HAND SANITIZING WIPES- ethyl alcohol cloth
N A Artis Inc

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 Ingredients

Ethyl Alcohol 75% v/v.

Purpose

Antiseptic

Warnings

For external use only.

Flammable, keep away from heat or flame.

Do not use

- on open skin wounds.

When using this product, keep out of eyes, ears and mouth. In case of contact with eyes,rinse thoroughly with water immediately.

Stop using and consult a doctor if irritation or rash occurs, these may be sighs of a serious condition.

- Discard used wipes immediately to prevent cross contamination.
- Children should be under adult supervision when using this product.
- Not recommended for infants.

Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center immediately.

Uses

- Decrease bacteria on the skin that could cause disease.
- For use when soap and water are not available.

Directions

- Open the pouch.
- Pull out wipe and close the pouch.
- Unfold and wipe hands, fingers and wrists thoroughly and allow to dry.
- For dirty hands, use first wipe to clean hands, then discard wipe; sanitize with second wipe.
- Discard after single use.
- Make sure to close the pouch firmly after use to retain moisture.

Other Information

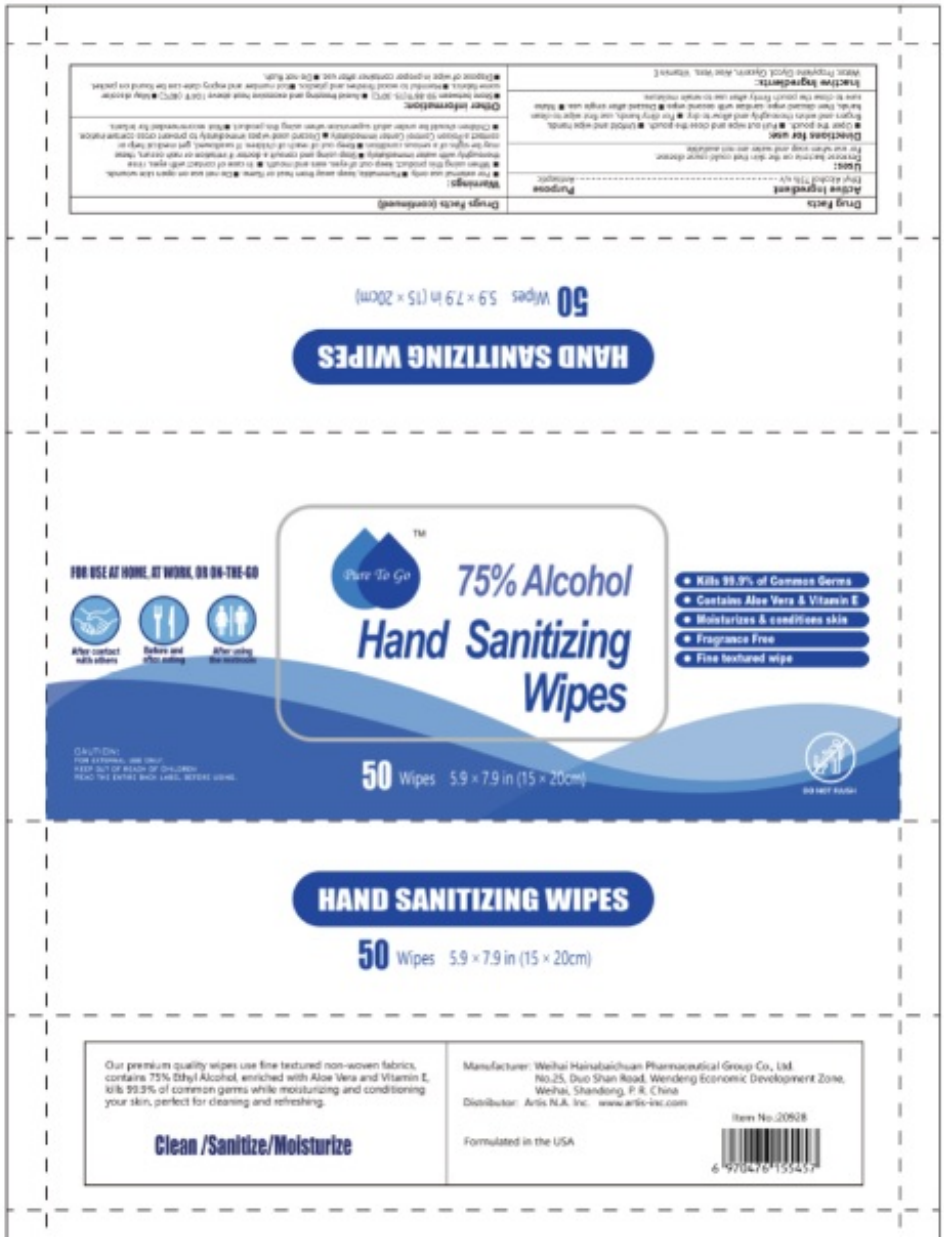
- Store between 59-86□(15-30□)
- Avoid freezing and excessive heat above 104□ (40□)
- May discolor some fabrics.

- Harmful to wood finishes and plastics.
- Lot number and expiry date can be found on packet.
- Dispose of wipe in proper container after use.
- Do not flush.

Inactive Ingredients

Water, Propylene Glycol, Glycerin, Aloe Vera, Vitamin E

Image



HAND SANITIZING WIPES

ethyl alcohol cloth

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:79798-001

Route of Administration	TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	75 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
.ALPHA.-TOCOPHEROL (UNII: H4N855PNZ1)	
ALOE (UNII: V5VD430YW9)	
GLYCERIN (UNII: PDC6A3C0OX)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:79798-001-01	5 in 1 PACKET	07/23/2020	
1		3.6 mL in 1 NOT APPLICABLE; Type 0: Not a Combination Product		
2	NDC:79798-001-02	10 in 1 PACKET	07/23/2020	
2		3.6 mL in 1 NOT APPLICABLE; Type 0: Not a Combination Product		
3	NDC:79798-001-03	20 in 1 PACKET	07/23/2020	
3		3.6 mL in 1 NOT APPLICABLE; Type 0: Not a Combination Product		
4	NDC:79798-001-04	50 in 1 PACKET	07/23/2020	
4		3.6 mL in 1 NOT APPLICABLE; Type 0: Not a Combination Product		
5	NDC:79798-001-05	80 in 1 PACKET	07/23/2020	
5		3.6 mL in 1 NOT APPLICABLE; Type 0: Not a Combination Product		
6	NDC:79798-001-06	80 in 1 CANISTER	07/23/2020	
6		3.6 mL in 1 NOT APPLICABLE; Type 0: Not a Combination Product		
7	NDC:79798-001-07	160 in 1 CANISTER	07/23/2020	
7		3.6 mL in 1 NOT APPLICABLE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	07/23/2020	

Labeler - N A Artis Inc (081009519)

Revised: 7/2020 N A Artis Inc