ALCOHOL- alcohol liquid
H-E-B

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.

HEB Hand Sanitizer
826

Active ingredient

Ethyl Alcohol 62%

Purpose

Antiseptic

Uses

- for handwashing to decrease bacteria on the skin that could cause disease
- recommended for repeated use

Warnings

For exterenal use only-hands

Flammable, keep away from fire or flame.

When using this product

- keep out of eyes. In case of contact with eyes, flush thoroghly with water.
- avoid contact with broken skin
- do not inhale or ingest

Stop use and ask a doctor if

- irritation and redness develop
- 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

- wet hands thoroughly with product and allow to dry without wiping
- for children under 6, use only under adult supervision
- not recommended for infants

Other information

- do not store above 105°F
- may discolor some fabrics
- harmful to wood finishes and plastics

Inactive ingredients

water, carbomer, fragrance, glycerin, isopropyl myristate, Aloe barbadensis leaf juice, tocopheryl acetate, blue 1, yellow 5

Questions?

1-888-287-1915

*Effective at eliminating 99.99% of many common harmful germs and bacteria in as little as 15 seconds.

SDS-MO-15036 SDA-WI-2486 DSP-MO-28 DSP-MO-34

DISTRIBUTED BY: H-E-B, SAN ANTONIO, TX 748204

PRINCIPAL DISPLAY PANEL

HILL COUNTRY FARE

HAND

SANITIZER

WITH ALOE

KILLS 99.99% OF GERMS

12 FL OZ (354 mL)

ALCOHOL
alcohol liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:37808-826
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	62 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
CARBOMER INTERPOLYMER TYPE A (ALLYL SUCROSE CROSSLINKED) (UNII: 59TL3WG5CO)	
GLYCERIN (UNII: PDC6A3C0OX)	
TOCOPHEROL (UNII: R0ZB2556P8)	
WATER (UNII: 059QF0KO0R)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	
ALOE (UNII: V5VD430YW9)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:37808-826-32	355 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/19/2019	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	07/19/2019	

Labeler - H-E-B (007924756)

Registrant - Vi Jon, Inc. (088520668)

Establishment

Name	Address	ID/FEI	Business Operations
Vi Jon, Inc.		088520668	manufacture(37808-826)

Establishment

Name	Address	ID/FEI	Business Operations
Vi Jon, Inc		150931459	manufacture(37808-826)