Study	Respondents	No. of respondents	No. of re- sponses/re- spondent	Avg. burden/ response (in hrs.)	Total burden (in hrs.)
Prevalence	Households	100	1	0.75	75
	Daycare Centers	10	1	0.25	2.5
Longitudinal	Pregnant Women	300	3.5	0.167	175.35
	Infants	300	7	0.524	1,100.40
Total					1,353.25

2. Weekly and Annual Morbidity and Mortality Reports—In 1878 Congress authorized the U.S. Marine Hospital Service (later re-named the U.S. Public Health Service) to collect morbidity reports on cholera, smallpox, plague, and yellow fever from U.S. consuls overseas; this information was to be used for instituting quarantine measures to prevent the introduction and spread of these diseases in the United States. In 1879, a specific Congressional appropriation was made for the collection and publication of reports of these notifiable diseases. The authority for weekly reporting and publication was expanded by Congress in 1893 to include data from state and municipal authorities throughout the U.S. To increase the uniformity of the data, Congress enacted a law in 1902

directing the Surgeon General of the Public Health Service to provide forms for the collection and compilation of data and for the publication of reports at the national level.

In 1961, responsibility for the collection of data on nationally notifiable diseases and deaths in 121 U.S. cities was transferred from the National Office of Vital Statistics to CDC. For 37 years the MMWR has consistently served as CDC's main communication mode for disease outbreaks and trends in health and health behavior. In collaboration with the Council of State and Territorial Epidemiologists (CSTE), CDC has demonstrated the efficiency and effectiveness of computer transmission of data. The data collected electronically for publication in the MMWR provides

information which CDC and State epidemiologists use to detail and more effectively interrupt outbreaks. Reporting also provides the timely information needed to measure and demonstrate the impact of changed immunization laws or a new therapeutic measure. Users of data include, but are not limited to, congressional offices, state and local health agencies, health care providers, and other health related groups.

The dissemination of public health information is accomplished through the MMWR series of publications. The publications consist of the MMWR, the CDC Surveillance Summaries, the Recommendations and Reports, and the Annual Summary of Notifiable Diseases. The total cost to respondents is estimated to be \$48,100.

Respondents	No.of respondents	No. of re- sponses/re- spondent	Avg. burden/ response (in hrs.)	Total burden (in hrs.)
State and local health departments	178	52	.45	4,165
Total				4,165

Dated: January 29, 1998.

#### Wilma G. Johnson,

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

[FR Doc. 98–2677 Filed 2–3–98; 8:45 am] BILLING CODE 4163–18–P

# DEPARTMENT OF HEALTH AND HUMAN SERVICES

# Administration for Children and Families

# Submission for OMB Review; Comment Request

Title: Child Support Enforcement Program Financial Report, ACF–396. OMB No.: 0970–0014.

Description: Used by the States to report expenditures and estimates made

under title IV–D for the purposes of enforcing the support obligations owed by absent parents to their children and the spouse (or former spouse) with whom such children are living; locating absent parents; establishing paternity; and assuring that assistance in obtaining support will be available to all children for whom such assistance is required.

Respondents: Federal Government; and State, Local or Tribal Government.

#### ANNUAL BURDEN ESTIMATES

Instrument	Number of re- spondents	Number of responses per respondent	Average bur- den hours per response	Total burden hours
OCSE-396, Parts 1 & 2	54	4	4.25	918
	54	2	2.0	216

Estimated Total Annual Burden Hours: 1.134.

Additional Information: Copies of the proposed collection may be obtained by writing to The Administration for

Children and Families, Office of Information Services, Division of Information Resource Management Services, 370 L'Enfant Promenade, S.W., Washington, D.C. 20447, Attn: ACF Reports Clearance Officer.

OMB Comment: OMB is required to make a decision concerning the collection of information between 30 and 60 days after publication of this document in the **Federal Register**. Therefore, a comment is best assured of having its full effect if OMB receives it within 30 days of publication. Written comments and recommendations for the proposed information collection should be sent directly to the following: Office of Management and Budget, Paperwork Reduction Project, 725 17th Street, N.W., Washington, D.C. 20503, Attn: Ms. Wendy Taylor.

Dated: January 29, 1998.

#### **Bob Sargis**,

Acting Reports Clearance Officer. [FR Doc. 98–2633 Filed 2–3–98; 8:45 am] BILLING CODE 4184–01–M

# DEPARTMENT OF HEALTH AND HUMAN SERVICES

# Administration for Children and Families

### Submission for OMB Review; Comment Request

*Title:* Refugee Resettlement Program Estimates: CMA, ORR-1. *OMB No.:* 0970–0030.

Description: ORR reimburses, to the extent of available appropriations, certain non-Federal costs for the provision of cash and medical assistance to refugees, along with allowable expenses in the administration of the Refugee Resettlement Program. ORR needs sound State estimates of likely expenditures for refugee cash, medical, and administrative (CMA) expenditures so that it can anticipate Federal costs in upcoming quarters. If Federal costs are anticipated to exceed budget allocations, ORR must take steps to reduce Federal expenses, such as limiting the number of months of eligibility for Refugee Cash Assistance (RCA) and Refugee Medical Assistance (RMA).

To meet the need for reliable State estimates of anticipated expenses, ORR has developed a single-page form in which States estimate the average number of recipients for each category of assistance, the average unit cost over the next 12 months, and the expense for the overall administration of the program. This form, the ORR-1 (formerly Form FSA-601) must be submitted prior to the beginning of each Federal fiscal year. Without this information, ORR would be sent out of compliance with the intent of its legislation and otherwise unable to estimate program costs adequately.

In addition, the ORR–1 serves as the State's application for reimbursement of its CMA expenses. Submission of this form is thus required by section 412(a)(4) of the Immigration and Nationality Act which provides that "no grant or contract may be awarded under this section unless an appropriate proposal and application . . . are submitted to, and approved by, the appropriate administering official."

*Respondents:* State, Local or Tribal Government.

#### ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total bur- den hours
ORR-1	24	1	.5	.24

Estimated Total Annual Burden Hours: 24.

Additional Information: Copies of the proposed collection may be obtained by writing to The Administration for Children and Families, Office of Information Services, Division of Information Resource Management Services, 370 L'Enfant Promenade, S.W., Washington, D.C. 20447, Attn: ACF Reports Clearance Officer.

OMB Comment: OMB is required to make a decision concerning the collection of information between 30 and 60 days after publication of this document in the **Federal Register**. Therefore, a comment is best assured of having its full effect if OMB receives it within 30 days of publication. Written comments and recommendations for the proposed information collection should be sent directly to the following: Office of Management and Budget, Paperwork Reduction Project, 725 17th Street, N.W., Washington, D.C. 20503, Attn: Ms. Wendy Taylor.

Dated: January 29, 1998.

#### **Bob Sargis**,

Acting Reports Clearance Officer.
[FR Doc. 98–2710 Filed 2–3–98; 8:45 am]
BILLING CODE 4184–01–M

# DEPARTMENT OF HEALTH AND HUMAN SERVICES

**Food and Drug Administration** 

[Docket No. 98F-0053]

BP Chemicals, Inc.; Filing of Food Additive Petition

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

**ACTION:** Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that BP Chemicals, Inc., has filed a petition proposing that the food additive regulations be amended to provide for the safe use of nitrile rubber modified acrylonitrile-methyl acrylate copolymers as beverage containers. **DATES:** Written comments on the petitioner's environmental assessment by March 6, 1998.

ADDRESSES: Submit written comments to the Dockets Management Branch (HFA–305), Food and Drug Administration, 12420 Parklawn Dr., rm. 1–23, Rockville, MD 20857.

FOR FURTHER INFORMATION CONTACT: Richard H. White, 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:** 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 8B4564) has been filed by BP Chemicals, Inc., c/o The Weinberg Group, Inc., 1220 19th Street NW., suite 300, Washington, DC 20036–2400. The petition proposes to amend the food additive regulations in § 177.1480 *Nitrile rubber modified acrylonitrile*-