PRODUCT INFORMATION

Bimeda®

Flunazine®

(flunixin meglumine)

Equine Paste

SYRINGE CONTAINS FLUNIXIN MEGLUMINE EQUIVALENT TO 1500 mg FLUNIXIN FOR ORAL USE IN HORSES ONLY

ANADA 200-581, Approved by FDA

CAUTION

Federal law restricts this drug to use by or on the order of a licensed veterinarian.

INDICATIONS

For the alleviation of inflammation and pain associated with musculoskeletal disorders in the horse.

PACKAGING

LIST NO.	UNIT PACKAGE	CASE SIZE	PALLET
1FLU015	30 g	48 (4 x 12)	3,072

BENEFITS

- Controls inflammatory response
- NSAID approved for oral administration
- Non-narcotic
- Non-steroidal; no steroidal side effects
- Active up to 36 hours
- **Apple-flavored**
- **Safe:** Approved by FDA

WARNING

Do not use in horses intended for human consumption.

Bimeda, Inc.
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Oakbrook Terrace, IL 60181 USA
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See reverse side for additional information

TECHNICAL INFORMATION

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KEEP OUT OF REACH OF CHILDREN

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ACTIVITY:

Flunixin meglumine is a potent, nonnarcotic, nonsteroidal, analgesic agent with anti-inflammatory and antipyretic activity. It is significantly more potent than pentazocine, meperidine, and codeine as an analgesic in the rat yeast paw test. Oral studies in the horse show onset of flunixin activity occurs within 2 hours of administration. Peak response occurs between 12 and 16 hours and duration of activity is 24 to 36 hours.

CONTRAINDICATIONS:

There are no known contraindications to this drug when used as directed.

PRECAUTIONS:

The effect of flunixin meglumine on pregnancy has not been determined. Studies to date show there is no detrimental effect on stallion spermatogenesis with or following the recommended dose of flunixin meglumine.

SIDE EFFECTS:

During field studies with flunixin meglumine, no significant side effects were reported.

DOSAGE AND ADMINISTRATION:

The recommended dose of flunixin meglumine is 0.5 mg per lb of body weight once daily. The Flunazine® Equine Paste syringe, calibrated in twelve 250-lb weight increments, delivers 125 mg of flunixin for each 250 lbs (see dosage table). One syringe will treat a 1000-lb horse once daily for 3 days, or three 1000-lb horses one time.

DOSAGE:

Syring Mark *	Horse Weight (lbs)	Flunazine® Equine Paste Delivered (g)	mg Flunixin Delivered
0	---	---	---
250	250	2.5	125
500	500	5.0	250
750	750	7.5	375
1000	1000	10.0	500

** Use dial edge nearest syringe barrel to mark dose.*

The paste is orally administered by inserting the nozzle of the syringe through the interdental space, and depositing the required amount of paste on the back of the tongue by depressing the plunger.

Treatment may be given initially by intravenous or intramuscular injection of Flunazine® Injectable Solution, followed by Flunazine® Equine Paste on Days 2 to 5. Flunixin meglumine treatment should not exceed 5 consecutive days.

TOXICITY:

No toxic effects were observed in rats given oral flunixin meglumine 2 mg/kg per day for 42 days. Higher doses produced ulceration of the gastrointestinal tract. The emetic dose in dogs is between 150 and 250 mg/kg. Flunixin was well tolerated in monkeys dosed daily with 4 mg/kg for 56 days. No adverse effects occurred in horses dosed orally with 1.0 or 1.5 mg/lb for 5 consecutive days.

STORAGE: Store at 20˚C - 25˚C (68˚F - 77˚F); excursions permitted between 15˚C - 30˚C (between 59˚F - 86˚F).

APPLE FLAVORED

HOW SUPPLIED:

Contains: 12 - Flunazine® (flunixin meglumine) Equine Paste Syringes 30 g each (syringe contains flunixin meglumine equivalent to 1500 mg flunixin).

To obtain an SDS or for assistance, contact Bimeda, Inc. at 1-888-524-6332.

TAKE TIME  OBSERVE LABEL DIRECTIONS

www.BimedaUS.com

PENN Veterinary Supply, Inc. | 53 Industrial Circle • Lancaster • PA 17601 • 1-800-233-0210 • www.pennvet.com