HOME BOOV-EL GUM SCENTED HAND SANITIZER- benzalkonium chloride gel
SHENZHEN LANTERN SCIENCE CO., LTD

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.

PURPOSE

Antibacterial

To decrease bacteria on skin that could cause disease

ACTIVE INGREDIENT

BENZALKONIUM CHLORIDE.....0.1%

DOSAGE AND ADMINISTRATION

Rub a dime sized drop into hands

For children under 6 use unde adult supervision

WARNINGS

For external use only - hands

Keep out of eyes.

Avoid contact with broken skin.

Do not inhale or ingest

If swallowed get medical help or contact a poison control center right away

KEEP OUT OF REACH OF CHILDREN

Keep out of reach of children

For children under 6, use under adult supervision

INDICATIONS AND USAGE

To decrease bacteria on skin that could cause disease.

INACTIVE INGREDIENTS

WATER (AQUA/EAU)
GLYCERIN
COCETH-7
PPG-1-PEG-9 LAURYL GLYCOL ETHER
CARBOMER
PEG-40 HYDROGENATED CASTOR OIL
FRAGRANCE (PARFUM)
SODIUM HYDROXIDE

PACKAGE LABEL

HOME BOOV-EL GUM SCENTED HAND SANITIZER

benzalkonium chloride gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:54860-020
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	1 mg in 100 mg

Inactive Ingredients	
Ingredient Name	**Strength**
COCETH-7 (UNII: 58Y261JLH5)	
CARBOMER 1342 (UNII: 809Y72KV36)	
GLYCERIN (UNII: PDC6A3C0OX)	
PPG-1-PEG-9 LAURYL GLYCOL ETHER (UNII: 5R8J43K25L)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
CASTOR OIL (UNII: D5340Y2I9G)	
WATER (UNII: 059QF0KO0R)	

Product Characteristics

Color	red (RED 40) , red (RED 33) , blue (BLUE 1) , yellow (YELLOW 5)	**Score**	
Shape		**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:54860-020-01	60 mg in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/01/2016	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	03/01/2016	

Labeler - SHENZHEN LANTERN SCIENCE CO., LTD (421222423)

Establishment

Name	Address	ID/FEI	Business Operations
SHENZHEN LANTERN SCIENCE CO., LTD		421222423	manufacture(54860-020)

Revised: 2/2016 SHENZHEN LANTERN SCIENCE CO., LTD