

51ST ANNUAL CONFERENCE

PROCEEDINGS

American Association of Bovine Practitioners



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Phoenix, Arizona
September 13-15, 2018

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PROCEEDINGS
of the
FIFTY-FIRST ANNUAL CONFERENCE
AMERICAN ASSOCIATION OF BOVINE PRACTITIONERS

September 13-15, 2018

Phoenix, Arizona

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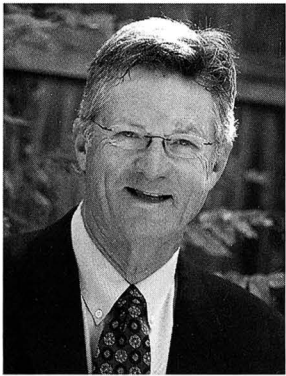
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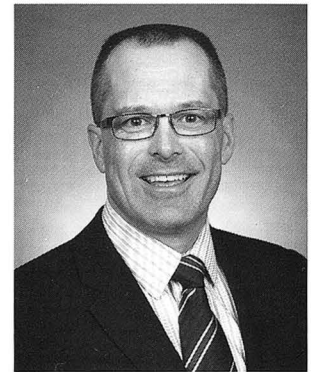
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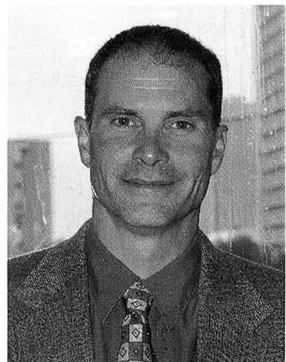
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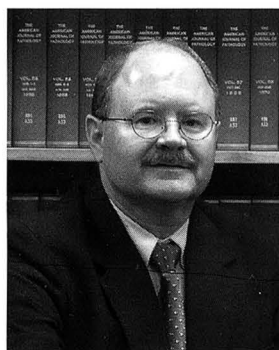
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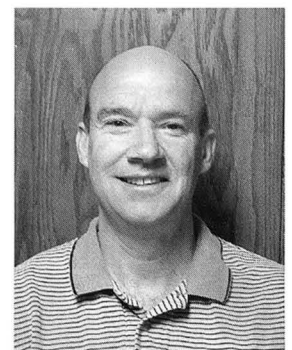
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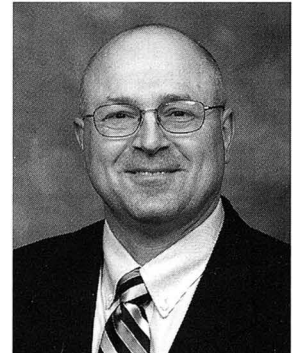
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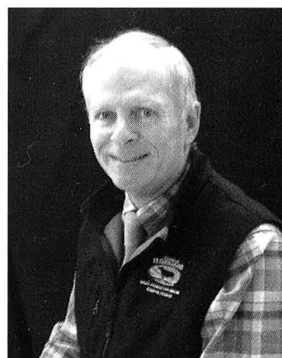
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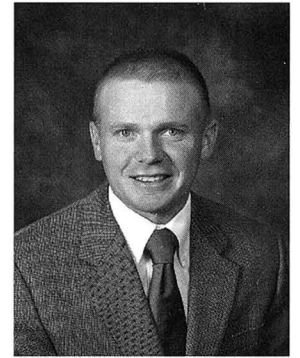
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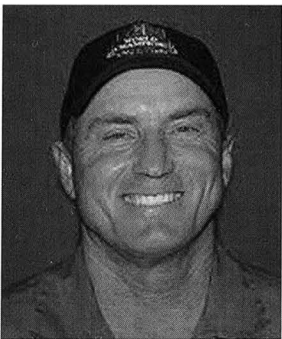
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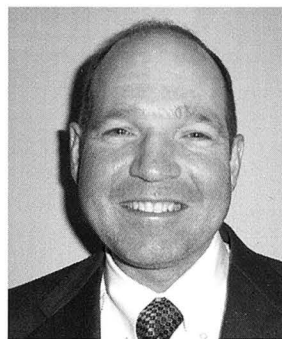
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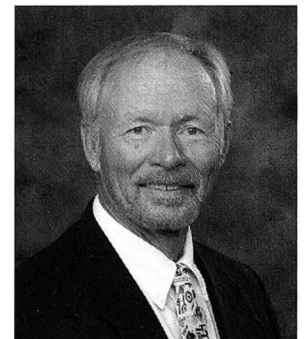
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Injectable
Baytril® 100
 (enrofloxacin)

100 mg/mL Antimicrobial Injectable Solution
 For Subcutaneous use In Beef Cattle, Non-Lactating Dairy Cattle
 For Intramuscular Or Subcutaneous Use In Swine
 Not For Use In Female Dairy Cattle 20 Months Of Age Or Older
 Or In Calves To Be Processed For Veal

BRIEF SUMMARY:
 Before using Baytril® 100, please consult the product insert, a summary of which follows:

CAUTION:
 Federal (U.S.A.) law restricts this drug to use by or on the order of a licensed veterinarian.
 Federal (U.S.A.) law prohibits the extra-label use of this drug in food-producing animals.
 To assure responsible antimicrobial drug use, enrofloxacin should only be used as a second-line drug for colibacillosis in swine following consideration of other therapeutic options.

INDICATIONS:
Cattle - Single-Dose Therapy: Baytril® 100 is indicated for the treatment of bovine respiratory disease (BRD) associated with *Mannheimia haemolytica*, *Pasteurella multocida*, *Histophilus somni* and *Mycoplasma bovis* in beef and non-lactating dairy cattle; and for the control of BRD in beef and non-lactating dairy cattle at high risk of developing BRD associated with *M. haemolytica*, *P. multocida*, *H. somni* and *M. bovis*.

Cattle - Multiple-Day Therapy: Baytril® 100 is indicated for the treatment of bovine respiratory disease (BRD) associated with *Mannheimia haemolytica*, *Pasteurella multocida* and *Histophilus somni* in beef and non-lactating dairy cattle.

Swine: Baytril® 100 is indicated for the treatment and control of swine respiratory disease (SRD) associated with *Actinobacillus pleuropneumoniae*, *Pasteurella multocida*, *Haemophilus parasuis*, *Streptococcus suis*, *Bordetella bronchiseptica* and *Mycoplasma hyopneumoniae*. Baytril® 100 is indicated for the control of colibacillosis in groups or pens of weaned pigs where colibacillosis associated with *Escherichia coli* has been diagnosed.

Use within 30 days of first puncture and puncture a maximum of 30 times with a needle or 4 times with a dosage delivery device. Any product remaining beyond these parameters should be discarded.

RESIDUE WARNINGS:

Cattle: Animals intended for human consumption must not be slaughtered within 28 days from the last treatment. This product is not approved for female dairy cattle 20 months of age or older, including dry dairy cows. Use in these cattle may cause drug residues in milk and/or in calves born to these cows. A withdrawal period has not been established for this product in pre-ruminating calves. Do not use in calves to be processed for veal.

Swine: Animals intended for human consumption must not be slaughtered within 5 days of receiving a single-injection dose.

HUMAN WARNINGS:
For use in animals only. Keep out of the reach of children. Avoid contact with eyes. In case of contact, immediately flush eyes with copious amounts of water for 15 minutes. In case of dermal contact, wash skin with soap and water. Consult a physician if irritation persists following ocular or dermal exposures. Individuals with a history of hypersensitivity to quinolones should avoid this product. In humans, there is a risk of user photosensitization within a few hours after excessive exposure to quinolones. If excessive accidental exposure occurs, avoid direct sunlight. For customer service or to obtain product information, including a Safety Data Sheet, call 1-800-633-3796. For medical emergencies or to report adverse reactions, call 1-800-422-9874.

PRECAUTIONS:
 The effects of enrofloxacin on cattle or swine reproductive performance, pregnancy and lactation have not been adequately determined. The long-term effects on articular joint cartilage have not been determined in pigs above market weight. Subcutaneous injection in cattle and swine, or intramuscular injection in swine, can cause a transient local tissue reaction that may result in trim loss of edible tissue at slaughter. Baytril® 100 contains different excipients than other Baytril® products. The safety and efficacy of this formulation in species other than cattle and swine have not been determined. Quinolone-class drugs should be used with caution in animals with known or suspected Central Nervous System (CNS) disorders. In such animals, quinolones have, in rare instances, been associated with CNS stimulation which may lead to convulsive seizures. Quinolone-class drugs have been shown to produce erosions of cartilage of weight-bearing joints and other signs of arthropathy in immature animals of various species. See Animal Safety section for additional information.

ADVERSE REACTIONS:
 No adverse reactions were observed during clinical trials.

ANIMAL SAFETY:
 In feeder calves, clinical signs including depression, incoordination, muscle fasciculation and inappetance have been observed at higher than approved label dosages. In swine subcutaneous safety studies, incidental lameness of short duration and musculoskeletal stiffness have been observed at higher than approved label dosages. In swine intramuscular safety studies, transient decreases in feed and water consumption were observed after each treatment. Mild, transient, post-treatment injection site swellings were observed in pigs receiving the 37.5 mg/kg BW dose. Injection site inflammation was found on post-mortem examination in all enrofloxacin-treated groups.

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HOW LONG DOES IT LAST?

THAT'S THE WRONG QUESTION.

The question to ask is how fast does the treatment kill BRD-causing bacteria.

Single-dose Baytril® 100 (enrofloxacin) Injectable delivers effective, therapeutic levels of drug in the lung tissues in 1-2 hours*¹ and kills 97% of BRD-causing bacteria in 1-2 hours.*^{2,3} The sooner bacteria are killed, the faster a calf will feel better and get back to work eating and gaining weight. You know the drill. You turn to the one you trust.

*The clinical significance of *in vitro* data has not been demonstrated. Federal law restricts this drug to use by or on the order of a licensed veterinarian. Extra-label use of this product in food-producing animals is prohibited.

¹Davis JL, Foster DM, Papich MG. (2007). Pharmacokinetics and tissue distribution of enrofloxacin and its active metabolite ciprofloxacin in calves. *J Vet Pharmacol Ther.* 30(6):564-571.

²Blondeau JM, Borsos S, Blondeau LD, et al. (2005). The killing of clinical isolates of *Mannheimia haemolytica* (MH) by enrofloxacin (ENR) using minimum inhibitory and mutant prevention drug concentrations and over a range of bacterial inocula. In: ASM Conference on Pasteurellaceae; 23-26 October 2005; Kohala Coast, Big Island, Hawaii: American Society of Microbiology; Abstract B12.

³Blondeau JM, Borsos SD, Hesje CH, et al. Comparative killing of bovine isolates of *Mannheimia haemolytica* by enrofloxacin, florfenicol, tilmicosin and tulathromycin using the measured minimum inhibitory concentration (MIC) and mutant prevention concentration (MPC) drug values. In: International Meeting of Emerging Diseases and Surveillance (IMED); Vienna, Austria: International Society for Infectious Diseases. February 23-25, 2007. Figures 8-10.

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BL171579



Baytril® 100
(enrofloxacin)



100 mg/mL Antimicrobial Injectable Solution



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