WET WIPES- wet wipes cloth Shandong Tricol Marine Biological 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.

Uses

- 1. Hand sanitizer to help reducebacteria on the skin.
- 2. For use when soap and water are not available.

For external use only. Flammable. Keep away from heat or flame.

Do not use on open skin wounds.

Do not use in children less than 2months of age.

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

Stop use and ask a doctor irritation or rash occours.

Keep out of reach of children. If swallowed, get medical help, or contact a Poison Control Center right away.

Directions

- 1.open resealable label and pull one sheet from pack.
- 2. Thoroughly clean hands or affected area and discard in receptacle. Do not flush.
- 3. Allow to dry without wiping.
- 4.Be sure to reseal label completely to retain moisture.

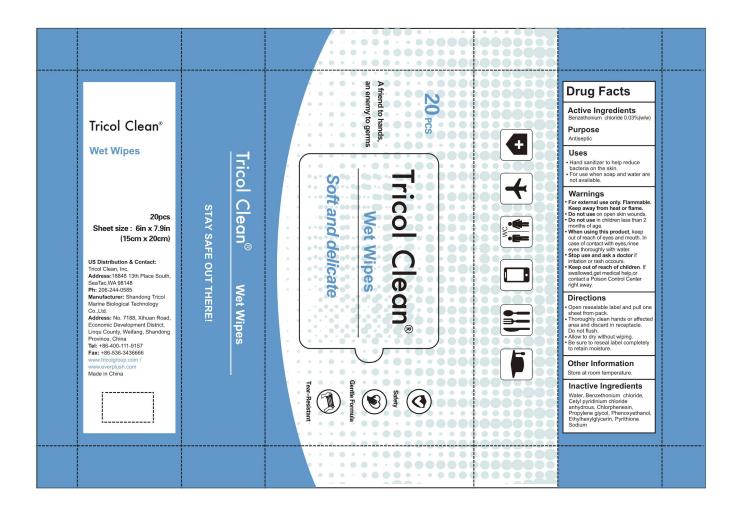
Other Information

Store at room temperature.

Water, Benzethonium chloride, Cetylpyridinium chloride anhydrous, Chlorphenesin, Propylene glycol, Phenoxyethanol, Ethylhexylglycerin, Pyrithione Sodium

Benzethonium chloride

Antiseptic



wet wipes cloth

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:54156-001	
Route of Administration	TOPICAL			

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
BENZETHO NIUM CHLO RIDE (UNII: PH41D05744) (BENZETHONIUM - UNII:1VU15B70BP)	BENZETHONIUM CHLORIDE	0.03 g in 100 g		

Inactive Ingredients		
Ingredient Name	Strength	
WATER (UNII: 059QF0KO0R)		
PYRITHIONE SODIUM (UNII: 6L3991491R)		
CHLORPHENESIN (UNII: 1670 DAL4SZ)		
CETYLPYRIDINIUM CHLORIDE ANHYDROUS (UNII: 6BR7T22E2S)		
PHENO XYETHANO L (UNII: HIE492ZZ3T)		
PROPYLENE GLYCOL (UNII: 6 DC9 Q16 7 V3)		
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)		

l	Packaging			
l	# Item Code	Package Description	Marketing Start Date	Marketing End Date
l	1 NDC:54156-001-01	20 in 1 PACKET	11/05/2020	
	1	0.03 g in 1 PATCH; 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	part333E	11/05/2020		

Labeler - Shandong Tricol Marine Biological Technology Co.,Ltd (541569370)

Establishment				
Name	Address	ID/FEI	Business Operations	
Shandong Tricol Marine Biological Technology Co.,Ltd		541569370	manufacture(54156-001)	

Revised: 11/2020 Shandong Tricol Marine Biological Technology Co.,Ltd