

Cefenil® RTU

(cefotiofur hydrochloride sterile suspension)

For intramuscular and subcutaneous use in cattle and intramuscular use in swine. This product may be used in lactating dairy cattle. Not for use in calves to be processed for veal.

CAUTION: Federal (USA) law restricts this drug to use by or on the order of a licensed veterinarian. Federal law prohibits extra-label use of this drug in cattle and swine for disease prevention purposes; at unapproved doses, frequencies, durations, or routes of administration; and in unapproved major food producing species/production classes.

DESCRIPTION

CEFENIL® RTU (cefotiofur hydrochloride sterile suspension) is a ready to use formulation that contains the hydrochloride salt of cefotiofur, which is a broad spectrum cephalosporin antibiotic. Each mL of this ready-to-use sterile suspension contains cefotiofur hydrochloride equivalent to 50 mg cefotiofur, 5.73 mg aluminum monostearate, 1.03 mg sorbitan monooleate and medium chain triglycerides.

Structure:

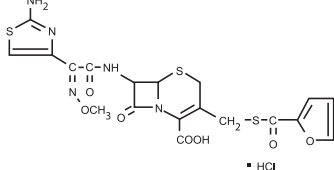


Figure 1

Chemical Name of Cefotiofur Hydrochloride: 5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid, 7-[[2-amino-4-thiazolyl(methoxyimino)acetyl]amio]-3-[[[2-furanylcarbonyl]thio]methyl]-8-oxo-, hydrochloride salt [6R-[6a,7b(2Z)]]

INDICATIONS

Swine: CEFENIL RTU is indicated for treatment/control of swine bacterial respiratory disease (swine bacterial pneumonia) associated with *Actinobacillus (Haemophilus) pleuropneumoniae*, *Pasteurella multocida*, *Salmonella choleraesuis* and *Streptococcus suis*.

Cattle: CEFENIL RTU is indicated for treatment of the following bacterial diseases:

- Bovine respiratory disease (BRD, shipping fever, pneumonia) associated with *Mannheimia haemolytica*, *Pasteurella multocida* and *Histophilus somni*.
- Acute bovine interdigital necrobacillosis (foot rot, pododermatitis) associated with *Fusobacterium necrophorum* and *Bacteroides melaninogenicus*.
- Acute metritis (0 to 14 days post-partum) associated with bacterial organisms susceptible to cefotiofur.

DOSE AND ADMINISTRATION

Shake for 90 seconds to ensure complete resuspension before using.

Swine: Administer intramuscularly at a dosage of 1.36 to 2.27 mg cefotiofur equivalents/lb (3.0 to 5.0 mg/kg) BW (1 mL of sterile suspension per 22 to 37 lb BW). Treatment should be repeated at 24 h intervals for a total of three consecutive days.

Cattle: - For bovine respiratory disease and acute bovine interdigital necrobacillosis: administer by intramuscular or subcutaneous administration at the dosage of 0.5 to 1.0 mg cefotiofur equivalents/lb (1.1 to 2.2 mg/kg) BW (1 to 2 mL sterile suspension per 100 lb BW). Administer daily at 24 h intervals for a total of three consecutive days. Additional treatments may be administered on Days 4 and 5 for animals which do not show a satisfactory response (not recovered) after the initial three treatments. In addition, for BRD only, administer intramuscularly or subcutaneously 1.0 mg cefotiofur equivalents/lb (2.2 mg/kg) BW every other day on Days 1 and 3 (48 h interval). Do not inject more than 15 mL per injection site.

Selection of dosage level (0.5 to 1.0 mg/lb) and regimen/duration (daily or every other day for BRD only) should be based on an assessment of the severity of disease, pathogen susceptibility and clinical response.

- For acute post-partum metritis: administer by intramuscular or subcutaneous administration at the dosage of 1.0 mg cefotiofur equivalents/lb (2.2 mg/kg) BW (2 mL sterile suspension per 100 lb BW). Administer at 24 h intervals for five consecutive days. Do not inject more than 15 mL per injection site.

CONTRAINDICATIONS

As with all drugs, the use of CEFENIL RTU is contraindicated in animals previously found to be hypersensitive to the drug.

WARNINGS

NOT FOR HUMAN USE. KEEP OUT OF REACH OF CHILDREN.

Penicillins and cephalosporins can cause allergic reactions in sensitized individuals. Topical exposures to such antimicrobials, including cefotiofur, may elicit mild to severe allergic reactions in some individuals. Repeated or prolonged exposure may lead to sensitization. Avoid direct contact of the product with the skin, eyes, mouth, and clothing. Persons with a known hypersensitivity to penicillin or cephalosporins should avoid exposure to this product. In case of accidental eye exposure, flush with water for 15 minutes. In case of accidental skin exposure, wash with soap and water. Remove contaminated clothing. If allergic reaction occurs (e.g., skin rash, hives, difficult breathing), seek medical attention. The safety data sheet contains more detailed occupational safety information. To report suspected adverse drug events, for technical assistance or to obtain a copy of the safety data sheet (SDS), contact Norbrook at 1-866-591-5777. For additional information about adverse drug experience reporting for animal drugs, contact FDA at 1-888-FDA-VETS or online at www.fda.gov/reportmalae.

RESIDUE WARNINGS:

Swine: When used according to label indications, dosage, and route of administration, treated swine must not be slaughtered for 4 days following the last treatment.

Use of dosages in excess of those indicated or by unapproved routes of administration may result in illegal residues in edible tissues.

Cattle: When used according to label indications, dosage and route of administration, treated cattle must not be slaughtered for 3 days following the last treatment. When used according to label indications, dosage and route of administration, a milk discard time is not required. Use of dosages in excess of those indicated or by unapproved routes of administration, such as intramammary, may result in illegal residues in edible tissues and/or milk. A withdrawal period has not been established in pre-ruminating calves. Do not use in calves to be processed for veal.

PRECAUTIONS

The effects of cefotiofur on cattle and swine reproductive performance, pregnancy, and lactation have not been determined.

Swine: Areas of discoloration associated with the injection site at time periods of 11 days or less may result in trim-out of edible tissues at slaughter. The safety of cefotiofur has not been demonstrated for pregnant swine or swine intended for breeding.

Cattle: Following intramuscular or subcutaneous administration in the neck, areas of discoloration at the site may persist beyond 11 days resulting in trim loss of edible tissues at slaughter. Following intramuscular administration in the rear leg, areas of discoloration at the injection site may persist beyond 28 days resulting in trim loss of edible tissues at slaughter.

CLINICAL PHARMACOLOGY

Swine: Cefotiofur administered as either cefotiofur sodium or cefotiofur hydrochloride is metabolized rapidly to desfuroylcefotiofur, the primary metabolite. Administration of cefotiofur to swine as either the sodium or hydrochloride salt provides effective concentrations of cefotiofur and desfuroylcefotiofur metabolites in plasma above the MIC₉₀ for the labeled pathogens: *Actinobacillus pleuropneumoniae*, *Pasteurella multocida*, *Streptococcus suis* and *Salmonella choleraesuis* for the 24 hour (h) period between the dosing intervals. The MIC₉₀ for *Salmonella choleraesuis* (1.0 µg/mL) is higher than the other three pathogens and plasma concentrations exceed this value for the entire dosing interval only after the 2.27 mg/lb (5.0 mg/kg) body weight (BW) dose.

Comparative Bioavailability Summary

Comparable plasma concentrations of cefotiofur administered as cefotiofur hydrochloride sterile suspension or cefotiofur sodium sterile solution were demonstrated after intramuscular administration of 2.27 mg cefotiofur equivalents/lb (5.0 mg/kg) BW. See Table 1 and Figure 2.

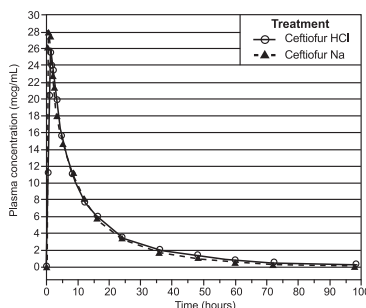
Table 1. Swine plasma concentrations and related parameters * of cefotiofur and desfuroylcefotiofur metabolites after cefotiofur hydrochloride sterile suspension, 50 mg/mL, or cefotiofur sodium sterile powder, 50 mg/mL, administered at 2.27 mg/lb cefotiofur equivalents/lb (5.0 mg/kg) BW IM.

	Cefotiofur hydrochloride	Cefotiofur sodium
C _{max} µg/mL:	26.1 ± 5.02	29.2 ± 5.01
t _{max} h:	0.66 – 2.0 (range)	0.33 – 2.0 (range)
AUC _{0-100h} µg·h/mL:	321 ± 50.2	314 ± 55.1
t _{1/2} h:	16.2 ± 1.55	14.0 ± 1.23
C _{24h} µg/mL:	3.45 ± 0.431	3.53 ± 0.791
C _{72h} µg/mL:	0.518 ± 0.126	0.407 ± 0.0675
t _{0.2} h:	93.8 ± 7.98	85.0 ± 7.71

Definitions:

- C_{max} - maximum plasma concentration in µg/mL.
 - t_{max} - the time after initial injection to when C_{max} occurs, measured in hours.
 - AUC_{0-100h} - the area under the plasma concentration vs. time curve from time of injection to the limit of quantitation of the assay (0.15 µg/mL).
 - t_{1/2} - the plasma half life of the drug in hours.
 - C_{24h} - the concentration of drug at 24 h after administration.
 - C_{72h} - the concentration of drug at 72 h after administration.
 - t_{0.2} - the time (in hours) plasma concentrations remain above 0.2 µg/mL.
- * Due to significant period effect and significant sequence effect in this study, data from period 1 only were used to evaluate these parameters.

Figure 2. Swine plasma concentrations of cefotiofur and desfuroylcefotiofur metabolites after cefotiofur hydrochloride sterile suspension, 50 mg/mL, or cefotiofur sodium sterile powder, 50 mg/mL, were administered intramuscularly at 2.27 mg cefotiofur equivalents/lb (5.0 mg/kg) BW.



Concentrations of total cefotiofur in the lungs of pigs administered radiolabeled cefotiofur at 2.27 or 3.41 mg cefotiofur equivalents/lb (5.0 or 7.5 mg/kg) BW 12 h after the last of three daily intramuscular injections at 24 h intervals averaged 3.66 and 5.63 µg/g.

Cattle: Cefotiofur administered as either cefotiofur sodium or cefotiofur hydrochloride is metabolized rapidly to desfuroylcefotiofur, the primary metabolite. Administration of cefotiofur to cattle as either the sodium or hydrochloride salt provides effective concentrations of cefotiofur and desfuroylcefotiofur metabolites in plasma above the MIC₉₀ for the bovine respiratory disease (BRD) label pathogens *Mannheimia haemolytica*, *Pasteurella multocida* and *Histophilus somni* for at least 48 h. The relationship between plasma concentrations of cefotiofur and desfuroylcefotiofur metabolites above the MIC₉₀ in plasma and efficacy has not been established for the treatment of bovine interdigital necrobacillosis (foot rot) associated with *Fusobacterium necrophorum* and *Bacteroides melaninogenicus*.

Comparative Bioavailability Summary

The comparability of plasma concentrations of cefotiofur following administration of cefotiofur hydrochloride sterile suspension or cefotiofur sodium sterile solution was demonstrated after intramuscular or subcutaneous administration of cefotiofur hydrochloride and intramuscular administration of cefotiofur sodium at 1.0 mg cefotiofur equivalents/lb (2.2 mg/kg) BW. See Table 2 and Figure 3.

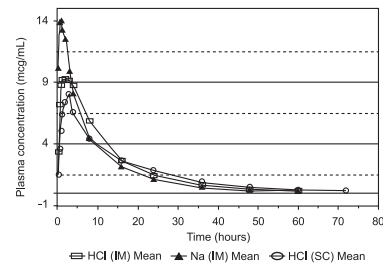
Table 2. Cattle plasma concentrations and related parameters of cefotiofur and desfuroylcefotiofur metabolites after cefotiofur hydrochloride sterile suspension, 50 mg/mL, administered intramuscularly or subcutaneously at 1.0 mg cefotiofur equivalents/lb (2.2 mg/kg) BW and cefotiofur sodium sterile powder, 50 mg/mL, administered intramuscularly at 1.0 mg cefotiofur equivalents/lb (2.2 mg/kg) BW.

	Cefotiofur hydrochloride	SC	Cefotiofur sodium
C _{max} µg/mL:	11.0 ± 1.69	8.56 ± 1.89	14.4–16.5
t _{max} h:	1–4 (range)	1–5 (range)	0.33–3.0
t _{0.2} h:	60.5 ± 6.27	51.0 ± 6.53	50.7–50.9
AUC _{0-100h} µg·h/mL:	160 ± 30.7	95.4 ± 17.8	115–142
t _{1/2} h:	12.0 ± 2.63	11.5 ± 2.57	9.50–11.1
C _{24h} µg/mL:	1.47 ± 0.380	0.926 ± 0.257	0.86–1.16
C _{48h} µg/mL:	0.340 ± 0.110	0.271 ± 0.086	0.250–0.268

Definitions:

- C_{max} - maximum concentration of drug in plasma in µg/mL.
- t_{max} - the time after initial injection to when C_{max} occurs, measured in hours.
- t_{0.2} - the time (in hours) plasma drug concentrations remain above 0.2 µg/mL.
- AUC_{0-100h} - the area under the plasma drug concentrations vs. time curve from time of injection to the limit of quantitation of the assay (0.15 µg/mL).
- t_{1/2} - the drug half life in plasma expressed in hours.
- C_{24h} - the plasma drug concentration 24 h after administration.
- C_{48h} - the plasma drug concentration 48 h after administration.

Figure 3. Cattle plasma concentrations of cefotiofur and desfuroylcefotiofur metabolites after cefotiofur hydrochloride sterile suspension, 50 mg/mL, was administered either intramuscularly or subcutaneously or cefotiofur sodium sterile powder, 50 mg/mL, was administered intramuscularly at 1.0 mg cefotiofur equivalents/lb (2.2 mg/kg) BW.



Total residues of cefotiofur were measured in the lungs of cattle administered radiolabeled cefotiofur at 1.0 mg cefotiofur equivalents/lb (2.2 mg/kg) BW at 24 h intervals for five consecutive days. Twelve h after the fifth injection of cefotiofur hydrochloride, total cefotiofur concentrations in the lung averaged 1.15 µg/g, while total cefotiofur concentrations in the lung 8 h after the fifth cefotiofur sodium injection averaged 1.18 µg/g.

CLINICAL MICROBIOLOGY

CEFENIL RTU is a ready to use formulation that contains the hydrochloride salt of cefotiofur, which is a broad spectrum cephalosporin antibiotic active against gram-positive and gram-negative bacteria including β-lactamase-producing strains. Like other cephalosporins, cefotiofur is bactericidal, *in vitro*, resulting in inhibition of cell wall synthesis.

Swine: Studies with cefotiofur have demonstrated *in vitro* and *in vivo* activity against gram-negative pathogens, including *Actinobacillus (Haemophilus) pleuropneumoniae*, *Pasteurella multocida*, *Salmonella choleraesuis*, and the gram-positive pathogen *Streptococcus suis*, all of which can be associated with swine bacterial respiratory disease – SRD (swine bacterial pneumonia). A summary of the minimum inhibitory concentration (MIC) values from SRD pathogens isolated from clinical field effectiveness studies is found in Table 3. Historic diagnostic laboratory MIC values for SRD pathogens from the US and Canada are found in Table 4.

Cattle: Studies with cefotiofur have demonstrated *in vitro* and *in vivo* activity against *Mannheimia haemolytica*, *Pasteurella multocida* and *Histophilus somni*, the three major pathogenic bacteria associated with bovine respiratory disease (BRD, pneumonia, shipping fever), and against *Fusobacterium necrophorum* and *Bacteroides melaninogenicus*, two of the major pathogenic anaerobic bacteria associated with acute bovine interdigital necrobacillosis (foot rot, pododermatitis). A summary of the MIC values for BRD and foot rot pathogens isolated from clinical field effectiveness studies is found in Table 3. Historic diagnostic MIC values for BRD and foot rot pathogens from the US and Canada are found in Table 4.

Antimicrobial Susceptibility

Summaries of MIC data are presented in Tables 3 and 4. Testing followed Clinical and Laboratory Standards Institute (CLSI) Guidelines.

Table 3. Cefotiofur MIC Values of Bacterial Isolates from Clinical Field Studies in the USA

Animal	Organism	Number Tested	Date Tested	MIC ₉₀ * (µg/mL)	MIC Range (µg/mL)	
Bovine	<i>Mannheimia haemolytica</i>	461	1988-1992	0.06	≤0.03-0.13	
	<i>Mannheimia haemolytica</i>	42	1993	0.015	≤0.003-0.03	
	<i>Pasteurella multocida</i>	318	1988-1992	0.06	≤0.03-0.25	
	<i>Pasteurella multocida</i>	48	1993	≤0.003	≤0.003-0.015	
	<i>Histophilus somni</i>	109	1988-1992	0.06	≤0.03-0.13	
	<i>Histophilus somni</i>	59	1993	≤0.0019	no range	
	<i>Fusobacterium necrophorum</i>	17	1994	≤0.06	no range	
	Swine	<i>Actinobacillus pleuropn.</i>	83	1993	≤0.03	≤0.03-0.06
		<i>Pasteurella multocida</i>	74	1993	≤0.03	≤0.03-0.06
<i>Streptococcus suis</i>		94	1993	0.25	≤0.03-1.0	
<i>Salmonella choleraesuis</i>		50	1993	1.0	1.0-2.0	
beta-hemolytic <i>Streptococcus</i> spp.		24	1993	≤0.03	≤0.03-0.06	
<i>Actinobacillus suis</i>		77	1998	0.0078	0.0019-0.0078	
<i>Haemophilus parasuis</i>		76	1998	0.06	0.0039-0.25	

*Minimum inhibitory concentration (MIC) for 90% of the isolates

Table 4. Cefiofur MIC Values of Bacterial Isolates from Diagnostic Laboratories* in the USA and Canada

Animal	Organism	Number Tested	Date Tested	MIC ^{**} (µg/mL)	MIC Range (µg/mL)	
Bovine	<i>Mannheimia haemolytica</i>	110	1997-1998	0.06	≤0.03-0.25	
	<i>Mannheimia haemolytica</i>	139	1998-1999	≤0.03	≤0.03-0.5	
	<i>Mannheimia haemolytica</i>	209	1999-2000	≤0.03	≤0.03-0.12	
	<i>Mannheimia haemolytica</i>	189	2000-2001	≤0.03	≤0.03-0.12	
	<i>Pasteurella multocida</i>	107	1997-1998	≤0.03	≤0.03-0.25	
	<i>Pasteurella multocida</i>	181	1998-1999	≤0.03	≤0.03-0.5	
	<i>Pasteurella multocida</i>	208	1999-2000	≤0.03	≤0.03-0.12	
	<i>Pasteurella multocida</i>	259	2000-2001	≤0.03	≤0.03-0.12	
	<i>Histophilus somni</i>	48	1997-1998	≤0.03	≤0.03-0.25	
	<i>Histophilus somni</i>	87	1998-1999	≤0.03	≤0.03-0.125	
	<i>Histophilus somni</i>	77	1999-2000	≤0.03	≤0.03-0.06	
	<i>Histophilus somni</i>	129	2000-2001	≤0.03	≤0.03-0.12	
	<i>Bacteroides fragilis</i> group	29	1994	16.0	≤0.06->16.0	
	<i>Bacteroides</i> spp., non- <i>fragilis</i> group	12	1994	16.0	0.13->16.0	
	<i>Peptostreptococcus anaerobius</i>	12	1994	2.0	0.13-2.0	
	Swine	<i>Actinobacillus pleuropn.</i>	97	1997-1998	≤0.03	no range
		<i>Actinobacillus pleuropn.</i>	111	1998-1999	≤0.03	≤0.03-0.25
		<i>Actinobacillus pleuropn.</i>	126	1999-2000	≤0.03	≤0.03-0.06
		<i>Actinobacillus pleuropn.</i>	89	2000-2001	≤0.03	≤0.03-0.06
		<i>Pasteurella multocida</i>	114	1997-1998	≤0.03	≤0.03-1.0
<i>Pasteurella multocida</i>		147	1998-1999	≤0.03	≤0.03-0.5	
<i>Pasteurella multocida</i>		173	1999-2000	≤0.03	≤0.03-0.06	
<i>Pasteurella multocida</i>		186	2000-2001	≤0.03	≤0.03-0.12	
<i>Streptococcus suis</i>		106	1997-1998	0.5	≤0.03-4.0	
<i>Streptococcus suis</i>		142	1998-1999	0.25	≤0.03-1.0	
<i>Streptococcus suis</i>		146	1999-2000	0.06	≤0.03-4.0	
<i>Streptococcus suis</i>		167	2000-2001	0.06	≤0.03-4.0	
<i>Salmonella choleraesuis</i>		96	1999-2000	1.0	0.03->4.0	
<i>Salmonella choleraesuis</i>		101	2000-2001	1.0	0.5-2.0	

*The following *in vitro* data are available but their clinical significance is unknown.

**Minimum inhibitory concentration (MIC) for 90% of the isolates.

Based on the pharmacokinetic studies of cefiofur in swine and cattle after a single intramuscular injection of 1.36 to 2.27 mg cefiofur equivalents/lb (3.0 to 5.0 mg/kg) BW (swine) or 0.5 to 1.0 mg cefiofur equivalents/lb (1.1 to 2.2 mg/kg) BW (cattle) and the MIC and disk (30 µg) diffusion data, the following breakpoints are recommended by CLSI.

Zone Diameter (mm)	MIC (µg/mL)	Interpretation
≥ 21	≤ 2.0	(S) Susceptible
18-20	4.0	(I) Intermediate
≤ 17	≥ 8.0	(R) Resistant

A report of "Susceptible" indicates that the pathogen is likely to be inhibited by generally achievable blood levels. A report of "Intermediate" is a technical buffer zone and isolates falling into this category should be retested. Alternatively the organism may be successfully treated if the infection is in a body site where drug is physiologically concentrated. A report of "Resistant" indicates that the achievable drug concentrations are unlikely to be inhibitory and other therapy should be selected. Standardized procedures¹ require the use of laboratory control organisms for both standardized diffusion techniques and standardized dilution techniques. The 30 µg cefiofur sodium disk should give the following zone diameters and the cefiofur sodium standard reference powder (or disk) should provide the following MIC values for the reference strain. Cefiofur sodium disks or powder reference standard is appropriate for both cefiofur salts.

Table 5. Acceptable quality control ranges for cefiofur against Clinical and Laboratory Standards Institute recommended American type Culture Collection (ATCC) reference strains

Organism name (ATCC No.)	Zone diameter* (mm)	MIC range (µg/mL)
<i>Escherichia coli</i> (25922)	26-31	0.25-1.0
<i>Staphylococcus aureus</i> (29213)	-	0.25-1.0
<i>Staphylococcus aureus</i> (25923)	27-31	-
<i>Pseudomonas aeruginosa</i> (27853)	14-18	16.0-64.0
<i>Actinobacillus pleuropneumoniae</i> (27090)	34-42**	0.004-0.015***
<i>Histophilus somni</i> (700025)	36-46**	0.0005-0.004***

*All testing performed using a 30 µg disk.

**Quality control ranges are applicable only to tests performed by disk diffusion test using a chocolate Mueller-Hinton agar, incubated in 5-7% CO₂ for 20-24 hours.

***MIC quality control ranges are applicable only to tests performed by broth microdilution procedures using veterinary fastidious medium (VFM).

CLINICAL EFFICACY

Cattle: In addition to demonstrating comparable plasma concentrations, the following clinical efficacy data are provided.

A clinical study was conducted to evaluate the efficacy of cefiofur hydrochloride administered subcutaneously for the treatment of the bacterial component of BRD under natural field conditions. When uniform clinical signs of BRD were present, 60 cattle (111 to 207 kg) were randomly assigned to one of the following treatment groups: negative control or cefiofur hydrochloride at 0.5 or 1.0 mg cefiofur equivalents/lb (1.1 or 2.2 mg/kg) BW. Treatments were administered daily for three consecutive days. Cattle were evaluated daily and animals that died or were euthanized were necropsied and the lung lesions scored. On Day 15, all surviving animals were euthanized and necropsied and the lung lesions scored. Mortality rates were 65%, 10% and 5% for negative controls, 0.5 mg cefiofur equivalents/lb and 1.0 mg cefiofur equivalents/lb, (1.1 or 2.2 mg/kg) BW, respectively. Mortality rates for both cefiofur hydrochloride treatment groups were lower than for negative controls (P < 0.0001). Rectal temperatures 24 h after third treatment were 104.0°F, 103.1°F and 102.8°F for negative controls, 0.5 mg/lb and 1.0 mg/lb (1.1 or 2.2 mg/kg) BW, respectively. The temperatures for both cefiofur hydrochloride treatment groups were lower than the negative controls (P ≤ 0.05). Cefiofur hydrochloride administered subcutaneously for three consecutive days at 0.5 or 1.0 mg cefiofur equivalents/lb (1.1 or 2.2 mg/kg) BW is an effective treatment for the bacterial component of BRD. A three-location clinical field study was conducted to evaluate the efficacy of cefiofur hydrochloride administered intramuscularly daily for three days or every other day (Days 1 and 3) for the treatment of the bacterial component of naturally occurring BRD. When uniform signs of BRD were present, 360 beef crossbred cattle were randomly assigned to one of the following treatment groups: negative control, cefiofur sodium at 0.5 mg cefiofur equivalents/lb (1.1 mg/kg) BW daily for three days, cefiofur hydrochloride at 1.0 mg cefiofur equivalents/lb (2.2 mg/kg) BW daily for three days, or cefiofur hydrochloride at 1.0 mg cefiofur equivalents/lb BW on Days 1 and 3 (every other day). All treatments were administered intramuscularly. All cefiofur treatment groups (hydrochloride and sodium) and treatment regimens (every day and every other day) significantly (P<0.05) reduced Day 4 rectal temperature as compared to the negative control. Clinical success on Days 10 and 28 and mortality to Day 28 were not different for the cefiofur groups (hydrochloride and sodium) and treatment regimens (every day and every other day). The results of this study demonstrate that daily and every other day (Days 1 and 3) intramuscular administration of cefiofur hydrochloride are effective treatment regimens for the bacterial component of BRD. An eight location study was conducted under natural field conditions to evaluate the efficacy of cefiofur hydrochloride for the treatment of acute post-partum metritis (0 to 14 days post-partum). When clinical signs of acute post-partum metritis (rectal temperature ≥103°F and fetid vaginal discharge) were observed, 361 lactating dairy cows were assigned randomly to treatment or negative control. Cattle were dosed either subcutaneously or intramuscularly, daily for five consecutive days. On days 1, 5 and 9 after the last day of dose administration, cows were evaluated for clinical signs of acute post-partum metritis. A cure was defined as rectal temperature <103°F and lack of fetid discharge. Cure rate for the 1.0 mg cefiofur equivalents/lb (2.2 mg/kg) BW dose group was significantly improved relative to cure rate of the negative control on day 9. The results of this study demonstrate that cefiofur hydrochloride administered daily for five consecutive days at a dose of 1.0 mg cefiofur equivalents/lb (2.2 mg/kg) BW is an effective treatment for acute post-partum metritis.

ANIMAL SAFETY

Swine: Results from a five-day tolerance study in normal feeder pigs indicated that cefiofur sodium was well tolerated when administered at 57 mg cefiofur equivalents/lb (125 mg/kg) (more than 25 times the highest recommended daily dosage of 2.27 mg/lb (5.0 mg/kg)) BW for five consecutive days. Cefiofur administered intramuscularly to pigs produced no overt adverse signs of toxicity.

To determine the safety margin in swine, a safety/toxicity study was conducted. Five barrows and five gilts per group were administered cefiofur sodium intramuscularly at 0, 2.27, 6.81 and 11.36 mg cefiofur equivalents/lb (0, 5, 15, 25 mg/kg) BW for 15 days. This is 0, 1, 3 and 5 times the highest recommended dose of 2.27 mg/lb (5.0 mg/kg) BW/day and 5 times the recommended treatment length of 3 days. There were no adverse systemic effects observed, indicating that cefiofur has a wide margin of safety when injected intramuscularly into feeder pigs at the highest recommended dose of 2.27 mg cefiofur equivalents/lb (5.0 mg/kg) BW daily for 3 days or at levels up to 5 times the highest recommended dose for 5 times the recommended length of treatment.

A separate study evaluated the injection site tissue tolerance of cefiofur hydrochloride in swine when administered intramuscularly in the neck at 1.36 and 2.27 mg cefiofur equivalents/lb (3.0 to 5.0 mg/kg) BW. Animals were necropsied at intervals to permit evaluations at 12 h, and 3, 5, 7, 9, 11, 15, 20, and 25 days after last injection. Injection sites were evaluated grossly at necropsy. No apparent changes (swelling or inflammation) were observed clinically after 12 h post-injection. Areas of discoloration associated with the injection site were observed at time periods less than 11 days after last injection.

Cattle: Results from a five-day tolerance study in feeder calves indicated that cefiofur sodium was well tolerated at 25 times (25 mg cefiofur equivalents/lb (55 mg/kg) BW) the highest recommended dose of 1.0 mg cefiofur equivalents/lb (2.2 mg/kg) BW for five consecutive days. Cefiofur administered intramuscularly had no adverse systemic effects. In a 15-day safety/toxicity study, five steer and five heifer calves per group were administered cefiofur sodium intramuscularly at 0 (vehicle control), 1, 3, 5 and 10 times the highest recommended dose of 1.0 mg cefiofur equivalents/lb (2.2 mg/kg) BW to determine the safety factor. There were no adverse systemic effects indicating that cefiofur sodium has a wide margin of safety when injected intramuscularly into the feeder calves at 10 times (10 mg cefiofur equivalents/lb (22 mg/kg) BW) the recommended dose for three times (15 days) the recommended length of treatment of three to five days. Local tissue tolerance to intramuscular injection of cefiofur hydrochloride was evaluated in the following study. Results from a tissue tolerance study indicated that cefiofur hydrochloride was well tolerated and produced no systemic toxicity in cattle when administered intramuscularly in the neck and rear leg at a dose of 1.0 mg cefiofur equivalents/lb (2.2 mg/kg) BW at each injection site. This represents a total dose per animal of 2.0 mg cefiofur equivalents/lb (4.4 mg/kg) BW. Clinically noted changes (local swelling) at injection sites in the neck were very infrequent (2/48 sites) whereas noted changes in rear leg sites were more frequent (21/48 sites). These changes in the rear leg injection sites were generally evident on the day following injection and lasted from 1 to 11 days. At necropsy, injection sites were recognized by discoloration of the subcutaneous tissues and muscle that resolved in approximately 7 to 15 days in the neck and 19 to 28 days in the rear leg. Results from another tissue tolerance study indicated that cefiofur hydrochloride was well tolerated and produced no systemic toxicity to cattle when administered subcutaneously at 0.5 or 1.0 mg cefiofur equivalents/lb (1.1 or 2.2 mg/kg) BW at 24 h intervals for 5 days. Mild and usually transient, clinically visible or palpable reactions (local swelling) were localized at the injection site. At necropsy, injection sites were routinely recognized by edema, limited increase in thickness and color changes of the subcutaneous tissue and/or fascial surface of underlying muscle. The fascial surface of the muscle was visibly affected in most cases through 9.5 days after injection. Underlying muscle mass was not involved. There were no apparent differences in tissue response to administration of cefiofur hydrochloride at 0.5 or 1.0 mg cefiofur equivalents/lb (1.1 or 2.2 mg/kg) BW.

TISSUE RESIDUE DEPLETION

Swine: A pivotal tissue residue decline study was conducted in swine. In this study, pigs received 2.27 mg of cefiofur per lb body weight (5 mg of cefiofur per kg body weight) per day for three consecutive days. Cefiofur residues in tissues were less than the tolerances for cefiofur residues in tissues such as kidney, liver and muscle by 4 days after dosing. These data collectively support a 4-day pre-slaughter withdrawal period in swine when used according to label directions.

Cattle: Two pivotal tissue residue decline studies were conducted in cattle. In the first study, cattle received an intramuscular injection of 1.0 mg of cefiofur per lb body weight (2.2 mg of cefiofur per kg body weight) for five consecutive days. Cefiofur residues in tissues were less than the tolerances for cefiofur residues in tissues such as kidney, liver and muscle by 3 days after dosing. In the second study, cattle received a subcutaneous injection of 1.0 mg of cefiofur per lb body weight (2.2 mg of cefiofur per kg body weight) for five consecutive days. Cefiofur residues in tissues were less than the tolerances for cefiofur residues in tissues such as kidney, liver and muscle by 3 days after dosing. These data collectively support a 3-day pre-slaughter withdrawal period in cattle when used according to label directions. In addition, two blood-level bioequivalence studies were conducted in cattle (one using subcutaneous administration and one using intramuscular administration). Blood concentrations of cefiofur (measured as cefiofur free acid equivalents) were greater than the analytical method's limit of quantification through 12 hours after administration, and these data demonstrated bioequivalence between Cefenil® RTU and the referenced listed new animal drug. These data support a zero-day milk discard time in lactating dairy cows.

STORAGE CONDITIONS

Do not store above 30°C (86°F). Shake well before using. Protect from freezing. Contents should be used within 42 days after the first dose is removed.

HOW SUPPLIED

CEFENIL RTU is available in 100 mL and 250 mL vials.

¹ Clinical and Laboratory Standards Institute (CLSI). Performance Standards for Antimicrobial Disk and Dilution Susceptibility Tests for Bacteria Isolated from Animals; Approved Standard – Second Edition. NCCLS document M31-A2. CLSI, 940 West Valley Road, Suite 1400, Wayne, Pennsylvania 19087-1898, 2002.

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