VETAMEG- flunixin meglumine paste Aspen Veterinary Resources

VetaMeg® Equine Paste (flunixin meglumine paste)

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

DESCRIPTION: Each 30-g syringe of VetaMeg Equine Paste contains flunixin meglumine equivalent to 1500 mg flunixin.

INDICATIONS: VetaMeg Equine Paste is recommended for the alleviation of inflammation and pain associated with musculoskeletal disorders in the horse.

ACTIVITY: Flunix in 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 flunix in activity occurs withing 2 hours of administration. Peak response occurs between 12 and 16 hours and duration of activity is 24-36 hours.

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

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

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.

To report suspected adverse drug events, for technical assistance or to obtain a copy of the Safety Data Sheet (SDS), contact Aspen Veterinary Resources at info@aspenveterinaryresources.com. For additional information about adverse drug experience reporting for animal drugs, contact FDA at 1-888-FDA-VETS or online at www.fda.gov/reportanimalae.

DOSAGE AND ADMINISTRATION: The recommended dose of flunixin meglumine is 0.5 mg per lb of body weight once daily. The VetaMeg 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	Horse Weight	VetaMeg⊺	Equine Paste Delevered mg Flunixin
Mark*	(lbs)	(g)	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 VetaMeg Injectable Solution, followed by VetaMeg 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.

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.

Approved by FDA under ANADA # 200-581

NDC # 46066-503-01



VETAMEG® EQUINE PASTE

Net Wt 30 g



(flunixin meglumine paste)

Apple Flavored

Syringe contains flunixin meglumine equivalent to 1500 mg FLUNIXIN For oral use in horses only.

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.

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.

See product information sheet for additional information.

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

Approved by FDA under ANADA # 200-581

Manufactured for:

Aspen Veterinary Resources®, Ltd.

Liberty, MO 64068

www.aspenveterinaryresources.com

1VFT021 8VET0260 A375B Rev. 06/23



VETAMEG

flunixin meglumine paste

Product Information

Product Type PRESCRIPTION ANIMAL DRUG Item Code (Source) NDC:46066-503

Route of Administration ORAL

Active Ingredient/Active Moiety

Ingredient Name Basis of Strength Strength FLUNIXIN MEGLUMINE (UNII: 8Y3JK0JW3U) (FLUNIXIN - UNII:356IB1O400) FLUNIXIN MEGLUMINE 1500 mg in 30 g

Packaging				
	# Item Code	Package Description	Marketing Start Date	Marketing End Date
П	1 NDC:46066-503-01	30 a in 1 SYRINGE		

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
ANADA	ANADA200581	02/26/2015		

Labeler - Aspen Veterinary Resources (627265361)

Registrant - Bimeda Inc. (060492923)

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

Revised: 3/2024 Aspen Veterinary Resources