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):

☐ **A)** 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 *Pasteurella* 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 *Anaplasma marginale* 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 Calves, Beef, and Nonlactating Dairy Cattle: For treatment of bacterial enteritis caused by *Escherichia coli* and bacterial pneumonia caused by *Pasteurella multocida* organisms susceptible to chlortetracycline.

Drug level: _____ g/ton in order to provide 10 mg/lb body weight / 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.

Withdrawal Period and Residue Warnings: No withdrawal period is required when used according to labeling. A withdrawal period has not been established for this product in pre-ruminating calves. Do not use in calves to be processed for veal.

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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.

	Drug(s) and Dose Range(s) [all listed doses are 90% DM basis]	Specifications*
☐	5 to 40 grams per ton monensin to provide 50 to 480 mg per head per day (RUMENSIN™) [NADA 95-735]	Growing beef steers and heifers fed in confinement for slaughter
☐	10 to 40 grams per ton monensin to provide 0.14 to 0.42 mg per pound of bodyweight per day, up to 480 mg per head per day (RUMENSIN™) [NADA 95-735]	Growing beef steers and heifers fed in confinement for slaughter
☐	10 to 200 grams per ton monensin to provide 0.14 to 1.0 mg per pound bodyweight per day (RUMENSIN™) [NADA 95-735]	Beef calves 2 months of age and older
☐	7.14 to 40 grams per ton monensin to provide 50 to 480 mg per pound bodyweight per day (RUMENSIN™) [NADA 95-735]	Growing beef steers and heifers fed in confinement for slaughter
☐	Other FDA-approved, conditionally approved, or indexed combinations:	

*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: _____

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