SUNBURN RELIEF- lidocaine hcl 0.5% gel
Old East Main Co.

Dollar General 005.003/005AE
Sunburn Relief Gel

Active ingredient

Lidocaine HCl 0.5%

Purpose

External analgesic

Uses

for the temporary relief of pain and itching associated with

- minor burns
- sunburn
- minor cuts
- scrapes
- insect bites
- minor skin irritations

Warnings

For external use only

Do not use

in large quantities, particularly over raw surfaces or blistered areas

When using this product

avoid contact with the eyes

Stop use and ask a doctor if

condition worsens, or if 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 of age and older: apply to affected area not more than 3 to 4 times daily
- children under 2 years of age: ask a doctor

Inactive ingredients

water, propylene glycol, glycerin, Aloe Barbadensis Leaf Juice, sodium hydroxide, isopropyl alcohol, polysorbate 80, carbomer, phenoxyethanol, benzyl alcohol, menthol, disodium EDTA, blue 1, yellow 5

Adverse reaction

Distributed By

Old East Main Co.

100 MISSION RIDGE,

GOODLETTSVILLE, TN 37072

100% Satisfction Guaranteed! (888)309-9030

principal display panel

DG™|health

SUNBURN RELIEF

PAIN RELIEVING GEL

with ALOE

Helps relieve the pain, discomfort and itchiness resulting from sunburn

NET WT 8 OZ (226 g)

SUNBURN RELIEF

lidocaine hcl 0.5% gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:55910-177
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
LIDOCAINE HYDROCHLORIDE (UNII: V13007Z41A) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE HYDROCHLORIDE ANHYDROUS	5 mg in 1 g

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
GLYCERIN (UNII: PDC6A3C0OX)	

ALOE VERA LEAF (UNII: ZY81Z83H0X)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
MENTHOL (UNII: L7T10EIP3A)	
EDETATE DISODIUM ANHYDROUS (UNII: 8NLQ36F6MM)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:55910-177-34	226 g in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	12/30/2019	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	12/30/2019	

Labeler - Old East Main Co. (068331990)

Registrant - Nice-Pak Products, LLC (119091520)

Establishment

Name	Address	ID/FEI	Business Operations
Nice-Pak Products, LLC		119091514	manufacture(55910-177)

Revised: 3/2026 Old East Main Co.