DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Proposed Information Collection Activity; Intergovernmental Reference Guide (IRG) (OMB No.: 0970–0209)

AGENCY: Office of Child Support Enforcement, Administration for Children and Families, HHS.

ACTION: Request for public comment.

SUMMARY: The Administration for Children and Families (ACF), Office of Child Support Enforcement (OCSE), is requesting the Office of Management and Budget (OMB) to approve the Intergovernmental Reference Guide (IRG), with content revisions, for an additional three years. The IRG contains state and tribal child support information that assists child support enforcement (CSE) agencies in the administration of their respective programs. The current OMB approval expires on January 31, 2022.

DATES: *Comments due within 60 days of publication.* In compliance with the requirements of the Paperwork Reduction Act of 1995, the ACF is soliciting public comment on the specific aspects of the information collection described above.

ADDRESSES: Copies of the proposed collection of information can be obtained and comments may be forwarded by emailing *infocollection@acf.hhs.gov.* Alternatively, copies can also be obtained by writing to the Administration for Children and Families, Office of Planning, Research and Evaluation, 330 C Street SW, Washington, DC 20201, Attn: ACF Reports Clearance Officer. All requests, emailed or written, should be identified by the title of the information collection.

SUPPLEMENTARY INFORMATION:

Description: The IRG is a centralized and automated repository of state and tribal profiles that contains high-level descriptions of each CSE program. These profiles provide state, tribal, and foreign country CSE agencies with an effective and efficient method for updating and accessing information needed to process intergovernmental child support cases. Proposed revisions to the state profile include content changes and organizational updates. Proposed revisions to the tribal profile are only organizational, no content changes are proposed.

Respondents: State and Tribal Child Support Enforcement Agencies.

ANNUAL BURDEN ESTIMATES

Information collection instrument	Total number of annual respondents	Number of annual responses per respondent	Average annual burden hour per response	Annual burden hours
IRG: State Profile Guide (states and territories)	54	18	0.3	292
IRG: Tribal Profile Guide	62	18	0.3	335

Estimated Total Annual Burden Hours: 627.

Comments: The Department specifically requests comments 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) the quality, utility, and clarity of the information to be collected; and (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.

Consideration will be given to comments and suggestions submitted within 60 days of this publication.

Authority: 42 U.S.C. 652(a)(7); 42 U.S.C. 666(f); 45 CFR 301.1; 45 CFR 303.7; and 45 CFR 309.120.

Mary B. Jones,

ACF/OPRE Certifying Officer.

[FR Doc. 2021–16244 Filed 7–29–21; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2020–N–2340]

Matthew Hebert: Final Debarment Order

AGENCY: Food and Drug Administration, Health and Human Services (HHS).

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is issuing an order under the Federal Food, Drug, and Cosmetic Act (FD&C Act) debarring Matthew Hebert for a period of 5 years from importing articles of food or offering such articles for importation into the United States. FDA bases this order on a finding that Mr. Hebert was convicted of a felony count under Federal law for conduct relating to the importation into the United States of an article of food. Mr. Hebert was given notice of the proposed debarment and an opportunity to request a hearing within the timeframe prescribed by regulation. As of May 5, 2021 (30 days after receipt of the notice), Mr. Hebert has not responded. Mr. Hebert's failure to respond and request a hearing constitutes a waiver of his right to a hearing concerning this matter.

DATES: This order is applicable July 30, 2021.

ADDRESSES: Submit applications for termination of debarment to the Dockets Management Staff (HFA–305), Food and Drug Administration, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852, 240–402–7500, or at *https://www.regulations.gov.*

FOR FURTHER INFORMATION CONTACT: Jaime Espinosa (ELEM–4029), Division of Enforcement, Office of Strategic Planning and Operational Policy, Office of Regulatory Affairs, Food and Drug Administration, 12420 Parklawn Dr., Rockville, MD 20857, 240–402–8743, or at *debarments@fda.hhs.gov.*

SUPPLEMENTARY INFORMATION:

I. Background

Section 306(b)(1)(C) of the FD&C Act (21 U.S.C. 335a(b)(1)(C)) permits debarment of an individual from importing an article of food or offering such an article for import into the United States if FDA finds, as required by section 306(b)(3)(A) of the FD&C Act, that the individual has been convicted of a felony for conduct relating to the importation into the United States of any food. On December 11, 2020, Mr. Hebert was convicted, as defined in section 306(*l*)(1)(A) of the FD&C Act, in the U.S. District Court for the Northern

District of Texas-Dallas Division, when the court accepted Mr. Hebert's plea of guilty and entered judgment against him for the offense of introduction of misbranded food into interstate commerce with intent to defraud and mislead, in violation of sections 301(a) and 303(a)(2) of the FD&C Act (21 U.S.C. 331(a) and 333(a)(2)).

FDA's finding that the debarment is appropriate is based on the felony conviction referenced herein. The factual basis for this conviction is as follows: As contained in the superseding indictment, filed on January 5, 2016, Mr. Hebert was a co-owner of USP Labs with primarily responsibilities over product packaging design. As contained in the factual résumé submitted as part of Mr. Hebert's plea agreement on March 11, 2019, and the factual resumes submitted as part of plea agreements with his codefendants, one of Mr. Hebert's codefendants instructed a Chinese company to have 2 metric tons of ground cynanchum auriculatum root powder shipped internationally to S.K. Laboratories in California for inclusion in USP Labs' dietary supplement products, using the false name ''cynanchum auriculatum root extract.'' USP Labs sent false labels to retailers and wholesalers listing ''cynanchum auriculatum (root) extract'' as an ingredient in OxyElite Pro ''Advanced Formula'' (which went on sale in or around August 2013), even though that ingredient was not present in the product. Beginning in or around August 2013, Mr. Hebert, USP Labs, and others working at USP Labs and S.K. Laboratories, did knowingly, and with the intent to defraud and mislead, cause the shipment of a misbranded food, namely the OxyElite Pro ''Advanced Formula'' dietary supplement, in interstate commerce. Specifically, on or about October 4, 2013, with intent to defraud and mislead, Mr. Hebert caused the shipment of misbranded OxyElite Pro ''Advanced Formula'' in interstate commerce. The labeling for OxyElite Pro ''Advanced Formula'' falsely declared cynanchum auriculatum (root) extract as an ingredient, when in fact OxyElite Pro ''Advanced Formula'' contained imported cynanchum auriculatum powder but no cynanchum auriculatum (root) extract.

As a result of this conviction, FDA sent Mr. Hebert, by certified mail on March 29, 2021, a notice proposing to debar him for a period of 5 years from importing articles of food or offering such articles for import into the United States. The proposal was based on a finding under section 306(b)(1)(C) of the FD&C Act that Mr. Hebert's felony conviction of ''introduction of misbranded food into interstate commerce with intent to defraud and mislead'' in violation of sections 301(a) and 303(a)(2) of the FD&C Act constitutes conduct relating to the importation into the United States of an article of food because Mr. Hebert caused the shipment of a misbranded food in interstate commerce, and the food was misbranded because its labeling falsely declared cynanchum auriculatum (root) extract as an ingredient, when in fact the imported ingredient was cynanchum auriculatum powder, not cynanchum auriculatum root extract.

The proposal was also based on a determination, after consideration of the relevant factors set forth in section 306(c)(3) of the FD&C Act, that Mr. Hebert should be subject to a 5-year period of debarment. The proposal also offered Mr. Hebert an opportunity to request a hearing, providing Mr. Hebert 30 days from the date of receipt of the letter in which to file the request, and advised Mr. Hebert that failure to request a hearing constituted a waiver of the opportunity for a hearing and of any contentions concerning this action. Mr. Hebert failed to respond within the timeframe prescribed by regulation and has, therefore, waived his opportunity for a hearing and waived any contentions concerning his debarment (21 CFR part 12).

II. Findings and Order

Therefore, the Assistant Commissioner, Office of Human and Animal Food Operations, under section 306(b)(1)(C) of the FD&C Act, under authority delegated to the Assistant Commissioner, finds that Mr. Matthew Hebert has been convicted of a felony count under Federal law for conduct relating to the importation into the United States of an article of food and that he is subject to a 5 year period of debarment.

As a result of the foregoing finding, Mr. Hebert is debarred for a period of 5 years from importing articles of food or offering such articles for import into the United States, effective (see **DATES**). Pursuant to section 301(cc) of the FD&C Act, the importing or offering for import into the United States of an article of food by, with the assistance of, or at the direction of Matthew Hebert is a prohibited act.

Any application by Mr. Hebert for termination of debarment under section 306(d)(1) of the FD&C Act should be identified with Docket No. FDA–2020–N–2340 and sent to the Dockets Management Staff (see **ADDRESSES**). The public availability of information in these submissions is governed by 21 CFR 10.20(j).

Publicly available submissions will be placed in the docket and will be viewable at *https://www.regulations.gov* or at the Dockets Management Staff (see **ADDRESSES**) between 9 a.m. and 4 p.m., Monday through Friday, 240–402–7500.

Dated: July 23, 2021.

Lauren K. Roth,

Acting Principal Associate Commissioner for Policy.

[FR Doc. 2021–16211 Filed 7–29–21; 8:45 am]

BILLING CODE 4164–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Resources and Services Administration

Notice of a Supplemental Award to the University of Arkansas System Telehealth Focused Rural Health Research Center

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

ACTION: Notice.

SUMMARY: HRSA announces a supplemental award in the amount of $100,000 annually through fiscal year 2024. This funding will support the University of Arkansas Telehealth Focused Rural Health Research Center (TF RHRC), a current HRSA-funded cooperative agreement. The supplemental request is for the remaining period of performance of the current cooperative agreement, subject to the availability of funds and successful performance of the activities in a given budget year.

FOR FURTHER INFORMATION CONTACT: For further information regarding this request, please contact Nicole Hewitt, (*nhewitt@hrsa.gov*), (301) 443–3893.

SUPPLEMENTARY INFORMATION:

Intended Recipient of Award: University of Arkansas System.

Amount of Award Increase Non-Competitive Award: $100,000 annually.

Period of Supplemental Funding: 9/1/2021–08/31/2024.

CFDA Number: 93.155.

Authority: Section 711(b)(5) of the Social Security Act (42 U.S.C. 912(b)(5)).

Justification: The TF RHRC program is authorized by Section 711(b)(5) of the Social Security Act (42 U.S.C. 912(b)(5)), as amended. This program is within HRSA's Federal Office of Rural Health Policy's (FORHP) Office for the Advancement of Telehealth (OAT), which administers grants, cooperative