

ADVENTURE HAND SANITIZER - FRAGRANCE 60% - isopropyl alcohol spray
NICETY SOLUTIONS LLC

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.

Adventure Hand Sanitizer 60% Isopropyl Alcohol Fragrance

Active ingredient

Isopropyl Alcohol 60% v/v

Purpose

Antimicrobial

Use

For hand sanitizing to decrease bacteria on skin.

Warnings

For external use only

Flammable, keep away from fire and flame

Do not use

in the eyes

Keep out of reach of children

If swallowed, get medical help or contact an Poison Control Center right away.

Stop use and ask a doctor if

redness or irritation develops and persists for more than 72 hours

Directions

Spray hands thoroughly. Rub hands together until dry.

Inactive ingredients

Water, Propanediol, Aloe Vera Powder, Panthenol, Hydroxyethylcellulose

Label

Adventure Hand Sanitizer

By the makers of Shower Pouch

Remove odors

Kills germs on the go

1.3 fl oz [38 ml]

ADVENTURE HAND SANITIZER



BY THE MAKERS OF

S H O W E R
P O U C H®

FRAGRANCE



REMOVES ODOR ► KILLS GERMS ► ON THE GO

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Questions or comments

(310) 571-8848 | info@nicetysolutions.com



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www.theshowerpouch.com

Distributed by:

Nicety Solutions LLC
Los Angeles, CA 90504



PN: HSF60I-050320

ADVENTURE HAND SANITIZER - FRAGRANCE 60 %

isopropyl alcohol spray

Product Information

Product Type

HUMAN OTC DRUG

Item Code (Source)

NDC:77371-126

Route of Administration		TOPICAL			
Active Ingredient/Active Moiety					
Ingredient Name				Basis of Strength	Strength
ISOPROPYL ALCOHOL (UNII: ND2M416302) (ISOPROPYL ALCOHOL - UNII:ND2M416302)				ISOPROPYL ALCOHOL	0.6 mL in 1 mL
Inactive Ingredients					
Ingredient Name					Strength
WATER (UNII: 059QF0KO0R)					
ALOE VERA LEAF (UNII: ZY81Z83H0X)					
PANTHENOL (UNII: WV9CM0O67Z)					
HYDROXYETHYL CELLULOSE, UNSPECIFIED (UNII: T4V6TWG28D)					
PROPANEDIOL (UNII: 5965N8W85T)					
Packaging					
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:77371-126-01	38 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	05/05/2020		
Marketing Information					
Marketing Category		Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final		part333E	05/05/2020		
Labeler - NICETY SOLUTIONS LLC (030405122)					

Revised: 5/2020

NICETY SOLUTIONS LLC