



Deracin (chlortetracycline) Veterinary Feed Directive for use in Calves, Beef Cattle and Nonlactating Dairy Cattle

Client:	Veterinarian:		
Busines Home Address	Address:		
Phone #	#: Phone #:		
	ate number of animals to be treated:		
	of animals:		
	structions and/or other animal identifications:		
	an, Drug Level in Medicated Feed, and Duration of Use (select one and specify the additional required information): A) For Growing Cattle (over 400 lbs): For the reduction of the incidence of liver abscesses.		
	Drug level: g/ton in order to provide 70 mg / head / day Duration of use: days		
	B) For Beef Cattle: For the control of bacterial pneumonia associated with shipping fever complex caused by <i>Pasteurella</i> spp. susceptible to chlortetracycline. Drug level: g/ton in order to provide 350 mg / head / day Duration of use: days		
	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		
	D) For Beef Cattle (over 700 lbs.): Control of active infection of anaplasmosis caused by Anaplasma marginale susceptible to chlortetracycline. Drug level: g/ton in order to provide 0.5 mg/lb body weight / day Duration of use: days		
	E) For Beef and Nonlactating Dairy Cattle: As an aid in control of active infection of anaplasmosis caused by <i>Anaplasma marginale</i> susceptible to chlortetracycline when delivered in a free-choice feed. Drug level: g/ton in order to provide 0.5 to 2.0 mg/lb body weight / day Duration of use: days		
	F) 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> organisms susceptible to chlortetracycline. Drug Concentration:		
	☐ Complete Feed g/ton (500 to 4,000 g/ton to provide 10 mg/lb body weight / day)		
	☐ Top Dress g/ton (4,000 to 20,000 g/ton to provide 10 mg/lb body weight / day) Duration of Feeding: days (Feed for not more than 5 days)		

Caution: Use of feed containing this Veterinary Feed Directive (VFD) drug in a manner other than as directed on the labeling (extra-label use) is not permitted.









Combin	nation (Jse:			
		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) n combination with any other animal drugs.			
		FD authorizes the use of the VFD drug(s) cited in this order in the following FDA-approved, conditionally approved, or ed combination(s) in medicated feed that contains the VFD drug(s) as a component.			
		Drug(s) and Dose Range(s) [all listed doses are 90% DM basis]	Specifications*		
		12.9 to 90.8 g/ton decoquinate to provide 22.7 mg per 100 lb body weight day (Deccox®) [ANADA 200-622]	Calves, beef, and non-lactating dairy cattle.		
		90.9 to 535.7 g/ton decoquinate to provide 22.7 mg per 100 lb body weight day (Deccox®) [ANADA 200-622]	Calves, beef, and non-lactating dairy cattle.		
		Other FDA-approved, conditionally approved, or indexed combination:			
	*for complete information see the approved Type C medicated feed label This VFD authorizes the use of the VFD drug(s) cited in this order in any FDA-approved, conditionally approved, or indexed combination(s) in medicated feed that contains the VFD drug(s) as a component.				
VFD Issuance Date: VFD Expiration Date: Month/Day/Year (Not to exceed 6 months from issuance date)					
Veterina	rian's siç	gnature:			
Original – Veterinarian Copy – Supplier All parties must retain a copy of this VFD for 2 years after issuance					