DEPARTMENT OF HEALTH AND HUMAN SERVICES

Agency for Health Care Policy and Research

Agency for Toxic Substances and Disease Registry

The Agency for Toxic Substances and Disease Registry announces the following meeting.

Name: Expert Panel on Pediatric Environmental Medicine.

Times and Dates: 3 p.m.–5 p.m., September 17, 1997; 9 a.m.–5 p.m., September 18, 1997; 8 a.m.–3 p.m., September 19, 1997.

Place: The Atlanta Marriott Norcross, 475 Technology Parkway, Norcross, Georgia 30092, telephone 770/263–8558.

Status: Open to the public, limited only by the space available. The meeting room accommodates approximately 50 people.

Purpose: The Agency for Toxic Substances and Disease Registry (ATSDR) is developing age-specific clinical practice guidelines in environmental medicine for children. The purpose of the expert panel workshop is to provide a forum for ATSDR to solicit individual expert consultation on issues of science and public health practice for the consideration in the development medical guidance. The guidelines initially proposed for development are: (1) general health assessment templates for children who have suffered environmental exposures and (2) a module on assessing growth and development in environmentally exposed children.

Matters To Be Considered: Participants will discuss their suggestions for pediatricians and environmental specialists confronted with situations in which children of various ages need environmental medicine evaluations; appropriate criteria for referral to specialists in environmental medicine or other specialties; how referrals should be arranged and what initial tests should be done. Guidance will be developed that (1) is age-specific; (2) is reasonable, quick, and easy-to-use; and (3) includes suggestions for basic standards of pediatric environmental medical care.

Agenda items are subject to change as priorities dictate.

Contact Person For More Information: Christine Rosheim, Health Education Specialist, Division of Health Education and Promotion, ATSDR, M/S E–33, 1600 Clifton Road NE, Atlanta, Georgia 30333, telephone 404/639–6205 or fax 404/ 639–6207.

Dated: August 28, 1997.

Carolyn J. Russell,

Director, Management Analysis and Services Office, Centers for Disease Control and Prevention (CDC).

[FR Doc. 97–23572 Filed 9–4–97; 8:45 am]

BILLING CODE 4163–70–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

The National Center for Environmental Health (NCEH) of the Centers for Disease Control and Prevention (CDC) announces the following meetings

Name: Teleconference meetings of the Ad Hoc Group for Early Hearing Detection and Intervention (EHDI), formerly known as the Ad Hoc Group for Universal Newborn Hearing Screening (UNHS).

Times and Dates: 2 p.m.–3 p.m., October 7, 1997; 2 p.m.–3 p.m., November 4, 1997; 2 p.m.–3 p.m., December 2, 1997; 2 p.m.–3 p.m., January 6, 1998; 2 p.m.–3 p.m., February 3, 1998; 2 p.m.–3 p.m., March 3, 1998.

Place: National Center for Environmental Health, Division of Birth Defects and Developmental Disabilities, Room 2103A, Building 101, 4770 Buford Highway, NE, Atlanta, Georgia 30341, telephone 770/488–7401.

Status: Open for participation by anyone with an interest in EHDI. All participants in the monthly conference calls are, by definition, members of the Ad Hoc Group for Early Hearing Detection and Intervention. Persons wishing to participate must E-mail or fax their request. The e-mail address is ehdi@cdc.gov; the fax number is 770/488–7361. Within one week of each teleconference, participants will be notified of the toll-free teleconference phone number, a caller code and the agenda. Each participant will have the responsibility to call in to connect to the conference call.

Purpose: This meeting will provide a forum for persons associated with EHDI programs to report and review relevant activities. Each conference call will be comprised of a series of scheduled presentations. Each presentation will be followed by a brief question and answer period. The agenda for the conference call will be determined by the Division of Birth Defects and Developmental Disabilities in collaboration with the Office on Disability and Health, NCEH (pending approval); the National Institute on Deafness and Communicative Disorders, National Institutes of Health; the Bureau of Maternal and Child Health, Health Resources and Services Administration; Office of Special Education and Rehabilitative Services, Department of Education; and others interested in early hearing detection programs.

Suggestions and feedback are invited by the conference call planners. Participants requesting to be on the agenda or to make written comments can send their requests or comments to the E-mail address or fax numbers noted above.

Matters To Be Discussed: Topics to be discussed during the meetings include progress on State and National activities to implement EHDI programs; progress on establishing State and National data systems on EHDI; and guidelines for establishing screening, diagnosis, and intervention protocols.

Contact Person for More Information: June Holstrum, Ph.D., Division of Birth Defects and Developmental Disabilities, NCEH, CDC, 4770 Buford Highway, NE, M/S F–15, Atlanta, Georgia 30341, telephone 770/488–7401, fax 770/488–7361.

Dated: August 28, 1997.

Carolyn J. Russell

Director, Management Analysis and Services Office, Centers for Disease Control and Prevention (CDC).

[FR Doc. 97–23578 Filed 9–4–97; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Proposed Information Collection Activity; Comment Request Proposed Projects

Title: Revised Form OCSE–100, State Plan for Child Support Collection and Establishment of Paternity Under Title IV–D of the Social Security Act.

OMB No.: 0970–0017.

Description: The State plan preprint and amendments serve as a contract with OCSE in outlining the activities the States will perform as required by law in order for States to receive federal funds to meet the costs of these activities. Due to enactment of HR2105, technical amendments for PRWORA, we are updating our State plan by revising 5 preprint pages. We are requesting approval of the revised State plan preprint pages for Section 2.1, Establishing Paternity and Securing Support, Section 2.5, Services to Individuals Not Receiving Title IV–A and IV–E Foster Care Assistance, Section 2.12–16 State Law Authorizing Suspension of Licenses, Section 2.12–20, Adoption of Uniform State Laws, and Section 3.16, Cooperation by Applicants for and Recipients of Part A Assistance. The information collected on the State plan pages is necessary to enable OCSE to monitor compliance with the requirements in Title IV–D of the Social Security Act and implementing regulations.

Respondents: States, Guam, Virgin Islands, Puerto Rico and District of Columbia.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours.
State Plan	54	5	.717	193

Estimated Total Annual Burden Hours: 193.

In compliance with the requirements of section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, the Administration for Children and Families is soliciting public comment on the specific aspects of the information collection described above. Copies of the proposed collection of information can be obtained and comments may be forwarded by writing to the Administration for Children and Families, Office of Information Services, Division of Information Resource Management Services, 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.

The Department specifically requests comments on: (a) Whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information shall have practical utility; (b) the accuracy of the agency's estimate of the burden of the proposed collection of information; (c) the quality, utility, and clarity of the information to be collected; and (d) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques or other forms of information technology. Consideration will be given to comments and suggestions submitted within 60 days of this publication.

Dated: August 29, 1997.

Bob Sargis,

Acting Reports Clearance Officer.

[FR Doc. 97–23649 Filed 9–4–97; 8:45 am]

BILLING CODE 4184–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 90F–0142]

Olin Corp.; Filing of Food Additive Petition; Amendment

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is amending the filing notice for a food additive petition filed by Olin Corp., to indicate that the petitioned additive, polyurethane resins derived from the reactions of toluene diisocyanate or 4,4′-methylenebis (cyclohexylisocyanate) with carboxylic acid-modified polypropylene glycol and with triethylamine and ethylenediamine as a component of adhesives for articles intended to contact food is more appropriately identified as polyurethane resins derived from the reactions of toluene diisocyanate or 4,4′-methylenebis (cyclohexylisocyanate) with fumaric acid-modified polypropylene glycol or fumaric acid-modified tripropylene glycol, triethylamine, and ethylenediamine as a component of adhesives for articles intended to contact food.

DATES: Written comments on the petitioner's environmental assessment by October 6, 1997.

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: Daniel N. Harrison, Center for Food Safety and Applied Nutrition (HFS–215), Food and Drug Administration, 200 C St. SW., Washington, DC 20204, 202–418–3084.

SUPPLEMENTARY INFORMATION: In a notice published in the **Federal Register** of May 10, 1990 (55 FR 19667), FDA announced that a food additive petition (FAP OB4201) had been filed by Olin Corp., 120 Long Ridge Rd., Stamford, CT 06904, proposing that § 175.105 *Adhesives* (21 CFR 175.105) be amended to provide for the safe use of polyurethane resins derived from the reaction of toluene diisocyanate or 4,4′-methylenebis (cyclohexylisocyanate) with carboxylic acid-modified polypropylene glycol and with triethylamine and ethylenediamine as a component of adhesives for articles intended to contact food.

Upon further review of the petition, the agency has determined that the petition specifically requests the use of polyurethane resins derived from the reaction of toluene diisocyanate or 4,4′-methylenebis (cyclohexylisocyanate) with fumaric acid-modified polypropylene glycol or fumaric acid-modified tripropylene glycol, triethylamine, and ethylenediamine as a component of adhesives for articles intended to contact food. Therefore, FDA is amending the filing notice of May 10, 1990, to state that the petitioner requests that the food additive regulations be amended to provide for the safe use of polyurethane resins derived from the reaction of toluene diisocyanate or 4,4′-methylenebis (cyclohexylisocyanate) with fumaric acid-modified polypropylene glycol or fumaric acid-modified tripropylene glycol, triethylamine, and ethylenediamine as a component of adhesives for articles intended to contact food.

The potential environmental impact of this action is being reviewed. To encourage public participation consistent with regulations promulgated under the National Environmental Policy Act (40 CFR 1501.4(b)), the agency is placing the environmental assessment submitted with the petition that is the subject of this notice on public display at the Dockets Management Branch (address above) for public review and comment. Interested persons may, on or before October 6, 1997, submit to the Dockets Management Branch (address above) written comments. Two copies of any comments are to be submitted, except that individuals may submit one copy. Comments are to be identified with the docket number found in brackets in the heading of this document. Received comments may be seen in the office above between 9 a.m. and 4 p.m., Monday through Friday. FDA will also place on public display any amendments to, or comments on, the petitioner's environmental assessment without further announcement in the **Federal Register**. If, based on its review, the agency finds that an environmental impact statement is not required and this petition results in a regulation, the notice of availability of the agency's finding of no significant impact and the evidence supporting that finding will be published with the regulation in the