TABLE 1.—ESTIMATED ANNUAL REPORTING BURDEN[1]

No. of Respondents	Annual Frequency per Response	Total Annual Responses	Hours per Response	Total Hours
2,000	1	2,000	.5	1,000

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

The burden estimate is based on FDA's experience with the 1998 survey mentioned in the previous paragraph.

Dated: August 11, 2000.

William K. Hubbard,

Senior Associate Commissioner for Policy, Planning, and Legislation.

[FR Doc. 00–21007 Filed 8–17–00; 8:45 am]

BILLING CODE 4160–01–F

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 00C–1444]

FEM, Inc.; Filing of Color Additive Petition

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that FEM, Inc., has filed a petition proposing that the color additive regulations be amended to eliminate the limitation on the amount of silver used as a color additive in fingernail polish.

FOR FURTHER INFORMATION CONTACT: James C. Wallwork, Center for Food Safety and Applied Nutrition (HFS–215), Food and Drug Administration, 200 C St. SW., Washington, DC 20204, 202–418–3078.

SUPPLEMENTARY INFORMATION: Under the Federal Food, Drug, and Cosmetic Act (sec. 721(d)(1) (21 U.S.C. 379e(d)(1))), notice is given that a color additive petition (CAP 0C0272) has been filed by FEM, Inc., 1521 Laguna St. #210, Santa Barbara, CA 93101. The petition proposes to amend the color additive regulations in § 73.2500 *Silver* (21 CFR 73.2500) to eliminate the limitation on the amount of silver used as a color additive in fingernail polish.

The agency has determined under 21 CFR 25.32(r) that this action is of a 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: August 1, 2000.

Alan M. Rulis,

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

[FR Doc. 00–21012 Filed 8–17–00; 8:45 am]

BILLING CODE 4160–01–F

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 93F–0360]

Cognis Corporation; Withdrawal of Food Additive Petition

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 3B4400) proposing that the food additive regulations be amended to provide for the safe use of pentaerythritol mixed esters of $C_{16–18}$ fatty acids as a dispersant for titanium dioxide in polyethylene, polypropylene, and polystyrene intended for contact with food.

FOR FURTHER INFORMATION CONTACT: Vivian M. Gilliam, Center for Food Safety and Applied Nutrition (HFS–215), Food and Drug Administration, 200 C St. SW., Washington, DC 20204, 202–418–3094.

SUPPLEMENTARY INFORMATION: In a notice published in the **Federal Register** of October 29, 1993 (58 FR 58172), FDA announced that a food additive petition (FAP 3B4400) had been filed by Henkel Corporation, 300 Brookside Ave., Ambler, PA 19002–3498. The petition proposed to amend the food additive regulations to provide for the safe use of pentaerythritol mixed esters of $C_{16–18}$ fatty acids as a dispersant for titanium dioxide in polyethylene, polypropylene, and polystyrene intended for contact with food. Henkel Corporation has since changed its name to Cognis Corporation. Cognis Corporation has now withdrawn the petition without prejudice to a future filing (21 CFR 171.7).

Dated: August 1, 2000.

Alan M. Rulis,

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

[FR Doc. 00–21008 Filed 8–17–00; 8:45 am]

BILLING CODE 4160–01–F

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 98F–1193]

Troy Corporation; Withdrawal of Food Additive Petition

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 7B4533) proposing that the food additive regulations be amended to provide for the safe use of 3–iodo–2–propynyl butyl carbamate as a fungicidal additive for wood products intended to contact food.

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: In a notice published in the **Federal Register** of December 24, 1998 (63 FR 71295), FDA announced that a food additive petition (FAP 7B4533) had been filed by Troy Corporation, c/o S.L. Graham & Associates, 1801 Peachtree Lane, Bowie, MD 20721. The petition proposed to amend the food additive regulations in § 178.3800 *Preservatives for wood* (21 CFR 178.3800) to provide for the safe use of 3-iodo-2-propynyl butyl carbamate as a fungicidal additive for wood products intended to contact food. Troy Corporation has now withdrawn the petition without prejudice to a future filing (21 CFR 171.7).

Dated: July 26, 2000.

Alan M. Rulis,

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

[FR Doc. 00–21057 Filed 8–17–00; 8:45 am]

BILLING CODE 4160–01–F