

Federal Communications Commission.

Marlene H. Dortch,
Secretary.

[FR Doc. 2017-17131 Filed 8-14-17; 8:45 am]

BILLING CODE 6712-01-P

FEDERAL COMMUNICATIONS COMMISSION

[OMB 3060-0208]

Information Collection Being Reviewed by the Federal Communications Commission

AGENCY: Federal Communications Commission.

ACTION: Notice and request for comments.

SUMMARY: As part of its continuing effort to reduce paperwork burdens, and as required by the Paperwork Reduction Act of 1995 (PRA), the Federal Communications Commission (FCC or Commission) invites the general public and other Federal agencies to take this opportunity to comment on the following information collections. Comments are requested concerning: Whether the proposed collection of information is necessary for the proper performance of the functions of the Commission, including whether the information shall have practical utility; the accuracy of the Commission's burden estimate; ways to enhance the quality, utility, and clarity of the information collected; ways to minimize the burden of the collection of information on the respondents, including the use of automated collection techniques or other forms of information technology; and ways to further reduce the information collection burden on small business concerns with fewer than 25 employees.

The FCC may not conduct or sponsor a collection of information unless it displays a currently valid Office of Management and Budget (OMB) control number. No person shall be subject to any penalty for failing to comply with a collection of information subject to the PRA that does not display a valid OMB control number.

DATES: Written comments should be submitted on or before October 16, 2017. If you anticipate that you will be submitting comments, but find it difficult to do so within the period of time allowed by this notice, you should advise the contacts below as soon as possible.

ADDRESSES: Direct all PRA comments to Cathy Williams, FCC, via email: PRA@fcc.gov and to Cathy.Williams@fcc.gov.

FOR FURTHER INFORMATION CONTACT: For additional information about the information collection, contact Cathy Williams at (202) 418-2918.

SUPPLEMENTARY INFORMATION: As part of its continuing effort to reduce paperwork burdens, and as required by the PRA, 44 U.S.C. 3501-3520, the FCC invites the general public and other Federal agencies to take this opportunity to comment on the following information collections. Comments are requested concerning: Whether the proposed collection of information is necessary for the proper performance of the functions of the Commission, including whether the information shall have practical utility; the accuracy of the Commission's burden estimate; ways to enhance the quality, utility, and clarity of the information collected; ways to minimize the burden of the collection of information on the respondents, including the use of automated collection techniques or other forms of information technology; and ways to further reduce the information collection burden on small business concerns with fewer than 25 employees.

OMB Control Number: 3060-0208.

Title: Section 73.1870, Chief

Operators.

Form Number: Not applicable.

Type of Review: Extension of a currently approved collection.

Respondents: Business and other for-profit; Not-for-profit institutions.

Number of Respondents and

Responses: 18,498 respondents; 36,996 responses.

Estimated Time per Response: 0.166-26 hours.

Frequency of Response:

Recordkeeping requirement; Third party disclosure requirement.

Total Annual Burden: 484,019 hours.

Total Annual Cost: None.

Obligation to Respond: Required to obtain or retain benefits. The statutory authority for this collection of information is contained in Sections 154(i) of the Communications Act of 1934, as amended.

Nature and Extent of Confidentiality: There is no need for confidentiality with this collection of information.

Privacy Impact Assessment(s): No impact(s).

Needs and Uses: The information collection requirements contained in 47 CFR 73.1870 require that the licensee of an AM, FM, or TV broadcast station designate a chief operator of the station. Section 73.1870(b)(3) requires that this designation must be in writing and posted with the station license. Section 73.1870(c)(3) requires that the chief

operator, or personnel delegated and supervised by the chief operator, review the station records at least once each week to determine if required entries are being made correctly, and verify that the station has been operated in accordance with FCC rules and the station authorization. Upon completion of the review, the chief operator must date and sign the log, initiate corrective action which may be necessary and advise the station licensee of any condition which is repetitive. The posting of the designation of the chief operator is used by interested parties to readily identify the chief operator. The review of the station records is used by the chief operator, and FCC staff in investigations, to ensure that the station is operating in accordance with its station authorization and the FCC rules and regulations.

Federal Communications Commission.

Marlene H. Dortch,

Secretary, Office of the Secretary.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Agency for Healthcare Research and Quality

Supplemental Evidence and Data Request on Mobile Health Technology for Diabetes

AGENCY: Agency for Healthcare Research and Quality, HHS.

ACTION: Request for supplemental evidence and data submissions.

SUMMARY: The Agency for Healthcare Research and Quality is seeking scientific information submissions from the public. Scientific information is being solicited to inform our review of *Mobile Health Technology for Diabetes*, which is currently being conducted by the AHRQ's Evidence-based Practice Centers (EPC) Program. Access to published and unpublished pertinent scientific information will improve the quality of this review.

DATES: *Submission Deadline* on or before September 14, 2017.

ADDRESSES:

Email submissions: SEADS@epc-src.org.

Print submissions:

Mailing Address: Portland VA Research Foundation, Scientific Resource Center, ATTN: Scientific Information Packet Coordinator, P.O. Box 69539, Portland, OR 97239

Shipping Address (FedEx, UPS, etc.): Portland VA Research Foundation,

Scientific Resource Center, ATTN:
Scientific Information Packet
Coordinator, 3710 SW U.S. Veterans
Hospital Road, Mail Code: R&D 71,
Portland, OR 97239

FOR FURTHER INFORMATION CONTACT:
Ryan McKenna, Telephone: 503–220–
8262 ext. 51723 or Email: [SEADS@epc-
src.org](mailto:SEADS@epc-src.org).

SUPPLEMENTARY INFORMATION: The
Agency for Healthcare Research and
Quality (AHRQ) has commissioned the
Evidence-based Practice Centers (EPC)
Program to complete a review of the
evidence for Mobile Health Technology
for Diabetes. AHRQ is conducting this
systematic review pursuant to Section
902(a) of the Public Health Service Act,
42 U.S.C. 299a(a).

The EPC Program is dedicated to
identifying as many studies as possible
that are relevant to the questions for
each of its reviews. In order to do so,
we are supplementing the usual manual
and electronic database searches of the
literature by requesting information
from the public (e.g., details of studies
conducted). We are looking for studies
that report on *Mobile Health Technology
for Diabetes*, including those that
describe adverse events. The entire
research protocol, including the key
questions, is also available online at:
[http://www.effectivehealthcare.ahrq.gov/
index.cfm/search-for-guides-reviews-
and-reports/?pageaction=display
roduct&productid=2484](http://www.effectivehealthcare.ahrq.gov/index.cfm/search-for-guides-reviews-and-reports/?pageaction=displayproduct&productid=2484).

This is to notify the public that the
EPC Program would find the following
information on *Mobile Health
Technology for Diabetes* helpful:

- A list of completed studies that
your organization has sponsored for this
indication. In the list, please *indicate
whether results are available on
ClinicalTrials.gov along with the
ClinicalTrials.gov trial number.*

- *For completed studies that do not
have results on ClinicalTrials.gov,*
please provide a summary, including
the following elements: Study number,
study period, design, methodology,
indication and diagnosis, proper use
instructions, inclusion and exclusion
criteria, primary and secondary

outcomes, baseline characteristics,
number of patients screened/eligible/
enrolled/lost to follow-up/withdrawn/
analyzed, effectiveness/efficacy, and
safety results.

- *A list of ongoing studies that your
organization has sponsored for this
indication.* In the list, please provide the
ClinicalTrials.gov trial number or, if the
trial is not registered, the protocol for
the study including a study number, the
study period, design, methodology,
indication and diagnosis, proper use
instructions, inclusion and exclusion
criteria, and primary and secondary
outcomes.

- Description of whether the above
studies constitute ALL Phase II and
above clinical trials sponsored by your
organization for this indication and an
index outlining the relevant information
in each submitted file.

Your contribution will be very
beneficial to the EPC Program. Materials
submitted must be publicly available or
able to be made public. Materials that
are considered confidential; marketing
materials; study types not included in
the review; or information on
indications not included in the review
cannot be used by the EPC Program.
This is a voluntary request for
information, and all costs for complying
with this request must be borne by the
submitter.

The draft of this review will be posted
on AHRQ’s EPC Program Web site and
available for public comment for a
period of 4 weeks. If you would like to
be notified when the draft is posted,
please sign up for the email list at:
[https://www.effectivehealthcare.ahrq.
gov/index.cfm/join-the-email-list1/](https://www.effectivehealthcare.ahrq.gov/index.cfm/join-the-email-list1/).

*The systematic review will answer the
following questions. This information is
provided as background. AHRQ is not
requesting that the public provide
answers to these questions.*

The Guiding Questions

I. Which specific mobile health
technology (mHealth) technologies for
diabetes self-management have been
researched?

II. What are the characteristics (e.g.,
interoperability, functions,

acceptability/usability, connection to
electronic health records) of these
specific mHealth technologies?

III. What patient outcomes are
associated with the use of these specific
mHealth technologies?

IV. What are the harms and costs
associated with these specific mHealth
technologies?

Sharon B. Arnold,

Deputy Director.

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**DEPARTMENT OF HEALTH AND
HUMAN SERVICES**

**Administration for Children and
Families**

**Submission for OMB Review;
Comment Request**

Title: Head Start Program Information
Report.

OMB No.: 0970–0427.

Description: The Office of Head Start
within the Administration for Children
and Families, United States Department
of Health and Human Services, is
proposing to renew authority to collect
information using the Head Start
Program Information Report (PIR),
monthly enrollments, contacts,
locations, and reportable conditions. All
information is collected through a single
system, the Head Start Enterprise
System (HSES). The PIR provides
information about Head Start and Early
Head Start services received by the
children and families enrolled in Head
Start programs. The information
collected in the PIR is used to inform
the public about these programs, to
make periodic reports to Congress about
the status of children in Head Start
programs as required by the Head Start
Act, and to assist the administration and
training/technical assistance of Head
Start programs.

Respondents: Head Start and Early
Head Start program grant recipients.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
Head Start Program Information Report (PIR)	3,267	1	4	13,068
Grantee Monthly Enrollment Reporting	2,049	12	0.05	1,229
Contacts, Locations & Reportable Conditions	3,267	1	0.25	817

*Estimated Total Annual Burden
Hours:* 15,114.

Additional Information: Copies of the
proposed collection may be obtained by

writing to the Administration for
Children and Families, Office of