

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. 2017-01741 Filed 1-25-17; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[60Day-17-0006; Docket No. CDC-2017-
0004]

Proposed Data Collection Submitted for Public Comment and Recommendations

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

ACTION: Notice with comment period.

SUMMARY: The Centers for Disease
Control and Prevention (CDC), as part of
its continuing efforts to reduce public
burden and maximize the utility of
government information, invites the
general public and other Federal
agencies to take this opportunity to
comment on proposed and/or
continuing information collections, as
required by the Paperwork Reduction
Act of 1995. This notice invites
comment on an extension request for
the information collection titled
“Statements in Support of Application
of Waiver of Inadmissibility.” Approved
under Office of Management and Budget
(OMB) Control Number 0920-0006, this
information collection allows CDC to
review Class A medical waiver
applications for prospective immigrants
to the United States. CDC assists DHS/
USCIS in determining whether or not a
prospective immigrant with a Class A
mental health designation may be
admitted into the United States.

DATES: Written comments must be
received on or before March 27, 2017.

ADDRESSES: You may submit comments,
identified by Docket No. CDC-2017-
0004 by any of the following methods:

- *Federal eRulemaking Portal:*
Regulations.gov. Follow the instructions
for submitting comments.

- *Mail:* Leroy A. Richardson,
Information Collection Review Office,
Centers for Disease Control and

Prevention, 1600 Clifton Road NE., MS-
D74, Atlanta, Georgia 30329.

Instructions: All submissions received
must include the agency name and
Docket Number. All relevant comments
received will be posted without change
to *Regulations.gov*, including any
personal information provided. For
access to the docket to read background
documents or comments received, go to
Regulations.gov.

Please note: All public comment
should be submitted through the
Federal eRulemaking portal
(*Regulations.gov*) or by U.S. mail to the
address listed above.

FOR FURTHER INFORMATION CONTACT: To
request more information on the
proposed project or to obtain a copy of
the information collection plan and
instruments, contact the Information
Collection Review Office, Centers for
Disease Control and Prevention, 1600
Clifton Road NE., MS-D74, Atlanta,
Georgia 30329; phone: 404-639-7570;
Email: omb@cdc.gov.

SUPPLEMENTARY INFORMATION:

Under the Paperwork Reduction Act
of 1995 (PRA) (44 U.S.C. 3501-3520),
Federal agencies must obtain approval
from the Office of Management and
Budget (OMB) for each collection of
information they conduct or sponsor. In
addition, the PRA also requires Federal
agencies to provide a 60-day notice in
the **Federal Register** concerning each
proposed collection of information,
including each new proposed
collection, each proposed extension of
existing collection of information, and
each reinstatement of previously
approved information collection before
submitting the collection to OMB for
approval. To comply with this
requirement, we are publishing this
notice of a proposed data collection as
described below.

Comments are invited 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)
ways to enhance the quality, utility, and
clarity of the information to be
collected; (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; and (e) estimates of capital
or start-up costs and costs of operation,
maintenance, and purchase of services
to provide information. Burden means
the total time, effort, or financial
resources expended by persons to
generate, maintain, retain, disclose or
provide information to or for a Federal
agency. This includes the time needed
to review instructions; to develop,
acquire, install and utilize technology
and systems for the purpose of
collecting, validating and verifying
information, processing and
maintaining information, and disclosing
and providing information; to train
personnel and to be able to respond to
a collection of information, to search
data sources, to complete and review
the collection of information; and to
transmit or otherwise disclose the
information.

Proposed Project

Statements in Support of Application
of Waiver of Inadmissibility (OMB
Control No. 0920-0006; Expires 8/31/
2017)—Extension—National Center for
Emerging and Zoonotic Infectious
Diseases (NCEZID), Centers for Disease
Control and Prevention (CDC).

Background and Brief Description

Section 212(a)(1) of the Immigration
and Nationality Act states that aliens
with specific health related conditions
are ineligible for admission into the
United States. The Attorney General
may waive application of this
inadmissibility on health-related
grounds if an application for waiver is
filed and approved by the consular
office considering the application for
visa. CDC uses this application
primarily to collect information to
establish and maintain records of waiver
applicants in order to notify the U.S.
Citizenship and Immigration Services
when terms, conditions and controls
imposed by waiver are not met.

CDC is requesting approval from OMB
to collect this data for another three
years. Based on a review of the number
of waivers processed by CDC over the
last three years, CDC does not request a
change in the amount of burden.

Respondents must mail these
documents to CDC, and this entails an
additional cost. CDC estimates that
respondents will spend approximately
\$15 per year on postal fees, for a total
of \$3,000 annually.

ESTIMATE OF ANNUALIZED BURDEN HOURS

Type of respondent	Form	Number of respondents	Number of responses per respondent	Average burden per response (in hours)	Total burden hours
Physician	CDC 4.422-1	200	1	10/60	33
Physician	CDC 4.422-1a or letter	200	1	20/60	67
Total	100

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Chief, Information Collection Review Office,
Office of Scientific Integrity, Office of the
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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Office of the Secretary

[Document Identifier: 0990-0419-60D]

Agency Information Collection Activities; Proposed Collection; Public Comment Request

AGENCY: Office of the Secretary, HHS.

ACTION: 60-Day notice.

SUMMARY: In compliance with section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, the Office of the Secretary (OS), Department of Health and Human Services, announces plans to submit an Information Collection Request (ICR), described below, to the Office of Management and Budget (OMB). The ICR is for extending the use of the approved information collection assigned OMB control number 0990-

0419, which expires on June 30, 2017.

Prior to submitting the ICR to OMB, OS seeks comments from the public regarding the burden estimate, below, or any other aspect of the ICR.

DATES: Comments on the ICR must be received on or before March 27, 2017.

ADDRESSES: Submit your comments to *Information.CollectionClearance@hhs.gov* or by calling (202) 690-5683.

SUPPLEMENTARY INFORMATION: When submitting comments or requesting information, please include the document identifier 0990-0419-60D for reference.

Information Collection Request Title: Acquisition Regulation Clause Patent Rights and Rights in Data.

OMB No.: 0990-0419.

Abstract: The Department of Health and Human Services; Office of the Assistant Secretary for Financial Resources and Office of Grants and Acquisition Policy and Accountability, Division of Acquisition, is requesting an approval by OMB for an extension of a previously approved information collection request, 0990-0419—Acquisition Regulation Clause Patent Rights and Rights in Data. HHS found that systematically, over a period of several years, when Determination of Exceptional Circumstances (DEC) were

executed, additional legal protection for the patent and data rights of third parties beyond those covered by FAR 27.306 were necessary. A DEC is executed consistent with the policy and objectives of the Bayh-Dole Act, 35 U.S.C. 200, *et seq.*, to ensure that subject inventions made under contracts and subcontracts (at all tiers) are used in a manner to promote free competition and enterprise without unduly encumbering future research and discovery; to encourage maximum participation of small business firms in federally supported research and development efforts; to promote collaboration between commercial concerns and nonprofit organizations including universities; to ensure that the Government obtains sufficient rights in federally supported inventions to meet its needs; to protect the public against nonuse or unreasonable use of inventions; and in the case of fulfilling the mission of the U.S. Department of Health and Human Services, to ultimately to benefit the public health.

Likely Respondents: Administrative, technical, legal and management personnel.

The total annual burden hours estimated for this ICR are summarized in the table below.

TOTAL ESTIMATED ANNUALIZED BURDEN—HOURS

Information collection	Type of respondent and hours for each	Number of respondents	Number of responses per respondent	Average burden per response (hours)	Total burden hours
(a)	Technical (4)	63	1	8	504
	Legal (2)				
	Management (2)				
(b)	Technical (8)	63	1	12	756
	Legal (2)				
	Management (2)				
(c)	Technical (8)	63	3	12 (36)	2268
	Legal (3)				
	Management (1)				
(d)	Technical (8)	63	3	14 (42)	2646
	Legal (4)				
	Management (2)				
(e)	Technical (6)	63	1	10	630
	Legal (2)				
	Management (2)				