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)

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: 0A5MM307FC)	
ALPHA-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
POLYOXYL 40 HYDROGENATED CASTOR OIL (UNII: 7YC686GQ8F)	
SODIUM LAUROYL SARCOSINATE (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)

Revised: 10/2017 Walgreen Company