
SYRINGE CONTAINS FLUNIXIN MEGLUMINE EQUIVALENT TO 1500 mg FLUNIXIN

FOR ORAL USE IN HORSES ONLY

KEEP OUT OF REACH OF CHILDREN

WARNING: Do not use in horses intended for human consumption.

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

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.

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

DOSE: 0.5 mg per pound of body weight per day for up to 5 days. Each calibration on the syringe' doses 250 lbs of body weight. Administer orally 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. *Use dial edge nearest syringe barrel to mark dose.

ACTIVITY: Flunixin meglumine is a potent, nonnarcotic, nonsteroidal, analgesic agent with 1 antiinflammatory 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 1 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.

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

See product information sheet for additional information.

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 TABLE

Syringe Mark*	Horse Weight (lbs)	Flunazine [®] Equine Paste Delevered (g)	mg Flunixin Delivered
0			
250	250	2.5	125
500	500	5.0	250
750	750	7.5	375

* 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.





Product Informati	on							
Product T ype		PRESCRIPTION ANIMAL DRUG		Item Code (Source)		ce)	NDC:61133-6007	
Route of Administrati	on	ORAL						
Active Ingredient/	Active Moi	ety						
Ingredient Name					Basis of Strength			rength
FLUNIXIN MEGLUMINE (UNII: 8 Y3JK0JW3U) (FLUNIXIN - UNII:356 IB 10400) FLUNIXIN MEGLUMINE								
FLUNIXIN MEGLUMIN	E (UNII: 8 Y3JK(0JW3U) (FLUNIXIN - UNI	II:356 IB 1O 400)	F	LUNIXIN M	IEGLUMINE	1500 m	ig in 30 ε
				-				
Packaging		DJW3U) (FLUNIXIN - UNI Kage Description	II:356 IB 10 400) Marke tin	-			2 1500 m eting End	
Packaging # Item Code		kage Description		-				
Packaging # Item Code	Pack	kage Description		-				g in 30 g I Date
Packaging # Item Code 1 NDC:61133-6007-1	Pack 30 g in 1 S	kage Description		-				
Packaging # Item Code 1 NDC:61133-6007-1 Marketing Info	Pack 30 g in 1 S rmation	Kage Description SYRINGE	Marketin	g Start	Date	Mark	eting End	l Date
Packaging	Pack 30 g in 1 S rmation	kage Description SYRINGE on Number or Monogra	Marketin	g Start	Date eting Start	Mark		l Date

Labeler - Bimeda Inc. (060492923)

Registrant - Bimeda Inc. (060492923)

Establishment

Name	Address	ID/FEI	Business Operations
Bimeda-MTC Animal Health		256232216	manufacture

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Bimeda Inc.