

**DEPARTMENT OF HEALTH AND HUMAN SERVICES****Health Resources and Services Administration****Agency Information Collection Activities: Proposed Collection: Comment Request; Information Collection Request Title: Survey of Eligible Users of the National Practitioner Data Bank, OMB No. 0915–0366—Reinstatement With Change**

**AGENCY:** Health Resources and Services Administration (HRSA), Department of Health and Human Services.

**ACTION:** Notice.

**SUMMARY:** In compliance with the requirement for opportunity for public comment on proposed data collection projects of the Paperwork Reduction Act of 1995, HRSA announces plans to submit an Information Collection Request (ICR), described below, to the Office of Management and Budget (OMB). Prior to submitting the ICR to OMB, HRSA seeks comments from the public regarding the burden estimate, below, or any other aspect of the ICR.

**DATES:** Comments on this ICR should be received no later than March 12, 2021.

**ADDRESSES:** Written comments and recommendations for the proposed information collection should be sent within 30 days of publication of this notice to [www.reginfo.gov/public/do/PRAMain](http://www.reginfo.gov/public/do/PRAMain). Find this particular information collection by selecting “Currently under Review—Open for Public Comments” or by using the search function.

**FOR FURTHER INFORMATION CONTACT:** To request more information on the proposed project or to obtain a copy of the data collection plans and draft instruments, email [paperwork@hrsa.gov](mailto:paperwork@hrsa.gov) or call Lisa Wright-Solomon, the HRSA

Information Collection Clearance Officer at (301) 443–1984.

**SUPPLEMENTARY INFORMATION:** When submitting comments or requesting information, please include the information request collection title for reference.

*Information Collection Request Title:* Survey of Eligible Users of the National Practitioner Data Bank, OMB No. 0915–0366—Reinstatement with Change.

*Abstract:* HRSA plans to survey National Practitioner Data Bank (NPDB) users. The purpose of this survey is to assess the overall satisfaction of the eligible users of the NPDB. This survey will evaluate the effectiveness of the NPDB as a flagging system, source of information, and its use in decision making. Furthermore, this survey will collect information from organizations and individuals who query the NPDB to understand and improve their user experience. This survey is a reinstatement of the 2012 NPDB survey with some changes.

A 60-day Notice published in the **Federal Register** on October 16, 2020, vol. 85, No. 201; pp. 65833–34. There were no comments.

*Need and Proposed Use of the Information:* The survey will collect information regarding the participants’ experiences of querying and reporting to the NPDB, perceptions of health care practitioners with reports, impact of NPDB reports on organizations’ decision-making, and satisfaction with various NPDB products and services.

The survey will also be administered to health care practitioners that use the self-query service provided by the NPDB. The self-queriers will be asked about their experiences of querying, the impact of having reports in the NPDB on their careers and health care organizations’ perceptions, and their satisfaction with various NPDB products and services. Understanding self-

queriers’ satisfaction and their use of the information is an important component of the survey.

Proposed changes to this ICR include the following:

1. In the proposed entity survey, there are 37 modules and 258 questions. From the previous 2012 survey, there are 15 deleted questions and 13 new questions in addition to proposed changes to 12 survey questions.

2. In the proposed self-query survey, there are 22 modules and 88 questions. From the previous 2012 survey, there are five deleted questions and five new questions in addition to proposed changes to two survey questions.

*Likely Respondents:* Health care entities and health care practitioners who are eligible users of the NPDB will be asked to complete a web-based survey. Data gathered from the survey will be compared with previous survey results. This survey will provide HRSA with the information necessary for research purposes and for improving the usability and effectiveness of the NPDB.

*Burden Statement:* Burden in this context means the time expended by persons to generate, maintain, retain, disclose, or provide the information requested. 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. The total annual burden hours estimated for this ICR are summarized in the table below.

*Total Estimated Annualized Burden Hours:*

| Form name                             | Number of respondents | Number of responses per respondent | Total responses | Average burden per response (in hours) | Total burden hours |
|---------------------------------------|-----------------------|------------------------------------|-----------------|--|--------------------|
| NPDB Users Entities Respondents ..... | 15,000                | 1                                  | 15,000          | 0.25                                   | 3,750              |
| NPDB Self-Query Respondents .....     | 2,000                 | 1                                  | 2,000           | 0.10                                   | 200                |
| Total .....                           | 17,000                | .....                              | 17,000          | .....                                  | 3,950              |

HRSA specifically requests comments on (1) the necessity and utility of the proposed information collection for the proper performance of the agency’s functions, (2) the accuracy of the estimated burden, (3) ways to enhance the quality, utility, and clarity of the information to be collected, and (4) the

use of automated collection techniques or other forms of information

technology to minimize the information collection burden.

**Maria G. Button,**

*Director, Executive Secretariat.*

[FR Doc. 2021–02757 Filed 2–9–21; 8:45 am]

**BILLING CODE 4165–15–P**