PharmGate Animal Health

Sequential VFD ID Number

Pennchlor (chlortetracycline) Veterinary Feed Directive for use in Cattle

Client: _____ Veterinarian: _____

Business or Home Address: _____ Address: _____

_____ _____

Phone #: _____ Phone #: _____

Approximate number of animals to be treated: _____

Location of animals: _____

Special Instructions and/or other animal identifications:

Indication, Drug Level in Medicated Feed, and Duration of Use (select one and specify the additional required information):

☐ 1) Growing Cattle (over 400 lb): For the reduction of the incidence of liver abscesses.
Drug level:_____g/ton (to achieve 70 mg/head/day)
Duration of use:_____days

☐ 2) Beef Cattle: Control of bacterial pneumonia associated with shipping fever complex caused by *Pasteurella spp.* susceptible to chlortetracycline.
Drug level:_____g/ton (to achieve 350 mg/head/day)
Duration of use:_____ days

☐ 3) Beef Cattle (under 700 lb): Control of active infection of anaplasmosis caused by *Anaplasma marginale* susceptible to chlortetracycline.
Drug level:_____g/ton (to achieve 350 mg/head/day)
Duration of use:_____ days

☐ 4) Beef Cattle (over 700 lb): Control of active infection of anaplasmosis caused by *Anaplasma marginale* susceptible to chlortetracycline.
Drug level:_____g/ton (to achieve 0.5 mg/lb BW/day)
Duration of use:_____ days

☐ 5) Calves, Beef and Non-Lactating Dairy Cattle: Treatment of bacterial enteritis caused by *Escherichia coli* and bacterial pneumonia caused by *Pasteurella multocida* organisms susceptible to chlortetracycline.
Drug level:_____g/ton (to achieve 10 mg/lb BW/day)
Duration of use: _____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.

Warning: A withdrawal period has not been established for this product in pre-ruminating calves. Do not use in calves to be processed for veal.

Indication 1: No withdrawal period required. Indications 2, 3, 4: Withdraw 48 hrs prior to slaughter. Indication 5: Withdraw 24 hrs prior to slaughter.

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Combination 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 any other animal drugs.

☐ 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))

	Drug(s) and Dose Range(s)	Specifications*
□	12.9 to 90.8 g/ton decoquinate to provide 22.7 mg/100 lb body weight per day decoquinate	Calves, beef and non-lactating dairy cattle
□	90.9 to 535.7 g/ton decoquinate to provide 22.7 mg/100 lb body weight per day decoquinate	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)

Veterinarian's signature: _____

Copy – Supplier **Copy – Client** **Original – Veterinarian**