

**ADVENTURE HAND SANITIZER FRAGRANCE- 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.*

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**Adventure Hand Sanitizer 60% Ethyl Alcohol Fragrance**

***Active ingredient***

Ethyl 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, Fragrance, Aloe Vera Powder, Panthenol, Hydroxyethylcellulose

**Label**

Adventure Hand Santitizer

By the makers of Shower Pouch

Remove odors

Kills germs on the go

1.3 fl oz [38 ml]

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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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# Drug Facts

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## **Questions or comments**

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

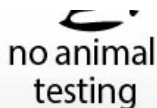


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HSF60E-050420

**Distributed by:**  
Nicety Solutions LLC  
Los Angeles, CA 90504





## ADVENTURE HAND SANITIZER FRAGRANCE

alcohol spray

### Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:77371-122
Route of Administration	TOPICAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	0.6 mL in 1 mL

### Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
PANTHENOL (UNII: WW9CM0O67Z)	
HYDROXYETHYL CELLULOSE, UNSPECIFIED (UNII: T4V6TWG28D)	
PROPANEDIOL (UNII: 5965N8W85T)	

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:77371-122-01	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)

**Registrant** - NICETY SOLUTIONS LLC (030405122)