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

Solarcaine After Sun Pain Relief

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

L50210A | B6526

CONTAINS NO CFCs WHICH
DEPLETE THE OZONE LAYER

Solarcaine After sun pain relief

SOLARCAINE BURN PAIN RELIEF 4OZ

solarcaine burn pain relief 4oz spray

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:65197-512
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 80 (UNII: 6OZP39ZG8H)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	

ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:65197-512-04	113 g in 1 CAN; Type 0: Not a Combination Product	11/15/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	11/15/2024	

Labeler - WellSpring Pharmaceutical Corporation (110999054)

Revised: 6/2024

WellSpring Pharmaceutical Corporation