## Veterinary Feed Directive for Use in Beef Cattle

Phone #: Phone #: FAX or email: (optional)	iber, li
Address:	
Phone #:       Phone #:       FAX or email: (optional)         FAX or email: (optional)       FAX or email: (optional)       FAX or email: (optional)         Indication, Drug Level in Medicated Feed, and Duration of Use: (specify additional required information)       For reduction of incidence of liver abscesses associated with Fusobacterium necrophorum and Arcanobacterium pyogenes.         Drug Level:      g/ton (8 to 10 g/ton)       Duration of Feeding:      days         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.         Approximate number of Cattle to be treated:	
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Premise or Location of cattle:         Special Instructions and/or other animal identifications:         Affirmation of Intent (for combination VFD drugs): check the appropriate box:         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 drein 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, 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*         Monensin at 10 to 40 g/ton to provide 0.14 to 0.42 mg/lb bodyweight/day       For use in cattle fed in confinement for slaughter for the last 28 to 42 days on fer shared day	***
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Monensin at 10 to 40 g/ton to provide 0.14 to 0.42 mg/lb bodyweight/day Ractopamine at 9.8 to 24.6 g/ton to provide 90 to 430 mg/head/day	
bodyweight/day Ractopamine at 9.8 to 24.6 g/ton to provide 90 to 430 mg/head/day     mg/head/day     slaughter for the last 28 to 42 days on ference of the last 28 to 42 days on	
Ractopamine at 9.8 to 24.6 g/ton to provide 90 to 430 mg/head/day	اممم
mg/head/day	ea.
(augustical by Dymansis $M$ and Astansis $\Lambda$ ANADA 200 EQE)	
(supplied by Rumensin and Actogain, ANADA 200-585)	
Monensin at 10 to 40 g/ton to provide 0.14 to 0.42 mg /lb bodyweight/day For use in cattle fed in confinement for slaughter for the last 28 to 42 days on fe	aad
bodyweight/day Ractopamine at 8.2 to 24.6 g/ton to provide 70 to 430	ea.
mg/head/day	
(supplied by Rumensin <sup>™</sup> and Actogain <sup>®</sup> , ANADA 200-585)	
	For use in cattle fed in confinement for
<ul> <li>bodyweight/day</li> <li>Ractopamine up to 800 g/ton as a top dress to provide 70 to</li> </ul>	ea.
400 mg/head/day	
(supplied by Rumensin <sup>™</sup> and Actogain <sup>®</sup> , ANADA 200-585)	
Monensin at 10 to 40 g/ton to provide 0.14 to 0.42 mg/lb bodyweight/day For use in cattle fed in confinement for slaughter for the last 28 to 42 days on fe	aad
bodyweight/day Ractopamine at 9.8 to 24.6 g/ton to provide 90 to 430	eu.
mg/head/day	
Melengestrol acetate, at the rate of 0.125 to 1.0 mg/lb (top	
dress or mixed at bunk with other Type C feed) to provide 0.25- 0.5 mg/head/day. (supplied by Rumensin <sup>™</sup> and Actogain <sup>®</sup> and	
melengestrol acetate, MGA <sup>®</sup> , ANADA 200-583)	
Other FDA-approved, conditionally approved, or indexed	
combination.	
*for complete information see the approved Type C medicated feed label.	
<ul> <li>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.</li> </ul>	d
Warning: No withdrawal period is required.	
Date of VFD Issuance:(dd/mm/yyyy) Date of VFD Expiration:(dd/mm/yyyy)	

Veterinarian's signature: \_

(Cannot exceed 6 months after issuance)

Color Z Original – Veterinarian