invoke the exemption to be 194 in the future.

*Estimated Time per Response:* 15 minutes for those adding a tariff rule to use a combination of tariff rates and NRAs, and 1 hour for recordkeeping requirements. For those using NRAs exclusively, one hour to publish an NRA rules tariff.

*Total Annual Burden:* Of the 194 new NVOCCs estimated to file a rule or prominent notice in their respective tariffs, we estimate that 3% (6) will use NRAs exclusively. The burden is calculated as follows: 188 × .25 hours = 47 hours and 6 × 1 hour = 6 hours (3% using NRAs exclusively). Recording keeping requirements for the total number of NVOCCs that have invoked the exemption thus far is 2,349 × 1 hour = 2,349. Total annual burden is estimated to be 2,402 hours.

**William Cody,**

*Secretary.*

[FR Doc. 2022–06621 Filed 3–29–22; 8:45 am]

**BILLING CODE 6730–02–P**

## FEDERAL MEDIATION AND CONCILIATION SERVICE

### Notice of Stakeholder Survey for Qualitative Feedback on Agency Service Delivery

**AGENCY:** Federal Mediation and Conciliation Service (FMCS).

**ACTION:** 60-Day notice and request for comments.

**SUMMARY:** The Federal Mediation and Conciliation Service (FMCS), invites the general public and other Federal Agencies to take this opportunity to comment on the following information collection request, Stakeholder Survey for Qualitative Feedback on Agency Service Delivery. This information collection request was previously approved by the Office of Management Budget (OMB) and FMCS is requesting a revision of a currently approved collection. This collection was developed as part of a Federal Government-wide effort to streamline the process for seeking feedback from the public on service delivery.

**DATES:** Comments must be submitted on or before May 31, 2022.

**ADDRESSES:** You may submit comments, identified by Stakeholder Survey for Qualitative Feedback on Agency Service Delivery, through one of the following methods:

- *Email: register@fmcs.gov;*
- *Mail:* Office of the General Counsel, One Independence Square, 250 E St. SW, Washington, DC 20427. Please note that at this time, mail is sometimes delayed. Therefore, we encourage emailed comments.

**FOR FURTHER INFORMATION CONTACT:** David Thaler, 980–812–0051, *dthaler@fmcs.gov.*

**SUPPLEMENTARY INFORMATION:** Copies of the agency questions are available here.

#### I. Information Collection Request

*Agency:* Federal Mediation and Conciliation Service.

*Form Number:* OMB No. 3076–0017.

*Type of Request:* Revision of a currently approved collection.

*Affected Entities:* Federal government and private sector.

*Frequency:* This survey is completed once.

*Abstract:* This information collection provides a means to garner qualitative customer and stakeholder feedback in an efficient, timely manner, in accordance with the Administration's commitment to improving service delivery. This feedback will provide insights into customer or stakeholder perceptions, experiences, and expectations. The surveys will provide notice of issues with service, or focus attention on areas where communication, training or changes in operations might improve delivery of products or services. The surveys are not statistical surveys that yield quantitative results that can be generalized to the population of study. These collections will allow for ongoing, collaborative, and actionable communications between the Agency and its customers and stakeholders. It will also allow feedback to contribute directly to improve program management. Responses will be assessed to plan and inform efforts to improve or maintain the quality of service offered to the public. Collecting this information is critical for ensuring quality service offered to the public.

*Burden:* FMCS receives approximately 7,100 responses per year and the time required is approximately one minute.

#### II. Request for Comments

FMCS solicits comments to:

i. Evaluate whether the proposed collections of information are necessary for the proper performance of the functions of the agency, including whether the information will have practical utility.

ii. Enhance the accuracy of the agency's estimates of the burden of the proposed collection of information.

iii. Enhance the quality, utility, and clarity of the information to be collected.

iv. Minimize the burden of the collections of information on those who are to respond, including the use of appropriate automated, electronic collection technologies or other forms of information technology.

#### III. The Official Record

The official records are electronic records.

#### List of Subjects

Labor-Management Relations.

Dated: March 25, 2022.

**Anna Davis,**

*Acting General Counsel.*

[FR Doc. 2022–06658 Filed 3–29–22; 8:45 am]

**BILLING CODE 6732–01–P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Administration for Children and Families

### Submission for OMB Review; Administration for Native Americans Project Outcome Assessment Survey (OMB #0970–0379)

**AGENCY:** Administration for Native Americans, Administration for Children and Families, HHS.

**ACTION:** Request for public comment.

**SUMMARY:** The Administration for Children and Families (ACF) is requesting a 3-year extension of the form Administration for Native Americans (ANA) Project Outcome Assessment Survey (OMB #0970–0379, expiration 6/30/2022). There are minor changes and updates requested to the form.

**DATES:** *Comments due within 30 days of publication.* OMB must make a decision about 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.

**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.* Find this particular information collection by selecting ''Currently under 30-day Review—Open for Public Comments'' or by using the search function. You can also obtain copies of the proposed collection of information by emailing *infocollection@acf.hhs.gov.* Identify all emailed

requests by the title of the information collection.

**SUPPLEMENTARY INFORMATION:**

*Description:* The information collected by the Project Impact Assessment Survey is needed for two main reasons: (1) To collect crucial information required to report on the ANA established Government Performance and Results Act measures, and (2) to properly abide by ANA's congressionally mandated statute (42 U.S.C. 2992 *et seq.*) found within the Native American Programs Act of 1974, as amended, which states that ANA will evaluate projects assisted through ANA grant dollars ''including evaluations that describe and measure the impact of such projects, their effectiveness in achieving stated goals, their impact on related programs, and their structure and mechanisms for delivery of services.'' The information collected with this survey will fulfill ANA's statutory requirement and will also serve as an important planning and performance tool for ANA.

Updates to this information collection address the Indian Community Economic Enhancement Act of 2020 (Pub. L. 116–261). It also addresses the flexibilities and assistance offered under COVID–19 recovery assistance.

*Respondents:* Tribal Governments, Native American nonprofit organizations, and Tribal Colleges and Universities.

BURDEN ESTIMATES

| Instrument | Number of respondents | Number of responses per respondent | Average burden hours per response | Total burden hours |
|---|---|---|---|---|
| ANA Project Outcome Assessment Survey | 85 | 1 | 6 | 510 |

*Estimated Total Burden Hours:* 510.

*Authority:* 42 U.S.C. 2992.

**Mary B. Jones,**

*ACF/OPRE Certifying Officer.*

[FR Doc. 2022–06652 Filed 3–29–22; 8:45 am]

**BILLING CODE 4184–34–P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Food and Drug Administration

**[Docket No. FDA–2012–N–0280]**

### Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Financial Disclosure by Clinical Investigators

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

**ACTION:** Notice.

**SUMMARY:** The Food and Drug Administration (FDA, Agency, or we) is announcing that a proposed collection of information has been submitted to the Office of Management and Budget (OMB) for review and clearance under the Paperwork Reduction Act of 1995.

**DATES:** Submit written comments (including recommendations) on the collection of information by April 29, 2022.

**ADDRESSES:** To ensure that comments on the information collection are received, OMB recommends that written comments be submitted to *https://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. The OMB control number for this information collection is 0910–0396. Also include the FDA docket number found in brackets in the heading of this document.

**FOR FURTHER INFORMATION CONTACT:** Amber Sanford, Office of Operations, Food and Drug Administration, Three White Flint North, 10A–12M, 11601 Landsdown St., North Bethesda, MD 20852, 301–796–8867, *PRAStaff@fda.hhs.gov.*

**SUPPLEMENTARY INFORMATION:** In compliance with 44 U.S.C. 3507, FDA has submitted the following proposed collection of information to OMB for review and clearance.

### Financial Disclosure by Clinical Investigators

*OMB Control Number 0910–0396—Extension*

Respondents to this collection are sponsors of marketing applications that contain clinical data from studies covered by the regulations. These sponsors represent pharmaceutical, biologic, and medical device firms. Respondents are also clinical investigators who provide financial information to the sponsors of marketing applications.

In the **Federal Register** of December 2, 2021 (86 FR 68500), FDA published a 60-day notice requesting public comment on the proposed collection of information. Although one comment was received, it was not responsive to the four collection of information topics solicited.

Table 1 shows information that is the basis of the estimated number of respondents in tables 2 through 4.

TABLE 1—ESTIMATED NUMBER OF APPLICATIONS, CLINICAL TRIALS, AND INVESTIGATORS SUBJECT TO THE REGULATION BY TYPE OF APPLICATION [1]

| Application type | Total number of applications | Number of applications affected | Number of trials | Number of investigators |
|---|---|---|---|---|
| Drugs: | | | | |
| New drug application (NDA), new molecular entity (NME) | 55 | 55 | 3 to 10 | 3 to 100. |
| NDA non-NME | 78 | 37 | 3 to 10 | 3 to 100. |
| NDA efficacy supplement | 196 | 119 | 1 to 3 | 10 to 30. |
| Abbreviated new drug application (ANDA) | 821 | 1 | 1.1 | 2. |
| ANDA supplement | 10,894 | 1 | 1 | 2. |
| CBER Biologics: | | | | |
| Biologics license application (BLA) | 10 | 10 | 3 to 10 | 3 to 100. |
| BLA efficacy supplement | 30 | 30 | 1 to 3 | 10 to 30. |
| CDER Biologics: | | | | |