TABLE 4.—ESTIMATED ANNUAL RECORDKEEPING BURDEN (NONREGISTERED UNLICENSED MIXER-FEEDERS)[1]

21 CFR Section	No. of Recordkeepers	Annual Frequency per Recordkeeping	Total Annual Records	Hours per Recordkeeper	Total Hours
225.142	45,000	4	180,000	1	180,000
225.158	45,000	1	45,000	4	180,000
225.180	45,000	32	1,440,000	.12	172,000
225.202	45,000	260	11,700,000	.33	3,861,000
Total					4,393,000

[1]There are no capital costs or operating and maintenance costs associated with this collection of information.

The estimate of the times required for record preparation and maintenance is based on agency communications with industry. Other information needed to finally calculate the total burden hours (i.e., number of recordkeepers, number of medicated feeds being manufactured, etc.) is derived from agency records and experience.

Dated: November 13, 2007.

Jeffrey Shuren,

Assistant Commissioner for Policy.

[FR Doc. E7–22587 Filed 11–16–07; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 2007N–0278]

Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Voluntary Registration of Cosmetic Product Establishments

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 Office of Management and Budget (OMB) for review and clearance under the Paperwork Reduction Act of 1995.

DATES: Fax written comments on the collection of information by December 19, 2007.

ADDRESSES: To ensure that comments on the information collection are received, OMB recommends that written comments be faxed to the Office of Information and Regulatory Affairs, OMB, Attn: FDA Desk Officer, FAX: 202–395–6974, or e-mailed to *baguilar@omb.eop.gov*. All comments should be identified with the OMB control number 0910–0027. Also include the FDA docket number found in brackets in the heading of this document.

FOR FURTHER INFORMATION CONTACT: Jonna Capezzuto, Office of the Chief Information Officer (HFA–250), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–4659.

SUPPLEMENTARY INFORMATION: In compliance with 44 U.S.C. 3507, FDA has submitted the following proposed collection of information to OMB for review and clearance.

Voluntary Registration of Cosmetic Product Establishments—(OMB Control Number 0910–0027)—Extension

The Federal Food, Drug, and Cosmetic Act (the act) provides FDA with the responsibility for assuring consumers that cosmetic products in the United States are safe and properly labeled. Cosmetic products that are adulterated under section 601 of the act (21 U.S.C. 361) or misbranded under section 602 of the act (21 U.S.C. 362) may not be distributed in interstate commerce. To assist FDA in carrying out its responsibility to regulate cosmetics, FDA has developed the Voluntary Cosmetic Registration Program (VCRP). In 21 CFR part 710, FDA requests that establishments that manufacture or package cosmetic products register with the agency on Form FDA 2511 entitled ''Registration of Cosmetic Product Establishment.'' The term ''Form FDA 2511'' refers to both the paper and electronic versions of the form. The electronic version of Form FDA 2511 is available on FDA's VCRP Web site at *http://www.cfsan.fda.gov/~dms/cos-regn.html*. FDA's online registration system, intended to make it easier to participate in the VCRP, was made available industry-wide on December 1, 2005. The agency strongly encourages electronic registration of Form FDA 2511 because it is faster and more convenient. A registering facility will receive confirmation of electronic registration, including a registration number, by e-mail, usually within 7 business days. The online system also allows for amendments to past submissions. Submission of the paper version of Form FDA 2511 remains an option as described in *http://www.cfsan.fda.gov/~dms/cos-reg2.html*. However, due to the high volume of online participation, the VCRP is allocating its limited resources primarily to electronic registrations.

Because registration of cosmetic product establishments is not mandatory, voluntary registration provides FDA with the best information available about the locations, business trade names, and types of activity (manufacturing or packaging) of cosmetic product establishments. FDA places the registration information in a computer database and uses the information to generate mailing lists for distributing regulatory information and for inviting firms to participate in workshops on topics in which they may be interested. FDA also uses the information for estimating the size of the cosmetic industry and for conducting onsite establishment inspections. Registration is permanent, although FDA requests that respondents submit an amended Form FDA 2511 if any of the originally submitted information changes.

In the **Federal Register** of July 19, 2007 (72 FR 39626), FDA published a 60-day notice requesting public comment on the information collection provisions. No comments were received.

FDA estimates the burden of this information collection as follows:

TABLE 1.—ESTIMATED ANNUAL REPORTING BURDEN[1]

21 CFR Part	Form	No. of Respondents	Annual Frequency per Response	Total Annual Responses	Hours per Response	Total Hours
710	FDA 2511	135	1	135	0.2	27

[1]There are no capital costs or operating and maintenance costs associated with this collection of information.

FDA bases its estimate on its review of the registrations received over the past 3 fiscal years. The total annual responses (averaged over fiscal years 2004 through 2006) is 9 times the previous total reported in 2004 (for fiscal years 2000 through 2003) due to increased participation by cosmetic companies, because of a renewed industry commitment to the program, and implementation of the online registration system on December 1, 2005. Due to the ease of online registration, FDA estimates that the hours per response have declined from 0.4 hours to 0.2 hours. Thus, the total estimated hour burden for this information collection is 27 hours, which is 4.5 times the previous level reported in 2004.

Dated: November 13, 2007.

Jeffrey Shuren,

Assistant Commissioner for Policy.

[FR Doc. E7–22588 Filed 11–16–07; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 2006F–0058]

ARCH Chemicals, Inc.; Withdrawal of Food Additive Petition FAP 6B4764

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing the withdrawal, without prejudice to a future filing, of a food additive petition (FAP 6B4764) proposing that the food additive regulations be amended to provide for the safe use of poly (iminoimidocarbonyliminoimidocarbonyliminohexamethylene) hydrochloride (CAS Reg No. 32289–58–0) as an antimicrobial agent in the manufacture of food-contact paper and paperboard.

FOR FURTHER INFORMATION CONTACT: Elizabeth S. Furukawa, Center for Food Safety and Applied Nutrition (HFS–275), Food and Drug Administration, 5100 Paint Branch Pkwy., College Park, MD 20740–3835, 301–436–1216, e-mail: *Elizabeth.Furukawa@fda.hhs.gov*.

SUPPLEMENTARY INFORMATION: In a notice published in the **Federal Register** of February 15, 2006 (71 FR 7975), FDA announced that a food additive petition (FAP 6B4764) had been filed by ARCH Chemicals, Inc., 1955 Lake Park Dr., suite 100, Smyrna, GA 30080. The petition proposed to amend the food additive regulations in 21 CFR 176.170 *Components of paper and paperboard in contact with aqueous and fatty foods* and 21 CFR 176.180 *Components of paper and paperboard in contact with dry food* to provide for the safe use of poly (iminoimidocarbonyliminoimido carbonyliminohexamethylene) hydrochloride (CAS Reg. No. 32289–58–0) as an antimicrobial agent in the manufacture of food-contact paper and paperboard. ARCH Chemicals, Inc., has now withdrawn the petition without prejudice to a future filing (21 CFR 171.7).

Dated: November 9, 2007.

Laura M. Tarantino,

Director, Office of Food Additive Safety, Center for Food Safety and Applied Nutrition.

[FR Doc. E7–22536 Filed 11–16–07; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[FDA 225–07–7001]

Memorandum of Understanding Between the Food and Drug Administration and the Association of American Feed Control Officials

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is providing notice of a memorandum of understanding (MOU) between FDA and the Association of American Feed Control Officials (AAFCO). The purpose of this MOU is to facilitate FDA's collaboration with AAFCO in the AAFCO New and Modified Ingredient Definition Process by clarifying the responsibilities of FDA and AAFCO in defining feed ingredients, in providing mechanisms for resolving disputes that may arise, and in providing mechanisms for modifying the ingredient definition process when required.

DATES: The agreement became effective August 30, 2007.

FOR FURTHER INFORMATION CONTACT: Sharon Benz, Division of Animal Feeds (HFV–220), Center for Veterinary Medicine, Food and Drug Administration, 7519 Standish Pl., Rockville, MD 20855, 240–453–6864.

SUPPLEMENTARY INFORMATION: In accordance with 21 CFR 20.108(c), which states that all written agreements and MOUs between FDA and others shall be published in the **Federal Register**, the agency is publishing notice of this MOU.

Dated: November 12, 2007.

Randall W. Lutter,

Deputy Commissioner for Policy.

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