

CVS HEALTH ANTISEPTIC WOUND WASH- lidocaine hydrochloride, benzalkonium chloride rinse

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

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Active Ingredients

Benzalkonium Cl 0.13% w/w

Lidocaine HCl 2.5% w/w

Purpose

First aid antiseptic

Pain reliever

Uses

First aid to help prevent bacterial contamination or skin infections, and temporary relief of pain and itching associated with minor:

cuts

scrapes

burns

sunburns

skin irritations

Warnings

For external use only

Ask doctor before use if you have

deep or puncture wounds

animal bites

serious burns

When using this product

do not use 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.

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 to 3 times daily.

may be covered with a sterile bandage

if bandaged, let dry first.

Other Information

Protect from excessive heat.

Questions or comments

1-800-635-3696

Inactive Ingredients

Camphor, Deionized Water, Propylene Glycol



CVS HEALTH ANTISEPTIC WOUND WASH

lidocaine hydrochloride, benzalkonium chloride rinse

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:69842-925
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LIDOCAINE HYDROCHLORIDE (UNII: V13007Z41A) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE HYDROCHLORIDE	0.025 g in 1 g
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.0013 g in 1 g

Inactive Ingredients

Ingredient Name	Strength
CAMPHOR (SYNTHETIC) (UNII: 5TJD82A1ET)	
WATER (UNII: 059QF0K00R)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69842-925-01	177 g in 1 BOTTLE; Type 0: Not a Combination Product	03/01/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part348	03/01/2017	

Labeler - CVS Pharmacy (062312574)

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