ALLAPURE ADVANCED FORMULA HAND SANITIZER- benzalkonium chloride liquid
Allapure, 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.

Allapure Advanced Formula Hand Sanitizer

Drug Facts

Active Ingredient:

Benzalkonium Chloride 0.13%

Purpose

Antiseptic/Hand & Skin Sanitizer

Uses:

Allapure *Advanced Hand Sanitizer & Wound Care is revolutionary wound care and disease protection. It helps decrease bacteria on the skin from cuts, lacerations, incisions and abrasions. Allapure* Advanced Hand Sanitizer & Wound Care promotes healing and is gentle on delicate wound tissue. Recommended for repeated use.

Warnings:

Do not freeze. For external use only.

Do not use

in ears, eyes or mouth.

When using this product,

avoid contact with the eyes. In case of contact, flush eyes with water.

Stop use and ask a doctor

if redness or irritation develop and persist for more than 72 hours.

Keep out of reach of children.

Children should be supervised when using this product.

Directions:

Apply liberally to the palms of the hands or areas of damaged skin. Rub into skin until dry. Repeat if neccesary.

Inactive Ingredients:

1-Octadecyldimethyl(3-Triethoxysilylpropyl) ammonium chloride, Aloe Barbadensis leaf extract, Aqua, C12-C15 Pareth 12, Caprylyl Glucoside, Polyhexanide, Phenoxyethanol, Triethanolamine.

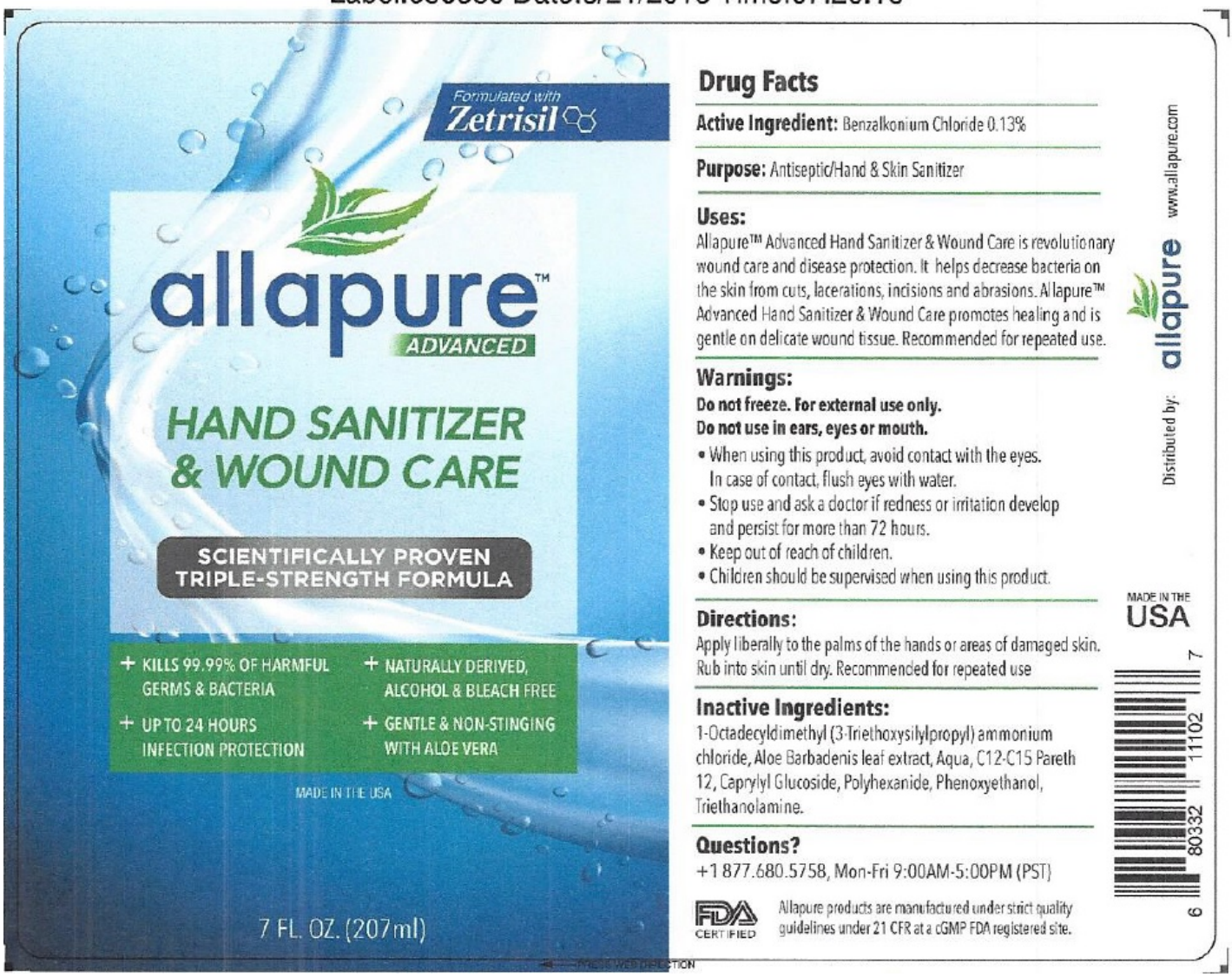
Questions?

+1 877.680.5758, Mon-Fri 9:00AM-5:00PM (PST)

Allapure Advanced Formula Hand Sanitizer 1.7oz (72552-001-01)

Allapure Advanced Formula Hand Sanitizer 7oz (72552-001-07)

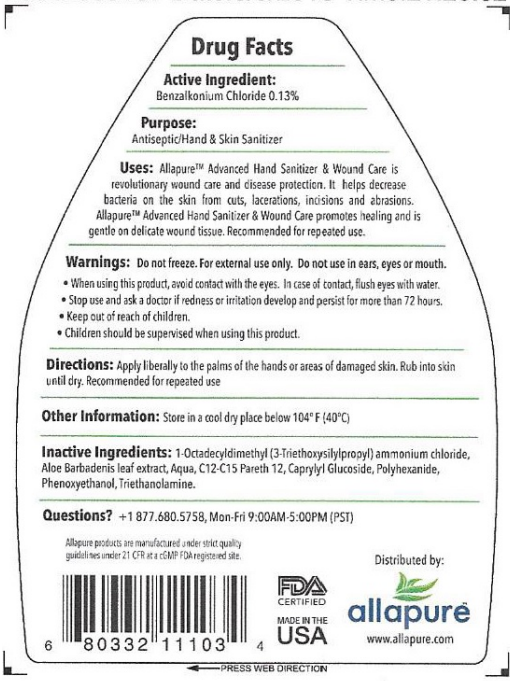
Label:656380 Date:8/21/2018 Time:07:20:16



Allapure Advanced Formula Hand Sanitizer 8.25oz (72552-001-08)

Label:655116 Date:8/9/2018 Time:20:45:42

Label:655157 Date:8/9/2018 Time:21:20:02



ALLAPURE ADVANCED FORMULA HAND SANITIZER

benzalkonium chloride liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:72552-001
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**
ALOE VERA LEAF (UNII: ZY81Z83H0X)	

WATER (UNII: 059QF0KO0R)	
C12-15 PARETH-12 (UNII: 131316X18L)	
CAPRYLYL GLUCOSIDE (UNII: V109WUT6RL)	
POLIHEXANIDE (UNII: 322U039GMF)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
TROLAMINE (UNII: 9O3K93S3TK)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72552-001-01	50 mL in 1 BOTTLE; Type 0: Not a Combination Product	09/24/2018	
2	NDC:72552-001-07	207 mL in 1 BOTTLE; Type 0: Not a Combination Product	09/24/2018	
3	NDC:72552-001-08	245 mL in 1 BOTTLE; Type 0: Not a Combination Product	09/24/2018	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	09/24/2018	

Labeler - Allapure, LLC (081227305)

Revised: 1/2019 Allapure, LLC