Your customers demand quality and a reliable supply.

Zoetis helps you deliver proven products, consistently.

You need a trusted ally.

Zoetis is a long-time, trusted and reliable animal health supplier that delivers day-in and day-out to help ensure you and your customers are able to keep business moving forward.

1. Zoetis is a reliable supplier.
   - Zoetis has a history of reliable supply, and has brought you DRAXXIN® (tulathromycin injection) Injectable Solution for more than 15 years.
   - We have high standards for production, quality control and manufacturing performance.

2. We deliver a competitive offering.
   - Our comprehensive portfolio allows you to offer a wide-array of animal health products from a single source provider.

3. We provide unmatched support to help you help your customers.
   - The sales and channel teams are ready to serve you and troubleshoot if necessary.
   - We offer unmatched technical support, protocol development and training from our technical veterinary team.
   - Our industry leadership and visibility in the market helps promote Zoetis products and support your sales initiatives.

IMPORTANT SAFETY INFORMATION: DRAXXIN has a pre-slaughter withdrawal time of 18 days in cattle. Do not use in female dairy cattle 20 months of age or older. Do not use in animals known to be hypersensitive to the product. See Brief Summary of Prescribing Information on the following page.
Zoetis offers all of that, plus a product with 15 years of clinical performance. Because your customers need to have confidence in what you are selling.

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First, always ask if a Zoetis representative has been in contact recently:

Doctor, have you recently utilized DRAXXIN from Zoetis?

If YES, confirm and discuss benefits.

If NO, engage/detail the customer as the primary contact.

Confidence

I believe DRAXXIN will continue to be an important tool for managing Bovine Respiratory Disease (BRD) for your customers.

Neutral invitation

Let’s take a look at some basic information about DRAXXIN...

It’s the customer’s decision

...so you can decide whether DRAXXIN is a solution you’ll consider for your patients presenting with signs of Bovine Respiratory Disease (BRD).

Brief Summary of Prescribing Information for Cattle.

Antibiotic of the subclass triamilide. Each mL of DRAXXIN contains 100 mg of tulathromycin/mL of infectious bovine keratoconjunctivitis (IBK) associated with Moraxella bovis.

DRAXXIN Injectable Solution is indicated for the treatment of bovine respiratory disease (BRD) associated with Pasteurella haemolytica, Pasteurella multocida, Histophilus somni, and Mycoplasma bovis.

DRAXXIN Injectable Solution is indicated for the treatment of foot rot (interdigital necrobacillosis) associated with Fusobacterium necrophorum and Porphyromonas levii.

DRAXXIN Injectable Solution is indicated for the treatment of bovine foot rot (interdigital necrobacillosis) associated with Haemophilus somnus and Moraxella bovis.

DRAXXIN Injectable Solution is indicated for the treatment of bovine foot rot (interdigital necrobacillosis) associated with Moraxella bovis.

DRAXXIN Injectable Solution is indicated for the treatment of bovine foot rot (interdigital necrobacillosis) associated with Pasteurella multocida, Histophilus somni, and Mycoplasma bovis.

Foot Rot – DRAXXIN Injectable Solution is indicated for the treatment of bovine foot rot (interdigital necrobacillosis) associated with Fusobacterium necrophorum and Porphyromonas levii.

Suckling Calves, Dairy Calves, and Veal Calves

BRD – DRAXXIN Injectable Solution is indicated for the treatment of BRD associated with M. haemolytica, P. multocida, M. somni, and M. bovis.

DOSEAGE AND ADMINISTRATION

Inject subcutaneously as a single dose in the neck at a dosage of 2.5 mg/kg (1.1 mL/100 lb) body weight (BW). Do not inject more than 10 mL per injection site.

Table 1. DRAXXIN Cattle Dosing Guide

<table>
<thead>
<tr>
<th>Animal Weight (Pounds)</th>
<th>Dose Volume (mL)</th>
</tr>
</thead>
<tbody>
<tr>
<td>100</td>
<td>1.1</td>
</tr>
<tr>
<td>200</td>
<td>2.3</td>
</tr>
<tr>
<td>300</td>
<td>3.4</td>
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<tr>
<td>800</td>
<td>9.1</td>
</tr>
<tr>
<td>900</td>
<td>10.2</td>
</tr>
<tr>
<td>1000</td>
<td>11.4</td>
</tr>
</tbody>
</table>

CONTRAINDICATIONS

The use of DRAXXIN Injectable Solution is contraindicated in animals previously found to be hypersensitive to the drug.

WARNINGS

FOR USE IN ANIMALS ONLY.

NOT FOR HUMAN USE.

KEEP OUT OF REACH OF CHILDREN.

NOT FOR USE IN CHICKENS OR TURKEYS.

RESIDUE WARNINGS

Cattle intended for human consumption must not be slaughtered within 18 days from the last treatment. This drug is not approved for use in female dairy cattle 50 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.

PRECAUTIONS

The effects of DRAXXIN on bovine reproductive performance, pregnancy, and lactation have not been determined. Subcutaneous injection can cause a transient local tissue reaction that may result in trim loss of edible tissue at slaughter.

Antibiotic of the subclass triamilide. Each mL of DRAXXIN contains 100 mg of tulathromycin/mL.