Food and Drug Administration, HHS

§ 556.275 Fenbendazole.
(a) Acceptable daily intake (ADI). The ADI for total residues of fenbendazole is 40 micrograms per kilogram of body weight per day.
(b) Tolerances—(1) Cattle—(i) Liver (the target tissue). The tolerance for parent fenbendazole (the marker residue) is 0.8 part per million (ppm).
   (ii) Muscle. The tolerance for parent fenbendazole (the marker residue) is 0.4 ppm.
   (iii) Milk. The tolerance for fenbendazole sulfone metabolite (the marker residue in cattle milk) is 0.6 ppm.
(2) Swine—(1) Liver (the target tissue). The tolerance for parent fenbendazole (the marker residue) is 6 ppm.
   (ii) Muscle. The tolerance for parent fenbendazole (the marker residue) is 2 ppm.
   (3) Turkeys—(1) Liver (the target tissue). The tolerance for fenbendazole sulfone (the marker residue) is 6 ppm.
   (ii) Muscle. The tolerance for fenbendazole sulfone (the marker residue) is 2 ppm.
(4) Goats—(1) Liver (the target tissue). The tolerance for parent fenbendazole (the marker residue) is 0.8 ppm.
   (ii) Muscle. The tolerance for parent fenbendazole (the marker residue) is 0.4 ppm.

§ 556.277 Fenprostalene.

A tolerance for marker residue of fenprostalene in cattle is not needed. The safe concentrations for the total residues of fenprostalene in the uncooked edible tissues of cattle are 10 parts per billion in muscle, 20 parts per billion in liver, 40 parts per billion in kidney, 40 parts per billion in fat, and 100 parts per billion in the injection site. As used in this section “tolerance” refers to a concentration of a marker residue in the target tissue selected to monitor for total residues of the drug in the target animal, and “safe concentrations” refer to the concentrations of total residues considered safe in edible tissues.

§ 556.230 Erythromycin.
Tolerances for residues of erythromycin in food are established as follows:
(a) 0.1 part per million in uncooked edible tissues of beef cattle and swine.
(b) Zero in milk.
(c) 0.025 part per million in uncooked eggs.
(d) 0.125 part per million (negligible residue) in uncooked edible tissues of chickens and turkeys.

§ 556.240 Estradiol and related esters.
No residues of estradiol, resulting from the use of estradiol or any of the related esters, are permitted in excess of the following increments above the concentrations of estradiol naturally present in untreated animals:
(a) In uncooked edible tissues of heifers, steers, and calves:
   (1) 120 parts per trillion for muscle.
   (2) 480 parts per trillion for fat.
   (3) 360 parts per trillion for kidney.
   (4) 240 parts per trillion for liver.
(b) [Reserved]

§ 556.260 Ethopabate.
Tolerance for residues of ethopabate converted to metaphenetidine are established in the edible tissues of chickens as follows:
(a) 1.5 parts per million in uncooked liver and kidney.
(b) 0.5 part per million in uncooked muscle.

§ 556.273 Famphur.
Tolerances are established for residues of famphur including its oxygen analog in or on meat, fat, or meat by-products of cattle at 0.1 part per million.

§ 556.277 Fenprostalene.
a tolerance for marker residue of fenprostalene in cattle is not needed. The safe concentrations for the total residues of fenprostalene in the uncooked edible tissues of cattle are 10 parts per billion in muscle, 20 parts per billion in liver, 40 parts per billion in kidney, 40 parts per billion in fat, and 100 parts per billion in the injection site. As used in this section “tolerance” refers to a concentration of a marker residue in the target tissue selected to monitor for total residues of the drug in the target animal, and “safe concentrations” refer to the concentrations of total residues considered safe in edible tissues.