

CATTLYST- laidlomycin propionate potassium granule
Phibro Animal Health

Cattlyst®
50G

Cattlyst 50G

Laidlomycin propionate potassium

Type A Medicated Article

Net Wt 50 LB (22.68 kg)

For further manufacturing only.

Active Drug Ingredient

Laidlomycin propionate potassium, 50 grams per pound, in a carrier suitable for incorporation in feed.

Weight Gain and Feed Efficiency

For increased rate of weight gain and improved feed efficiency in growing beef steers and heifers fed in confinement for slaughter when fed a complete Type C medicated feed containing 5 g/ton laidlomycin propionate potassium to provide 30 to 75 mg laidlomycin propionate potassium per head per day.

For improved feed efficiency in growing beef steers and heifers fed in confinement for slaughter when fed a complete Type C medicated feed containing 5 to 10 g/ton laidlomycin propionate potassium to provide 30 to 150 mg laidlomycin propionate potassium per head per day.

Not for use in animals intended for breeding.

Mixing And Feeding Instructions

Read all directions carefully before mixing and feeding.

Type B Medicated Feed

Thoroughly mix the following amounts of Cattlyst 50G to make one ton of Type B medicated feed containing the concentrations laidlomycin propionate potassium shown in TABLE 1. When using small quantities of Cattlyst 50G (e.g. two lb/ton or less), perform an intermediate blending step to ensure adequate mixing.

Type B liquid medicated feed manufactured from Cattlyst 50G must have a dry matter content between 62 and 75%, pH between 6.0 to 8.0, and concentration of laidlomycin propionate potassium of 100 to 2,000 g/ton. The label of the Type B liquid medicated

feed must carry daily, and prior to use, recirculating/agitation directions and an expiration of 21 days from date of manufacture. The Type C medicated feed manufactured from this Type B liquid medicated feed must be fed within 7 days from date of manufacture.

TABLE 1 - Type B Medicated Feed

Pounds of Cattlyst 50G per ton of Type B medicated feed	Laidlomycin propionate potassium concentration in Type B medicated feed (g/ton)
2	100
4	200
20	1,000
40	2,000

Type C Medicated Feed

Thoroughly mix the following amounts of Cattlyst 50G to make one ton of Type C Medicated Feed that provides 5-10 grams of laidlomycin propionate potassium per ton of feed on a 90% dry matter basis (TABLE 2). An intermediate blending step should be performed to ensure adequate mixing. For example, use one part Cattlyst 50G Type A Medicated Article with nine parts of feed ingredients to prepare an intermediate blend before preparing a Type C Medicated Feed.

TABLE 2 - Type C Medicated Feed

Pounds of Cattlyst 50G per ton of complete Type C medicated feed	Pounds of a 1,000 g/ton Type B medicated feed per ton of complete Type C medicated feed	Laidlomycin propionate potassium concentration in complete Type C medicated feed (g/ton)
0.1	10	5
0.15	15	7.5
0.20	20	10

Feeding Directions

Feed the Type C medicated feed as the sole ration.

Animal Safety Warning

Do not allow horses or other equines access to feeds containing laidlomycin propionate potassium (Cattlyst 50G). The safety of laidlomycin propionate potassium (Cattlyst 50G) in unapproved species has not been established.

User Safety Warnings

When mixing and handling Cattlyst 50G use appropriate protective clothing, eye, face, and respiratory protection. If accidental eye or skin contact occurs, rinse immediately

with water. Operators should wash thoroughly with soap and water after handling Cattlyst 50G.

Keep container closed between uses.

Store below 25°C (77°F), excursions permitted to 37°C (99°F).

Restricted Drug (California) - Use Only as Directed. Not for human use.

WITHDRAWAL PERIODS

No withdrawal period is required when used according to labeling.

Approved by FDA under NADA # 141-025

Questions/Comments? For technical assistance or to obtain a Safety Data Sheet(s), contact Phibro Animal Health Corporation at 888-403-0074 or www.pahc.com. To report side effects, contact Phibro Animal Health Corporation at 1-888-475-7355. For additional information about reporting side effects, contact FDA at 1-888-FDA-VETS or <http://www.fda.gov/reportanimalae>

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Distributed by:
Phibro Animal Health Corporation
Teaneck, NJ 07666

Net Wt 50 LB (22.68 kg)

176-9014-03

TAKE TIME



**OBSERVE LABEL
DIRECTIONS**

Cattlyst 50 Label

Label

Cattlyst 50G
(laidlomycin propionate potassium
Type A medicated article)

Phibro

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Cattlyst 50G
(laidlomycin propionate potassium
Type A medicated article)

Phibro

Cattlyst 50G
(laidlomycin propionate potassium
Type A medicated article)

recommend no print on short lap (except full coverage color)

Cattlyst 50G
(laidlomycin propionate potassium
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Net Wt 50 LB (22.68 kg)

Active Drug Ingredient:
Laidlomycin propionate potassium, 50 grams per pound, in a carrier suitable for incorporation in feed.

Indications for Use:
For increased rate of weight gain and improved feed efficiency in growing beef steers and heifers fed in confinement for slaughter when fed a complete Type C medicated feed containing 5 gram laidlomycin propionate potassium to provide 30 to 75 mg laidlomycin propionate potassium per head per day.
For improved feed efficiency in growing beef steers and heifers fed in confinement for slaughter when fed a complete Type C medicated feed containing 5 to 10 gram laidlomycin propionate potassium to provide 30 to 100 mg laidlomycin propionate potassium per head per day.
Not for use in animals intended for breeding.

Mixing Directions:
Read all directions carefully before mixing.
To Prepare Type B Medicated Feed:
Thoroughly mix the following amount of Cattlyst 50G to make one ton of Type B medicated feed containing the concentrations of laidlomycin propionate potassium shown in TABLE 1. When using small quantities of Cattlyst 50G (e.g., two-bale or less), perform an intermediate blending step to ensure adequate mixing.
Type B liquid medicated feed manufactured from Cattlyst 50G must have a dry matter content between 62 and 75% and between 6.0 to 6.5 laid concentration of laidlomycin propionate potassium of 100 to 2,000 g/ton. The label of the Type B liquid medicated feed must carry date, and prior to use, resubmitting/expiration date and an expiration of 21 days from date of manufacture. The Type C medicated feed manufactured from the Type B liquid medicated feed for the seller. Keep from date of manufacture.

TABLE 1 - Type B Medicated Feed

Pounds of Cattlyst 50G per ton of Type B medicated feed	Laidlomycin propionate potassium concentration in Type B medicated feed (g/ton)
2	100
4	200
20	1,000
40	2,000

To Prepare Type C Medicated Feed:
Thoroughly mix the following amount of Cattlyst 50G to make one ton of Type C Medicated Feed that provides 5 - 10 grams of laidlomycin propionate potassium per head per day in 100 to 2,000 g/ton. An intermediate blending step should be performed to ensure adequate mixing. For example, mix one part Cattlyst 50G Type A Medicated Article with nine parts of feed ingredients to prepare an intermediate blend before preparing a Type C Medicated Feed.

TABLE 2 - Type C Medicated Feed

Pounds of Cattlyst 50G per ton of complete Type C medicated feed	Pounds of a 1,000 g/ton Type B medicated feed per ton of complete Type C medicated feed	Laidlomycin propionate potassium concentration in complete Type C medicated feed (g/ton)
5.1	10	5
8.15	15	7.5
16.2	30	15

Feeding Directions:
Feed the Type C medicated feed on the sole ration.

Animal Safety Warnings:
Do not allow horses or other equine species to feed containing laidlomycin propionate potassium (Cattlyst 50G). The safety of laidlomycin propionate potassium (Cattlyst 50G) to companion equine has not been established.

User Safety Warnings:
When mixing and handling Cattlyst 50G use appropriate protective clothing, eye, face, and respiratory protection. If accidental contact with contact occurs, rinse immediately with water. Operators should wear breathing and eye and water after handling Cattlyst 50G. Keep container closed between uses.
Store below 30°C (87°F), moisture permitted to 33°C (91°F).
Restrict drug (Cattlyst) - Use only as directed. Not for human use.

WITHDRAWAL PERIODS
No withdrawal period is required when used according to labeling.

Approved for sale under #114-025

Supplier/Manufacturer: For technical assistance or to order a CATTLYST 50G Data Sheet, contact Phibro Animal Health Corporation at 800-863-6114 or visit our web site at www.phibro.com.
Phibro Animal Health Corporation, 1500 45th Street, Fort Dodge, IA 50501. For additional information, visit www.phibro.com.
Phibro Animal Health Corporation is a subsidiary of Phibro Animal Health Corporation, an affiliate.

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Net Wt 50 LB (22.68 kg)

Z Reference No.	FD Reference No.	Description	Colors
5148010	1006030	BAG CATTLYST 50G	Black
2148010	1006030	Canister	Black
6670 50	50	Quantity	Black
1007000	41005	Additional Information	Black

CATTLYST

laidlomycin propionate potassium granule

Product Information

Product Type	OTC TYPE A MEDICATED ARTICLE ANIMAL DRUG	Item Code (Source)	NDC:66104-2000
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LAILOMYCIN PROPIONATE POTASSIUM (UNII: 05TAA9I0Z8) (LAILOMYCIN - UNII:MI3S6ZF5QE)	LAILOMYCIN	50 g in 0.45 kg

Product Characteristics

Color	white (white to off white)	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:66104-2000-1	22.68 kg in 1 BAG		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
NADA	NADA141025	01/01/2009	

Labeler - Phibro Animal Health (006989008)

Revised: 1/2026

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