Sequential VFD ID Number

Veterinary Feed Directive for use in Calves, Beef Cattle and Nonlactating Dairy Cattle Chlortetracycline

Client: Business or Home Address:		JOE PRODUCER	Veterinarian:	DR JD EXAMPLE		
		222 SECOND DRIVE	Address:	111 FIRST LANE		
Phone Number:		XXX-XXX-XXXX	Phone Number:	XXX-XXX-XXXX		
		r of animals to be treated: 100 222 SECOND LANE				
		nd/or other animal identifications:				
FEED 4 OI	UNCES PER I	HEAD DAILY TO PROVIDE 750 MG PER 1	500 ANIMAL			
Indication	ns, Drug Lev	vel in Medicated Feed, and Duratio	n of Use (select one and spec	ify the additional required in	formation):	
	A. For Gro	wing Cattle (over 400 lbs): For the r Drug level: Duration of use:	g/ton in order to provide 70 n			
	 B. For Beef Cattle: For the control of bacterial pneumonia associated with shipping fever complex caused by <i>Pasteurella spp</i>. susceptible to chlortetracycline. Drug level:g/ton in order to provide 350 mg / head/ day 					
		Drug level: Duration of use:		ing / neau/ day		
	C. For Beef Cattle (under 700 lbs.): Control of active infection of anaplasmosis caused by <i>Anaplasma marginale</i> susceptible to chlortetracycline. Drug level: g/ton in order to provide 350 mg / head/ day Duration of use: days					
Ø	D. For Beef Cattle (over 700 lbs.): Control of active infection of anaplasmosis caused by <i>Anaplasma marginale</i> susceptible to chlortetracycline. Drug level: 6000 g/ton in order to provide 0.5 mg/lb body weight / day Duration of use: 180 days					
	E. For Calves, Beef, and Nonlactating Dairy Cattle; For treatment of bacterial enteritis caused by <i>Escherichia coli</i> and bacterial pneumonia caused by <i>Pasteurella multocida</i> susceptible to chlortetracycline. Drug level: g/ton in order to provide 10 mg/lb body weight / day Duration of use: days (1 to 5 days)					
Caution: l	Use of feed	containing this Veterinary Feed Di	rective (VFD) drug in a manno	er other than as directed on t	he labeling (extra-label use) is not permitted.	
		For use	in Dry Feeds Only. Not for U	se in Liquid feed Supplement	s.	
	Residue	e Warnings: Zero-day withdrawal p	period. A withdrawal period h Do not use in calves to be		his product in pre-ruminating calves.	
Combina	tion Use:					
	This VFD only authorizes the use of the VFD drug(s) cited in this order and is not intended to authorize the use of such drug(s) in combination with a other animal drugs.				ne use of such drug(s) in combination with any	
	This VFD authorizes the use of the VFD drug(s) cited in this order in the following FDA-approved, conditionally approved, or indexed combination(s) in medicated feed that contains the VFD drug(s) as a component. (List the specific approved combination(s))					
Ø		outhorizes the use of the VFD drug(s contains the VFD drug(s) as a compo	•	A-approved, conditionally approved, or indexed combination(s) in medicated		
VFD Issua	nce Date:	01-01-xxxx	VFD Expiration Date	e: 6-01-XXXX		
			•	Mon	th/Day/Year	
Veterinari	ian's Signat	ure: Digitally signed by Veterinarian		(Not to exceed 6 month	is from issuance date)	
Original	– Veterina	arian	Copy – Supplier		Copy – Client	

All parties must retain a copy of this veterinary feed directive for 2 years after issuance