V-MAX 50- virginiamycin powder Phibro Animal Health

V-MaxTM 50 (Virginiamycin) TYPE A MEDICATED ARTICLE

To be mixed in cattle feed

Active Drug Ingredient

Virginiamycin: 11%

(Contains 50 g virginiamycin activity per lb)

Caution

CAUTION: Federal law restricts medicated feed containing this veterinary feed directive (VFD) drug to use by or on the order of a licensed veterinarian.

Inert ingredients:

Processed grain by-products, roughage products, calcium carbonate, carboxymethylcellulose, mineral oil

Important: Must be diluted in feed before use

Follow Directions on Back Panel

V-Max is a registered trademark of Phibro Animal Health Corporation

Phibro Animal Health Corporation, Teaneck, NJ 07666

Approved by FDA under NADA # 140-998

8812000

101-9141-06

NET CONTENTS: 50 lb (22.7 kg)

Distributed by:

PHIBRO ANIMAL HEALTH, INC.

Teaneck, NJ 07666, USA

(50 g virginiamycin activity per lb)

For use in complete feeds for cattle fed in confinement for slaughter as specified below Directions for Use

for slaughter	mg/hd/day	complete feed (90% dry matter basis)		
Reduction of incidence of	85-240	13.5-16.0		
liver abscesse				
Caution: Not for use in animals intended for breeding.				

Mixing Directions

Preparation of Type B Medicated Feeds for cattle fed in confinement for slaughter—Thoroughly mix the following amounts of V-Max 50 Type A Medicated Article to make 1 ton of Type B Medicated Feed to provide the concentrations shown in Table 1. An intermediate blending step, consistent with the mixing equipment specifications, should be performed to ensure adequate mixing.

Table 1. Type B Medicated Feed

lb of V-Max 50 Type A Medicated Article per ton of supplement	Virginiamycin concentration in Type B Medicated Feed (g/ton)	
5	250	
10	500	
20	1,000	
40	2,000	
100	5,000	
200	10,000	

Preparation of Type C Medicated Feed for cattle fed in confinement for slaughter:

From Type B Medicated Feed: The Type B Medicated Feed must be diluted to a Type C Medicated Feed before being fed. Prepare a Type B Medicated Feed as described above. Thoroughly mix the V-Max 50 Type B Medicated Feed to make 1 ton of Type C Medicated Feed to provide 13.5 16.0 g of virginiamycin per ton of complete feed on a 90% dry matter basis using the examples provided in Table 2.

Table 2. Type C Medicated Feed prepared from V-Max 50 Type B Medicated Feed

lb of a 500g/ton Type B Medicated Feed per ton of complete feed	lb of a 1,000 g/ton Type B Medicated Feed per ton of complete feed	lb of a 5,000 g/ton Type B Medicated Feed per ton of complete feed	lb of a 10,000 g/ton Type B Medicated Feed per ton of complete feed	Virginiamycin concentration in complete feed 90% dry matter basis (g/ton)
54	27	5.4	2.7	13.5
64	32	6.4	3.2	16.0

From Type A Medicated Article: V-Max 50 Type A Medicated Article must be diluted to a Type C Medicated Feed before being fed. Thoroughly mix the Type A Medicated Article to make 1 ton of Type C Medicated Feed to provide 13.5 – 16.0 g of virginiamycin per ton of complete feed on a 90% dry matter basis using the examples provided in Table 3.

An intermediate blending step, consistent with the blending equipment specifications, should be performed to ensure adequate mixing.

Table 3. Type C Medicated Feed prepared from V-Max 50 Type A Medicated Article

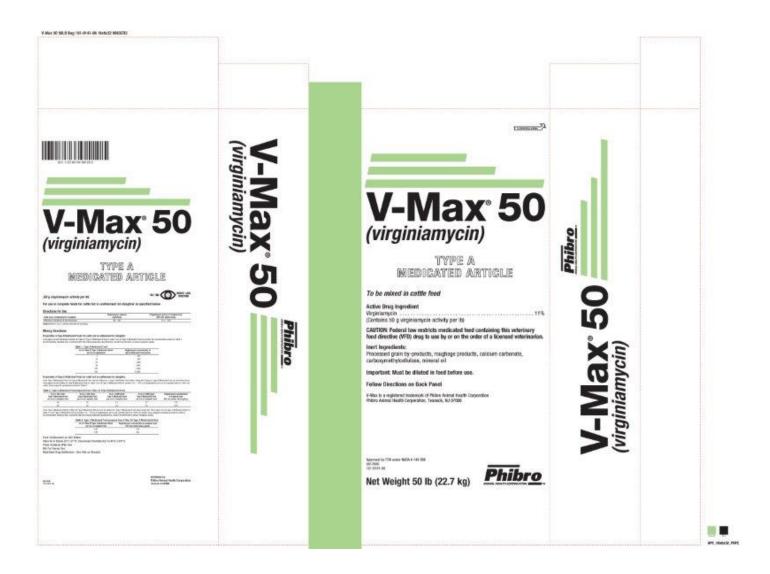
	Virginiamycin concentration in complete feed 90% dry matter basis (g/ton)
0.27	13.5
0.32	16.0

Feed continuously as sole ration

Store at or Below 25°C/77F°, Excursions Permitted Up To 40°C (104°F)
Close container after use
NOT FOR HUMAN USE
RESTRICTED DRUG (CALIFORNIA) - USE ONLY AS DIRECTED

V-Max 50

V-Max 50 Label



V-MAX 50

virginiamycin powder

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VFD TYPE A MEDICATED ARTICLE ANIMAL **Product Type** DRUG

Item Code (Source)

NDC:66104-9601

Route of Administration

ORAL

Active Ingredient/Active Moiety

Ingredient Name

Basis of Strength

Strength

VIRGINIAMYCIN (UNII: C49WS9N75L) (VIRGINIAMYCIN - UNII:C49WS9N75L)

VIRGINIAMYCIN

227 g in 0.45 kg

Inactive Ingredients

Ingredient Name	Strength
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CALCIUM CARBONATE (UNII: H0G9379FGK)

MINERAL OIL (UNII: T5L8T28FGP)

CARBOXYMETHYLCELLULOSE SODIUM (UNII: K6790BS311)

Packaging				
# Item Code	Package Description	Marketing Start Date	Marketing End Date	
1 NDC:66104-9601-1	22.7 kg in 1 BAG			
Marketing Information				
Marketing Category	Application Number or Mo Citation	onograph Marketing S Date	Start Marketing End Date	
NADA	NADA140998	04/01/2010		

Labeler - Phibro Animal Health (006989008)

Registrant - Phibro Animal Health (006989008)

Revised: 11/2023 Phibro Animal Health