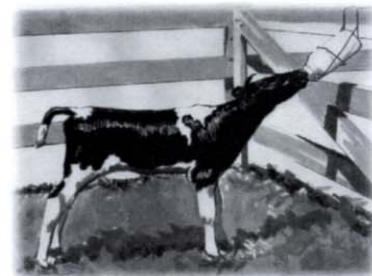


FrontLine®

MARKETING INFORMATION FOR TODAY'S FEED PROFESSIONAL



New Antibiotic Regulations for Chlortetracycline, Oxytetracycline and Neo-Oxy

The FDA has amended the usage of Chlortetracycline and Oxytetracycline in calf milk replacers, as well as the labeling for Neomycin/Oxytetracycline. The details for each drug are as follows:

Chlortetracycline

There will be only two choices of CTC drug level:

Drug Level	Claim	Use & Warning Statements
20 grams/ton	For increased weight gain and improved feed efficiency in replacement calves up to 250 pounds	A withdrawal period has not been established for this product in pre-ruminating calves. Do not use in calves to be processed for veal.
2000 grams/ton	For treatment of bacterial enteritis caused by <i>E. coli</i> and bacterial pneumonia caused by <i>P. multocida</i> organisms susceptible to chlortetracycline.	Treat for not more than 5 days. A withdrawal period has not been established for this product in pre-ruminating calves. Do not use in calves to be processed for veal.

All labeling must be changed upon receipt of Type A medicated articles bearing the revised labeling.

Neomycin/Oxytetracycline

The FDA is requiring a change in the warning statement labeling of milk replacer precuts containing Neomycin/Oxytetracycline. The labeling will read as follows:

WARNING: Withdraw this feed 30 days prior to slaughter. A withdrawal period has not been established for this product in pre-ruminating calves. Do not use in calves to be processed for veal.

Milk Products, Inc. has already received Type A drug in our inventory with this warning. Therefore, tag changes will be made as we reprint labeling.

Oxytetracycline (Oxy)

There will be two choices of Oxy available:

Drug Level	Claim	Use & Warning Statements
10-20 grams/ton	For increased rate of gain and improved feed efficiency.	None
2000 grams/ton	Treatment of bacterial enteritis caused by <i>E. coli</i> susceptible to oxytetracycline.	Feed continuously for 7 to 14 days. Withdraw 5 days before slaughter.

All labeling must be changed upon receipt of Type A medicated articles bearing the revised labeling.