

XTRACARE WET WIPES- benzethonium chloride swab
China Ningbo Shangge Cosmetic Technology Corp

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.

XtraCare Wet Wipes

Active Ingredient

Purpose

Benzethonium Chloride 0.3% Antibacterial

Uses

hand sanitizer to reduce bacteria on the skin.

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

XtraCare Wet Wipes

Antibacterial

Fresh Citrus Scent

Pocket Size

Kills 99.9% of Germs

Warnings

For external use only.

Do not use if you are allergic to any of the ingredients.

When using this product do not get into eyes. If eye contact occurs, rinse thoroughly with water.

Stop use and ask a doctor if irritation and redness develops, and continues for more than 72 hours.

Directions

for adults and children 2 years and older

- apply to hands
- allow to dry without wiping
- ask a doctor before using on children under 2 years

Inactive Ingredients

water (aqua), ethyl alcohol, propylene glycol, PEG-40 hydrogenated castor oil, fragrance, disodium EDTA, aloe barbadensis leaf juice, methylparaben, propylparaben

Questions/comments? 1-855-345-5575

DISTRIBUTED BY

REJOICE INTERNATIONAL INC

NORTHVILLE, MI 48168 USA

MADE IN CHINA





Drug Facts

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XTRACARE WET WIPES

benzethonium chloride swab

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:58503-062
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZETHONIUM CHLORIDE (UNII: PH41D05744) (BENZETHONIUM - UNII:1VU15B70BP)	BENZETHONIUM CHLORIDE	3 mg in 100 mg

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
ALCOHOL (UNII: 3K9958V90M)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
POLYOXYL 40 HYDROGENATED CASTOR OIL (UNII: 7YC686GQ8F)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:58503-062-01	18 in 1 PACKAGE	03/13/2014	
1		100 mg in 1 APPLICATOR; 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	03/13/2014	

Labeler - China Ningbo Shangge Cosmetic Technology Corp (529287434)

Establishment

Name	Address	ID/FEI	Business Operations
China Ningbo Shangge Cosmetic Technology Corp		529287434	manufacture(58503-062)

