf extract, Denatonium Benzoate, Phenoxyethanol, Etyhexylglycerin, Yellow 5(CI 19140),

Blue 1(CI 42090).

Package Label - Principal Display Panel

PROJECT SPECS.	PHOTOGRAPHY	ILLUSTRATION	REVISION	INKS
Project Name: SCENT THEORY Gel Hand Sanitizer File Name: ST_12oz_Aloe_GelHandSanitizer_355ml_V3.ai Production: James King Date: 05.18.2020	☐ Custom ☐ Supplier ☒ Stock/Royalty Free ☐ Stock/Rights Mngd. ☐ Photo/Model Release ☐ N/A	☐ Custom ☐ Supplier ☐ Stock/Royalty Free ☐ Stock/Rights Mngd. ☒ N/A	3	BLACK PMS 360 C INDICATES BACKED WITH WHITE INDICATES KNOCKOUT TO BOTTLE

355ml hand sanitizer wrap label / size:190*75mm



Front Back

HAND SANITIZER

alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:51706-903
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	62 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
DENATONIUM BENZOATE (UNII: 4YK5Z54AT2)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
CARBOMER INTERPOLYMER TYPE A (ALLYL SUCROSE CROSSLINKED) (UNII: 59TL3WG5CO)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
GLYCERIN (UNII: PDC6A3C0OX)	
WATER (UNII: 059QF0KO0R)	
TROLAMINE (UNII: 9O3K93S3TK)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:51706-903-01	355 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/27/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	05/27/2020	

Labeler - Landy International (545291775)

Establishment

Name	Address	ID/FEI	Business Operations
Landy International		545291775	manufacture(51706-903)

Revised: 3/2022 Landy International