HAND SANITIZING WIPES- benzalkonium chloride swab
HANGZHOU GUOGUANG TOURING COMMODITY 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.

001

Active ingredient

Benzalkonium chloride 0.13%

Purpose

Antibacterial

Use

For hand sanitizing to decrease bacteria on the skin

recommended for repeated use

Warnings

For external use only.
When useing this product avoid contact with eyes.In case of eye contact,flush eyes with water.
Stop use and ask a doctor if irritation or redness develops.
Keep out of reach of children.
If swallowed, get medical help or contact a Poison Control Center right away.

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Directions

Take wipe and rub thoroughly over all surfaces of both hands,Rub hands together briskly to dry .Dispose of wipe.

Purified Water ,Glycerin,Aloe Barbadensis Leaf Juice,Anionic surfactant.

HAND SANITIZING WIPES

benzalkonium chloride swab

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:61312-001
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.13 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
GLYCERIN (UNII: PDC6A3C0OX)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
SODIUM DODECYLBENZENESULFONATE (UNII: 554127163Y)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:61312-001-01	10 in 1 BOTTLE	01/02/2018	
1		0.00468 g in 1 POUCH; Type 0: Not a Combination Product		
2	NDC:61312-001-02	25 in 1 BOTTLE	01/02/2018	
2		0.00468 g in 1 POUCH; Type 0: Not a Combination Product		
3	NDC:61312-001-03	40 in 1 BOTTLE	01/02/2018	
3		0.00468 g in 1 POUCH; Type 0: Not a Combination Product		
4	NDC:61312-001-04	60 in 1 BOTTLE	01/02/2018	
4		0.00468 g in 1 POUCH; Type 0: Not a Combination Product		
5	NDC:61312-001-05	100 in 1 BOTTLE	01/02/2018	
5		0.00468 g in 1 POUCH; Type 0: Not a Combination Product		
6	NDC:61312-001-06	120 in 1 BOTTLE	01/02/2018	
6		0.00468 g in 1 POUCH; Type 0: Not a Combination Product		
7	NDC:61312-001-07	200 in 1 BOTTLE	01/02/2018	
7		0.00468 g in 1 POUCH; Type 0: Not a Combination Product		
8	NDC:61312-001-08	240 in 1 BOTTLE	01/02/2018	
8		0.00468 g in 1 POUCH; Type 0: Not a Combination Product		
9	NDC:61312-001-09	250 in 1 BOTTLE	01/02/2018	

9		0.00468 g in 1 POUCH; Type 0: Not a Combination Product		
10	NDC:61312-001-10	800 in 1 BOTTLE	01/02/2018	
10		0.00468 g in 1 POUCH; Type 0: Not a Combination Product		
11	NDC:61312-001-11	1200 in 1 BOTTLE	01/02/2018	
11		0.00468 g in 1 POUCH; Type 0: Not a Combination Product		
12	NDC:61312-001-12	1500 in 1 BOTTLE	01/02/2018	
12		0.00468 g in 1 POUCH; 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	03/14/2013	

Labeler - HANGZHOU GUOGUANG TOURING COMMODITY CO., LTD. (526890634)

Establishment

Name	Address	ID/FEI	Business Operations
HANGZHOU GUOGUANG TOURING COMMODITY CO., LTD.		526890634	manufacture(61312-001)

Revised: 1/2018 HANGZHOU GUOGUANG TOURING COMMODITY CO., LTD.