# ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per re- spondent	Average burden hours per response	Total bur- den hours
Lien	53,254	1	0.25	13,313

#### *Estimated Total Annual Burden Hours:* 13,313.

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, SW., Washington, DC 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: August 18, 1997.

#### Bob Sargis,

Acting Reports Clearance Officer. [FR Doc. 97–22285 Filed 8–21–97; 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:* Administrative Subpoena. *OMB No.:* 0970–0152. *Description: Respondents:* Individuals and Households; not-for-profit institutions;

business or other for-profit; and State, Local or Tribal Govt. Annual Burden Estimates:

Instrument	Number of respondents	Number of responses per re- spondent	Average burden hours per response	Total bur- den hours
Subpoena	15,391	1	0.5	7,696

#### *Estimated Total Annual Burden Hours:* 7,696.

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, SW., Washington, DC 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, NW., Washington, DC 20503, Attn: Ms. Wendy Taylor.

Dated: August 18, 1997.

#### Bob Sargis,

Acting Reports Clearance Officer. [FR Doc. 97–22286 Filed 8–21–97; 8:45 am] BILLING CODE 4184–01–M

#### DEPARTMENT OF HEALTH AND HUMAN SERVICES

#### Food and Drug Administration

#### Advisory Committee; Notice of Meeting

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

## ACTION: Notice.

This notice announces a forthcoming meeting of a public advisory committee of the Food and Drug Administration (FDA). At least one portion of the meeting will be closed to the public.

*Name of Committee:* General Hospital and Personal Use Devices Panel of the Medical Devices Advisory Committee.

*General Function of the Committee:* To provide advice and recommendations to the agency on FDA

regulatory issues. *Date and Time:* The meeting will be held on September 15, 1997, 10 a.m. to

5:30 p.m., and September 15, 1997, 10 a.m. to a.m. to 12 m.

*Location:* Holiday Inn, Walker/ Whetstone Rooms, Two Montgomery Village Ave., Gaithersburg, MD.

*Contact Person:* Martha T. O'Lone, Center for Devices and Radiological Health (HFZ-480), Food and Drug Administration, 9200 Corporate Blvd., Rockville, MD 20850, 301-443-8913, or FDA Advisory Committee Information Line, 1-800-741-8138 (301-443-0572 in the Washington, DC area), code 12520. Please call the Information Line for upto-date information on this meeting.

Agenda: On September 15 and 16, 1997, the committee will discuss and make recommendations on the draft guidance entitled "Testing for Skin Sensitization to Chemicals in Latex Products." Single copies of this draft guidance are available to the public from the Division of Small Manufacturers Assistance, 1350 Piccard Dr., Rockville, MD 20851, 1-800-638-2041, or on the Internet using the World Wide Web (WWW) (http:// www.fda.gov/cdrh/draftgui.html).

Procedure: On September 15, 1997, from 10:30 a.m. to 5:30 p.m., and September 16, 1997, from 8 a.m. to 12 m., the meeting is open to the public. Interested persons may present data, information, or views, orally or in writing, on issues pending before the committee. Written submissions may be made to the contact person by August 9, 1997. Oral presentations from the public will be scheduled between approximately 11 a.m. and 12 m. on September 15, 1997. Time allotted for each presentation may be limited. Those desiring to make formal oral presentations should notify the contact