Respondents	Number of respondents	Number of responses/respondents	Avg. burden/response (in hrs.)	Total burden (in hrs.)
Pediatricians	900	1	0.33	297

Dated: June 4, 1999.

Nancy Cheal,

Acting Associate Director for Policy, Planning and Evaluation Centers for Disease Control and Prevention (CDC).

[FR Doc. 99–14832 Filed 6–10–99; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 99F–1719]

Angus Chemical Co.; Filing of Food Additive Petition

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that Angus Chemical Co. has filed a petition proposing that the food additive regulations be amended to provide for the safe use of 4-(Diiodomethylsulfonyl) toluene as a slimicide in the manufacture of food-contact paper and paperboard.

FOR FURTHER INFORMATION CONTACT: Mark A. Hepp, Center for Food Safety and Applied Nutrition (HFS–215), Food and Drug Administration, 200 C St. SW., Washington, DC 20204, 202–418–3098.

SUPPLEMENTARY INFORMATION: Under the Federal Food, Drug, and Cosmetic Act (sec. 409(b)(5) (21 U.S.C. 348(b)(5))), notice is given that a food additive petition (FAP 9B4668) has been filed by Angus Chemical Co., c/o Phillip A. Johns, 10900 Silent Wood Pl., North Potomac, MD 20878–4829. The petition proposes to amend the food additive regulations in § 176.300 *Slimicides* (21 CFR 176.300) to provide for the safe use of 4-(Diiodomethylsulfonyl) toluene as a slimicide in the manufacture of food-contact paper and paperboard.

The agency has determined under 21 CFR 25.32(q) that this action is of the type that does not individually or cumulatively have a significant effect on the human environment. Therefore, neither an environmental assessment nor an environmental impact statement is required.

Dated: May 24, 1999.

Alan M. Rulis,

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

[FR Doc. 99–14839 Filed 6–10–99; 8:45 am]

BILLING CODE 4160–01–F

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 99N–1174]

Dietary Supplements; Center for Food Safety and Applied Nutrition Strategy; Public Meeting; Correction

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is correcting a document that appeared in the **Federal Register** of May 13, 1999 (64 FR 25889). The document announced a public meeting to solicit comments that will assist the Center for Food Safety and Applied Nutrition to develop an overall strategy for achieving effective regulation of dietary supplements under the Dietary Supplement Health and Education Act. The document published with an incorrect date for the submission of written comments. This document corrects that error.

FOR FURTHER INFORMATION CONTACT: Naomi Kulakow, Center for Food Safety and Applied Nutrition (HFS–165), Food and Drug Administration, 200 C St. SW., Washington, DC 20204, 202–205–8682, FAX 202–260–8957, e-mail ''nkulakow@bangate.fda.gov''.

In FR Doc. 99–12039, appearing on page 25889 in the **Federal Register** of Thursday, May 13, 1999, the following corrections are made:

1. On page 25889, in the third column, under the ''Dates'' caption, ''May 28, 1999.'' is corrected to read ''August 20, 1999.''

2. On page 25890, in the third column, under the ''Comments'' section, in the second line, ''May 28, 1999,'' is corrected to read ''August 20, 1999,''.

Dated: June 7, 1999.

William K. Hubbard,

Associate Commissioner for Policy Coordination.

[FR Doc. 99–14840 Filed 6–10–99; 8:45 am]

BILLING CODE 4160–01–F

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[FDA 225–98–4001]

Memorandum of Understanding Between the Food and Drug Administration and States of Iowa

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 the FDA and the State of Iowa Department of Public Health. The purpose of the MOU is to establish policies, procedures, and responsibilities for the billing and collection of mammography facility inspection fees under the Mammography Quality Standards Act.

DATES: The agreement became effective July 14, 1998.

FOR FURTHER INFORMATION CONTACT: Lireka P. Joseph, Center for Devices and Radiological Health (HFZ–200), Food and Drug Administration, 2094 Gaither Rd., Gaithersburg, MD 20850, 301–443–2845.

SUPPLEMENTARY INFORMATION: In accordance with 21 CFR 20. 108(c), which states that all written agreements and MOU's between FDA and others shall be published in the **Federal Register**, the agency is publishing notice of this MOU.

Dated: June 4, 1999.

William K. Hubbard,

Associate Commissioner for Policy Coordination.

BILLING CODE 4160–01–F