SANIGUARD-SF- alcohol liquid
Inopak. Ltd

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.

Inopak SaniGuard-SF

Drug Facts

Active ingredient

Alcohol 70% v/v

Purpose

Antiseptic

Uses

- to decrease bacteria on the skin after changing diapers, assisting ill persons or before contact with a person under medical care or treatment
- Recommended for repeated use

Warnings

For external use only.

FLAMMABLE, keep away from heat or flames.

When using this product keep away from eyes. In case of eye contact, flush eyes with water.

Stop use and ask a doctor if irritation and redness develop or if condition persists for more than 72 hours.

Keep out of reach of children. If swallowed, contact a physician or poison control center.

Directions

- apply liberally to hands to cover all surfaces.
- rub hands together until dry.
- supervise children under 6 years old.
- not recommended for infants.

Inactive ingredients Water, PEG-10 Dimethicone, Glycerin, Isopropyl Myristate, Polyquaternium-11, Disodium EDTA, Aloe Barbadensis Leaf Juice, Tocopheryl Acetate

Manufacturd for

Inopak, LTD. Ringwood, NJ 07456 • 1-800-762-7725 • www.inopak.com

SaniGuard SF

foam hand sanitizer

alcohol 70%

Compliant with CDC Hand Hygiene Guidlines

Does not contain DEA/MEA, triclosan, parabens, formaldehyde, dyes, fragrance

NET CONTENTS: 33.8 FL. OZ. (1000 ML)

5068-FL1000 5068-FL Rev. 6.22

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NSF

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Manufactured for

INOPAK

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SANIGUARD-SF

alcohol liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:58575-150
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**
EDETATE DISODIUM (UNII: 7FLD91C86K)	
ALPHA-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	
PEG-10 DIMETHICONE (600 CST) (UNII: 8PR7V1SVM0)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
POLYQUATERNIUM-11 (1000000 MW) (UNII: 0B44BS5IJS)	
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:58575-150-02	60 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/01/2011	04/20/2021
2	NDC:58575-150-80	800 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/01/2011	
3	NDC:58575-150-10	1000 mL in 1 BAG; Type 0: Not a Combination Product	05/01/2011	04/20/2021
4	NDC:58575-150-12	1250 mL in 1 BAG; Type 0: Not a Combination Product	05/01/2011	04/20/2021
5	NDC:58575-150-37	3840 mL in 1 BAG; Type 0: Not a Combination Product	05/01/2011	04/20/2021
6	NDC:58575-150-18	540 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/01/2011	
7	NDC:58575-150-03	50 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/20/2021	
8	NDC:58575-150-01	1000 mL in 1 POUCH; Type 0: Not a Combination Product	04/20/2021	
9	NDC:58575-150-42	3800 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/20/2021	

Marketing Information

Marketing	Application Number or Monograph	Marketing Start	Marketing End

Category	Citation	Date	Date
OTC monograph not final	part333E	05/01/2011	

Labeler - Inopak. Ltd (194718243)

Revised: 6/2022 Inopak. Ltd