GOOD NEIGHBOR PHARMACY ALOE VERA- lidocaine hydrochloride gel
Amerisourcebergen Drug Corporation

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.

Good Neighbor Pharmacy Aloe Vera

Drug Facts

Active Ingredients

Lidocaine Hydrochloride 0.5%

Pain Reliever

Temporary relief of pain and itching due to sunburn, minor burns, insect bites, cuts, scrapes

For external use only

Do not use in large quantities, particularly over raw or blistered areas.

When using this product

• avoid contact with eyes.

Rinse with water if contact occurs.

Stop use and ask a doctor if

• condition worsens

or clear up and occur again within a few days.

• if symptoms persist for more than 7 days.

Keep out of reach of children.

If product is swallowed, get medical help or contact a Poison Control Center right away.

• adults and children 2 years and older: apply to affected area not more than 3 - 4 times a day. • children under 2 years of age: consult a physician

Aloe Barbadensis Leaf Juice, Carbomer, Diazolidinyl Urea, Disodium EDTA, FD&C Blue #1, FD&C Yellow #5, Glycerin, Isopropyl Alcohol (0.05% v/v), Menthol, Polysorbate 80, Propylene Glycol, Triethanolamine, Water.

This product is not manufactured or distributed by Schering-Plough

Healthcare Products, Inc., owner of the Solarcaine® trademark.

Distributed By

AmerisourceBergen

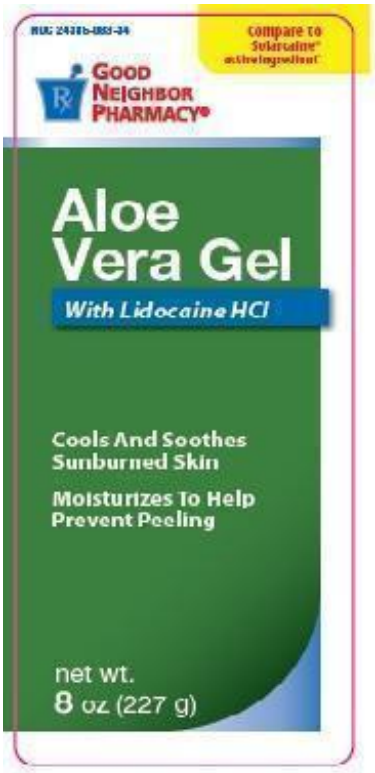
1300 Morris Drive

Chesterbrook, PA 19087

Visit us at

www.goodneighborpharmacy.com

Good Neighbor Pharmacy Aloe Vera Gel



GOOD NEIGHBOR PHARMACY ALOE VERA

lidocaine hydrochloride gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:24385-083
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
LIDOCAINE HYDROCHLORIDE (UNII: V13007Z41A) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE HYDROCHLORIDE	500 g in 100000 mg

Inactive Ingredients	
Ingredient Name	**Strength**
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
CARBOMER 940 (UNII: 4Q93RCW27E)	
DIAZOLIDINYL UREA (UNII: H5RIZ3MPW4)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	

FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
GLYCERIN (UNII: PDC6A3C0OX)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
MENTHOL (UNII: L7T10EIP3A)	
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
TROLAMINE (UNII: 9O3K93S3TK)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:24385-083-34	500 mg in 1 BOTTLE; 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	part348	11/28/1988	

Labeler - Amerisourcebergen Drug Corporation (007914906)

Establishment

Name	Address	ID/FEI	Business Operations
Product Quest Mfg, LLC		927768135	manufacture(24385-083) , label(24385-083)

Revised: 7/2015 Amerisourcebergen Drug Corporation