GENRX HAND SANITIZING- benzalkonium chloride spray
PureTek Corporation

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 ingredient

Benzalkonium Chloride 0.13%

Purpose

Antiseptic

Uses

■ a topical antiseptic that protects against germs

Warnings

For external use only

Do not use on

■ deep or puncture wounds ■ animal bites ■ serious burns

When using this product

■ do not get into eyes

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
■ allow to dry without wiping
■ repeat as necessary

Other information

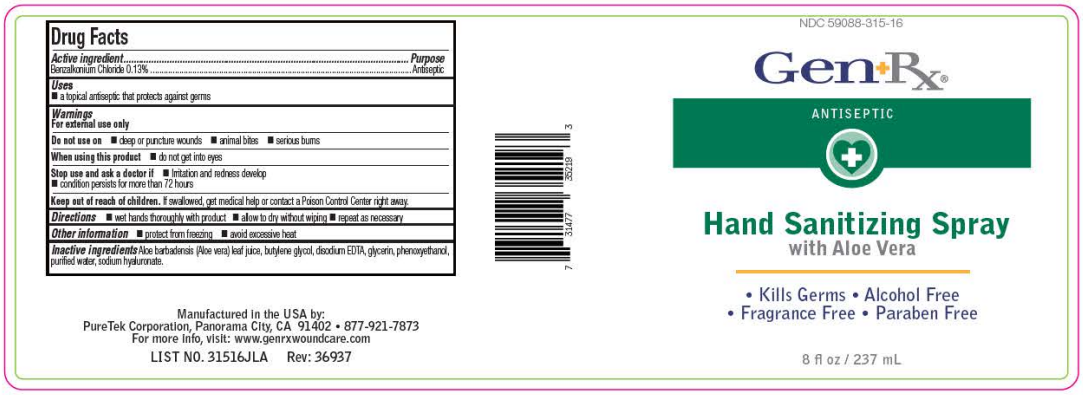
■ protect from freezing ■ avoid excessive heat

Inactive ingredients

Aloe barbadensis (Aloe vera) leaf juice, butylene glycol, disodium EDTA, glycerin, phenoxyethanol,

purified water, sodium hyaluronate.

GenRx Hand Sanitizing Spray (8 oz label)



GENRX HAND SANITIZING

benzalkonium chloride spray

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:59088-315
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	1.3 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
EDETATE DISODIUM (UNII: 7FLD91C86K)	
GLYCERIN (UNII: PDC6A3C0OX)	
WATER (UNII: 059QF0KO0R)	
HYALURONATE SODIUM (UNII: YSE9PPT4TH)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:59088-315-16	237 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	03/18/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	03/18/2020	

Labeler - PureTek Corporation (785961046)

Revised: 3/2020 PureTek Corporation