hundred thousand dollars ($1,300,000) to the Commission to be used for equitable relief, including restitution, and any attendant expenses for the administration of such equitable relief. To facilitate the payment of redress, Part VI of the proposed order requires Wacoal America to provide to the Commission a searchable electronic file containing the name and contact information of all consumers who purchased the iPants garments directly from respondent from January 1, 2011, through the date of entry of the order. Part VIII of the proposed order requires respondent to comply with the provisions of Appendix A to the order, which sets out the methods for notifying consumers who may be entitled to file a claim for consumer redress.

Part IX of the proposed order is triggered whenever the human clinical testing requirement in either Part II or Part III applies. Part IX of the proposed order requires the company to secure and preserve all underlying or supporting data and documents generally accepted by experts in the field as relevant to an assessment of the test. There is an exception for a ''Reliably Reported'' test defined as a test published in a peer-reviewed journal that was not conducted, controlled, or sponsored by any proposed respondent or supplier. Also, the published report must provide sufficient information about the test for experts in the relevant field to assess the reliability of the results.

Part X of the proposed order contains recordkeeping requirements for advertisements and substantiation relevant to any representation covered by the proposed order. Parts XI, XII and XIII of the proposed order require respondent to provide copies of the order to its personnel; to notify the Commission of changes in corporate structure that might affect compliance obligations under the order; and to file compliance reports with the Commission. Part XIV provides that the order will terminate after twenty (20) years, with certain exceptions.

The purpose of this analysis is to facilitate public comment on the proposed order, and it is not intended to constitute an official interpretation of the complaint or and proposed order or to modify the proposed order's terms in any way.

By direction of the Commission.

**Donald S. Clark,**

*Secretary.*

[FR Doc. 2014–23681 Filed 10–3–14; 8:45 am]

**BILLING CODE 6750–01–P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Announcement of Availability of the Report on Carcinogens, Thirteenth Edition

**AGENCY:** National Toxicology Program (NTP), National Institute of Environmental Health Sciences (NIEHS); National Institutes of Health (NIH).

**ACTION:** Availability of the Report on Carcinogens, Thirteenth Edition (13th RoC).

**SUMMARY:** The Department of Health and Human Services released the 13th RoC to the public on October 2, 2014. The report is available on the RoC Web site at: *http://ntp.niehs.nih.gov/go/roc13* or electronically from the Office of the RoC (see **ADDRESSES** below).

**DATES:** The 13th RoC is available to the public on October 2, 2014.

**ADDRESSES:** Dr. Ruth Lunn, Director, Office of the RoC, NTP, NIEHS, P.O. Box 12233, MD K2–14, Research Triangle Park, NC 27709; telephone: (919) 316–4637; FAX: (301) 480–2970; *lunn@niehs.nih.gov.*

**FOR FURTHER INFORMATION CONTACT:** Questions or comments concerning the 13th RoC should be directed to Dr. Ruth Lunn (telephone: (919) 316–4637 or *lunn@niehs.nih.gov*).

**SUPPLEMENTARY INFORMATION:**

#### Background Information on the RoC

The RoC is a congressionally mandated document that identifies and discusses agents, substances, mixtures, or exposure circumstances (collectively referred to as ''substances'') that may pose a hazard to human health by virtue of their carcinogenicity. Substances are listed in the report as either *known* or *reasonably anticipated to be human carcinogens.* The listing of a substance in the RoC indicates a potential hazard, but does not establish the exposure conditions that pose a cancer hazard to individuals in their daily lives. For each listed substance, the RoC provides information from cancer studies that support the listing as well as information about potential sources of exposure and current federal regulations to limit exposures. Each edition of the RoC is cumulative, that is, it lists newly reviewed substances in addition to substances listed in the previous edition. Information about the RoC is available on the RoC Web site (*http://ntp.niehs.nih.gov/go/roc13*) or by contacting Dr. Lunn (see **ADDRESSES** above).

The NTP prepares the RoC on behalf of the Secretary of Health and Human Services. For the 13th RoC, the NTP followed an established, multi-step process with multiple opportunities for public input, and used established criteria to evaluate the scientific evidence on each candidate substance under review (*http://ntp.niehs.nih.gov/go/rocprocess*).

#### New Listings to the 13th RoC

The 13th RoC contains 243 listings, some of which consist of a class of structurally related chemicals or agents. There are three new listings and one revised listing in this edition. The revised listing is for *ortho*-toluidine, which was previously listed as *reasonably anticipated to be a human carcinogen* and is now listed as *known to be a human carcinogen.* The new listings in the 13th RoC are three substances—1-bromopropane, cumene, and pentachlorophenol and by-products of its synthesis—each listed as *reasonably anticipated to be a human carcinogen.*

Dated: September 26, 2014.

**Linda S. Birnbaum,**

*Director, National Institute of Environmental Health Sciences, and National Toxicology Program.*

[FR Doc. 2014–23748 Filed 10–3–14; 8:45 am]

**BILLING CODE 4140–01–P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Administration for Children and Families

### Submission for OMB Review; Comment Request

*Title:* Temporary Assistance for Needy Families/National Directory of New Hires Match Results Report

*OMB No.:* 0970–0311

*Description:* Section 453(j)(3) of the Social Security Act (the Act) allows for matching between the National Directory of New Hires (maintained by the Federal Office of Child Support Enforcement (OCSE) and State TANF Agencies for purposes of carrying out responsibilities under programs funded under part A of Title IV of the Act. To assist OCSE and the Office of Family Assistance (OFA) in measuring savings to the TANF program attributable to the use of NDNH data matches, the State TANF Agencies have agreed to provide OCSE with a written description of the performance outputs and outcomes attributable to the State TANF Agency's use of NDNH match results. This information will help OCSE

demonstrate how the NDNH supports the OCSE's mission and strategic goals.

*Respondents:* State TANF Agencies

ANNUAL BURDEN ESTIMATES

| Instrument | Number of respondents | Number of responses per respondent | Average burden hours per response | Total burden hours |
|---|---|---|---|---|
| TANF/NDNH Match Results Report | 12 | 4 | 0.17 | 8.16 |

*Estimated Total Annual Burden Hours:* 8.16

*Additional Information:* Copies of the proposed collection may be obtained by writing to the Administration for Children and Families, Office of Planning, Research and Evaluation, 370 L'Enfant Promenade, SW., Washington, DC 20447, Attn: ACF Reports Clearance Officer. All requests should be identified by the title of the information collection. Email address: *infocollection@acf.hhs.gov.*

*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, Fax: 202–395–7285, Email: *OIRA_SUBMISSION@OMB.EOP.GOV,* Attn: Desk Officer for the Administration for Children and Families.

**Robert Sargis,**

*Reports Clearance Officer.*

[FR Doc. 2014–23670 Filed 10–3–14; 8:45 am]

**BILLING CODE 4184–01–P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Administration for Children and Families

### Submission for OMB Review; Comment Request

*Title:* Refugee Assistance Program Estimates: CMA—ORR–1

*OMB No.:* 0970–0030

Description: The ORR–1, Cash and Medical Assistance (CMA) Program Estimates, is the application for grants under the CMA program. The application is required by the Office of Refugee Resettlement (ORR) program regulations at 45 CFR 400.11(b). The regulation specifies that States must submit, as their application for this program, estimates of the projected costs they anticipate incurring in providing cash and medical assistance for eligible recipients and the costs of administering the program. Under the CMA program, States are reimbursed for the costs of providing these services and benefits for eight months after an eligible recipient arrives in this country. The eligible recipients for these services and benefits are refugees, Amerasians, Cuban and Haitian Entrants, asylees, Afghans and Iraqi with Special Immigrant Visas, and victims of a severe form of trafficking. States that provide services for unaccompanied refugee minors also provide an estimate for the cost of these services for the year for which they are applying for grants.

*Respondents:* Respondents are the 47 States and the District of Columbia that participate in the Refugee Resettlement program.

ANNUAL BURDEN ESTIMATES

| Instrument | Number of respondents | Number of responses per respondent | Average burden hours per response | Total burden hours |
|---|---|---|---|---|
| ORR–1, Cash and Medical Assistance Program Estimates | 47 | 1 | 0.60 | 28.20. |

*Estimated Total Annual Burden Hours:* 28.20

**Additional Information**

Copies of the proposed collection may be obtained by writing to the Administration for Children and Families, Office of Planning, Research and Evaluation, 370 L'Enfant Promenade SW., Washington, DC 20447, Attn: ACF Reports Clearance Officer. All requests should be identified by the title of the information collection. Email address: *infocollection@acf.hhs.gov.*

**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, *Fax:* 202–395–7285, *Email: OIRA_SUBMISSION@OMB.EOP.GOV*, Attn: Desk Officer for the Administration for Children and Families.

**Robert Sargis,**

*Reports Clearance Officer.*

[FR Doc. 2014–23672 Filed 10–3–14; 8:45 am]

**BILLING CODE 4184–01–P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### National Institutes of Health

### National Institute of Neurological Disorders and Stroke; Notice of Closed Meetings

Pursuant to section 10(d) of the Federal Advisory Committee Act, as amended (5 U.S.C. App.), notice is hereby given of the following meetings.

The meetings will be closed to the public in accordance with the provisions set forth in sections 552b(c)(4) and 552b(c)(6), Title 5 U.S.C., as amended. The grant applications and the discussions could disclose confidential trade secrets or commercial property such as patentable materials,