SUBCHAPTER E—ANIMAL DRUGS, FEEDS, AND RELATED PRODUCTS

PART 500—GENERAL

Subpart A [Reserved]

Subpart B—Specific Administrative Rulings and Decisions

Sec.
500.23 Thermally processed low-acid foods packaged in hermetically sealed containers.
500.24 Emergency permit control.
500.25 Anthelmintic drugs for use in animals.
500.26 Timed-release dosage form drugs.
500.27 Methylene blue-containing drugs for use in animals.
500.29 Gentian violet for use in animal feed.
500.30 Gentian violet for animal drug use.
500.35 Animal feeds contaminated with Salmonella microorganisms.
500.45 Use of polychlorinated biphenyls (PCB's) in the production, handling, and storage of animal feed.
500.46 Hexachlorophene in animal drugs.
500.50 Propylene glycol in or on cat food.

Subpart C—Animal Drug Labeling Requirements

500.51 Labeling of animal drugs; misbranding.
500.52 Use of terms such as "tonic", "tone", "toner", or "conditioner" in the labeling of preparations intended for use in or on animals.
500.55 Exemption from certain drug-labeling requirements.

Subpart D—Requirements for Specific Animal Drugs

500.65 Epinephrine injection 1:1,000 in 10-milliliter containers for emergency treatment of anaphylactoid shock in cattle, horses, sheep, and swine.

Subpart E—Regulation of Carcinogenic Compounds Used in Food-Producing Animals

500.80 Scope of this subpart.
500.82 Definitions.
500.84 Operational definition of "no residue".
500.86 Marker residue and target tissue.
500.88 Regulatory method.
500.90 Waiver of requirements.


SOURCE: 40 FR 13802, Mar. 27, 1975, unless otherwise noted.