

WALG AFTRSN BURN RELIEF- lidocaine hydrochloride spray

Walgreen Company

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.

WAG Aftrsn Burn Relief Spray 8oz

Product

WALGREENS AFTERSUN BURN RELIEF SPRAY

Active Ingredients

Lidocaine Hydrochloride 0.5%

Purpose

Topical Anesthetic

Uses:

- Temporary relief of pain and itching
- Helps to relieve and soothe pain from sunburn, minor burns, skin irritations, scrapes, insect bites.

Warnings:

For external use only

Do not use

in large quantities, particularly over raw surfaces or areas with blisters.

When using this product:

- Avoid contact with eyes.

Stop use and ask a doctor if:

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 or older: apply to affected area not more than 3-4 times daily.
- Children under 2 years of age: do not use; ask a doctor.

Inactive Ingredients

Deionized Water, Glycerin, Propanediol, Aloe Barbadensis Leaf Juice, Carbomer, Tocopherol Acetate, PEG-40 Hydrogenated Castor Oil, Sodium Lauroyl Sarcosinate, Disodium EDTA, Phenoxyethanol, Sodium Hydroxide

Questions or comments?

www.walgreens.com or call toll free 1-800-XXX-XXXX

PRINCIPAL DISPLAY PANEL

Walgreens
Sunburn Relief
CONTINUOUS SPRAY
LIDOCAINE 0.5% /
EXTERNAL ANALGESIC
PAIN RELIEF
LIDOCAINE
8 OZ (227 g)

Drug Facts	
Active ingredient Lidocaine 0.50%	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 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. • Do not puncture or incise skin. Contents under pressure. Do not store at temperature above 120°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 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: ask a doctor • To apply to face, spray on palm of hand and gently apply	
Inactive ingredients: Water, Glycerin, Propylolol, Aloe Barbadensis Leaf Juice, Carbomer, Tocopheryl Acetate, PEG-40 Hydrogenated Castor Oil, Sodium Lauryl Sarcosinate, Disodium EDTA, Phenoxethanol, Sodium Hydroxide	
Questions or Comments? 1-800-45-4733	

Walgreens

Sunburn Relief

CONTINUOUS SPRAY

LIDOCAINE 0.5% /
EXTERNAL ANALGESIC

PAIN RELIEF

LIDOCAINE

Provides fast relief from:

- Sunburn
- Cuts & scrapes
- Insect bites
- Minor burns



NET WT 8 OZ (227 g)

Walgreens Pharmacist Recommended
Walgreens Pharmacist Survey

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WALG AFTRSN BURN RELIEF

lidocaine hydrochloride spray

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0363-9919
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LIDOCAINE HYDROCHLORIDE (UNII: V13007Z41A) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE HYDROCHLORIDE ANHYDROUS	0.5 mg in 100 mg

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
CARBOXYPOLYMETHYLENE (UNII: 0A5MMB07FC)	
ALPHA-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
POLYOXYL 40 HYDROGENATED CASTOR OIL (UNII: 7YC686GQ8F)	
SODIUM LAUROYL SARCO SINATE (UNII: 632GS99618)	
EDETATE DISODIUM ANHYDROUS (UNII: 8NLQ36F6MM)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0363-9919-36	237 mg in 1 BOTTLE; Type 0: Not a Combination Product	11/30/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH NOT FINAL	part348	11/30/2017	

Labeler - Walgreen Company (008965063)

Registrant - Walgreen Company (008965063)

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