ADVENTURE HAND SANITIZER - FRAGRANCE 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 Fragrance

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 Santitizer By the makers of Shower Pouch Remove odors Kills germs on the go 1.3 fl oz [38 ml]

BY THE MAKERS OF SHOWER POUCH

TIZER

FRAGRANCE



REMOVES ODOR ► KILLS GERMS ► ON THE GO

1.3 fl oz (38mL)

Drug Facts	6
Active ingredient Purpose Ethyl Alcohol 80% v/v Antimicrobial	
Use For hand sanitizing to decrease bacteria on skin.	
Warnings: For external use only	4
Flammable, keep away from fire and flame	
Do not use in the eyes	0
Stop use and consult a physician if redness or irritation develops and persists for more than 72 hours.	com
Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.	pouch.
Directions Spray hands thoroughly. Rub hands together until dry.	@shc
Inactive ingredients Water, Propanediol, Fragrance, Aloe Vera Powder, Panthenol, Hydroxyethylcellulose.	Follow us @showerpouch www.theshowerpouch.com
Questions or comments (310) 571-8848 info@nicetysolutions.com	Fol
Distributed by: Nicety Solutions LLC Los Angeles, CA 90504	BN: HSF80E-050320
ADVENTURE HAND SANITIZER - FRAGRAM	NCE 80%
Product Information	
Product Type HUMAN OTC DBUG Item Co	de (Source) NDC:7737

 Product Type
 HUMAN OTC DRUG
 Item Code (Source)
 NDC:77371-124

 Route of Administration
 TOPICAL

	lient/Active Moiety		
	Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3	K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	0.8 mL in 1 mL
Inactive Ingr	edients		
	Ingredient Name		Strength
WATER (UNII: 059	QF0KO0R)		
ALOE VERA LEAF	(UNII: ZY81Z83H0X)		
PANTHENOL (UNI	: WV9CM0067Z)		
HYDROXYETHYL	CELLULOSE, UNSPECIFIED (UNII: T4V6TWG28	D)	
PROPANEDIOL (U	NII: 5965N8W85T)		
Packaging			
# Item Code	Package Description	Marketing Start Date	Marketing End Date
	38 mL in 1 BOTTLE, SPRAY; Type 0: Not a	05/05/2020	
1 NDC:77371- 124-01	Combination Product		
	Combination Product		
1 124-01	Combination Product Information		
1 124-01		oh Marketing Start Date	Marketing End Date
Marketing	Information Application Number or Monograp Citation		

Labeler - NICETY SOLUTIONS LLC (030405122)

Registrant - NICETY SOLUTIONS LLC (030405122)

Revised: 7/2022

NICETY SOLUTIONS LLC