HANNAFORD HAND SANITIZER - ethyl alcohol spray Health-Tech, Inc.

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.

Drug Facts

Active Ingredient

Ethyl alcohol

Purpose

Antiseptic

Uses

- For hand sanitizing to decrease bacteria on the skin.
- Recommended for repeated use.

Warnings

- For external use only.
- Do not use in eye area.

Flammable: Keep away from heat or flame.

When using this product

- Keep out of eyes. In case of eye contact, flush eyes thoroughly with water.
- Avoid contact with broken skin.

Stop use and ask a doctor if

Irritation or redness develops or if condition persists for more than 72 hours.

Keep out of reach of children

If swallowed, get medical help, or contact a Poison control Center immediately.

Directions

- Pump one to three sprays onto palms of hands
- Rub thoroughly over all surfaces of both hands
- Rub hands together briskly until dry
- Children under 6 should be supervised when using this product
- Not recommended for infants

Other information

• Store at 20^0 to 25^0 C (68^0 to 77^0 F)

- Prior to initial use, prime pump by depressing multiple times
- May discolor certain fabrics
- harmful to wood finishes and plastics

Inactive Ingredients

Aloe Vera, D and C Green 5, DI Water, FD and C Yellow 5, Fragrance, Propylene Glycol, Tocopheryl Acetate

Hannaford bottle label



MM9

Hannaford blister card

MM10





Drug Facts

Active ingredients Ethyl Alcohol 62% Purpose Antiseptic

Uses Sanitizes & moisturizes. Kills most household gerns. Fits perfectly in purse, pocket or car. Leaves hands feeling soft with Vitamin E & Aloe. 70+ Sprays per bottle. For hand sanitizing to decrease bacteria on the skin that may cause disease, when soap and water are not available.

Warning

Flammable. Keep away from fire or flame.

For External Use Only

When using this product do not use in or near the eyes. In case of contact, rinse eyes thoroughly with water. Avoid contact with broken skin.

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

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Directions

Pump one to three sprays onto palms of hands. Rub thoroughly over all surfaces of both hands. Rub hands together briskly until dry. Children under 6 should be supervised when using this product. Not recommended for infants.

Other information Store below 25° to 40°C (68° to 77°F) May discolor some fabrics. Harmful to wood finishes & plastics

Inactive ingredients

Aloe vera, D&C Green No. 5, DI Water, FD&C Yellow No. 5, Fragrance, Propylene Glycol, Tocopheryl Acetate

*Effective at eliminating 99.99% of many common harmful germs and bacteria.

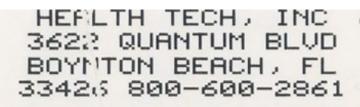
Distributed by: Hannaford Bros. Co., Scarborough, ME 04074.
For product questions or concerns contact us at 1-800-213-9040.
Please include UPC number and code from package.
www.hannaford.com

NDC 48871-002-01



This product carries our commitment to quality and value. If you're not completely satisfied, let us know - we'll doubly your money back.

Hannaford shipper label



HANNAFORD HAND SANITIZER SPRAY

12/6

CASE PACK ITEM: 69511 NDC 48871-002-01

CASE QTY



00041268169511

MM11

HANNAFORD HAND SANITIZER

ethyl alcohol spray

D-4-4	4	Tf		
Prod	HCL	THIO	amau	on

Product Type HUMAN OTC DRUG Item Code (Source) NDC:48871-002

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name		Basis of Strength	Strength
ı	alcohol (UNII: 3K9958 V90 M) (alcohol - UNII: 3K9958 V90 M)	alcohol	4.65 mL in 7.5 mL

inactive ingredients				
	Ingredient Name	Strength		

aloe vera leaf (UNII: ZY81Z83H0X)

ALPHA-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)

Propylene Glycol (UNII: 6DC9Q167V3)

water (UNII: 059QF0KO0R)

Inactive Ingredients

Product Characteristics

green (D and C green 5), yellow (FD and C yellow 5) Color Score

Shape	Size	
Flavor	Imprint Code	
Contains		

Packaging					
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:48871-002-01	6 in 1 CASE			
1		12 in 1 CARTON			
1		1 in 1 BLISTER PACK			
1		7.5 mL in 1 BOTTLE, SPRAY			

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part333	03/31/2010		

Labeler - Health-Tech, Inc. (084007889)

Establishment				
Name	Address	ID/FEI	Business Operations	
Health-Tech, Inc.		084007889	manufacture	

Revised: 3/2010 Health-Tech, Inc.