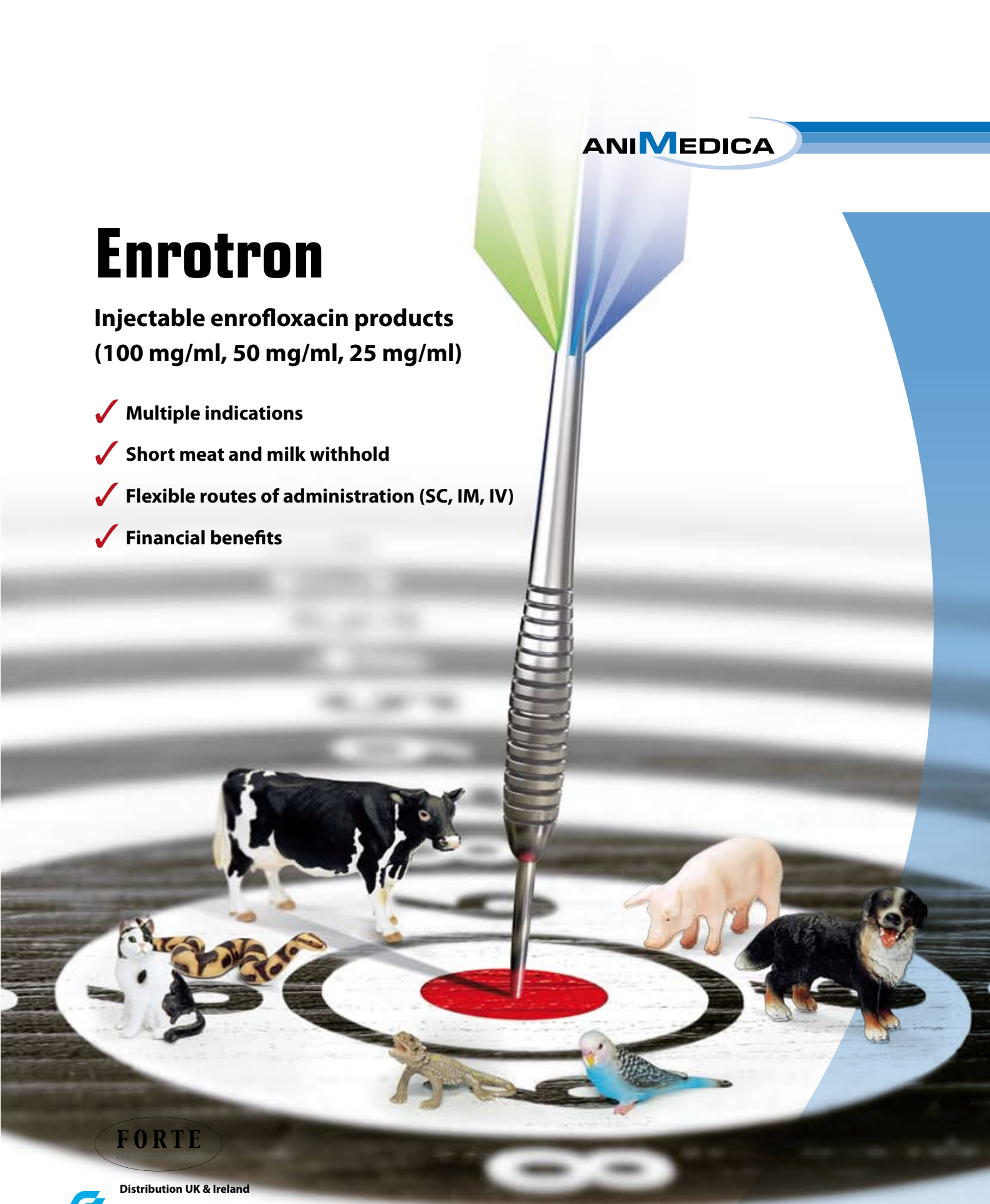
ANIMEDICA

Enrotron

Injectable enrofloxacin products (100 mg/ml, 50 mg/ml, 25 mg/ml)

- ✓ Multiple indications
- ✓ Short meat and milk withhold
- ✓ Flexible routes of administration (SC, IM, IV)
- ✓ Financial benefits



FORTE Healthcare Ltd

Distribution UK & Ireland

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Forte Healthcare

Enrotron 100 mg/ml

Solution for injection
for cattle and pigs

Enrotron 50 mg/ml

Solution for injection
for cattle, pigs, dogs and cats

Enrotron 25 mg/ml

Solution for injection
for dogs, cats and exotic animals

… COVERS ALL ASPECTS FOR AN EFFECTIVE ANTI-INFECTIVE TREATMENT.

What?

- Enrotron injectable products contain the active ingredient enrofloxacin, a well known veterinary fluoroquinolone, which is successfully used worldwide for the treatment of animals

Why?

- Enrotron is a broad spectrum antibiotic with concentration dependent activity against a wide range of gram-negative and gram-positive pathogens and mycoplasmas
- Rapidly and extensively absorbed from the injection site
- Easy to draw up and inject
- Short withdrawal period

When?

- **Dogs and cats:** infections of the alimentary, respiratory and urogenital tracts, skin, secondary wound infections and otitis externa
- **Exotic animals:** infections of the alimentary and respiratory tracts
- **Cattle:** respiratory and alimentary tract infections caused by bacteria or mycoplasma, secondary bacterial infections subsequent to viral infections, peracute/acute mastitis in lactating dairy cattle
- **Pigs:** diseases of the respiratory and alimentary tract of bacterial or mycoplasmal origin, multi-factorial diseases such as atrophic rhinitis and enzootic pneumonia

Enrotron 100 mg/ml

Cattle, S/C:	Meat & offal 10 days Milk 84 hours
Cattle, I/V:	Meat & offal 4 day Milk 72 hours
Pigs, I/M:	Meat & offal 10 days

Enrotron 50 mg/ml

Cattle, S/C:	Meat & offal 14 days
Pigs, I/M:	Meat & offal 10 days

Enrotron 100 mg/ml Solution for injection for cattle and pigs. Ingredients: Each ml contains: Active Substance: Enrofloxacin 100.0 mg. Excipients: 1-Butanol 30.0 mg. **Indications:** Cattle: Diseases of the respiratory and alimentary tract of bacterial or mycoplasmal origin (e.g. pasteurellosis, mycoplasmosis, coli-bacillosis, coli-septicaemia and salmonellosis) and secondary bacterial infections subsequent to viral infections (e.g., viral pneumonia), where clinical experience, supported where possible by sensitivity testing of the causal organism, indicates enrofloxacin as the drug of choice. Treatment of local signs (inflammation, milk quality and yield) associated with peracute/acute mastitis in lactating dairy cattle caused by E. coli, where herd history and previous sensitivity testing indicate enrofloxacin as the drug of choice. Pigs: Diseases of the respiratory and alimentary tract of bacterial or mycoplasmal origin (e.g. pasteurellosis, mycoplasmosis, coli-bacillosis, coli-septicaemia and salmonellosis) and multifactorial diseases such as atrophic rhinitis and enzootic pneumonia where clinical experience, supported where possible by sensitivity testing of the causal organism, indicates enrofloxacin as the drug of choice. **Contraindications:** Do not use the product for prophylaxis. Do not use in cases of known hypersensitivity to fluoroquinolones or to any of the excipients. Do not use when resistance / cross resistance to (fluoro)quinolones is known to occur. **Adverse reactions:** Local tissue reactions may occasionally occur at the injection site. Normal sterile precautions should be taken. **Use during pregnancy, lactation or lay:** There is no restriction on the use of this product during pregnancy and lactation. **Withdrawal Period(s):** Cattle: Subcutaneous use: Meat and offal: 10 days, Milk: 84 hours (7 milkings). Cattle: Intravenous use: Meat and offal: 4 days, Milk: 72 hours (6 milkings). Pigs: Intramuscular use: Meat and Offal 10 days.

Enrotron 50 mg/ml Solution for injection for cattle, pigs, dogs and cats. Ingredients: Each ml contains: Active Substance: Enrofloxacin 50.0 mg. Excipients: 1-Butanol 30.0 mg. **Indications:** Cattle: Diseases of the respiratory and alimentary tract of bacterial or mycoplasmal origin (e.g. pasteurellosis, mycoplasmosis, coli-bacillosis, coli-septicaemia and salmonellosis) and secondary bacterial infections subsequent to viral infections (e.g., viral pneumonia), where clinical experience, supported where possible by sensitivity testing of the causal organism, indicates enrofloxacin as the drug of choice. Pigs: Diseases of the respiratory and alimentary tract of bacterial or mycoplasmal origin (e.g. pasteurellosis, actinobacillosis, mycoplasmosis, coli-bacillosis, coli-septicaemia and salmonellosis) and multifactorial diseases such as atrophic rhinitis and enzootic pneumonia where clinical experience, supported where possible by sensitivity testing of the causal organism, indicates enrofloxacin as the drug of choice. Dogs and Cats: Treatment of bacterial infections of the alimentary, respiratory and urogenital tracts, skin, secondary wound infections and otitis externa where clinical experience, supported where possible by sensitivity testing of the causal organism, indicates enrofloxacin as the drug of choice. **Contraindications:** Do not treat dogs under 1 year of age with the product as damage to the articular cartilage may occur during the period of rapid growth, specifically in large breeds of dogs. As a precaution do not treat very large breeds of dog with the product until they are 18 months of age because of their longer growth period. Do not use in cats less than 8 weeks of age. Do not use Enrofloxacin for prophylaxis. Do not use in cases of known hypersensitivity to fluoroquinolones or to any of the excipients. Do not use when resistance / cross resistance to (fluoro)quinolones is known to occur. **Adverse reactions:** During the period of rapid growth, enrofloxacin may affect articular cartilage. Local tissue reactions may occur at the injection site. Normal sterile precautions should be taken. Dogs: Occasionally skin reactions have been seen after administration to kennelled greyhounds. **Use during pregnancy, lactation or lay:** There is no restriction on the use of this product during pregnancy and lactation. **Withdrawal Period(s):** Meat and Offal: Cattle: 14 days, Pigs: 10 days. Milk: Not permitted for use in lactating animals producing milk for human consumption.

Enrotron 25 mg/ml Solution for injection for dogs, cats and exotic animals. Ingredients: Each ml contains: Active Substance: Enrofloxacin 25.0 mg. Excipients: 1-Butanol 30.0 mg. **Indications:** The product is for use in dogs and cats in the treatment of bacterial infections of the alimentary, respiratory and urogenital tracts, skin, secondary wound infections and otitis externa where clinical experience, supported where possible by sensitivity testing of the causal organism, indicates enrofloxacin as the drug of choice. The product may also be used in exotic animals (small mammals, reptiles and avian species) for the treatment of bacterial infections of the alimentary and respiratory tracts where clinical experience, supported where possible by sensitivity testing of the causal organism, indicates enrofloxacin as the drug of choice. **Contraindications:** Do not use in dogs less than 1 year of age or in exceptionally large breeds of dog with a longer growth period under 18 months of age. Do not use for cats less than 8 weeks of age. Do not use in cases of known hypersensitivity to fluoroquinolones or to any of the excipients. Do not use when resistance / cross resistance to (fluoro) quinolones is known to occur. **Adverse reactions:** In dogs enrofloxacin may affect articular cartilage during the period of rapid growth. Occasionally skin reactions have been seen after administration to kennelled greyhounds. Local tissue reactions may occasionally occur at the injection site. Normal sterile recautions should be taken. Exotic Animals: Muscle bruising after injection in reptiles and birds has been reported occasionally. **Use during pregnancy, lactation or lay:** Do not use during pregnancy and lactation. **Withdrawal period:** Not applicable.

For animal treatment only – to be supplied only on veterinary prescription. Marketing authorisation holder: FORTE Healthcare Limited, Cougar Lane, Naul, Co. Dublin, Ireland. 12/2011.

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