

LICEFREEE AFTERLICE- pramoxine hci spray
Tec Laboratories Inc.

LiceFreee AfterLice Spray

Active ingredient

Pramoxine HCl 1%

For external use only

When using this product

- avoid contact with eyes

Stop use and ask a doctor if

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

Keep out of reach of children

- If swallowed, get medical help or contact a Poison Control Center right away.

Other Information

Store at 59-86°F (15-30°C)

Inactive ingredients

alcohol, benzethonium Cl, menthol, panthenol, polyoxyl 35 castor oil, polyquaternium-10, purified water, tea tree oil

Questions or comments?

Call **1-800-482-4464**.

Uses

for the temporary relief of itching associated with

- insect bites
- minor skin irritations

Directions

- Adults and children 2 years of age and older: spray on affected area not more than 3-4 times daily
- Children under 2 years; do not use; consult a doctor

Pain and itch relief spray for scalp irritation caused by lice bites and lice treatments.

Licefreee! AfterLice™

Pramoxine HCl Pain
Relieving Spray 1%

Relieve itch and
soothe scalp
irritation caused
by lice bites and
treatment

6 FL OZ
(177.4 mL)



Drug Facts

Active ingredient
Pramoxine HCl 1% External Analgesic

Purpose

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- insect bites ■ minor skin irritations

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TEC LABS®

Manufactured by
Tec Laboratories, Inc.
Albany, OR 97321
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LICEFREEE AFTERLICE

pramoxine hci spray

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:51879-260
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
PRAMOXINE HYDROCHLORIDE (UNII: 88AYB867L5) (PRAMOXINE - UNII:068X84E056)	PRAMOXINE HYDROCHLORIDE	10 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
TEA TREE OIL (UNII: VIF565UC2G)	
BENZETHONIUM CHLORIDE (UNII: PH41D05744)	
MENTHOL (UNII: L7T10EIP3A)	
POLYOXYL 35 CASTOR OIL (UNII: 6D4M1DAL6O)	
ALCOHOL (UNII: 3K9958V90M)	
WATER (UNII: 059QF0KO0R)	
PANTHENOL (UNII: WW9CM0067Z)	
POLYQUATERNIUM-10 (400 CPS AT 2%) (UNII: HB1401PQFS)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:51879-260-06	177.4 g in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/16/2023	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	10/16/2023	

Labeler - Tec Laboratories Inc. (083647792)

Registrant - Tec Laboratories Inc. (083647792)

Establishment

Name	Address	ID/FEI	Business Operations
Tec Laboratories Inc.		083647792	manufacture(51879-260)

Revised: 11/2024

Tec Laboratories Inc.