

DEPARTMENT OF HEALTH AND HUMAN SERVICES**Administration for Children and Families****Submission for OMB Review; Comment Request**

Title: Interstate Administrative Subpoena.

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 to be used in State child support enforcement programs to collect information for use in 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 in interstate child support cases. Tribal IV–D agencies are not required to use this form but may choose to do so.

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	53,488	1	0.50	26,744

Estimated Total Annual Burden Hours: 26,744.

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. 2013–19921 Filed 8–15–13; 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: Job Search Assistance (JSA) Strategies Evaluation.

OMB No.: New Collection.

Description: The Administration for Children and Families (ACF) is proposing an information collection activity as part of the Job Search Assistance (JSA) Strategies Evaluation. The proposed information collection consists of semi-structured interviews with key respondents involved with job search assistance programs in states and localities. Through this information collection and other study activities, ACF seeks to identify the types of job search assistance strategies that should be tested within the context of current TANF policies and requirements.

Respondents: State and local TANF administrators, program staff, and stakeholders such as researchers and policy experts.

ANNUAL BURDEN ESTIMATES

Instrument	Total number of respondents	Annual number of respondents	Number of responses per respondent	Average burden hours per response	Annual burden hours
Discussion Guide for Use with Researchers and Policy Experts	15	8	1	1	8
Discussion Guide for use with State and Local TANF Administrators	35	18	1	2.5	45
Discussion Guide for Use with Program Staff	50	25	1	2	50

Estimated Total Annual Burden Hours: 103.

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: OPRE Reports

Clearance Officer. All requests should be identified by the title of the information collection. Email address: OPREinfocollection@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,
ACF Reports Clearance Officer.
[FR Doc. 2013–19929 Filed 8–15–13; 8:45 am]
BILLING CODE 4184–09–P

**DEPARTMENT OF HEALTH AND
HUMAN SERVICES**

Food and Drug Administration
[Docket No. FDA–2009–N–0380]

**Agency Information Collection
Activities; Submission for Office of
Management and Budget Review;
Comment Request; Product
Jurisdiction: Assignment of Agency
Component for Review of Premarket
Applications**

AGENCY: Food and Drug Administration,
HHS.

ACTION: Notice.

SUMMARY: The Food and Drug
Administration (FDA) is announcing
that a proposed collection of
information has been submitted to the
Office of Management and Budget
(OMB) for review and clearance under
the Paperwork Reduction Act of 1995.

DATES: Fax written comments on the
collection of information by September
16, 2013.

ADDRESSES: To ensure that comments on
the information collection are received,
OMB recommends that written
comments be faxed to the Office of

Information and Regulatory Affairs,
OMB, Attn: FDA Desk Officer, FAX:
202–395–7285, or emailed to
oira_submission@omb.eop.gov. All
comments should be identified with the
OMB control number 0910–0523. Also
include the FDA docket number found
in brackets in the heading of this
document.

FOR FURTHER INFORMATION CONTACT:
Jonna Capezzuto, Office of Operations,
Food and Drug Administration, 1350
Piccard Dr., PI50–400B, Rockville, MD
20850, 301–796–3794,
Jonnalynn.capezzuto@fda.hhs.gov.

SUPPLEMENTARY INFORMATION: In
compliance with 44 U.S.C. 3507, FDA
has submitted the following proposed
collection of information to OMB for
review and clearance.

**Product Jurisdiction: Assignment of
Agency Component for Review of
Premarket Applications—(OMB Control
Number 0910–0523)—Extension**

This regulation relates to Agency
management and organization and has
two purposes. The first is to implement
section 503(g) of the Federal Food, Drug,
and Cosmetic Act (21 U.S.C. 353(g)), as
added by the Safe Medical Devices Act
of 1990 (Pub. L. 101–629), and amended
by the Medical Device User Fee and
Modernization Act of 2002 (Pub. L. 107–
250), by specifying how FDA will
determine the organizational component
within FDA assigned to have primary
jurisdiction for the premarket review
and regulation of products that are
comprised of any combination of: (1) A
drug and a device; (2) a device and a

biological product; (3) a biological
product and a drug; or (4) a drug, a
device, and a biological product. The
second purpose of this regulation is to
enhance the efficiency of Agency
management and operations by
providing procedures for classifying and
determining which Agency component
is designated to have primary
jurisdiction for any drug, device, or
biological product where such
jurisdiction is unclear or in dispute.

The regulation establishes a
procedure by which an applicant may
obtain an assignment or designation
determination. The regulation requires
that the request include the identity of
the applicant, a comprehensive
description of the product and its
proposed use, and the applicant's
recommendation as to which Agency
component should have primary
jurisdiction, with an accompanying
statement of reasons. The information
submitted would be used by FDA as the
basis for making the assignment or
designation decision. Most information
required by the regulation is already
required for premarket applications
affecting drugs, devices, biological
products, and combination products.
The respondents will be businesses or
other for-profit organizations.

In the **Federal Register** of May 2, 2013
(78 FR 25746), FDA published a 60-day
notice requesting public comment on
the proposed collection of information.
No comments were received.

FDA estimates the burden of this
collection of information as follows:

TABLE 1—ESTIMATED ANNUAL REPORTING BURDEN ¹

21 CFR Part	Number of respondents	Number of responses per respondent	Total annual responses	Average burden per response	Total hours
Part 3	59	1	59	24	1,416

¹ There are no capital costs or operating and maintenance costs associated with this collection of information.

These burden estimates are based on
the number of applications FDA
received over the past 2 fiscal years.

Dated: August 12, 2013.

Leslie Kux,
Assistant Commissioner for Policy.
[FR Doc. 2013–19916 Filed 8–15–13; 8:45 am]
BILLING CODE 4160–01–P

**DEPARTMENT OF HEALTH AND
HUMAN SERVICES**

National Institutes of Health

**National Cancer Institute; Notice of
Meeting**

Pursuant to section 10(a) of the
Federal Advisory Committee Act, as
amended (5 U.S.C. App.), notice is
hereby given of a meeting of the
President's Cancer Panel.

The meeting will be open to the
public, with attendance limited to space
available. Individuals who plan to
attend and need special assistance, such

as sign language interpretation or other
reasonable accommodations, should
notify the Contact Person listed below
in advance of the meeting.

Name of Committee: President's Cancer
Panel.

Date: October 11, 2013.

Time: 8:30 a.m. to 3:30 p.m.

Agenda: Cancer Communication for
Prevention: In the Digital Commons,
Opportunities Amongst the Challenges.

Place: National Institutes of Health, 9000
Rockville Pike, Building 31, C–Wing, 6th
Floor, Conference Room 10, Bethesda, MD
20892.

Contact Person: Abby B. Sandler, Ph.D.,
Executive Secretary, President's Cancer
Panel, Special Assistant to the Director, NCI