COMPANION HAND SANITIZER- hand sanitizer liquid
Neogen Corporation

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.

COMPANION HAND SANITIZER

DRUG FACTS

Active ingredients

Ethyl Alcohol 62%

PURPOSE

Antimicrobial

USES

- For hand sanitizing to decrease disease causing dermal bacteria
- Recommend for repeated use.

WARNINGS

Flammale. Keep away from fire or flame.

For External use only.

When using this product avoid contact with eyes. In case of eye contact, flush eyes with water.

Stop use and ask a doctor if irritation or redness develops, or if 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

Place 2 pumps of product into your palm. Rub hands together briskly until dry. Children under 6 years of age should be supervised when using this product.

OTHER INFORMATION

Store below 110°F (43°C). May discolor certain fabrics or surfaces.

INACTIVE INGREDIENTS

Water, DEA C8-18, Glycerin

COMPANION

Hand Sanitizer

- Kills 99.9% of harmful bacteria on contact.
- Foaming action.

Manufactured For: Neogen Corporation

944 Nandino Blvd.

Lexington, KY 40511 USA

800-477-8201 (USA/Canada) or 859-254-1221 L4849-1216

COMPANION HAND SANITIZER

hand sanitizer liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:59051-7100
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	338.82 g in 1 L

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
PEG-10 ACRYLATE/PERFLUOROHEXYLETHYL ACRYLATE COPOLYMER (UNII: D76Z87928N)	
2-(PERFLUOROHEXYL)ETHANOL (UNII: G2R5YO5N3V)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:59051-7100-2	16 in 1 PACKAGE, COMBINATION	06/05/2017	
1	NDC:59051-7100-1	0.2 L in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product		
2	NDC:59051-7100-3	3.78 L in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/05/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	06/05/2017	

Labeler - Neogen Corporation (042125879)

Registrant - Preserve International (117315289)

Establishment

Name	Address	ID/FEI	Business Operations
Preserve Inc.		808154199	manufacture(59051-7100) , api manufacture(59051-7100)

Revised: 7/2023 Neogen Corporation