BOLOR SANITIZING HAND WIPES- bolor sanitizing hand wipes solution Guangzhou Baihua Co., Ltd

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.

Active ingredient Purpose

Benzalkonium chloride 0.12% Antimicrobial

Purpose

Antimicrobial

Directions Take wipe and rub thoroughly over all surfaces of both hands. Wet hands thoroughly with wipe. Rub hands together briskly to dry your hands. Dispose of wipe. Do not flush.

Warnings For external use only

Inactive ingredients Aloe Barbadensis Leaf Extract, Centella AsiaOca Extract, Chlorhexidine Diacetate, Deionized Water, Ethylhexylglycerin, Glycerol, Lonicera Caprifolium Extract.

When using this product avoid contact with eyes. In case of contact, flush eyes with water

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away. Children under 6 years age should be supervised when using this product.

Drug Facts Active ingredient Purpose Benzalkonium chloride 0.12% Antimicrobial Uses Hand sanitizer to help reduce bacteria on skin Recommended for repeated use When using this product avoid contact with eyes. In case of contact, flush eyes with water. Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away. Children under 6 years age should be supervised when using this product. Stop use and ask a doctor if irritation and/or rash occurs and persists for more than 72 hours. Warnings For external use only Directions Take wipe and rub thoroughly over all surfaces of both hands. Wet hands thoroughly with wipe. Rub hands together briskly to dry your hands. Dispose of wipe. Do not flush. Inactive ingredients Aloe Barbadensis Leaf Extract, Centella AsiaΘca Extract, Chlorhexidine Diacetate, Deionized Water, Ethylhexylglycerin, Glycerol, Lonicera Caprifolium Extract.









bolor sanitizing hand wipes solution

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:77665-002

Route of Administration TOPICAL

Active Ingredient/Active Moiety Ingredient Name BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y) BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII: 7N6JUD5X6Y) BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII: 7N6JUD5X6Y)

Inactive Ingredients				
Ingredient Name	Strength			
CENTELLA ASIATICA (UNII: 7M867G6T1U)				
LONICERA CAPRIFOLIUM FLOWERING TOP (UNII: 1X0T378SXY)				
WATER (UNII: 059QF0KO0R)				
GLYCEROL FORMAL (UNII: 3L7GR2604E)				
ALOE BARBADENSIS LEAF JUICE (UNII: RUE8E6T4NB)				

Packaging					
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:77665-002- 02	60 in 1 PACKET	04/01/2025		
1	NDC:77665-002- 01	100 mg in 1 PACKET; Type 0: Not a Combination Product			
2	NDC:77665-002- 04	30 in 1 PACKET	04/01/2025		
2	NDC:77665-002- 03	100 mg in 1 PACKET; Type 0: Not a Combination Product			
3	NDC:77665-002- 06	10 in 1 PACKET	04/01/2025		
3	NDC:77665-002- 05	100 mg in 1 PACKET; Type 0: Not a Combination Product			

Marketing Information					
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
unapproved drug other		04/01/2025			

Labeler - Guangzhou Baihua Co., Ltd (545015588)

Establishment							
Name	Address	ID/FEI	Business Operations				
Guangzhou Baihua Co., Ltd		545015588	manufacture(77665-002)				

Revised: 5/2025 Guangzhou Baihua Co., Ltd