fees due, and to fulfill any applicable unsatisfied data requirements.

**V. Provisions for Disposition of Existing Stocks**

Existing stocks are those stocks of registered pesticide products which are currently in the United States and which were packaged, labeled, and released for shipment prior to the effective date of the cancellation action. EPA's existing stocks policy (56 FR 29362, June 26, 1991) provides that: "If a registrant requests to voluntarily cancel a registration where the Agency has identified no particular risk concerns, the registrant has complied with all applicable conditions of reregistration, conditional registration, and data call ins, and the registration is not subject to a Registration Standard, Label Improvement Program, or reregistration decision, the Agency will generally permit a registrant to sell or distribute existing stocks for 1 year after the cancellation request was received. Persons other than registrants will generally be allowed to sell, distribute, or use existing stocks until such stocks are exhausted."

Upon cancellation of the pesticides identified in Table 1, EPA anticipates allowing sale, distribution and use as described above. Exception to this general policy will be made in specific cases when more stringent restrictions on sale, distribution, or use of the products or their ingredients have already been imposed, as in a special review action, or where the Agency has identified significant potential risk concerns associated with a particular chemical.

**List of Subjects**

Environmental protection, Pesticides and pests.

Dated: January 14, 2010

**Richard P. Keigwin, Jr.,**

*Director, Pesticide Re-evaluation Division, Office of Pesticide Programs.*

[FR Doc. 2010–1476 Filed 1–25–10; 8:45 am]

**BILLING CODE 6560–50–S**

## FEDERAL COMMUNICATIONS COMMISSION

### Notice of Public Information Collection Being Reviewed by the Federal Communications Commission, Comments Requested

January 21, 2010.

**SUMMARY:** The Federal Communications Commission, as part of its continuing effort to reduce paperwork burden invites the general public and other Federal agencies to take this opportunity to comment on the following information collection(s), as required by the Paperwork Reduction Act of 1995, 44 U.S.C. 3501–3520. Comments are requested concerning (a) whether the proposed collection of information is necessary for the proper performance of the functions of the Commission, including whether the information shall have practical utility; (b) the accuracy of the Commission's burden estimate; (c) ways to enhance the quality, utility, and clarity of the information collected; (d) ways to minimize the burden of the collection of information on the respondents, including the use of automated collection techniques or other forms of information technology, and (e) ways to further reduce the information collection burden for small business concerns with fewer than 25 employees.

The FCC may not conduct or sponsor a collection of information unless it displays a currently valid control number. No person shall be subject to any penalty for failing to comply with a collection of information subject to the Paperwork Reduction Act (PRA) that does not display a valid control number.

**DATES:** Persons wishing to comments on this information collection should submit comments on or before March 29, 2010. If you anticipate that you will be submitting comments, but find it difficult to do so within the period of time allowed by this notice, you should advise the contact listed below as soon as possible.

**ADDRESSES:** Direct all PRA comments to Nicholas A. Fraser, Office of Management and Budget (OMB), via fax at (202) 395–5167, or via the Internet at Nicholas_A._Fraser@omb.eop.gov and to Judith B. Herman, Federal Communications Commission (FCC). To submit your PRA comments by e–mail send them to: PRA@fcc.gov.

**FOR FURTHER INFORMATION CONTACT:** Judith B. Herman, OMD, 202–418–0214. For additional information about the information collection(s) send an e–mail to PRA@fcc.gov or contact Judith B. Herman, 202–418–0214.

**SUPPLEMENTARY INFORMATION:**

OMB Control No: 3060–0295.

Title: Section 90.607(a)(1) and (b)(1), Supplemental Information To Be Furnished By Applicants For Facilities Under Subpart S.

Form No.: N/A.

Type of Review: Extension of a currently approved collection.

Respondents: Business or other for–profit; not–for–profit institutions, and state, local or tribal government.

Number of Respondents: 3,788 respondents; 3,788 responses.

Estimated Time Per Response: .25 hours.

Frequency of Response: One time reporting requirement.

Obligation to Respond: Required to obtain or retain benefits.

Total Annual Burden: 947 hours.

Privacy Act Impact Assessment: N/A.

Nature and Extent of Confidentiality: No questions of a confidential nature are asked.

Need and Uses: The Commission is submitting this information collection to the Office of Management and Budget (OMB) after this comment period in order to obtain the full three year clearance. There is a reduction in the number of respondents/responses and therefore, the total annual burden hours have been reduced.

This rule section requires the affected applicants to submit a list of any radio facilities they hold within 40 miles of the base station transmitter site being applied for. This information is used to determine if an applicant's proposed system is necessary in light of communications facilities it already owns. Such a determination helps the Commission to equitably distribute limited spectrum and prevents spectrum warehousing. The information is collected only once – upon initial license application.

Federal Communications Commission.

**Marlene H. Dortch,**

*Secretary, Office of the Secretary, Office of Managing Director.*

[FR Doc. 2010–1459 Filed 1–25–10; 8:45 am]

**BILLING CODE 6712–01–S**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Food and Drug Administration

**[Docket No. FDA–2009–N–0505]**

### Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Recordkeeping and Reporting Requirements for Human Food and Cosmetics Manufactured From, Processed With, or Otherwise Containing, Material From Cattle

**AGENCY:** Food and Drug Administration, HHS.

**ACTION:** Notice.

**SUMMARY:** The Food and Drug Administration (FDA) is announcing that a proposed collection of information has been submitted to the