

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.

Robert Sargis,
Reports Clearance Officer.
 [FR Doc. 2016-23951 Filed 10-3-16; 8:45 am]
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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Submission for OMB Review; Comment Request

Title: Interstate Administrative Subpoena and Notice of Interstate Lien. *OMB No.:* 0970-0152.
Description: Section 452(a)(11) of the Social Security Act requires the Secretary of the Department of Health and Human Services to promulgate a

form for administrative subpoenas and imposition of liens used by State child support enforcement (Title IV-D) agencies. The Interstate Administrative Subpoena is used to collect information for the establishment, modification and enforcement of child support orders in interstate cases. Section 454(9)(E) of the Social Security Act requires each State to cooperate with any other State in using the federal form for issuance of administrative subpoenas and imposition of liens in interstate child support cases. Tribal IV-D agencies are not required to use this form but may choose to do so. OMB approval of these forms is expiring in December 2016 and the Administration for Children and Families is requesting an extension of this form.

Respondents: State, local or Tribal agencies administering a child support enforcement program under title IV-D of the Social Security Act.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
Administrative Subpoena	31,344	1	0.50	15,672
Notice of Lien	1,916,891	1	0.25	479,223

Estimated Total Annual Burden Hours: 494,895.

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, 330 C Street SW., Washington, DC 20201. Attention 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, Email: OIRA_SUBMISSION@OMB.EOP.GOV. Attn:

Desk Officer for the Administration for Children and Families.

Robert Sargis,
Reports Clearance Officer.
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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Proposed Information Collection Activity; Comment Request

Proposed Projects

Title: Income Withholding Order/ Notice for Support (IWO). *OMB No.:* 0970-0154.
Description: The Income Withholding Order/Notice for Support (IWO) is the standard form that must be used to order and notify employers and income providers to withhold child support payments from an obligor's income. It also indicates where employers and other income providers must remit the payments and other information needed to withhold correctly. Child support agencies, courts, private attorneys, custodial parties, and others must use the IWO form to initiate

an income withholding order for support and give notice of income withholding. State child support agencies are required to have automated data processing systems containing current order and case information. State child support agencies providing services to custodial and/or noncustodial parties enter the terms of a child support order established by a tribunal into the state's automated system, which automatically populates the order information into the IWO form.

Employers and income providers also use the form to respond to the order/ notice with termination or income status information. Employers and other income providers may choose to receive the IWO form from child support agencies on paper or electronically, and may respond on paper or electronically to notify the sender of termination of employment or change in the income status.

The information collection activities pertaining to the IWO form are authorized by 42 U.S.C. 666(a)(1), (a)(8) and 666(b)(6), which require the use of the Income Withholding for Support (IWO) form to order income withholding for all child support orders. *Respondents:* Courts, private attorneys, custodial parties or their

representatives, employers, and other parties that provide income to noncustodial parents.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
Income withholding order/notice (Courts, private attorneys, custodial parties or their representatives).	3,699,790	1.00	5 minutes	308,316
Income withholding orders/termination of employment/income status (Employers and other income providers).	1,207,484	9.694	2 minutes	390,178
Electronic income withholding orders/termination of employment/income status (Employers and other income providers).	9,596	136.38	3 seconds	1,090
Estimated Total Annual Burden Hours	699,585

In compliance with the requirements of the Paperwork Reduction Act of 1995 (Pub. L. 104–13, 44 U.S.C. Chap 35), 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 Planning, Research and Evaluation, 330 C Street SW., Washington DC 20201. Attn: ACF Reports Clearance Officer. Email address: infocollection@acf.hhs.gov. 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.

Robert Sargis,

Reports Clearance Officer.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2016–N–2655]

Center for Devices and Radiological Health Veteran Amputee Devices; Public Workshop; Request for Comments

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice of public workshop; request for comments.

SUMMARY: The Food and Drug Administration (FDA) is announcing the following public workshop entitled “Center for Devices and Radiological Health Veteran Amputee Devices.” The purpose of this workshop is to engage all stakeholders involved in the research, development, and marketing of prosthetic limb medical devices used by veteran amputees. A specific goal is to engage veteran amputees who use prosthetic limb medical devices and hear their views on these devices so that these perspectives may be considered in the total product life cycle of prosthetic limb devices.

DATES: The public workshop will be held on October 31, 2016, from 9 a.m. to 4 p.m. Submit either electronic or written comments on the public workshop by November 30, 2016.

ADDRESSES: The public workshop will be held at FDA’s White Oak Campus, 10903 New Hampshire Ave., Bldg. 31 Conference Center, the Great Room (Rm. 1503), Silver Spring, MD 20993. Entrance for the public meeting participants (non-FDA employees) is through Building 1 where routine security check procedures will be performed. For parking and security information, please refer to <http://www.fda.gov/AboutFDA/WorkingatFDA/BuildingsandFacilities/WhiteOakCampusInformation/ucm241740.htm>.

You may submit comments as follows:

Electronic Submissions

Submit electronic comments in the following way:

- *Federal eRulemaking Portal:* <http://www.regulations.gov>. Follow the instructions for submitting comments. Comments submitted electronically, including attachments, to <http://www.regulations.gov> will be posted to the docket unchanged. Because your comment will be made public, you are solely responsible for ensuring that your comment does not include any confidential information that you or a third party may not wish to be posted, such as medical information, your or anyone else’s Social Security number, or confidential business information, such as a manufacturing process. Please note that if you include your name, contact information, or other information that identifies you in the body of your comments, that information will be posted on <http://www.regulations.gov>.

- If you want to submit a comment with confidential information that you do not wish to be made available to the public, submit the comment as a written/paper submission and in the manner detailed (see “Written/Paper Submissions” and “Instructions”).

Written/Paper Submissions

Submit written/paper submissions as follows:

- *Mail/Hand delivery/Courier (for written/paper submissions):* Division of Dockets Management (HFA–305), Food and Drug Administration, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852.

- For written/paper comments submitted to the Division of Dockets Management, FDA will post your comment, as well as any attachments, except for information submitted, marked and identified, as confidential, if submitted as detailed in “Instructions.”