subscale (C08) for one program that aims to improve mothers' belief in their parenting skills.

- Added items about confidence in future financial situations (C09a-C09b) for one program that aims to increase youths' belief in their ability to become self-sufficient.
- Added a question to ask if participants are covered by health insurance (C10b).
- Added the Center for Epidemiologic Studies Depression Scale Revised (CESD–R) (C22a) for use by a program that serves mothers with symptoms of depression. For this program, this more in-depth depression scale will be used instead of the shorter K–6 Distress Scale (C22).
- Added questions about whether the respondent is currently under court-ordered supervision (C38), whether any convictions were for felonies (C42), and if any of the reported incarcerations were for violating the terms of court-ordered supervision (C44) for a program that serves adults with criminal justice system involvement.
- Removed some items asking program group study participants about their satisfaction with the program.

We currently have OMB approval to provide a $40 gift card for follow-up survey completion. Based on new evidence, we propose to increase the first follow-up survey token of appreciation from $40 to $50 and to add a $5 prepaid gift card for the first follow-up survey sent with the advance letter. We would assess the effectiveness of the prepaid gift card using an experiment. If the experiment shows that prepaid tokens of appreciation are effective at increasing response rates or decreasing the treatment-control response rate differential, we will propose using it for remaining sample members for the first follow-up survey.

Second Follow-Up Survey (Instrument 4)

We propose changes to the second follow-up survey to match the changes proposed for the first follow-up survey. We propose introducing a $5 prepaid token of appreciation if the experiment for the first follow-up survey demonstrates it is effective. We do not propose changes to the previously approved $50 postpaid gift card for the second follow-up survey.

Respondents: Individuals enrolled in the NextGen Project.

Annual Burden Estimates

The annual burden estimates for the instruments we are requesting to revise are presented below. All currently approved materials under OMB # 0970–0545 and the associated burden can be found at: *https://www.reginfo.gov/public/do/PRAViewICR?ref_nbr=202012-0970-003.*

Instrument	Total number of respondents	Annual number of respondents	Number of responses per respondent	Average burden hours per response	Annual burden hours
PHASE 2					
First follow-up survey—participants	8,000	2,667	1	0.83	2,214
Second follow-up survey—participants	8,000	2,667	1	0.83	2,214

Total Annual Burden Hours, Phase 2: 4,428.

Authority: Section 413 of the Social Security Act, as amended by the FY 2017 Consolidated Appropriations Act, 2017 (Pub. L. 115–31).

Mary B. Jones,

ACF/OPRE Certifying Officer.

[FR Doc. 2022–06403 Filed 3–25–22; 8:45 am]

BILLING CODE 4184–09–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Submission for OMB Review; Measuring Human Trafficking Prevalence in Construction: A Field Test of Multiple Estimation Methods (New Collection)

AGENCY: Office of Planning, Research, and Evaluation, Administration for Children and Families, HHS.

ACTION: Request for public comment.

SUMMARY: The Administration for Children and Families (ACF) is proposing a new data collection activity for Measuring Human Trafficking Prevalence in Construction: A Field Test of Multiple Estimation Methods. This study will examine the labor trafficking and other labor exploitation experiences among individuals who work in construction. The goal of this study is to advance knowledge of promising methods for estimating human trafficking prevalence by field-testing two methods of prevalence estimation within the construction industry in Houston, Texas.

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 *OPREinfocollection@acf.hhs.gov.* Identify all requests by the title of the information collection.

SUPPLEMENTARY INFORMATION:

Description: The purpose of the proposed data collection activity is to estimate the prevalence of labor trafficking among construction workers in one location using two different sampling and estimation strategies. The proposed information collection activity is a one-time survey with up to 4,200 adults who worked in the construction industry in the selected geographic location in the 24 months prior to data collection. The construction worker survey will be offered in English and Spanish to workers identified through the following two sampling strategies: (1) Probability sample (*i.e.,* time location sample), and (2) a network sample. The survey instrument used for individuals recruited through the two different sampling strategies will be primarily the same and includes questions focused on the individuals' experiences with labor exploitation and trafficking; employment histories, including work after a natural disaster; social networks; and demographic data.

Respondents: English- and Spanish-speaking individuals who have worked in construction in Houston, Texas, in the 2 years prior to data collection will be invited to complete a survey.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents (total over request period)	Number of responses per respondent (total over request period)	Average burden per response (in hours)	Total burden (in hours)	Total burden (in hours)	Annual burden (in hours)
Construction Worker Survey	4,200	1	1	0.5	2,100	1,050

Estimated Total Annual Burden Hours: 1,050.

Authority: Section 105(d)(2) of the Trafficking Victims Protection Act of 2000 (Pub. L. 106–386) [22 U.S.C. 7103].

Mary B. Jones,

ACF/OPRE Certifying Officer.

[FR Doc. 2022–06415 Filed 3–25–22; 8:45 am]

BILLING CODE 4184–47–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2021–N–0709]

Prescription Drug User Fee Rates for Fiscal Year 2022; Correction

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice; correction.

SUMMARY: The Food and Drug Administration (FDA) is correcting a notice entitled ''Prescription Drug User Fee Rates for Fiscal Year 2022'' that appeared in the **Federal Register** of August 16, 2021. The document announced the Fiscal Year 2022 fee rates for the Prescription Drug User Fee Act. The document published with errors. The errors did not have an impact on the previously published user fee rates but are corrected in this document for clarity.

FOR FURTHER INFORMATION CONTACT: Misbah Tareen, Office of Financial Management, Food and Drug Administration, 4041 Powder Mill Rd., Rm. 61077A, Beltsville, MD 20705–4304, 301–796–3997.

SUPPLEMENTARY INFORMATION: In the **Federal Register** of August 16, 2021 (86 FR 45732), appearing on page 45736 in FR Doc. 2021–17505, the following corrections are made:

1. In the second column, in the last sentence of the third paragraph under ''D. FY 2022 Statutory Fee Revenue Adjustments for Operating Reserve'', ''both user fee funds available for obligation $126,873,636 and funds that are considered unavailable due to a lack of appropriations $98,850,995'' is corrected to read ''user fee funds considered unavailable due to a lack of appropriations $78,850,995, additional fee funds that are available for obligation but set aside for future year refunds as a matter of prudent operations $20,000,000, and carryover net of unavailable funds and the set-aside $126,873,636.''

2. The fourth footnote is corrected by removing the text and replacing it with: ''In recent PDUFA Annual Financial Reports, the category ''unavailable for use'' has been used to refer both to (1) fee funds that are considered unappropriated and (2) appropriated fee funds the Agency has maintained to provide for any refunds. FDA intends to discontinue use of the category ''unavailable for use'' in forthcoming reports to better reflect the difference between these line items and improve the clarity of its reporting. Although certain amounts have been maintained for future refunds as a matter of prudent operations, these amounts are considered appropriated and are available for obligation.''

3. In the second column, in the fifth paragraph under ''D. FY 2022, Statutory Fee Revenue Adjustments for Operating Reserve'', sentences 4 through 7 are corrected by removing the text and replacing it with ''FDA has decided to make an available operating reserve adjustment that is intended to increase the amount of available funds to approximately 8 weeks by the end of FY 2022, representing the low end of the 8- to 10-week range while mitigating the impact on fee amounts. FDA estimates the cost of operations per week is $22,144,672. Before the operating adjustment, the estimated end of year FY 2022 available operating reserve is $145,677,240, which equates to about 6½ weeks of available operating reserves. Adding the FY 2022 operating reserve adjustment of $39,402,923 to this amount is expected to provide approximately 8 weeks of available operating reserve, or $185,080,162 (including $20,000,000 in available fee funds maintained for any future refunds), and a total carryover of operating reserves (including unavailable funds) of $263,931,157.''

Dated: March 21, 2022.

Andi Lipstein Fristedt,

Deputy Commissioner for Policy, Legislation, and International Affairs, U.S. Food and Drug Administration.

[FR Doc. 2022–06427 Filed 3–25–22; 8:45 am]

BILLING CODE 4164–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2021–N–0336]

Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Quantitative Research on a Voluntary Symbol Depicting the Nutrient Content Claim ''Healthy'' on Packaged Foods

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA 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 (PRA).

DATES: Submit written comments (including recommendations) on the collection of information by April 27, 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 title of this information collection is ''Quantitative Research on a Voluntary Symbol Depicting the Nutrient Content Claim 'Healthy' on Packaged Foods.'' Also include the FDA docket number found in brackets in the heading of this document.

FOR FURTHER INFORMATION CONTACT: Ila S. Mizrachi, Office of Operations, Food and Drug Administration, Three White