Veterinary Feed Directive

Veterinarian:		Client:
Address:		Address:
		(business or home)
Phone:		Phone:
Fax or email (optional):		Fax or email (optional):
Drug(s) Name:	Drug(s) Level:	g/ton Duration of use:
		Number of reorders (refills) authorized (if permitted by the drug approval):
Indications for use (as approved):		
Caution (related to this medicated feed in	fany)*	
Oddaon (rolated to this medicated lood, r		
USE OF FEED CONTAINING THIS VETERINARY FEED DIRECTIVE (VFD) DRUG IN A MANNER OTHER THAN AS DIRECTED ON THE LABELING (EXTRALABEL USE) IS NOT PERMITTED		
Approximate Number of Animals:		
Premises:		
Other Identification (e.g., age, weight) (optional):		
Special Instructions (if any):		
Affirmation of intent (for combination VFD Drugs) (check one box)*: ("For VFD drugs for which there are no approved VFD combinations, only the first affirmation statement should be included on the VFD)		
☐ 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.		
		in this order in the following FDA-approved, conditionally approved, s the VFD drug(s) as a component:
Drug(s)		Drug Level(s) and any Special Instructions
☐ 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.		
	Withdrawal Time (if a be withdrawn	any): This VFD Feed must days prior to slaughter
VFD Date of Issuance:	(Month/Day/Year) VFD Exp	iration Date: (Month/Day/Year) (As specified in the approval; cannot exceed 6 months after issuance)
Veterinarian's Signature:		