### ESTIMATED ANNUALIZED BURDEN HOURS

Respondents	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hours)	Total
Patients Controls	Case questionnaire Control questionnaire	161 483	1 1	25/60 25/60	67 201 268

#### LeRoy Richardson,

Chief, Information Collection Review Office, Office of Scientific Integrity, Office of the Associate Director for Science, Office of the Director, Centers for Disease Control and Prevention.

[FR Doc. 2014-08014 Filed 4-9-14; 8:45 am]

BILLING CODE 4163-18-P

# **DEPARTMENT OF HEALTH AND HUMAN SERVICES**

#### Administration for Children and **Families**

## Submission for OMB Review; **Comment Request**

Title: ACF Program Instruction: Children's Justice Act.

OMB No.: 0970-0425.

Description: The Program Instruction, prepared in response to the enactment of the Children's Justice Act (CJA), Title II of Public Law 111-320, Child Abuse Prevention and Treatment Act Reauthorization of 2010, provides direction to the States and Territories to accomplish the purposes of assisting States in developing, establishing and operating programs designed to improve: (1) The assessment and investigation of suspected child abuse and neglect cases, including cases of suspected child sexual abuse and exploitation, in a manner that limits additional trauma to the child and the child's family; (2) the assessment and investigation of cases of suspected child abuse-related fatalities and suspected child neglect-related fatalities; (3) the investigation and prosecution of cases of child abuse and neglect, including child

sexual abuse and exploitation; and (4) the assessment and investigation of cases involving children with disabilities or serious health-related problems who are suspected victims of child abuse or neglect. This Program Instruction contains information collection requirements that are found in Public Law 111–320 at Sections 107(b) and 107(d), and pursuant to receiving a grant award. The information being collected is required by statute to be submitted pursuant to receiving a grant award. The information submitted will be used by the agency to ensure compliance with the statute; to monitor, evaluate and measure grantee achievements in addressing the investigation and prosecution of child abuse and neglect; and to report to Congress.

Respondents: State Governments.

### **ANNUAL BURDEN ESTIMATES**

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
Application and Annual Report	52	1	60	3,120

Estimated Total Annual Burden Hours: 3,120.

### **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. 2014-08065 Filed 4-9-14; 8:45 am] BILLING CODE 4184-01-P

**Food and Drug Administration** 

**HUMAN SERVICES** 

DEPARTMENT OF HEALTH AND

[Docket No. FDA-2010-N-0062]

**Agency Information Collection Activities; Proposed Collection; Comment Request; Exception From** General Requirements for Informed Consent

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

**ACTION:** Notice.

**SUMMARY:** The Food and Drug Administration (FDA) is announcing an opportunity for public comment on the proposed collection of certain information by the Agency. Under the Paperwork Reduction Act of 1995 (the PRA), Federal Agencies are required to publish notice in the Federal Register concerning each proposed collection of