April 22, at 5 p.m. Eastern Time. Members of the public are encouraged to submit registration requests via email in advance of the deadline. Seating for members of the public is limited and will be available on a first-come, first-served basis for those who register in advance. We will note when the limit of in-person attendees has been reached. If you have accessibility concerns and require assistance, contact *secretary@fmc.gov.*

FOR FURTHER INFORMATION CONTACT: Mr. Dylan Richmond, Designated Federal Officer of the National Shipper Advisory Committee, phone: (202) 523–5810; email: *drichmond@fmc.gov.*

SUPPLEMENTARY INFORMATION:

Background: The National Shipper Advisory Committee is a federal advisory committee. It operates under the provisions of the Federal Advisory Committee Act, 5 U.S.C. app., and 46 U.S.C. chapter 425. The Committee was established on January 1, 2021, when the National Defense Authorization Act for Fiscal Year 2021 became law. Public Law 116–283, section 8604, 134 stat. 3388 (2021). The Committee will provide information, insight, and expertise pertaining to conditions in the ocean freight delivery system to the Commission. Specifically, the Committee will advise the Federal Maritime Commission on policies relating to the competitiveness, reliability, integrity, and fairness of the international ocean freight delivery system. 46 U.S.C. 42502(b).

The Committee will hear from Commissioner Bentzel for an update on the Maritime Transportation Data Initiative. They will also receive updates from each of its subcommittees. The Committee will receive proposals for recommendations to the Federal Maritime Commission and may vote on these recommendations. These recommendations will also be available for the public to view in advance of the meeting on the NSAC's website, *https://www.fmc.gov/industry-oversight/national-shipper-advisory-committee/.*

Public Comments: Members of the public may submit written comments to NSAC at any time. Comments should be addressed to NSAC, c/o Dylan Richmond, Federal Maritime Commission, 800 North Capitol St NW, Washington, DC 20573 or *nsac@fmc.gov.*

The Committee will also take public comment at its meeting. If attending the meeting and providing comments, please note that in the registration request. Comments are most helpful if they address the Committee's objectives or their proposed recommendations. Comments at the meeting will be limited to 3 minutes each.

A copy of all meeting documentation, including meeting minutes, will be available at *www.fmc.gov* following the meeting.

By the Commission.

Dated: April 8, 2022.

William Cody,

Secretary.

[FR Doc. 2022–07896 Filed 4–12–22; 8:45 am]

BILLING CODE 6730–02–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifiers: CMS–10680 and CMS–10692]

Agency Information Collection Activities: Proposed Collection; Comment Request

AGENCY: Centers for Medicare & Medicaid Services, Health and Human Services (HHS).

ACTION: Notice.

SUMMARY: The Centers for Medicare & Medicaid Services (CMS) is announcing an opportunity for the public to comment on CMS' intention to collect information from the public. Under the Paperwork Reduction Act of 1995 (the PRA), federal agencies are required to publish notice in the **Federal Register** concerning each proposed collection of information (including each proposed extension or reinstatement of an existing collection of information) and to allow 60 days for public comment on the proposed action. Interested persons are invited to send comments regarding our burden estimates or any other aspect of this collection of information, including the necessity and utility of the proposed information collection for the proper performance of the agency's functions, the accuracy of the estimated burden, ways to enhance the quality, utility, and clarity of the information to be collected, and the use of automated collection techniques or other forms of information technology to minimize the information collection burden.

DATES: Comments must be received by June 13, 2022.

ADDRESSES: When commenting, please reference the document identifier or OMB control number. To be assured consideration, comments and recommendations must be submitted in any one of the following ways:

1. *Electronically.* You may send your comments electronically to *http://www.regulations.gov.* Follow the instructions for ''Comment or Submission'' or ''More Search Options'' to find the information collection document(s) that are accepting comments.

2. *By regular mail.* You may mail written comments to the following address: CMS, Office of Strategic Operations and Regulatory Affairs, Division of Regulations Development, Attention: Document Identifier/OMB Control Number: ____, Room C4–26–05, 7500 Security Boulevard, Baltimore, Maryland 21244–1850.

To obtain copies of a supporting statement and any related forms for the proposed collection(s) summarized in this notice, you may make your request using one of following:

1. Access CMS' website address at website address at *https://www.cms.gov/Regulations-and-Guidance/Legislation/PaperworkReductionActof1995/PRA-Listing.*

FOR FURTHER INFORMATION CONTACT: William N. Parham at (410) 786–4669.

SUPPLEMENTARY INFORMATION:

Contents

This notice sets out a summary of the use and burden associated with the following information collections. More detailed information can be found in each collection's supporting statement and associated materials (see **ADDRESSES**).

CMS–10680 Electronic Visit Verification Compliance Survey

CMS–10692 Home and Community Based Services (HCBS) Incident Management Survey

Under the PRA (44 U.S.C. 3501–3520), federal agencies must obtain approval from the Office of Management and Budget (OMB) for each collection of information they conduct or sponsor. The term ''collection of information'' is defined in 44 U.S.C. 3502(3) and 5 CFR 1320.3(c) and includes agency requests or requirements that members of the public submit reports, keep records, or provide information to a third party. Section 3506(c)(2)(A) of the PRA requires federal agencies to publish a 60-day notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension or reinstatement of an existing collection of information, before submitting the collection to OMB for approval. To comply with this requirement, CMS is publishing this notice.

Information Collection

1. *Title of Information Collection:* Electronic Visit Verification Compliance

Survey; *Type of Information Collection Request:* Extension without change of a currently approved collection; *Use:* The web-based survey will allow states to self-report their progress in implementing electronic visit verification (EVV) for personal care services (PCS) and home health care services (HHCS), as required by section 1903(l) of the Social Security Act. CMS will use the survey data to assess states' compliance with section 1903(l) of the Act and levy Federal Medical Assistance Percentage (FMAP) reductions where necessary as required by 1903(l) of the Act.

The survey will be disseminated to all 51 state Medicaid agencies (including the District of Columbia) and the Medicaid agencies of five US territories. States will be required to complete the survey in order to demonstrate that they are complaint with Section 1903(l) of the Act by reporting on their EVV implementation status for PCS provided under sections 1905(a)(24), 1915(c), 1915(i), 1915(j), 1915(k), and Section 1115 of the Act; and HHCS provided under 1905(a)(7) of the Act or under a demonstration project or waiver (*e.g.,* 1915(c) or 1115 of the Act).

The survey will be a live form, meaning states will have the ability to update their 1903(l) compliance status on a continuous basis. As FMAP reductions are assigned quarterly per 1903(l) of the Act, states who are not in compliance will be asked to review their survey information on a quarterly basis to ensure it is up-to-date and to update their survey responses as needed until they come into compliance. *Form Number:* CMS–10680 (OMB control number: 0938–1360); *Frequency:* On occasion; *Affected Public:* State, Local, or Tribal Governments; *Number of Respondents:* 56; *Number of Responses:* 336; *Total Annual Hours:* 504. (For questions regarding this collection contact Ryan Shannahan at 410–786–0295.)

2. *Type of Information Collection Request:* Extension without change of a currently approved collection; *Title of Information Collection:* Home and Community Based Services (HCBS) Incident Management Survey; *Use:* The Survey will be disseminated to all 51 state Medicaid agencies (including the District of Columbia) to assess incident management systems in 1915(c) waivers. States will be surveyed to identify methods and promising practices for identifying, reporting, tracking, and resolving incidents of abuse, neglect, and exploitation. The survey results will also be used to review the strengths and weaknesses of each state's incident management system and will inform guidance to help ensure compliance with sections 1902(a)(30(A) and 1915(c)(2)(A) of the Social Security Act. *Form Number:* CMS–10692 (OMB control number: 0938–1362); *Frequency:* Once and on occasion; *Affected Public:* State, Local, or Tribal Governments; *Number of Respondents:* 51; *Total Annual Responses:* 102; *Total Annual Hours:* 153. (For policy questions regarding this collection contact Ryan Shannahan at 410–786–0295.)

3.

Dated: April 8, 2022.

William N. Parham, III,

Director, Paperwork Reduction Staff, Office of Strategic Operations and Regulatory Affairs.

[FR Doc. 2022–07917 Filed 4–12–22; 8:45 am]

BILLING CODE 4120–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Request for Information: Technical Assistance Needs and Priorities on Implementation and Coordination of Early Childhood Development Programs in American Indian and Alaska Native Communities; Correction

AGENCY: Administration for Children and Families, HHS.

ACTION: Notice; correction.

SUMMARY: The Administration for Children and Families published a document in the **Federal Register** of March 22, 2022 concerning a request for information on technical assistance needs and priorities on implementation and coordination of early childhood development programs in American Indian and Alaska Native communities. The document contained incorrect dates.

FOR FURTHER INFORMATION CONTACT: Moushumi Beltangady at *Moushumi.beltangady@acf.hhs.gov* or 202–260–3613.

SUPPLEMENTARY INFORMATION:

Correction

In the **Federal Register** of March 22, 2022, in FR Doc. 2022–05962 (Vol. 87, No. 55) on page 16195, in the first column, final line, correct the **DATES** caption to read:

DATES: Send comments on or before May 20, 2022.

Kathleen D. Hamm,

Deputy Assistant Secretary for Early Childhood Development, Administration for Children and Families, U.S. Department of Health and Human Services.

[FR Doc. 2022–07840 Filed 4–12–22; 8:45 am]

BILLING CODE P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2022–N–0529]

Secura Bio, Inc.; Withdrawal of Approval of Relapsed or Refractory Follicular Lymphoma Indication for COPIKTRA

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that it is withdrawing approval of the relapsed or refractory follicular lymphoma indication for COPIKTRA (duvelisib) Capsules, approved under new drug application 211155, held by Secura Bio, Inc., 1995 Village Center Circle, Suite 128, Las Vegas, NV 89134. Secura Bio, Inc. voluntarily requested that the Agency withdraw approval of this indication and has waived its opportunity for a hearing.

DATES: Approval is withdrawn as of April 13, 2022.

FOR FURTHER INFORMATION CONTACT: Kimberly Lehrfeld, Center for Drug Evaluation and Research, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 51, Rm. 6226, Silver Spring, MD 20993–0002, 301–796–3137, *Kimberly.Lehrfeld@fda.hhs.gov.*

SUPPLEMENTARY INFORMATION: FDA approved COPIKTRA (duvelisib) Capsules for the treatment of adult patients with relapsed or refractory follicular lymphoma after at least two prior systemic therapies (the follicular lymphoma indication) on September 24, 2018, under the Agency's accelerated approval regulations, 21 CFR part 314, subpart H. As a condition of accelerated approval of COPIKTRA (duvelisib) Capsules for follicular lymphoma, the applicant was required to conduct a postmarketing trial to verify the clinical benefit of duvelisib for follicular lymphoma.

On November 22, 2021, FDA met with Secura Bio, Inc., to discuss the company's inability to conduct a