telephone 571–388–6570; or email *gsa.privacyact@gsa.gov*.

Pranjali Desai,

Director, Office of Information Management, General Services Administration.

[FR Doc. 2016–28084 Filed 11–21–16; 8:45 am]

BILLING CODE 6820–34–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[60Day–17–17CQ]

Proposed Data Collection Submitted for Public Comment and Recommendations—Zika Virus Associated Neurologic Illness Case Control Study; Correction

AGENCY: Centers for Disease Control and Prevention (CDC), Department of Health and Human Services (HHS).

ACTION: Notice; correction.

SUMMARY: The Centers for Disease Control and Prevention (CDC) published a document in the **Federal Register** of November 17, 2016, concerning request for comments on *Proposed Data Collection Submitted for Public Comment and Recommendations—Zika Virus Associated Neurologic Illness Case Control Study.* The document provided the incorrect agency identification number (60Day–17–17ZQ).

FOR FURTHER INFORMATION CONTACT: Leroy Richardson, 1600 Clifton Road, MS D–74, Atlanta, GA 30333; telephone (404) 639–4965; email: *omb@cdc.gov.*

Correction

In the **Federal Register** of November 17, 2016, in FR Doc. 2016–27692, on page 81143, in the first column (first heading), correct the agency identification number to read:

[60Day–17–17CQ; Docket No. CDC–2016–0107]

Dated: November 17, 2016.

Leroy A. 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. 2016–28072 Filed 11–21–16; 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: State Plan Child Support Collection and Establishment of Paternity Title IV–D, OCSE–100.

OMB No.: 0970–0017.

Description: The Office of Child Support Enforcement has approved a IV–D state plan for each state. Federal regulations require states to amend their state plans only when necessary to reflect new or revised federal statutes or regulations or material change in any state laws, regulations, policies, or IV–D agency procedures. The requirement for submission of a state plan and plan amendments for the Child Support Enforcement program is found in sections 452, 454, and 466 of the Social Security Act.

Respondents: State IV–D Agencies.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
State Plan (OCSE–100)	54	5	.5	135
State Plan Transmittal (OCSE–21–U4)	54	5	.25	67.5

Estimated Total Annual Burden Hours: 202.5.

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.

[FR Doc. 2016–28107 Filed 11–21–16; 8:45 am]

BILLING CODE 4184–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2015–E–4875]

Determination of Regulatory Review Period for Purposes of Patent Extension; ANAVIP

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) has determined the regulatory review period for ANAVIP and is publishing this notice of that determination as required by law. FDA has made the determination because of the submission of an application to the Director of the U.S. Patent and Trademark Office (USPTO), Department of Commerce, for the extension of a patent which claims that human biological product.