SOLARCAINE BURN PAIN RELIEF 4OZ- solarcaine burn pain relief 4oz spray
WellSpring Pharmaceutical Corporation

Solarcaine Burn Pain Relief Spray 4oz

Active Ingredients (w/w)

Lidocaine HCl 4%

Purpose

External Analgesic

Uses

temporarily relieves pain and itching due to:

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

Warnings

For external use only. When using this product

Do not use

in large quantities, particularly over raw surfaces or blistered areas

When using this product

- keep out of eyes
- use only as directed. Intentional misuse by deliberately concentrating and inhaling the contents can be harmful or fatal.
- do not puncture or incinerate. Contents under pressure. Avoid prolonged storage above 40°C (104°F).

Stop use and ask a doctor if

- condition gets worse
- symptoms last more than 7 days
- symptoms clear up and occur again in a few days

Keep out of reach of children

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

Directions

- shake well
- 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: do not use, ask a doctor
- to apply to face, spray in palm of hand and gently apply

Inactive ingredients

purified water, propylene glycol, aloe barbadensis leaf juice, polysorbate 80, ethylhexylglycerin, tocopheryl acetate (vitamin E acetate), 2-phenoxyethanol, sodium hydroxide

Questions?

1-844-241-5454

Distributed by:

Distributed by:

WellSpring Pharmaceutical Corporation
Sarasota, FL 34243

MONEY BACK GUARANTEE

Solarcaine is FSA/HSA eligible

PACKAGE LABEL. PRINCIPAL DISPLAY PANEL

#1 Pharmacist Recommended*

* Pharmacy Times/* U.S News & World

Report 2023-2024

DOT 2Q M5706

L50200A | B6522

CONTAINS NO CFCs WHICH
DEPLETE THE OZONE LAYER

Solarcaine 4%

SOLARCAINE BURN PAIN RELIEF 4OZ

solarcaine burn pain relief 4oz spray

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:65197-502
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
LIDOCAINE HYDROCHLORIDE (UNII: V13007Z41A) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE HYDROCHLORIDE ANHYDROUS	4 g in 100 g

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
POLYSORBATE 81 (UNII: 2MSF640LWM)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	

ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:65197-502-04	113 g in 1 CAN; Type 0: Not a Combination Product	01/01/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	01/01/2024	

Labeler - WellSpring Pharmaceutical Corporation (110999054)

Revised: 6/2023

WellSpring Pharmaceutical Corporation