Lubabegron Medicated Cattle Feed
Type B Medicated Feed
(lubabegron Type B medicated feed)

Do Not Feed Undiluted

Indications for Use

For reduction of ammonia gas emissions per pound of live weight and hot carcass weight in beef steers and heifers fed in confinement for slaughter during the last 14 to 91 days on feed.

Effectiveness has not been demonstrated when fed for less than 14 days.

Ammonia gas emissions were measured for individual animals or small groups of animals held in environmentally controlled facilities. Based on existing information, reliable predictions of the reduction of ammonia gas emissions cannot be made on a herd, farm, or larger scale.

Increased rate of weight gain, improved feed efficiency, and increased carcass leanness have not been demonstrated with this product.

ACTIVE DRUG INGREDIENT

Lubabegron (as lubabegron fumarate)[a]…..……...….45 to 720 g/ton[*]

GUARANTEED ANALYSIS

Crude Protein, not less than……..……..……	_____ %
Non-Protein Nitrogen (NPN)[1], not more than……..	_____ %
Crude Fat, not less than……...…….…..……...…….	_____ %
Crude Fiber, not more than…….…..……..……...……	_____ %
Calcium, not less than…….……..…….....……	_____ %
Calcium, not more than…….…….....	_____ %
Phosphorus, not less than…….……..…….	_____ %
Salt[1], not less than…….….....	_____ %
Salt[1], not more than…….….....	_____ %
Sodium[2], not less than…….….....	_____ %
Sodium[2], not more than…….……...……	_____ %
Potassium[1], not less than…….……..……...……..	_____ %
Vitamin A[1], not less than…….……..……...……..	_____ I.U./lb

[1] Guarantee required only when nutrient source is added except when the feed is intended, represented, or serves as a principal source of the nutrient.

[2] Sodium guarantee required only when total sodium exceeds that furnished by the maximum salt guarantee.

INGREDIENTS

Each ingredient must be named in accordance with the names and definitions adopted by the Association of American Feed Control Officials.

MIXING DIRECTIONS

Thoroughly mix lubabegron Type B medicated feed in a ton of complete cattle feed according to the table below to obtain the proper concentration in the Type C medicated feed. The following table gives examples of how some Type B medicated feed concentrations can be prepared:

Lubabegron Concentration in Type B Medicated Feed (g/ton)	**Resulting lubabegron Concentration in Type C Medicated Feed (g/ton)[a]**		
	1.25 g/ton	2.50 g/ton	4.54 g/ton
	lb of Type B medicated feed to make 1 ton of Type C medicated feed		
50	50.0	100.0	180.0
350	7.1	14.3	25.7
700	3.6	7.1	12.9

[a] Based on 90% Dry Matter Basis

CAUTION

Not approved for use in breeding animals because safety and effectiveness have not been evaluated in these animals. Do not allow horses or other equines access to feed containing Experior. A decrease in dry matter intake may be noticed in some animals.

WARNING

No withdrawal period is required when used according to labeling.

User Safety Warning: The active ingredient in Experior, lubabegron, is a beta-adrenergic agonist/antagonist. Individuals with cardiovascular disease should exercise special caution to avoid exposure. Not for human use. Keep out of reach of children. When mixing and handling Experior, use protective clothing, impervious gloves, protective eye wear, and a NIOSH-approved dust mask. Operators should wash thoroughly with soap and water after handling. If accidental eye contact occurs, immediately rinse thoroughly with water; if wearing contact lenses, rinse eyes first, then remove contact lenses and continue to rinse for 5-20 minutes. If irritation persists, seek medical attention. The safety data sheet contains more detailed occupational safety information. To report adverse drug events, access medical information, or obtain additional product information, call Elanco US Inc. at 1-800-428-4441. For additional information about adverse drug experience reporting for animal drugs, contact FDA at 1-888-FDA-VETS or

http://www.fda.gov/AnimalVeterinary/SafetyHealth.

Lot Number:_____

MANUFACTURED BY:
BLUE BIRD FEED MILL
Any town, USA 12345

Net Weight lb (kg) on bag or bulk

[a] Sourced from Experior™. Approved by FDA under NADA #XXX-XXX.
*The medicated feed label must state a single drug concentration.