

BARRIER WOUND CARE WITH PAIN RELIEF- povidone-iodine, lidocaine, and alcohol solution

Aurora Pharmaceutical, 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.

BARRIER® Wound Care Spray With Pain Relief

INDICATIONS

For topical use on animals as an aid in reducing pain, licking and chewing on wounds and surgical incisions. Kills up to 99% of surface germs in 15 seconds or less that can potentially cause an infection.

ACTIVE INGREDIENTS	PURPOSE
Povidone-Iodine (2% available iodine)	Antiseptic
Bitrex™ Denatonium Benzoate	Bitter Agent
Lidocaine 2% w/v	Pain Relief
Ethyl Alcohol – less than 80%	Antiseptic
Isopropyl Alcohol – less than 5%	Antiseptic

DIRECTIONS

Hold sprayer about 4–6 inches from area to be treated. Spray 1 or more times — allow to dry. Repeat as needed. Forms a highly visible coating that is durable.

WARNINGS

Do not use in the eyes. Do not use if you are allergic or sensitive to the active ingredients. Prolonged exposure may cause irritation. If swallowed, get medical help or contact a Poison Control Center right away. Avoid inhalation.

Store upright at 15°–25° C (59°–77° F). Brief excursions up to 40° C (104° F) are permitted.

**FLAMMABLE — USE WITH CAUTION
FOR EXTERNAL USE ONLY
KEEP OUT OF REACH OF CHILDREN
WET HAIR OR FUR IS FLAMMABLE
Limited Quantity**

MANUFACTURED IN THE USA

REORDER NO: 19001

MANUFACTURED BY:

Aurora Pharmaceutical, Inc.

NORTHFIELD, MINNESOTA 55057

1-888-215-1256

www.aurorapharmaceutical.com

IN 50-1082 06/2023



PRINCIPAL DISPLAY PANEL - 473 mL Bottle Label

NDC 51072-036-16

BARRIER®

**Wound Care Spray
With Pain Relief**

2% Available Iodine with Lidocaine

(Patent pending)

For Animal Use Only

16 fl. oz. (473 mL)

AURORA PHARMACEUTICAL®

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TAKE TIME
OBSERVE LABEL
DIRECTIONS

INDICACIONES: Para uso tópico en animales para ayudar a reducir el dolor, lametazo, y masticación en heridas e incisiones quirúrgicas. Mata hasta el 99% de los gérmenes superficiales en 15 segundos o menos, que potencialmente puede causar una infección.

INGREDIENTES ACTIVOS
Yodopovidona (Yodo al 2%)
Bitrex® Benzato de denatonio
Lidocaína 2% p/v
Alcohol etílico – menos del 80%
Alcohol isopropílico – menos del 5%

PROPOSITO
Antiséptico
Agente amargo
Anestésico local
Antiséptico
Antiséptico

INSTRUCCIONES DE USO: Mantener pulverizador a unas 4-6 pulgadas de la zona a tratar. Rocíar una o más veces, dejar a secar. Repetir según sea necesario. Forma una capa que es altamente visible y duradera.

ADVERTENCIAS: No utilice en los ojos. No utilice en caso de alergia o de sensibilidad a los ingredientes activos. La exposición prolongada puede causar irritación. En caso de ingestión, buscar ayuda médica o contactar al Centro de Control de Venenos (Poison Control Center) inmediatamente. Evitar la inhalación.

Almacenar verticalmente a 15°–25° C (59°–77° F). Excursiones breves hasta 40° C (104° F) son admisibles.

MANUFACTURED IN THE USA

INFLAMABLE — USAR CON CUIDADO SOLO PARA USO EXTERNO. MANTENER FUERA DEL ALCANCE DE LOS NIÑOS. EL PELO O PIEL MOJADO ES INFLAMABLE. Cantidad Limitada

BARRIER WOUND CARE WITH PAIN RELIEF
povidone-iodine, lidocaine, and alcohol solution

Product Information

Product Type	OTC ANIMAL DRUG	Item Code (Source)	NDC:51072-036
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
Povidone-Iodine (UNII: 85H0HZU99M) (IODINE - UNII:9679TC07X4)	IODINE	2 g in 100 mL
Lidocaine (UNII: 98PI200987) (Lidocaine - UNII:98PI200987)	Lidocaine	2 g in 100 mL
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	80 mL in 100 mL
ISOPROPYL ALCOHOL (UNII: ND2M416302) (ISOPROPYL ALCOHOL - UNII:ND2M416302)	ISOPROPYL ALCOHOL	5 mL in 100 mL

Product Characteristics

Color	brown (brown)	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:51072-036-16	473 mL in 1 BOTTLE		
2	NDC:51072-036-01	3790 mL in 1 BOTTLE		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		10/15/2010	

Labeler - Aurora Pharmaceutical, Inc. (832848639)

Establishment

Name	Address	ID/FEI	Business Operations
Aurora Pharmaceutical, Inc.		832848639	MANUFACTURE

Revised: 6/2024

Aurora Pharmaceutical, Inc.