PAIN RELIEF ROLL ON EVORA LABS- lidocaine liquid
Evora Worldwide

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.

Pain Releif Lidocaine 4%

Active ingredient **Purpose**

Lidocaine 4%..... Topical analgesic

Uses

For temporary relief of pain and itching

Warnings

For externa use only.

When using this product • use only as directed • do not bandage tightly • avoid contact with
eyes • do not apply to wounds or damaged skin • do not use in large quantities, particularly over raw surface or blistered areas.

Stop use and ask doctor if • condition worsens • symptoms persist for more than 7 days or clear up and occur again within a few days

Flammable • keep away from fire or flame

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 right away.

Directions

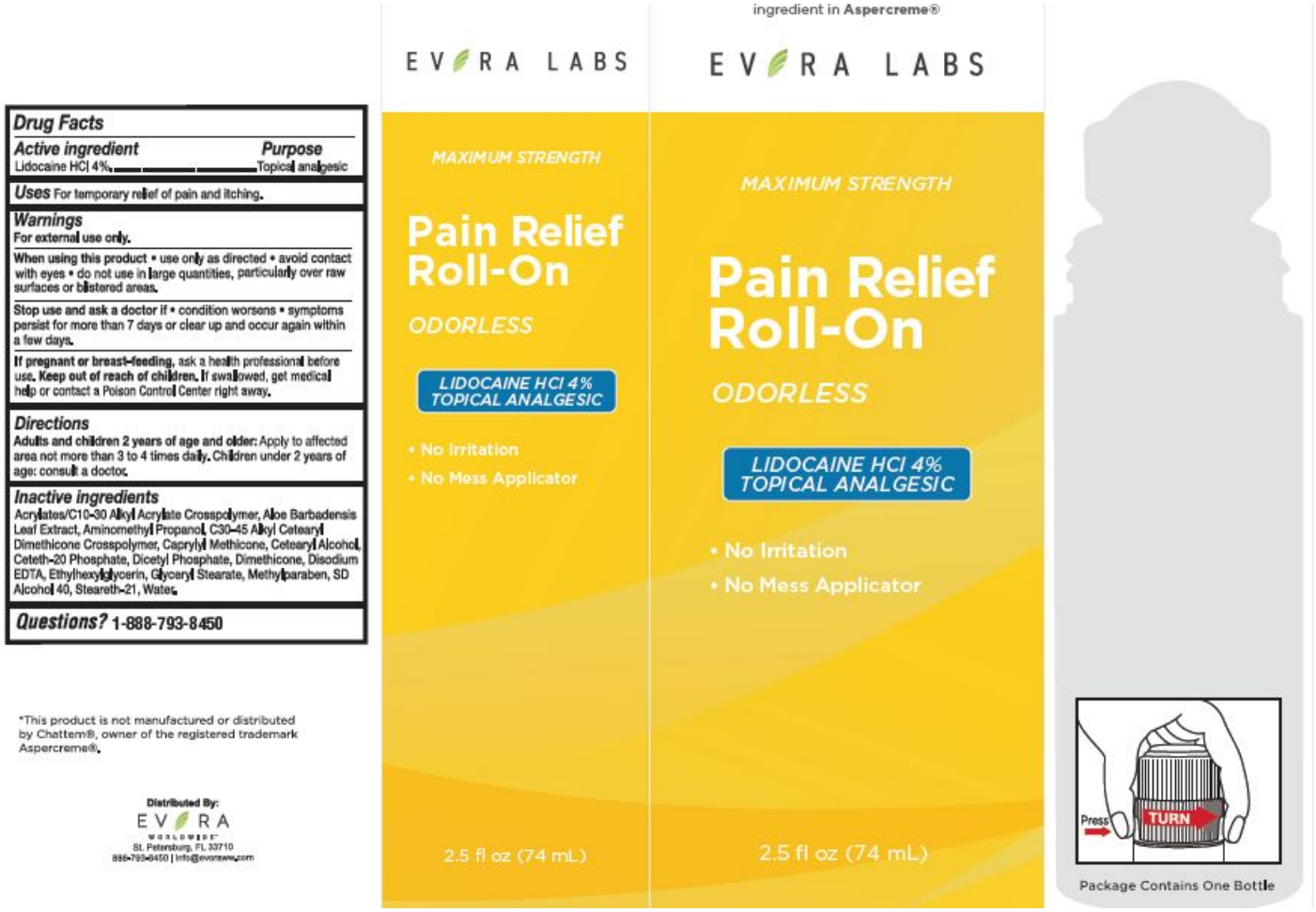
• Adults and children 2 years of age and older: apply to affected area not more than 3-4 times daily
• Children under 2 years old: consult a doctor

Inactive ingredients

Acrylates/C10-30 Alkyl Acrylate Crosspolymer
ALOE VERA LEAF
AMINOMETHYLPROPANOL
Caprylyl Methicone
CETOSTEARYL ALCOHOL
CETETH-20 PHOSPHATE
DIHEXADECYL PHOSPHATE
DIMETHICONE
EDETATE DISODIUM
Ethylhexylglycerin
GLYCERYL STEARATE
METHYLPARABEN
SD ALCOHOL 40
STEARETH-21
WATER

Questions?

1-888-793-8450



PAIN RELIEF ROLL ON EVORA LABS

lidocaine liquid

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:77375-007
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LIDOCAINE HYDROCHLORIDE (UNII: V13007Z41A) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE	4 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
CARBOMER INTERPOLYMER TYPE A (ALLYL SUCROSE CROSSLINKED) (UNII: 59TL3WG5CO)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	
MYRISTYL TRISILOXANE (UNII: J7960S4R1T)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
CETETH-20 PHOSPHATE (UNII: 921FTA1500)	
DIHEXADECYL PHOSPHATE (UNII: 2V6E5WN99N)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
ALCOHOL (UNII: 3K9958V90M)	
STEARETH-21 (UNII: 53J3F32P58)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:77375-007-02	1 in 1 CARTON	12/22/2020	
1		74 mL 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 not final	part348	12/22/2020	

Labeler - Evora Worldwide (081336028)

Establishment

Name	Address	ID/FEI	Business Operations
Inspec Solutions LLC		081030372	manufacture(77375-007)

Revised: 1/2022 Evora Worldwide