whether bagged or in bulk, shall be adequately labeled to assure that the feed can be properly used.

Subpart I—Records

§ 225.202 Formula, production, and distribution records.
Records shall be maintained identifying the formulation, date of mixing, and if not for own use, date of shipment. The records shall be adequate to facilitate the recall of specific batches of medicated feed that have been distributed. Such records shall be retained on the premises for 1 year following the date of last distribution.

(Approved by the Office of Management and Budget under control number 0910–0152)

[51 FR 7390, Mar. 3, 1986]

PART 226—CURRENT GOOD MANUFACTURING PRACTICE FOR TYPE A MEDICATED ARTICLES

Subpart A—General Provisions

Sec. 226.1 Current good manufacturing practice.
226.10 Personnel.

Subpart B—Construction and Maintenance of Facilities and Equipment
226.20 Buildings.
226.30 Equipment.

Subpart C—Product Quality Control
226.40 Production and control procedures.
226.52 Components.
226.58 Laboratory controls.

Subpart D—Packaging and Labeling
226.80 Packaging and labeling.

Subpart E—Records and Reports
226.102 Master-formula and batch-production records.
226.110 Distribution records.
226.115 Complaint files.


SOURCE: 40 FR 14031, Mar. 27, 1975, unless otherwise noted.