INSTANT HAND SANITIZER LEMON DROP - alcohol gel BB17, 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.

Active ingredients

Ethyl Alcohol 62% v/v

Purpose

Antimicrobial

KILLS MORE THAN 99.99% OF COMMON GERMS

WARNING:

Flammable, Keep away from fire or flame. **FOR EXTERNAL USE ONLY.** DO NOT USE IN THE EYES.

DISCONTINUE USE IF IRRITATION AND REDNESS DEVELOP. IF CONDITION PERSISTS FOR MORE THAN 72 HOURS, CONSULT A DOCTOR.

Keep out of reach of children.

DIRECTIONS:

Rub into skin until dry.

Other Information:

Store below 110 F.

INACTIVE INGREDIENTS:Water, Aloe barbadenis, Leaf Juice, Glycerin,

Propylene Glycol, Carbomer, Triethanolamine, Fragrance, Tocopheryl Acetate, FD&C Yellow NO.5



KILLS MORE THAN 99.99% OF COMMON GERMS

DRUG FACTS Active Ingredient: Ethyl Alcohol 62%

Purpose: Antimicrobial

WARNING: FLAMMABLE, KEEP AWAY FROM FIRE OR FLAME.

FOR EXTERNAL USE ONLY, DO NOT USE IN THE EYES. DISCON-TINUE USE IF IRRITATION AND REDNESS DEVELOP. IF CONDITION PERSISTS FOR MORE THAN 72 HOURS, CONSULT A DOCTOR.

KEEP OUT OF REACH OF CHILDREN.

Directions: Rub into skin until dry.

Other Information: Store below 110°F.

Inactive Ingredients: Water, Aloe Barbadensis Leaf Juice, Glycerin, Propylene Glycol, Carbomer,
Triethanolamine,
Fragrance, Tocopheryl
Acetate, FD&C Yellow NO.5

FOR BEST USE, PLEASE SHAKE ME



INSTANT HAND SANITIZER LEMON DROP

ethyl alcohol gel

Product Information

Product Type HUMAN OTC DRUG LABEL Item Code (Source) NDC:53603-1004

Route of Administration TOPICAL DEA Schedule

Active Ingredient/Active Moiety

Ingredient Name Basis of Strength Strength
Alcohol (Alcohol) Alcohol 62 mL in 100 mL

Inactive Ingredients Ingredient Name Strength WATER ALOE ALOE VERA LEAF GLYCERIN PROPYLENE GLYCOL CARBOMER HOMOPOLYMER TYPE C (ALLYL PENTAERYTHRITOL CROSSLINKED)

TROLAMINE	
.ALPHATO COPHEROL ACETATE	
FD&C YELLOW NO.5	

F	Packaging					
#	Item Code	Package Description	Marketing Start Date	Marketing End Date		
1	NDC:53603-1004-2	59.1 mL in 1 BOTTLE, SPRAY				

Marketing Information						
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date			
OTC monograph not final	part333E	12/06/2012				

Labeler - BB17, LLC (828378294)

Revised: 12/2012 BB17, LLC