TABLE 2.—ESTIMATED ANNUAL RECORDKEEPING BURDEN[1]

Standard No.	No. of Record-keepers	Annual Frequency per Record-keeping	Total Annual Records	Hours per Record-keeper	Total Hours
3, 4, and 6[2]	500	1	500	5	2,500

[1]There are no capital costs or operating and maintenance costs associated with this collection of information.
[2]The standards incorporate the best program management practices currently in use in the regulatory community. The recommended policies, procedures, and standard operating procedures contained in the various national standards are considered usual and customary management practices for State, local, and tribal agencies that regulate the retail segment of the food industry.

Dated: May 4, 2001.

William K. Hubbard,

Senior Associate Commissioner for Policy, Planning, and Legislation.

[FR Doc. 01–11618 Filed 5–8–01; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 99D–0239]

Agency Information Collection Activities; Announcement of OMB Approval; Request for Resolution of Scientific Disputes Concerning the Regulation of Medical Devices

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a collection of information entitled ''Request for Resolution of Scientific Disputes Concerning the Regulation of Medical Devices'' has been approved by the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995.

FOR FURTHER INFORMATION CONTACT: Peggy Schlosburg, Office of Information Resources Management (HFA–250), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–1223.

SUPPLEMENTARY INFORMATION: In the **Federal Register** of February 8, 2001 (66 FR 9585), the agency announced that the proposed information collection had been submitted to OMB for review and clearance under 44 U.S.C. 3507. An agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a currently valid OMB control number. OMB has now approved the information collection and has assigned OMB control number 0910–0467. The approval expires on April 30, 2004. A copy of the supporting statement for this information collection is available on the Internet at http://www.fda.gov/ohrms/dockets.

Dated: May 3, 2001.

William K. Hubbard,

Senior Associate Commissioner for Policy, Planning, and Legislation.

[FR Doc. 01–11583 Filed 5–8–01; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 01N–0179]

Purina Mills, Inc., et al.; Withdrawal of Approval of New Animal Drug Applications

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is withdrawing approval of 11 new animal drug applications (NADAs) listed below. In a final rule published elsewhere in this issue of the **Federal Register**, FDA is amending the animal drug regulations to remove the portions reflecting approval of the NADAs because the products are no longer manufactured or marketed.

DATES: Withdrawal of approval is effective May 21, 2001.

FOR FURTHER INFORMATION CONTACT: Pamela K. Esposito, Center for Veterinary Medicine (HFV–210), Food and Drug Administration, 7500 Standish Pl., Rockville, MD 20855, 301–827–5593.

SUPPLEMENTARY INFORMATION: The following sponsors have requested that FDA withdraw approval of the NADAs listed below because the products are no longer manufactured or marketed:

Sponsor	NADA No. Product (Drug)	21 CFR Cite Affected (Sponsor Drug Labeler Code)
Purina Mills, Inc., P.O. Box 66812, St. Louis, MO 63166–6812.	NADA 48–915 Purina® Bot Control (trichlorfon)	520.2520a (017800)
Golden Sun Feeds, Inc., 111 South Fifth St., Estherville, IA 51334.	NADA 97–567 Tylan® 10 Premix (tylosin phosphate).	558.625(b)(17) (021780)
.....	NADA 97–615 Swine Med-A-Mix TS 8000 Premix, Tylan® 5, 10, 20, 40 Sulfa-G (tylosin phosphate and sulfamethazine).	558.630(b)(4) and (b)(10) (021780)
Quali-Tech Products, Inc., 318 Lake Hazeltine Dr., Chaska, MN 55318–1093.	NADA 110–440 Hygromix Hygrowormer Hyanthelmix (hygromycin B).	558.274(a)(2), (a)(3), (a)(4), (c)(1)(i), and (c)(1)(ii) (016968)
Steris Laboratories, Inc., 620 North 51st Ave., Phoenix, AZ 85043–4705.	NADA 44–585 Oxytocin Injection	522.1680 (000402)
.....	NADA 45–578 Lidocaine Hydrochloride with Epi-nephrine Injection 2%.	522.1258 (000402)
.....	NADA 45–737 Sodium Pentobarbital Injection	522.1704(b) (000402)
.....	NADA 45–848 Phenylbutazone Injection	522.1720 (000402)
.....	NADA 110–349 Dexamethasone Injection	522.540(c)(2) (000402)