HAND SANITIZER - alcohol liquid Bullet Line, 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.

Drug Facts

Ethyl Alcohol (63%)

Antimicrobial

Hand sanitizer to help reduce bacteria on the skin

For external use only. Do not use in or near the eyes. Stop use and ask a doctor if irritation or rash appears and lasts. Keep out of reach of children. If swallowed contact a doctor or Poison Control Center right away. Flammable-keep away from open flame.

Water, Glycerin, Aloe Vera Leaf Extract, Citrus Peel Extract

Keep out of reach of children. If swallowed contact a doctor or Poison Control Center right away.

Wet hands thoroughly with product and allow to dry without wiping.

Drug Facts

Active ingredient Purpose Ethyl Alcohol (63%)...... Antimicrobial

Use - Hand sanitizer to help reduce bacteria on the skin.

Warnings - For external use only • Do not use in or near the eyes • Stop use and ask a doctor if irritation or rash appears and lasts.

 Keep out of reach of children. If swallowed contact a doctor or Poison Control Center right away
Flammable - keep away from open flame.

Directions - Wet hands thoroughly with product and allow to dry without wiping.

Inactive Ingredients - Water, Glycerin, Aloe Vera Leaf Extract, Citrus Peel Extract.





Mfg. for HLI, Portland, OR 97222 800-794-3620 ASI 62050 alcohol liquid

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:19392-130

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII: 3K9958V90M)	ALCOHOL	630 mL in 1000 mL

Inactive Ingredients				
Ingredient Name	Strength			
WATER (UNII: 059QF0KO0R)				
GLYCERIN (UNII: PDC6A3C0OX)				
ALOE VERA LEAF (UNII: ZY81Z83H0X)				
BITTER ORANGE OIL (UNII: 9TLV70SV6I)				

ı	Packaging					
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
		NDC:19392- 130-01	8 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/08/2014		

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part333E	10/08/2014		

Labeler - Bullet Line, LLC (122539042)

Revised: 12/2021 Bullet Line, LLC