SILVER ION ANTIBACTERIAL GEL- silver ion antibacterial gel gel Anson Bio-Technology 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.

75139-007 Silver Ion Antibacterial Gel

Main Active Ingredients and Content

Silver ions, 0.05%.

Glycerol, 5%.

Purpose

Antibacterial

Scope of Use

The product is suitable for antibacterial skin surface

Directions

After cleaning the skin surface, apply it directly to the skin surface for 5minandletitdry naturally without washing after use.

Warning

The product has antibacterial effect on staphylococcus aureus, escherichia coli and candida albicans.

Other Tips

Place away from light, seal and store at room temperature.

Precautions

See instructions for details

other ingredients

Carbomer, PEG 400, Sodium Hydroxide, Purified Water etc.

See instructions for details

Package Label - Principal Display Panel

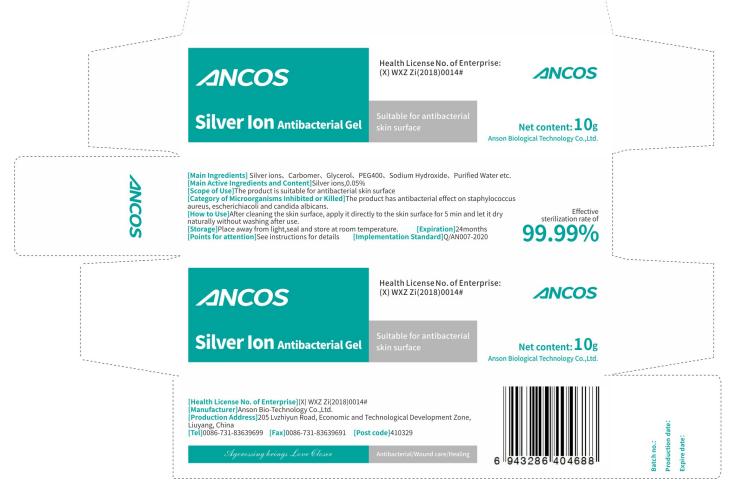
75139-007-01 3G



75139-007-02 5G



75139-007-03 10G



75139-007-04 20



75139-007-05 30G



SILVER ION ANTIBACTERIAL GEL

silver ion antibacterial gel gel

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:75139-007	
Route of Administration	TOPICAL			

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
GLYCERIN (UNII: PDC6A3C0OX) (GLYCERIN - UNII: PDC6A3C0OX)	GLYCERIN	5 g in 100 g	
SILVER CATION (UNII: 57N7B0K90A) (SILVER CATION - UNII:57N7B0K90A)	SILVER CATION	0.05 g in 100 g	

Inactive Ingredients		
Ingredient Name	Strength	
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)		
WATER (UNII: 059QF0KO0R)		

SODIUM HYDROXIDE (UNII: 55X04QC32I)	
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:75139-007- 02	5 g in 1 BOTTLE; Type 0: Not a Combination Product	04/28/2020	
2	NDC:75139-007- 01	3 g in 1 BOTTLE; Type 0: Not a Combination Product	04/28/2020	
3	NDC:75139-007- 05	30 g in 1 BOTTLE; Type 0: Not a Combination Product	04/28/2020	
4	NDC:75139-007- 04	20 g in 1 BOTTLE; Type 0: Not a Combination Product	04/28/2020	
5	NDC:75139-007- 03	10 g in 1 BOTTLE; Type 0: Not a Combination Product	04/28/2020	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part334	04/28/2020	

Labeler - Anson Bio-Technology Co.,Ltd. (547087171)

Establishment				
Name	Address	ID/FEI	Business Operations	
Anson Bio-Technology Co.,Ltd.		547087171	manufacture(75139-007)	

Revised: 4/2021 Anson Bio-Technology Co.,Ltd.