

SETT BEEF PRE-MIX No. 16

FOR SELF-FEEDING

Face Fly, Horn Fly, and Certain Internal Parasite Larvicide

Aids in preventing the breeding of horn flies and face flies in the manure of treated cattle; aids in the control of stomach worms (*Haemonchus*, *Ostertagia*, and *Trichostrongylus* spp.), hookworms (*Bunostomum* spp.), nodularworms (*Oesophagostomum* spp.), and largemouth bowelworms (*Chabertia* spp.) when used as directed.

ACTIVE INGREDIENT

Phenothiazine - - - - -	2.50%
INERT* INGREDIENTS - - - - -	97.50%
Total	100.00%

\* Refers only to ingredients which are not pesticides

GUARANTEED ANALYSIS

Calcium (Ca), Maximum - - - - -	13.00%
Calcium (Ca), Minimum - - - - -	11.00%
Phosphorus (P), Minimum - - - - -	6.00%
Salt (NaCl), Maximum - - - - -	20.00%
Salt (NaCl), Minimum - - - - -	18.00%
Iodine (I), Minimum - - - - -	0.007%
Magnesium (Mg), Minimum - - - - -	8.00%
Iron (Fe), Minimum - - - - -	0.20%
Copper (Cu), Minimum - - - - -	0.02%
Cobalt (Co), Minimum - - - - -	0.01%
Manganese (Mn), Minimum - - - - -	0.16%
Zinc (Zn), Minimum - - - - -	0.20%
Selenium (Se), Minimum - - - - -	0.0008%
Vitamin A, Minimum USP Units - - - - -	350,000
Vitamin D-3, Minimum USP Units - - - - -	200,000
Vitamin E, Minimum IU per lb. - - - - -	50

ACCEPTED  
with COMMENTS  
to EPA Letter Dated  
MAR 14 1983  
Under the Federal Insecticide,  
Fungicide, and Herbicide Act  
as amended, for the Pesticide  
registered under EPA Reg. No.  
40034-6

FEED INGREDIENTS

Monocalcium phosphate, dicalcium phosphate, calcium carbonate, salt, yeast culture, corn distillers dried grains with solubles, vitamin A acetate, D-activated animal sterol (source of vitamin D-3), vitamin E supplement, magnesium oxide, magnesium sulfate, potassium sulfate, copper oxide, cobalt carbonate, ethylenediamine dihydriodide (EDDI), iron sulfate, manganous oxide, zinc oxide, mineral oil, and sodium selenite.

KEEP OUT OF REACH OF CHILDREN



KEEP OUT OF REACH OF CHILDREN  
CAUTION  
See reverse of tag for additional precautionary statements.

PRECAUTIONARY STATEMENTS

HAZARDS TO HUMANS AND DOMESTIC ANIMALS

Do not administer to animals showing symptoms of acute respiratory conditions.

Individual animals are occasionally sensitive to phenothiazine. Sick, anemic, feverish, severely emaciated or generally weak animals, should be treated only under the advice of a veterinarian.

Do not treat lactating dairy animals. Do not administer within the last 4 weeks of pregnancy.

Do not treat animals with this product if they have been exposed to treatment with an organo-phosphate pesticide within several days since Phenothiazine may potentiate organo-phosphate toxicity. Occasionally, an animal may be sensitive to phenothiazine resulting in symptoms such as temporary blindness and loss of skin or hair. If toxic symptoms develop, take animals off pesticide and avoid exposure to sunlight for several days.

CAUTION. Harmful if swallowed, avoid contact with skin and eyes. Wash thoroughly with soap and water after handling and before eating or smoking. If in eyes, wash with plenty of water for 15 minutes. If irritation persists, see a physician. Not for human consumption.

ENVIRONMENTAL HAZARDS

This product is toxic to fish. Keep out of lakes, streams, and ponds. Do not contaminate water by cleaning equipment or disposal of wastes.

Net Wt. 50 lbs.

EPA Reg. No. 40034-6  
EPA Estab. No. 40034-OH-01

Manufactured By  
GOSSETT NUTRITION, INC.  
P.O. BOX 512  
MARION, OHIO 43302

**FEEDING DIRECTIONS:** It is a violation of Federal law to use this product in a manner inconsistent with its labeling.

Gossett BEEF PRE-MIX No.16 should be self-fed continuously as the sole source of salt and pre-mix from covered range feeders strategically placed near the cattle's drinking water or where they pass by them daily so they will have free access to them at all times.

For adequate control it is essential that the consumption of Phenothiazine be 2 grams per animal per day (3 oz. of Gossett Beef Pre-Mix No. 16).

If the daily intake is too high, add a little salt to cut down the consumption. If the intake is too low, add palatable protein such as cottonseed meal, soybean meal, etc., to increase the consumption.

Phenothiazine only destroys horn and face fly larvae. IT DOES NOT KILL OR REDUCE THE NUMBERS OF ADULT FACE OR HORN FLIES.

Since flies migrate from farm to farm, it is essential that a larvicide program be carried out on an area basis to assist in the reduction of adult face fly population. Therefore, the low level Phenothiazine program should be started as soon as the first flies appear in the spring. If the program is not started until the fly population has built up, it will take two to three weeks before horn fly population is appreciably reduced. During this two to three week period it is suggested that the animals be sprayed to cut down the horn fly population.

For internal parasite control, this product should be fed on a year round basis to cattle.

#### STORAGE AND DISPOSAL

Do not contaminate water, food, or feed by storage or disposal.

STORAGE: Store in a dry place. Keep separate from other feed products.

PESTICIDE DISPOSAL: Pesticide that cannot be used according to label instructions must be disposed of according to applicable Federal, State, or local procedures.

CONTAINER DISPOSAL: Completely empty bag by shaking and tapping sides and bottom to loosen clinging particles. Empty residue into application equipment. Then dispose of bags in a sanitary landfill or by incineration if allowed by state and local authorities.

U.S. ENVIRONMENTAL PROTECTION AGENCY  
OFFICE OF PESTICIDES PROGRAMS  
REGISTRATION DIVISION (WH-567)  
WASHINGTON, D.C. 20460

EPA REGISTRATION NO.	DATE OF ISSUANCE
TERM OF ISSUANCE	
NAME OF PESTICIDE PRODUCT	

NOTICE OF PESTICIDE:  REGISTRATION  
 REREGERISTRATION  
(Under the Federal Insecticide, Fungicide,  
and Rodenticide Act, as amended)

NAME AND ADDRESS OF REGISTRANT (Include ZIP code)

[Faint, illegible text]

161/87711  
18/1

NOTE: Changes in labeling formula differing in substance from that accepted in connection with this registration must be submitted to and accepted by the Registration Division prior to use of the label in commerce. In any correspondence on this product always refer to the above U.S. EPA registration number.

On the basis of information furnished by the registrant, the above named pesticide is hereby Registered/Reregistered under the Federal Insecticide, Fungicide, and Rodenticide Act.

A copy of the labeling accepted in connection with this Registration/Reregistration is returned herewith.

Registration is in no way to be construed as an indorsement or approval of this product by this Agency. In order to protect health and the environment, the Administrator, on his motion, may at any time suspend or cancel the registration of a pesticide in accordance with the Act. The acceptance of any name in connection with the registration of a product under this Act is not to be construed as giving the registrant a right to exclusive use of the name or to its use if it has been covered by others.

This product is conditionally registered in accordance with FIFRA Sec. 3(c)(7)(A) provided that you:

1. Submit and/or cite all data required for registration/reregistration of your product under FIFRA Sec. 3(c)(7) when the Agency requires all registrants of similar products to submit such data.

2. Make the labeling changes listed below before you release the product for shipment:

- Be sure to note the minimum type size requirements for the front panel signal word, the "KEEP OUT OF REACH OF CHILDREN", and the heading "STORAGE AND DISPOSAL". (Note: When submitting typed draft labeling, note the area of the front panel in square inches and the type sizes of the various labeling elements in parentheses next to each element.)

3. Submit five (5) copies of your final printed label a before you release the product for shipment. Refer to the 3-73 Enclosure for a further description of final printed labeling.

If these conditions are not complied with, the registration will be subject to cancellation in accordance with FIFRA Sec. 3(b). Your release for shipment of the product constitutes acceptance of these conditions.

ATTACHMENT IS APPLICABLE  
A stamped copy of the label is enclosed for your records.

SIGNATURE OF APPROVING OFFICIAL	DATE
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