#### BURN- lidocaine hydrochloride spray First Aid Only, Inc

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.

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Drug Facts

#### **Active Ingredients**

Lidocaine HCL 2.0%

#### Purpose

**Topical Pain Relief** 

#### Uses

temporary pain relief associated with minor burns

#### Warnings

#### For external use only.

#### Do not use

- In large quantities, particularly over raw or blistered area
- Near eyes, if this happens rinse thoroughly with water

**Stop use and ask doctor if** condition worsens or persists for more than 7 days or clears up and returns

**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; spray an even layer of burn spray over affected area not more than 3-4 times daily
- For children under 2 years of age, consult a physician.

#### **Inactive ingredients**

aloe vera, germaben II. Propylene glycol, purified water

#### **Burn Spray**

#### Pain relieving spray for use on minor burns.

M5082

3 fl. oz. (88.7 ml.)

-----FIRST AID ONLY

Distributed by First Aid Only, Inc.

Vancouver, WA 96882 USA

NDC 62985-5082-1

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BURN						
lidocaine hydrochloride spray						
Product Information						
Product Type	HUMAN OTC DRUG	Item Code (Source) ND		NDC:62985-5	DC:62985-5082	
Route of Administration	TOPICAL					
Active Ingredient/Active Moi	ety					
Ingredient Name			<b>Basis of Strength</b>		Strength	
Lidocaine Hydrochloride (UNII: V13007Z41A) (Lidocaine - UNII:98PI200987)			Lidocaine Hydrochloride Anhydrous		$20\ g\ in\ 1\ L$	
Inactive Ingredients						
	Ingredient Name			Strength		
aloe vera leaf (UNII: ZY81Z83H0X)						
propylene glycol (UNII: 6DC9Q167V3	)					
diazolidinyl urea (UNII: H5RIZ3MPW4	)					
water (UNII: 059QF0KO0R)						

Packaging								
# Item Code	2	Package Description	Marketing Start Date	Marketing End Date				
1 NDC:62985- 5082-1	0.088 Pro du	7 L in 1 BOTTLE, SPRAY; Type 0: Not a Combination	08/13/2014					
Marketing	Inform	nation						
Marketing Marketing Ca		nation Application Number or Monograph Citation	Marketing Start Date	Marketing End Dat				

## Labeler - First Aid Only, Inc (196551634)

## Registrant - Safetec of America, Inc. (874965262)

## Establishment

Name	Address	ID/FEI	Business Operations
Safetec of America, Inc.		874965262	MANUFACTURE(62985-5082)

Revised: 8/2014

First Aid Only, Inc