AIR RAID CARD ANTIBACTERIAL HAND SANITIZER- antibacterial hand sanitizer liquid
Jiangsu Manwei Pharmaceutical Co.,Ltd
Reference Label Set Id: a52591f0-7f38-280e-e053-2a95a90ad0bd

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.

Antibacterial Hand Sanitizer

Active Ingredient(s)

CHLOROXYLENOL 0.3%,SALICYLIC ACID 0.5%.

Purpose

Antiseptic, Hand Sanitizer

scope of application

applicable to hand hygiene

wet hands with water first, then take appropriate hand sanitizer in the hands to scrub,finally rinse with water.

Matters needing attention

This product is for external use only.

Do not use

It is forbidden to be taken orally. It is forbidden for people allergic to this product

Stop use and ask a doctor if irritation or rash occurs. These may be signs of a serious condition.

Do not use for children under 3 years old

Directions

wet hands with water first, then take appropriate hand sanitizer in the hands to scrub,finally rinse with water.

Other information

Store in a cool and ventilated place,

Inactive ingredients

PROPYLENE GLYCOL,

ETHYL BENZOATE,

EDETATE DISODIUM ANHYDROUS,

BENZYL ALCOHOL,

GLYCOSINE,

2-(METHYLAMINO)ETHANOL,
SODIUM CHLORIDE,
WATER,
SODIUM LAURETH-3 SULFATE,
POTASSIUM HYDROXIDE

Package Label - Principal Display Panel

AIR RAID CARD ANTIBACTERIAL HAND SANITIZER

antibacterial hand sanitizer liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:75086-117
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
CHLOROXYLENOL (UNII: 0F32U78V2Q) (CHLOROXYLENOL - UNII:0F32U78V2Q)	CHLOROXYLENOL	0.2 g in 100 mL
SALICYLIC ACID (UNII: O414PZ4LPZ) (SALICYLIC ACID - UNII:O414PZ4LPZ)	SALICYLIC ACID	0.2 g in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
SODIUM LAURETH-3 SULFATE (UNII: BPV390UAP0)	
WATER (UNII: 059QF0KO0R)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
POTASSIUM HYDROXIDE (UNII: WZH3C48M4T)	
METHYL BENZOATE (UNII: 6618K1VJ9T)	
EDETATE DISODIUM ANHYDROUS (UNII: 8NLQ36F6MM)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
GLYCOSINE (UNII: D5JUH3HNWF)	

2-(METHYLAMINO)ETHANOL (UNII: ZMQ4G4V497)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:75086-117-01	30 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/20/2020	
2	NDC:75086-117-02	60 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/20/2020	
3	NDC:75086-117-03	250 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/20/2020	
4	NDC:75086-117-04	300 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/20/2020	
5	NDC:75086-117-05	350 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/20/2020	
6	NDC:75086-117-06	500 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/20/2020	
7	NDC:75086-117-07	3800 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/20/2020	
8	NDC:75086-117-08	5000 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/20/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	04/20/2020	

Labeler - Jiangsu Manwei Pharmaceutical Co.,Ltd (554529430)

Establishment

Name	Address	ID/FEI	Business Operations
Jiangsu Manwei Pharmaceutical Co.,Ltd		554529430	manufacture(75086-117)

Revised: 5/2020 Jiangsu Manwei Pharmaceutical Co.,Ltd