Dated: July 12, 2022.

Sarah L. Stewart,

Deputy General Counsel, Federal Mine Safety and Health Review Commission.

[FR Doc. 2022–15277 Filed 7–15–22; 8:45 am]

BILLING CODE 6735–01–P

FEDERAL RESERVE SYSTEM

Change in Bank Control Notices; Acquisitions of Shares of a Bank or Bank Holding Company

The notificants listed below have applied under the Change in Bank Control Act (Act) (12 U.S.C. 1817(j)) and § 225.41 of the Board's Regulation Y (12 CFR 225.41) to acquire shares of a bank or bank holding company. The factors that are considered in acting on the applications are set forth in paragraph 7 of the Act (12 U.S.C. 1817(j)(7)).

The public portions of the applications listed below, as well as other related filings required by the Board, if any, are available for immediate inspection at the Federal Reserve Bank(s) indicated below and at the offices of the Board of Governors. This information may also be obtained on an expedited basis, upon request, by contacting the appropriate Federal Reserve Bank and from the Board's Freedom of Information Office at *https://www.federalreserve.gov/foia/request.htm.* Interested persons may express their views in writing on the standards enumerated in paragraph 7 of the Act.

Comments regarding each of these applications must be received at the Reserve Bank indicated or the offices of the Board of Governors, Ann E. Misback, Secretary of the Board, 20th Street and Constitution Avenue NW, Washington, DC 20551–0001, not later than August 2, 2022.

A. Federal Reserve Bank of Chicago (Colette A. Fried, Assistant Vice President) 230 South LaSalle Street, Chicago, Illinois 60690–1414:

1. *The Bell Family Voting Trust (''Trust''), Wisconsin Rapids, Wisconsin and Steven C. Bell and Paula J. Bell, both of Wisconsin Rapids, Wisconsin, Elizabeth Bell Killian, Spokane, Washington, Rebecca L. Kettleson, Wausau, Wisconsin and Margaret S. Bell, Chicago, Illinois, all co-trustees of the Trust;* to become members of the Bell Family Control Group, a group acting in concert, to acquire voting shares of WoodTrust Financial Corporation, and thereby indirectly acquire voting shares of WoodTrust Bank, both of Wisconsin Rapids, Wisconsin.

Board of Governors of the Federal Reserve System.

Michele Taylor Fennell,

Deputy Associate Secretary of the Board.

[FR Doc. 2022–15274 Filed 7–15–22; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

[OMB No. 0970–0036]

Submission for OMB Review; ORR–6 Performance Report

AGENCY: Office of Refugee Resettlement, Administration for Children and Families, HHS.

ACTION: Request for Public Comment.

SUMMARY: The Administration for Children and Families (ACF), Office of Refugee Resettlement (ORR) is requesting a renewal of the ORR–6 Performance Report (OMB #0970–0036, expiration 03/31/2023). ORR published a notice in the **Federal Register** on 8/12/2021 requesting comments within 60-days on revisions to the ORR–6.

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: ACF/ORR requests information from the ORR–6 Performance Report to determine effectiveness of state Cash and Medical Assistance (CMA) and Refugee Support Services programs. ORR uses state-by-state CMA utilization rates, derived from the ORR–6 Performance Report, to formulate program initiatives, priorities, standards, budget requests, and assistance policies. Federal regulations require state Refugee Resettlement, Replacement Designee agencies, and local governments submit statistical or programmatic information that the ORR Director determines to be required to fulfill their responsibility under the Immigration and Nationality Act (INA). The currently approved ORR–6 has been updated to add new data elements to better understand the meaning of existing data collection, and update the instructions and reformat some of the forms to provide clearer definitions and better distinguish the participation and performance results of different support services programs. In addition, some revisions are related to Afghanistan Supplemental Appropriations Act 2022, Additional Afghanistan Supplemental Appropriations Act 2022, and Additional Ukraine Supplemental Appropriations Act 2022

Respondents: State governments and Replacement Designees.

ANNUAL BURDEN ESTIMATES

Instrument	Total number of respondents	Total No. of responses per respondent	Average burden hours per response	Total burden hours	Annual burden hours
ORR–6 Performance Report	64	6	15	5,760	1,920

Estimated Total Annual Burden Hours: 1,920.

Authority: 8 U.S.C 1522 of the Immigration and Nationality Act (the Act) (Title IV, Sec. 412 of the Act), and 45 CFR 400.28(b).

Mary B. Jones,

ACF/OPRE Certifying Officer.

[FR Doc. 2022–15227 Filed 7–15–22; 8:45 am]

BILLING CODE 4184–45–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–1999–D–2955 (formerly 1999D–4071)]

International Cooperation on Harmonisation of Technical Requirements for Registration of Veterinary Medicinal Products; Impurities: Residual Solvents in New Veterinary Medicinal Products, Active Substances and Excipients (Revision 2); Draft Guidance for Industry; Availability

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice of availability.

SUMMARY: The Food and Drug Administration (FDA or Agency) is announcing the availability of a draft guidance for industry (GFI) #100 (VICH GL 18(R2)) entitled ''Impurities: Residual Solvents in New Veterinary Medicinal Products, Active Substances and Excipients (Revision 2).'' This draft guidance has been developed for veterinary use by the International Cooperation on Harmonisation of Technical Requirements for Registration of Veterinary Medicinal Products (VICH). The objective of this draft guidance is to recommend acceptable amounts for residual solvents in pharmaceuticals for the safety of the target animal as well as for the safety of residues in products derived from treated food-producing animals. This revision updates the listings and classification of solvents.

DATES: Submit either electronic or written comments on the draft guidance by September 16, 2022 to ensure that the Agency considers your comment on this draft guidance before it begins work on the final version of the guidance.

ADDRESSES: You may submit comments on any guidance at any time as follows:

Electronic Submissions

Submit electronic comments in the following way:

- *Federal eRulemaking Portal: https://www.regulations.gov.* Follow the instructions for submitting comments. Comments submitted electronically, including attachments, to *https://www.regulations.gov* will be posted to the docket unchanged. Because your comment will be made public, you are solely responsible for ensuring that your comment does not include any confidential information that you or a third party may not wish to be posted, such as medical information, your or anyone else's Social Security number, or confidential business information, such as a manufacturing process. Please note that if you include your name, contact information, or other information that identifies you in the body of your comments, that information will be posted on *https://www.regulations.gov.*
- If you want to submit a comment with confidential information that you do not wish to be made available to the public, submit the comment as a written/paper submission and in the manner detailed (see ''Written/Paper Submissions'' and ''Instructions'').

Written/Paper Submissions

Submit written/paper submissions as follows:

- *Mail/Hand Delivery/Courier (for written/paper submissions):* Dockets Management Staff (HFA–305), Food and Drug Administration, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852.
- For written/paper comments submitted to the Dockets Management Staff, FDA will post your comment, as well as any attachments, except for information submitted, marked and identified, as confidential, if submitted as detailed in ''Instructions.''

Instructions: All submissions received must include the Docket No. FDA–1999–D–2955 (formerly 1999D–4071) for ''Impurities: Residual Solvents in New Veterinary Medicinal Products, Active Substances and Excipients (Revision 2).'' Received comments will be placed in the docket and, except for those submitted as ''Confidential Submissions,'' publicly viewable at *https://www.regulations.gov* or at the Dockets Management Staff between 9 a.m. and 4 p.m., Monday through Friday, 240–402–7500.

- Confidential Submissions—To submit a comment with confidential information that you do not wish to be made publicly available, submit your comments only as a written/paper submission. You should submit two copies total. One copy will include the information you claim to be confidential with a heading or cover note that states ''THIS DOCUMENT CONTAINS CONFIDENTIAL INFORMATION.'' The Agency will review this copy, including the claimed confidential information, in its consideration of comments. The second copy, which will have the claimed confidential information redacted/blacked out, will be available for public viewing and posted on *https://www.regulations.gov.* Submit both copies to the Dockets Management Staff. If you do not wish your name and contact information to be made publicly available, you can provide this information on the cover sheet and not in the body of your comments and you must identify this information as ''confidential.'' Any information marked as ''confidential'' will not be disclosed except in accordance with 21 CFR 10.20 and other applicable disclosure law. For more information about FDA's posting of comments to public dockets, see 80 FR 56469, September 18, 2015, or access the information at: *https://www.govinfo.gov/content/pkg/FR-2015-09-18/pdf/2015-23389.pdf.*

Docket: For access to the docket to read background documents or the electronic and written/paper comments received, go to *https://www.regulations.gov* and insert the docket number, found in brackets in the heading of this document, into the ''Search'' box and follow the prompts and/or go to the Dockets Management Staff, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852, 240–402–7500.

You may submit comments on any guidance at any time (see 21 CFR 10.115(g)(5)).

Submit written requests for single copies of this draft guidance to the Policy and Regulations Staff (HFV–6), Center for Veterinary Medicine, 7500 Standish Pl., Rockville, MD 20855. Send one self-addressed adhesive label to assist that office in processing your request or include a fax number to which the draft guidance may be sent. See the **SUPPLEMENTARY INFORMATION** section for information on electronic access to the draft guidance.

FOR FURTHER INFORMATION CONTACT: Mai Huynh, Center for Veterinary Medicine (HFV–140), Food and Drug Administration, 7500 Standish Pl., Rockville, MD 20855, 240–402–0669, *mai.huynh@fda.hhs.gov.*

SUPPLEMENTARY INFORMATION:

I. Background

FDA is announcing the availability of a draft GFI #100 (VICH GL18(R2)) entitled ''Impurities: Residual Solvents in New Veterinary Medicinal Products, Active Substances and Excipients (Revision 2).'' Residual solvents in pharmaceuticals are defined in the guidance as organic volatile chemicals that are used or produced in the manufacture of active substances or