

HEALTHWISE PAIN RELIEF LIDOCAINE- lidocaine hydrochloride cream
Veridian Healthcare

HealthWise Pain Relief Lidocaine Cream

Active ingredient

Lidocaine HCl 4%

Purpose

Topical analgesic

Use

For the temporarily relief of pain

Warnings

For external use only. Avoid contact with eyes.

Do not use

- in large quantities, particularly over raw surfaces or blistered areas
- on puncture wounds
- for more than one week without consulting a doctor

When using this product

- use only as directed. Read and follow all directions and warnings on this carton.
- do not allow contact with the eyes
- do not bandage or apply local heat (such as heating pads) to the area of use

Stop use and ask a doctor if

- condition worsens
- symptoms persist for more than 7 days or clear up and occur again within a few days

If pregnant or breast-feeding,

ask a health professional before use.

Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center (800-222-1222) right away.

Directions

adults and children over 12 years:

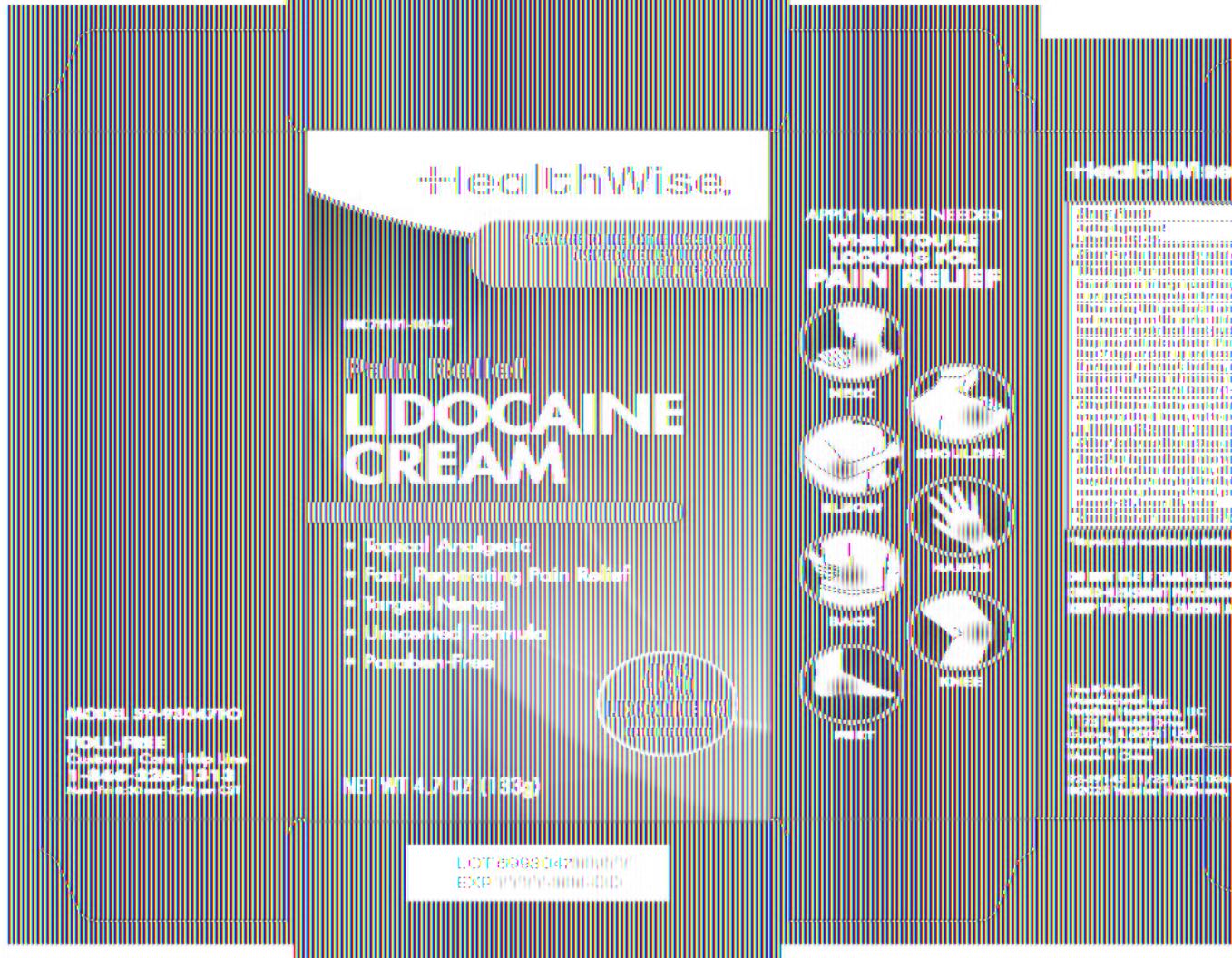
- apply a thin layer to affected area every 6 to 8 hours, not to exceed 3 applications in a 24 hour period

children 12 years or younger: ask a doctor***Inactive ingredients***

Inactive ingredients

aloe vera extract, cetostearyl alcohol, dimethicone, edetate disodium anhydrous, ethyl alcohol, ethylhexylglycerin, hydroxyethyl acrylate/sodium acryloyldimethyl taurate copolymer, iodopropynyl butylcarbamate, monostearin, phenoxyethanol, polyacrylic acid, potassium hydroxide, steareth-2, steareth-21, tween 60, water

PRINCIPAL DISPLAY PANEL



HEALTHWISE PAIN RELIEF LIDOCAINE

lidocaine hydrochloride cream

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:71101-107
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 100 g

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
ALCOHOL (UNII: 3K9958V90M)	
POLYACRYLIC ACID (250000 MW) (UNII: 9G2MAD7J6W)	
HYDROXYETHYL ACRYLATE/SODIUM ACRYLOYLDIMETHYL TAURATE COPOLYMER (45000 MPA.S AT 1%) (UNII: 86FQE96TZ4)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
EDETATE DISODIUM ANHYDROUS (UNII: 8NLQ36F6MM)	
GLYCERYL 1-STEARATE (UNII: 258491E1RZ)	
STEARETH-2 (UNII: V56DFE46J5)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
POLYSORBATE 60 (UNII: CAL22UVI4M)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
STEARETH-21 (UNII: 53J3F32P58)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
POTASSIUM HYDROXIDE (UNII: WZH3C48M4T)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
IODOPROPYNYL BUTYLCARBAMATE (UNII: 603P14DHEB)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71101-107-47	1 in 1 CARTON	01/01/2026	
1		133 g in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	01/01/2026	

Labeler - Veridian Healthcare (830437997)

Establishment

Name	Address	ID/FEI	Business Operations
Foshan Aqua Gel Biotech Co., Ltd.		529128763	manufacture(71101-107)

