EAR PAIN MD UNIT DOSE- lidocaine hydrochloride liquid
Eosera, Inc

Ear Pain MD® Unit Dose

DRUG FACTS

Active ingredient

Lidocaine HCL Monohydrate 4%

Purpose

Topical Analgesic

Use

for the temporary relief of pain.

Warnings

- Keep out of reach of children.

- For external use only.
- If swallowed, immediately call Poison Control 800-222-1222
- Avoid contact with the eyes.

Do not use

- in large quantities, particularly over raw skin surfaces or blistered areas.
- if you are allergic to any ingredient in this product.
- if you have an injury or perforation (hole) of the eardrum, including tubes in the ear.
- If conditions worsen, or if symptoms persist for more than 7 days or clear up and occur again within a few days, discontinue use of this product and consult a doctor.

Directions

Adults and children 2 years of age and older	Apply 2 to 3 drops to the affected area not more than 3 to 4 times daily.
For children under 2	DO NOT USE, consult a doctor.

Inactive ingredients

Aloe barbadensis (Aloe), Benzalkonium chloride, Benzyl alcohol, Hypromellose (HPMC), Water. May contain Hydrochloric acid and/or Sodium hydroxide to adjust pH.

Questions?

Call 844-732-7929

(M-F 9:00am – 5:00pm CST)

PRINCIPAL DISPLAY PANEL - 10 Vial Carton

eosera®
EAR
PAIN MD®

PAIN RELIEVING DROPS
#1
DOCTOR
RECOMMENDED*

With 4% LIDOCAINE

10
sigle use
vials

10 vials [0.5 mL each]

EAR PAIN MD UNIT DOSE

lidocaine hydrochloride liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:72429-0003
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 mL

Inactive Ingredients

Ingredient Name	Strength
Benzalkonium Chloride (UNII: F5UM2KM3W7)	
Benzyl Alcohol (UNII: LKG8494WBH)	
Sodium Acetate Anhydrous (UNII: NVG71ZZ7P0)	
ACETIC ACID (UNII: Q40Q9N063P)	
Water (UNII: 059QF0KO0R)	

Product Characteristics

Color	BLUE	**Score**	
Shape		**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72429-0003-9	10 in 1 CARTON	06/10/2024	
1		0.5 mL in 1 AMPULE; 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	06/10/2024	

Labeler - Eosera, Inc (079789050)

Establishment

Name	Address	ID/FEI	Business Operations
Eosera, Inc		079789050	MANUFACTURE(72429-0003)

Revised: 8/2024 Eosera, Inc