EHA- lidocaine hydrochloride lotion
Asclemed USA, Inc.

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.

Drug Facts

Active Ingredients

Lidocaine HCI 4%

Purpose

External Anesthetic

Uses

For temporary relief of pain and itching and minor skin irritations due to minor cuts and scrapes, sunburns, and minor burns.

Warning

For external use only.

Avoid contact with eyes

Stop using this product and ask doctor if

- symptoms last for more than seven days, or clear up and occur again within a few days
- if redness, irritation, swelling, pain or other symptoms increase

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Directions

For adults and children two-years or older, apply externally to the affected area. Do not use more than three to four times per day.

Inactive ingredients

Inactive ingredients: Aqua (Deionized Water), C13-14 Isoparaffin, Glyceryl Stearate, Helianthus Annuus (Sunflower) Seed Oil, Isopropyl Myristate, Laureth-7, M Polyacrylamide, Steric Acid.

Principal Display Panel - 88 mL Bottle Label

'Eha Lotion 4%

Pain Relief Lotion

For Professional Use Only

3 OZ (88 mL)

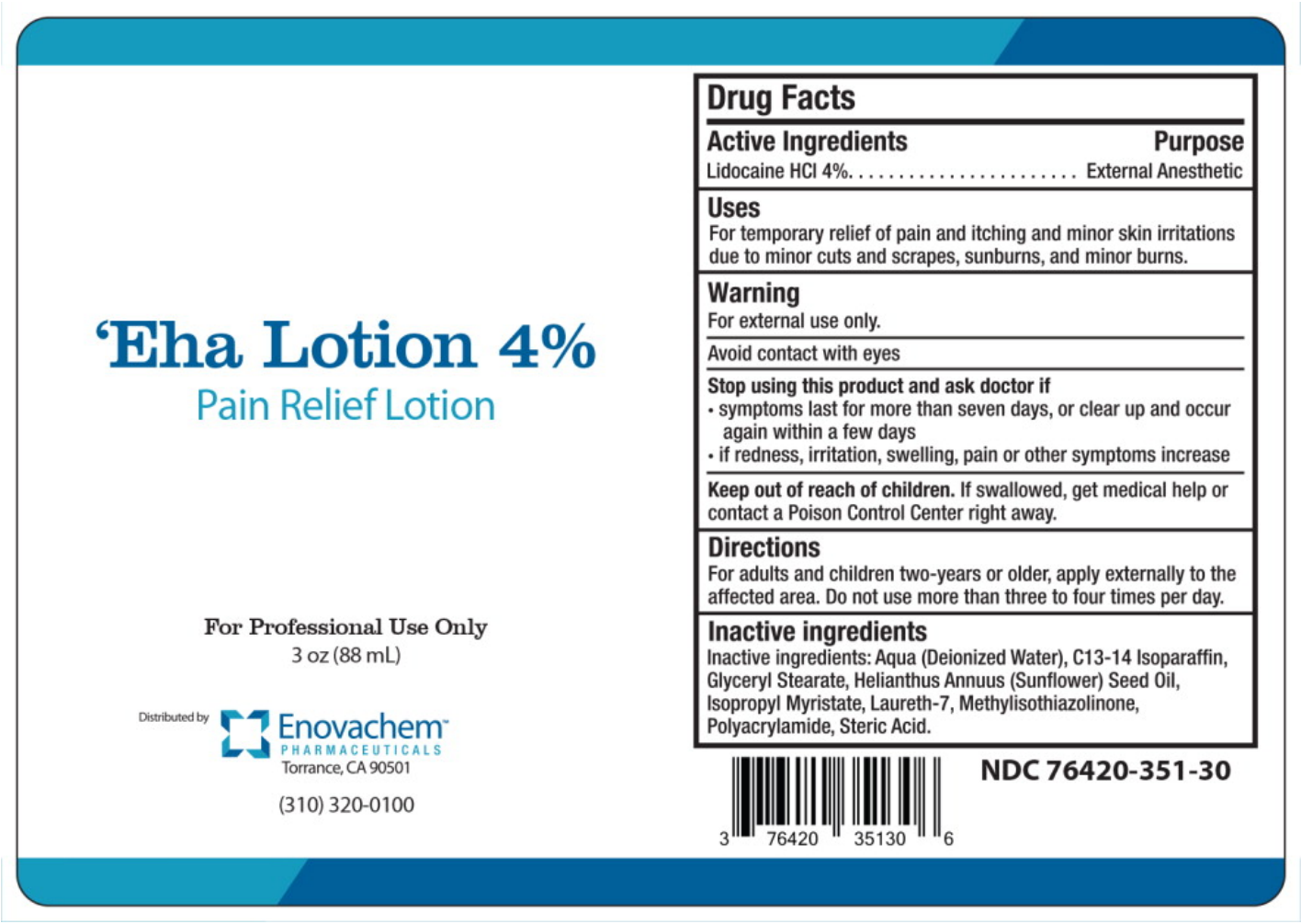
Distributed by

Enovachem™

PHARMACEUTICALS

Torrance, CA 90501

(310) 320-0100



EHA

lidocaine hydrochloride lotion

Product Information			
Product Type	HUMAN PRESCRIPTION DRUG	**Item Code (Source)**	NDC:76420-351
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 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
C13-14 ISOPARAFFIN (UNII: E4F12ROE70)	
GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4)	
HELIANTHUS ANNUUS FLOWERING TOP (UNII: BKJ0J3D1BP)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	
LAURETH-7 (UNII: Z95S6G8201)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	
STEARIC ACID (UNII: 4ELV7Z65AP)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:76420-351-30	88 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/07/2016	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		06/07/2016	

Labeler - Asclemed USA, Inc. (059888437)

Revised: 11/2018 Asclemed USA, Inc.