

**FLUNIXIN-S- flunixin meglumine injection, solution**  
**Norbrook Laboratories Limited**

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**Flunixin Injection –S**  
**(flunixin meglumine)**  
**50 mg/mL**  
**Veterinary**

ANADA 200-476, Approved by FDA.

**For intramuscular use in swine.**

**Not for use in breeding swine.**

**CAUTION**

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

**DESCRIPTION**

Each milliliter of FLUNIXIN INJECTION -S contains flunixin meglumine equivalent to 50 mg flunixin, 0.1 mg edetate disodium, 2.5 mg sodium formaldehyde sulfoxylate, 4.0 mg diethanolamine, 207.2 mg propylene glycol; 5.0 mg phenol as preservative, hydrochloric acid, water for injection q.s.

**PHARMACOLOGY**

Flunixin meglumine is a potent non-narcotic, non-steroidal, 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.

Flunixin is known to persist in inflammatory tissues<sup>1</sup> and is associated with anti-inflammatory properties which extend well beyond the period associated with detectable plasma drug concentrations<sup>2</sup>. Therefore, prediction of drug concentrations based upon estimated plasma terminal elimination half-life will likely underestimate both the duration of drug action and the concentration of drug remaining at the site of activity.

The pharmacokinetic profiles were found to follow a 2-compartmental model, although a deep (third) compartment was observed in some animals. The mean terminal elimination half-life ( $\beta$  half-life) of flunixin after a single intramuscular injection of flunixin injection (2.2 mg/kg) to pigs was between 3 and 4 hours. The mean observed maximum plasma concentration was 2944 ng/mL, achieved at a mean time of approximately 0.4 hours. The mean AUC(0-LOQ) was 6431 ng\*hr/mL. Following IM administration of flunixin, quantifiable drug concentration could be measured up to 18 hours post dose. The mean volume of distribution was 2003 mL/kg and the mean total clearance was 390 mL/hr/kg. The mean absolute bioavailability of flunixin following an intramuscular injection in the neck was 87%.

**INDICATION**

FLUNIXIN INJECTION -S is indicated for the control of pyrexia associated with swine respiratory disease.

**DOSE AND ADMINISTRATION**

The recommended dose for swine is 2.2 mg/kg (1 mg/lb; 2 mL per 100 lbs) body weight given by a single intramuscular administration. The injection should be given only in the neck musculature with a

maximum of 10 mL per site.

Note: Intramuscular injection may cause local tissue irritation and damage. In an injection-site irritation study, the tissue damage did not resolve in all animals by Day 28 post-injection. This may result in trim loss of edible tissue at slaughter.

## **CONTRAINDICATIONS**

There are no known contraindications to this drug in swine when used as directed. Do not use in animals showing hypersensitivity to flunixin meglumine. Use judiciously when renal impairment or gastric ulceration is suspected.

## **RESIDUE WARNINGS**

Swine must not be slaughtered for human consumption within 12 days of the last treatment.

## **PRECAUTIONS**

As a class, cyclo-oxygenase inhibitory NSAIDs may be associated with gastrointestinal, renal and hepatic toxicity. Sensitivity to drug-associated adverse events varies with the individual patient. Patients at greatest risk for adverse events are those that are dehydrated, on concomitant diuretic therapy, or those with existing renal, cardiovascular, and/or hepatic dysfunction. Concurrent administration of potentially nephrotoxic drugs should be carefully approached. NSAIDs may inhibit the prostaglandins that maintain normal homeostatic function. Such prostaglandin effects may result in clinically significant disease in patients with underlying or pre-existing disease that has not been previously diagnosed. Since many NSAIDs possess the potential to produce gastrointestinal ulceration, concomitant use of flunixin meglumine with other anti-inflammatory drugs, such as other NSAIDs and corticosteroids, should be avoided.

Not for use in breeding swine. The reproductive effects of FLUNIXIN INJECTION -S have not been investigated in this class of swine.

Intramuscular injection may cause local tissue irritation and damage. In an injection site irritation study, the tissue damage did not resolve in all animals by Day 28 post-injection. This may result in trim loss of edible tissue at slaughter.

## **SAFETY**

Flunixin was mildly irritating at the injection sites. No other flunixin-related changes (adverse reactions) were noted in swine administered a 1X (2.2 mg/kg; 1.0 mg/lb) dose for 9 days. Minimal toxicity manifested itself as statistically significant increased spleen weight at elevated doses (5X or higher daily for 9 days) with no change in normal microscopic architecture.

## **HOW SUPPLIED**

FLUNIXIN INJECTION -S, 50 mg/mL, is available in 100 mL and 250 mL multi-dose vials.

**Store between 2° and 30°C (36° and 86°F).**

## **REFERENCES**

1. Lees P, Higgins AJ. Flunixin inhibits prostaglandin E2 production in equine inflammation. *Res Vet Sci.* 1984; 37:347-349.
2. Odensvik K. Pharmacokinetics of flunixin and its effect on prostaglandin F2a metabolite concentrations after oral and intravenous administration in heifers. *J Vet Pharmacol Ther.* 1995;

18:254-259.

**Restricted Drug - California. Use Only as Directed.**

Made in the UK.

**Distributed by:**

Norbrook, Inc.  
9733 Loiret Blvd  
Lenexa, KS 66219  
USA

010670101

**Vial Label**

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Sterile Solution	250 mL
Veterinary	Multiple Dose Vial
	50 mg/mL

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Flunixin Injection –S

(flunixin meglumine)

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Not for use in breeding swine.

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<p><b>Description:</b> Each mL contains: flunixin meglumine equivalent to 50 mg flunixin, 0.1 mg edetate disodium, 2.5 mg sodium formaldehyde sulfoxylate, 4.0 mg diethanolamine, 207.2 mg propylene glycol; 5.0 mg phenol as preservative, hydrochloric acid, water for injection qs.</p> <p><b>Indication:</b> For the control of pyrexia associated with swine respiratory disease.</p> <p><b>Dosage and Administration:</b> For complete product information, please see the outer carton and package insert.</p> <p>Read accompanying directions carefully.</p> <p>000000L0?</p>	<p>Sterile Solution Veterinary</p> <p>250 mL Multiple Dose Vial 50 mg/mL</p> <p><b>Flunixin Injection -S</b> (flunixin meglumine)</p> <p>For intramuscular use in swine. Not for use in breeding swine.</p> <p><b>Caution:</b> Federal law restricts this drug to use by or on the order of a licensed veterinarian.</p> <p><b>Norbrook</b> </p>	<p><b>RESIDUE WARNINGS:</b> Swine must not be slaughtered for human consumption within 12 days of the last treatment.</p> <p><b>Store between 2° and 30° C (36° and 86° F).</b></p> <p>Restricted Drug - California. Use Only as Directed.</p> <p>ANADA 200-476, Approved by FDA.</p> <p>Made in the UK.</p> <p><b>Distributed by:</b> Norbrook, Inc. 9733 Loiret Blvd Lenexa, KS 66219</p> <p>Lot No.:</p> <p>Exp. Date:</p>
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**Carton Label**

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Sterile Solution	250 mL
Veterinary	Multiple Dose Vial
	50 mg/mL

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Flunixin Injection –S

(flunixin meglumine)



**Product Information**

<b>Product Type</b>	PRESCRIPTION ANIMAL DRUG	<b>Item Code (Source)</b>	NDC:55529-030
<b>Route of Administration</b>	INTRAMUSCULAR		

**Active Ingredient/Active Moiety**

<b>Ingredient Name</b>	<b>Basis of Strength</b>	<b>Strength</b>
flunixin meglumine (UNII: 8Y3JK0JW3U) (flunixin - UNII:356IB1O400)	flunixin	50 mg in 1 mL

**Inactive Ingredients**

<b>Ingredient Name</b>	<b>Strength</b>
sodium formaldehyde sulfoxylate (UNII: X4ZGP7K714)	2.5 mg in 1 mL
edetate disodium (UNII: 7FLD91C86K)	0.1 mg in 1 mL
diethanolamine (UNII: AZE05TDV2V)	4.0 mg in 1 mL
propylene glycol (UNII: 6DC9Q167V3)	207.2 mg in 1 mL
phenol (UNII: 339NCG44TV)	5.0 mg in 1 mL
hydrochloric acid (UNII: QTT17582CB)	
water (UNII: 059QF0K00R)	

**Packaging**

#	<b>Item Code</b>	<b>Package Description</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
1	NDC:55529-030-02	1 in 1 CARTON		
1		100 mL in 1 VIAL, GLASS		
2	NDC:55529-030-04	1 in 1 CARTON		
2		250 mL in 1 VIAL, GLASS		

**Marketing Information**

<b>Marketing Category</b>	<b>Application Number or Monograph Citation</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
ANADA	ANADA200476	10/05/2009	

**Labeler** - Norbrook Laboratories Limited (214580029)

Revised: 1/2017

Norbrook Laboratories Limited