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]

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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.

BILLING CODE 4160–01–S