BENZALKONIUM CHLORIDE AND LIDOCAINE HYDROCHLORIDE - benzalkonium chloride and lidocaine hydrochloride liquid CVS Pharmacy

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.

drug facts

Active ingredient Purpose Benzalkonium Cl 0.13% w/w......Antiseptic lidocaine HCl 2.5% w/w......Pain reliever

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Uses first aid to help prevent skin infection, and for temporary relief of pain and itching associated with minor - cuts - scrapes - burns

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

Directions - adults and children 2 years and older; clean the affected area; apply a small amount on the area 1-3 times daily; may be covered with a sterile bandage (let dry first) - children under 2 years, ask a doctor.

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Warnings For external use only.

Ask a doctor before use if you have - deep or puncture wounds - animal bites - serious burns

When using this product - do not use in or near the eyes - do not apply over large areas of the body or in large quantities - do not apply over raw surfaces or blistered areas.

Stop use and ask a doctor if - condition worsens - symptoms persist for more than 7 days, or clear up and occur again within a few days

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Other Information protect from excessive heat

Inactive Ingredients disodium EDTA, fragrance, nonoxynol-9, propylene glycol, purified water





benzalkonium chloride and lidocaine hydrochloride liquid

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:59779-255

Route of Administration TOPICAL

Active Ingredient/Active Moiety Ingredient Name Basis of Strength BENZALKO NIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM UNII:7N6 JUD5X6Y) BENZALKONIUM CHLORIDE in 1 mL LIDO CAINE HYDRO CHLORIDE (UNII: V13007Z41A) (LIDO CAINE - UNII:98 PI200987) LIDO CAINE HYDRO CHLORIDE 25 mg in 1 mL

Inactive Ingredients				
Ingredient Name	Strength			
WATER (UNII: 059QF0KO0R)				
PROPYLENE GLYCOL (UNII: 6 DC9 Q167V3)				
EDETATE DISO DIUM (UNII: 7FLD9 1C86K)				
NONOXYNOL-9 (UNII: 48Q180SH9T)				

ı	Packaging						
ı	# Item Code	Package Description	Marketing Start Date	Marketing End Date			
ı	1 NDC:59779-255-06	120 mL in 1 BOTTLE, DISPENSING					

Marketing Information						
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date			
OTC monograph final	part333A	07/08/2010				

Labeler - CVS Pharmacy (062312574)

Registrant - Pharma Pac, LLC (140807475)

Establishment						
Name	Address	ID/FEI	Business Operations			
Pharma Pac, LLC		140807475	manufacture			

Revised: 7/2010 CVS Pharmacy