7-SELECT AFTER SUN LIDICAINE HCL PAIN-RELIEVING WITH ALOE VERA- lidocaine hydrochloride gel
7-11

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.

7-Select After Sun Lidicaine HCl Pain-Relieving Gel with Aloe Vera

Active Ingredient

Lidocaine hydrochloride 0.5%

Purpose

External Analgesic

Uses

temporarily relieves pain and itching due to:

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

Warnings

- **For External Use Only**

Do not use

in large quantities, particularly over raw surfaces or blistered areas.

When using this product

- avoid contact with eyes. If contact occurs, rinse with water to remove.

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

- 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

Inactive Ingredients

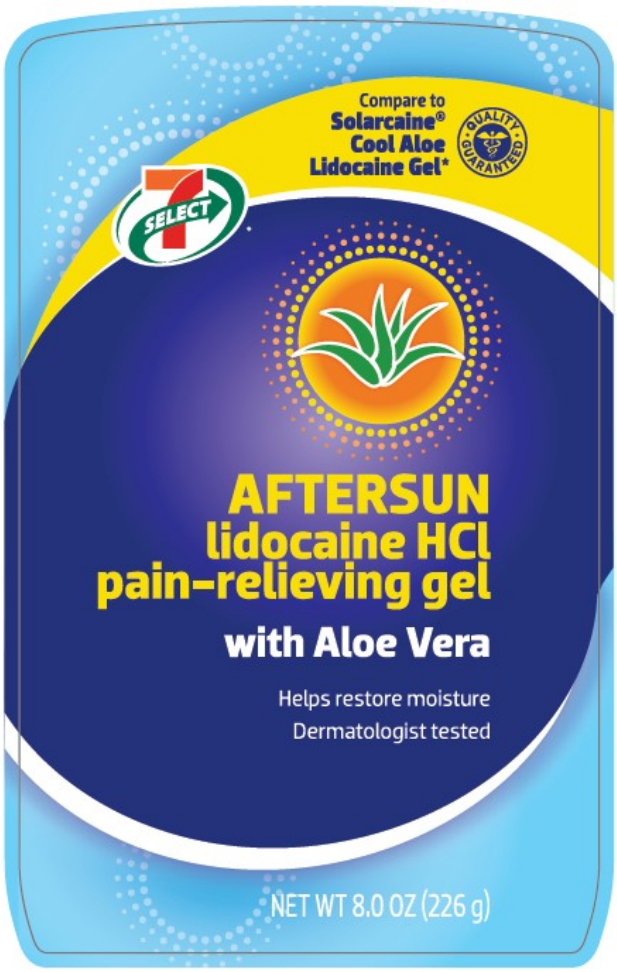
aloe barbadensis leaf juice, water, isopropyl alcohol, propylene glycol, glycerin, triethanolamine, carbomer, polysorbate 80, diazolidinyl urea, menthol, disodium EDTA, yellow 5, blue 1

Label

7-Select After Sun Lidicaine HCl Pain-Relieving Gel with Aloe Vera

8 OZ (226 g)

NDC 10202-950-80



7-SELECT AFTER SUN LIDICAINE HCL PAIN-RELIEVING WITH ALOE VERA

lidocaine hydrochloride gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:10202-950
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**

LIDOCAINE HYDROCHLORIDE (UNII: V13007Z41A) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE HYDROCHLORIDE ANHYDROUS	5 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
TROLAMINE (UNII: 9O3K93S3TK)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	
DIAZOLIDINYL UREA (UNII: H5RIZ3MPW4)	
WATER (UNII: 059QF0KO0R)	
MENTHOL (UNII: L7T10EIP3A)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
CARBOMER INTERPOLYMER TYPE A (55000 CPS) (UNII: 59TL3WG5CO)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
GLYCERIN (UNII: PDC6A3C0OX)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:10202-950-80	226 g in 1 BOTTLE; Type 0: Not a Combination Product	02/21/2019	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part348	02/21/2019	

Labeler - 7-11 (007347602)

Revised: 2/2021 7-11