

BISMUSAL - bismuth subsalicylate suspension
Durvet, 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.

BISMUSAL SUSPENSION Anti-Diarrheal Liquid

INDICATIONS

A palatable oral suspension for use as an aid in the control of simple diarrhea in cattle, horses and dogs.

CONTRAINDICATIONS

Not recommended for use in cats.

CAUTION

This product contains salicylate; do not administer with other salicylate-containing products, such as aspirin.

DOSAGE AND ADMINISTRATION

Administer orally after the first sign of diarrhea and after each loose bowel movement or as needed.

Dogs--1 to 3 tablespoonfuls

Calves-- 3 to 4 ounces

Foals-- 3 to 4 ounces

Cattle and Horses-- 6-10 fl. ozs.

This product may cause stools to darken. If diarrhea persists after using this product for 2 days, contact a veterinarian.

COMPOSITION

Bismuth Subsalicylate 1.75%
in a palatable aqueous suspension.
Flavoring and coloring added.

FOR ORAL USE ONLY

SHAKE WELL BEFORE USING

Store at room temperature not above 37°C (98.6°F).

Protect from freezing.

TAKE TIME OBSERVE LABEL DIRECTIONS

NDC 30798-334-35

FOR ANIMAL USE ONLY

KEEP OUT OF REACH OF CHILDREN

NET CONTENTS: 1 GALLON (3.785 L)

Manufactured for:
DURVET, INC.
 Blue Springs, Missouri 64014
 Iss. 2-02

NDC 30798-334-35

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B-0700-07 Iss. 2-02



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www.durvet.com


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Lot No. Exp. Date



BISMUSAL			
bismuth subsalicylate suspension			
Product Information			
Product Type	OTC ANIMAL DRUG	Item Code (Source)	NDC:30798-334
Route of Administration	ORAL		
Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
BISMUTH SUBSALICYLATE (UNII: 62TEY51RR1) (BISMUTH CATION - UNII:ZS9CD118YE)	BISMUTH SUBSALICYLATE	66.2 mg in 3.785 L	
Packaging			
#	Item Code	Package Description	Marketing Start Date Marketing End Date
1	NDC:30798-334-35	3.785 L in 1 JUG	
Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date

unapproved drug other	06/21/2002
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Labeler - Durvet, Inc. (056387798)

Revised: 2/2013

Durvet, Inc.