

Each mL contains: 50 mg flunixin (equivalent to 83 mg flunixin meglumine), 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.

RESIDUE WARNINGS:

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

USE WITHIN 28 DAYS OF FIRST PUNCTURE AND PUNCTURE A MAXIMUM OF 10 TIMES. WHEN USING A DRAW-OFF SPIKE OR NEEDLE WITH BORE DIAMETER LARGER THAN 18 GAUGE, DISCARD ANY PRODUCT REMAINING IN THE VIAL IMMEDIATELY AFTER USE.

NDC # 0061-1838-30

Sterile

100 mL
Multiple-Dose Vial
50 mg/mL

Banamine[®]-S
(flunixin meglumine injection)

Caution: Federal law restricts this drug to use by or on the order of a licensed veterinarian.
NADA #101-479, Approved by FDA.



SAMPLE ONLY

LOT
EXP

Varnish and
Ink free

Store at or below 25°C (77°F).
Do not freeze.

For intramuscular use in swine.
Not for use in breeding swine.
Before using this drug, read package insert for complete product information.

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