- Joan Ilardo, Director of Research Initiatives, Michigan State University, College of Human Medicine.
- Cheri Lattimer, Executive Director, National Transitions of Care Coalition.
- Cori McMahon, Vice President, Tridiuum.
- Alan Meade, Director of Rehabilitation Services, Holston Medical Group.
- Michael Minor, National Director, H.O.P.E. HHS Partnership, National Baptist Convention USA, Incorporated.
- Jina Ragland, Associate State Director of Advocacy and Outreach, AARP Nebraska.
- Morgan Reed, Executive Director, Association for Competitive Technology.
- Margot Savoy, Senior Vice President, American Academy of Family Physicians.
- Congresswoman Allyson Schwartz, Senior Advisor, FTI Consulting.
- Tia Whitaker, Statewide Director, Outreach and Enrollment, Pennsylvania Association of Community Health Centers.

II. Provisions of This Notice

In accordance with section 10(a) of the FACA, this notice announces a meeting of the APOE. The agenda for the February 3, 2022 meeting will include the following:

- Welcome and listening session with CMS leadership
- Recap of the previous (September 15, 2021) meeting
- CMS programs, initiatives, and priorities
- An opportunity for public comment
- Meeting summary, review of recommendations, and next steps

Individuals or organizations that wish to make a 5-minute oral presentation on an agenda topic should submit a written copy of the oral presentation to the DFO at the address listed in the **ADDRESSES** section of this notice by the date listed in the **DATES** section of this notice. The number of oral presentations may be limited by the time available. Individuals not wishing to make an oral presentation may submit written comments to the DFO at the address listed in the **ADDRESSES** section of this notice by the date listed in the **DATES** section of this notice.

III. Meeting Participation

The meeting is open to the public, but attendance is limited to registered participants. Persons wishing to attend this meeting must register at the website *https://www.eventbrite.com/e/apoe-february-3-2022-virtual-meeting-tickets-212590763697* or contact the DFO at the address or number listed in the **FOR FURTHER INFORMATION CONTACT** section of this notice by the date specified in the **DATES** section of this notice. This meeting will be held virtually. Individuals who are not registered in advance will be unable to attend the meeting.

IV. Collection of Information

This document does not impose information collection requirements, that is, reporting, recordkeeping, or third-party disclosure requirements. Consequently, there is no need for review by the Office of Management and Budget under the authority of the Paperwork Reduction Act of 1995 (44 U.S.C. Chapter 35).

The Administrator of the Centers for Medicare & Medicaid Services (CMS), Chiquita Brooks-LaSure, having reviewed and approved this document, authorizes Lynette Wilson, who is the Federal Register Liaison, to electronically sign this document for purposes of publication in the **Federal Register**.

Dated: January 11, 2022.

Lynette Wilson,

Federal Register Liaison, Centers for Medicare & Medicaid Services.

[FR Doc. 2022–00745 Filed 1–14–22; 8:45 am]

BILLING CODE 4120–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2021–N–1037]

Fresenius USA, Inc., et al.; Withdrawal of Approval of 216 Abbreviated New Drug Applications; Correction

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice; correction.

SUMMARY: The Food and Drug Administration (FDA) is correcting a notice that appeared in the **Federal Register** of October 22, 2021. The document announced the withdrawal of approval of 216 abbreviated new drug applications (ANDAs) from multiple applicants, as of November 22, 2021. The document was published with an incorrect date. In addition, the document indicated that FDA was withdrawing approval of ANDA 075941, Strontium Chloride SR–89 Injection, 1 millicurie/milliliter, held by Bio-Nucleonics, Inc., 1600 Market St., Suite 13200, Philadelphia, PA 19103, for repeated failure to submit annual reports. Before FDA withdrew the approval of this ANDA, the application holder informed FDA that it submitted annual reports for ANDA 075941. Therefore, FDA rescinds its withdrawal of approval of ANDA 075941. The approval of ANDA 075941 is still in effect.

FOR FURTHER INFORMATION CONTACT: James Hanratty, Center for Drug Evaluation and Research, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 75, Rm. 1671, Silver Spring, MD 20993–0002, 240–402–4718, *James.Hanratty@fda.hhs.gov.*

SUPPLEMENTARY INFORMATION:

Corrections

In the **Federal Register** of Friday, October 22, 2021 (86 FR 58675), in FR Doc. 2021–23075, the following corrections are made:

1. On page 58675, in the second column, correct the **DATES** section to read: **DATES:** Approval is withdrawn as of October 22, 2021.''

2. On page 58679, in the table, remove the entry for ANDA 075941.

Dated: January 12, 2022.

Lauren K. Roth,

Associate Commissioner for Policy.

[FR Doc. 2022–00831 Filed 1–14–22; 8:45 am]

BILLING CODE 4164–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2021–P–0885]

Determination That PEPCID (Famotidine) Tablet, 20 Milligrams and 40 Milligrams, Was Not Withdrawn From Sale for Reasons of Safety or Effectiveness

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA or Agency) has determined that PEPCID (famotidine) tablet, 20 milligrams (mg) and 40 mg, was not withdrawn from sale for reasons of safety or effectiveness. This determination means that FDA will not begin procedures to withdraw approval of abbreviated new drug applications (ANDAs) that refer to this drug product, and it will allow FDA to continue to approve ANDAs that refer to the product as long as they meet relevant legal and regulatory requirements.

FOR FURTHER INFORMATION CONTACT: Stacy Kane, Center for Drug Evaluation and Research, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 51, Rm. 6236, Silver Spring,