

ADVENTURE HAND SANITIZER - UNSCENTED 80% - 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 80% Ethyl Alcohol Unscented

Active ingredient

Ethyl Alcohol 80% 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

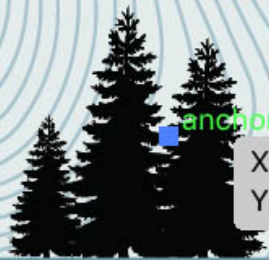
By the makers of Shower Pouch

Remove odors

Kills germs on the go

1.3 fl oz [38 ml]

ADVENTURE HAND SANITIZER



anchor

X: 194.83 pt

Y: 291.19 pt

BY THE MAKERS OF

S H O W E R
P O U C H ®

UNSCENTED



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Drug Facts

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Questions or comments

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



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HSU80E-050420

Distributed by:
Nicety Solutions LLC
Los Angeles, CA 90504





no animal
testing

manufactured
in USA

7 5 3 9 3 6 7 3 2 3 3 9

PN:

ADVENTURE HAND SANITIZER - UNSCENTED 80%

alcohol spray

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:77371-023
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	0.8 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:77371-023-01	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