

**RELIEF- pramoxine hydrochloride spray**  
**Elanco US 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.*

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**KEY INGREDIENTS:** Pramoxine Hydrochloride 1%, Omega-6 fatty acids and colloidal oatmeal.

**PRODUCT DESCRIPTION:** Relief Spray combines the soothing benefits of pramoxine and colloidal oatmeal in a moisturizing base that remains on the skin for anti-itch activity. It contains Omega-6 fatty acids for nourishment of the skin.

**INDICATIONS:** For temporary relief of itching, flaking and irritation.

**DIRECTIONS: Shake well before use.** Spray directly onto the affected areas. If necessary, part hair coat so that Relief Spray makes direct contact with the skin. Rub in well. May be used daily or as directed by your veterinarian.

**CAUTIONS:** For topical use on dogs, cats, and horses only. If redness or irritation persists or increases, discontinue use and contact your veterinarian. Avoid contact with eyes. If eye contact occurs, rinse thoroughly with water.

**STORAGE:** Store at room temperature. Protect from freezing.

Manufactured for Elanco US Inc.  
Greenfield, IN 46140 U.S.A.

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**DVM™**

**Elanco™**

90210026\_YL600283B

**Principal Display Panel - 237 mL Bottle Spray Label**

**Elanco™**

**Relief™**  
**SPRAY**

Colloidal oatmeal spray  
for temporary relief of itching and  
flaking on dogs, cats, and horses.

**1% Pramoxine HCl**

**WARNING: Keep out of reach of children.**

**Anti-itch**

NET CONTENTS 237mL (8 fl oz)



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## RELIEF

pramoxine hydrochloride spray

### Product Information

<b>Product Type</b>	OTC ANIMAL DRUG	<b>Item Code (Source)</b>	NDC:58198-0014
<b>Route of Administration</b>	TOPICAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>PRAMOXINE HYDROCHLORIDE</b> (UNII: 88AYB867L5) (PRAMOXINE - UNII:068X84E056)	PRAMOXINE HYDROCHLORIDE	10.8 mg in 1 mL

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:58198-0014-1	1 in 1 BOTTLE, SPRAY		
1		237 mL in 1 BOTTLE, SPRAY		

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
Unapproved drug other		02/15/2017	

**Labeler** - Elanco US Inc. (966985624)

Revised: 4/2023

Elanco US Inc.