Veterinary Feed Directive For Cattle

			OMYCIN® tracycline)	
Veterinari	ian:	•	Client:	
Address:			Business/Home Address: Phone:	
			ect one and specify additional required i	information)
	Growing Cattle (over 400 lb): For Drug Concentration: Duration of Feeding:	g/ton (to provide 7		
	Beef Cattle: Control of bacterial pneumonia associated with shipping fever complex caused by <i>Pasteurella spp.</i> susceptible to chlortetracycline.			
	Drug Concentration: Duration of Feeding:	days		
	chlortetracycline.		naplasmosis caused by <i>Anaplasma margi</i>	<i>inale</i> susceptible to
	Drug Concentration: g/ton (to provide 350 mg/head/day) Duration of Feeding: days			
	Beef Cattle (over 700 lb): Control of active infection of anaplasmosis caused by <i>Anaplasma marginale</i> susceptible to chlortetracycline.			
	Drug Concentration: g/ton (to provide 0.5 mg/lb body weight/day) Duration of Feeding: days			
5.	_	attle: As an aid in control owhen delivered in a free-c .0 mg/lb body weight/day)	f active infection of anaplasmosis caused hoice feed.	l by Anaplasma marginale
	6000 g/ton (to provide 0.5 to 2	.0 mg/lb body weight/day)		
	[Must use a FDA-approved proprie] 5000 g/ton (to provide 0.5 to 2)	-	in 21 CFR 558.128(e)(6).]	
	[Must use a FDA-approved proprie			
	700 g/ton (to provide 0.5 to 2.0 [Must use a FDA-approved proprie	tary formulation.]		
	Duration of Feeding:		pacterial enteritis caused by <i>Escherichia</i> c	coli and hacterial pneumonia
	caused by <i>Pasteurella multocida</i> ug Concentration:			on and bacterial pheumonia
	Complete Feed	_ g/ton (500 to 4,000 g/	ton to provide 10 mg/lb body weight/day	y) /da.x
Du	ration of Feeding:	g/ton (4,000 to 20,000 days (Feed for not more	g/ton to provide 10 mg/lb body weight/ e than 5 days)	udy)
•••••	USE OF FEED CONTAINING	THIS VETERINARY FEED	DIRECTIVE (VFD) DRUG IN A MANNER OTI TRA-LABEL USE) IS NOT PERMITTED.	HER THAN AS
	te number of Cattle to be treated: ructions and/or other animal ident		Premises or Location of Cattle:	
Special ilist	ructions and/or other animaridem	incations.		
Affirmation	of Intent (for combination VFD d	rugs): chock the appropria	to hav	
T	his VFD only authorizes the use of	the VFD drug(s) cited in th	is order and is not intended to authorize the	e use of such drug(s) in
T	ombination with any other animal his VFD authorizes the use of the \ ndexed combination(s) in medicate	VFD drug(s) cited in this ord	der in the following FDA-approved, conditio TD drug(s) as a component.	onally approved, or
	his VFD authorizes the use of the vombination(s) in medicated feed t	VFD drug(s) cited in this or	cific approved combination) der in any FDA-approved, conditionally app s) as a component.	roved, or indexed
WAR		uired. A withdrawal period	I has not been established for this product	in pre-ruminating calves.
,	D Issuance:(dd/m		Date of VFD Expiration:(Cannot exceed 6 months after issuance)	(dd/mm/yyyy)

Veterinarian's signature: _ WHITE Original - Veterinarian

CANARY Copy - Supplier

PINK Copy - Client