SUPPLEMENTARY INFORMATION:

Agenda: The meeting presentations will be heard, viewed, captioned, and recorded through an online teleconferencing platform. On July 14, 2020, the committee will discuss biologic license application (BLA) 761158, for belantamab mafodotin, submitted by GlaxoSmithKline Intellectual Property Development Ltd. England. The proposed indication (use) for this product is for the treatment of adults with relapsed or refractory multiple myeloma who have received at least four prior therapies including an anti-CD38 monoclonal antibody, a proteasome inhibitor, and an immunomodulatory agent.

FDA intends to make background material available to the public no later than 2 business days before the meeting. If FDA is unable to post the background material on its website prior to the meeting, the background material will be made publicly available on FDA's website at the time of the advisory committee meeting. Background material and the link to the online teleconference meeting room will be available at *https://www.fda.gov/AdvisoryCommittees/Calendar/default.htm.* Scroll down to the appropriate advisory committee meeting link. The meeting will include slide presentations with audio components to allow the presentation of materials in a manner that most closely resembles an in-person advisory committee meeting.

Procedure: Interested persons may present data, information, or views, orally or in writing, on issues pending before the committee. All electronic and written submissions submitted to the Docket (see **ADDRESSES**) on or before June 29, 2020, will be provided to the committee. Oral presentations from the public will be scheduled between approximately 11:30 a.m. and 12:30 p.m. Those individuals interested in making formal oral presentations should notify the contact person and submit a brief statement of the general nature of the evidence or arguments they wish to present, the names and addresses of proposed participants, and an indication of the approximate time requested to make their presentation on or before June 19, 2020. Time allotted for each presentation may be limited. If the number of registrants requesting to speak is greater than can be reasonably accommodated during the scheduled open public hearing session, FDA may conduct a lottery to determine the speakers for the scheduled open public hearing session. The contact person will notify interested persons regarding their request to speak by June 22, 2020.

For press inquiries, please contact the Office of Media Affairs at *fdaoma@fda.hhs.gov* or 301–796–4540.

FDA welcomes the attendance of the public at its advisory committee meetings and will make every effort to accommodate persons with disabilities. If you require accommodations due to a disability, please contact Yvette Waples (see **FOR FURTHER INFORMATION CONTACT**) at least 7 days in advance of the meeting.

FDA is committed to the orderly conduct of its advisory committee meetings. Please visit our website at *https://www.fda.gov/AdvisoryCommittees/AboutAdvisoryCommittees/ucm111462.htm* for procedures on public conduct during advisory committee meetings.

Notice of this meeting is given under the Federal Advisory Committee Act (5 U.S.C. app. 2).

Dated: June 15, 2020.

Lowell J. Schiller,

Principal Associate Commissioner for Policy.

[FR Doc. 2020–13345 Filed 6–19–20; 8:45 am]

BILLING CODE 4164–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2019–N–0163]

Hospira, Inc., et al.; Withdrawal of Approval of 12 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 February 12, 2019. The document announced the withdrawal of approval of 12 abbreviated new drug applications (ANDAs) from multiple applicants. The document erroneously included ANDA 077736 for Polyethylene Glycol 3350 Powder for Oral Solution, 17 grams/scoopful, held by Breckenridge Pharmaceutical, Inc. (Breckenridge). This document corrects that error.

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.

SUPPLEMENTARY INFORMATION: In the **Federal Register** of February 12, 2019 (84 FR 3467), in FR Doc. 2019–02032, the following correction is made:

1. On page 3467, in the table, the entry for ANDA 077736 is removed. The approval of ANDA 077736 was withdrawn effective November 2, 2018.

In the **Federal Register** of April 2, 2018 (83 FR 13994), FDA denied a hearing and issued an order withdrawing approval of multiple ANDAs for polyethylene glycol 3350, effective May 2, 2018. Breckenridge's ANDA 077736 was included in the April 2018 notice. In the **Federal Register** of July 30, 2018 (83 FR 36604), FDA subsequently published a notice granting a temporary stay of the effective date of the April 2018 notice, extending the withdrawal of approval of the ANDAs to November 2, 2018. Thus, the approval of ANDA 077736 was withdrawn effective November 2, 2018.

Dated: June 12, 2020.

Lowell J. Schiller,

Principal Associate Commissioner for Policy.

[FR Doc. 2020–13346 Filed 6–19–20; 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 Education Development Center for the Home Visiting Collaborative Improvement and Innovation Network 2.0 Cooperative Agreement

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

ACTION: Notice of a Supplemental Award to Education Development Center for Home Visiting Collaborative Improvement and Innovation Network 2.0 Cooperative Agreement.

SUMMARY: HRSA announces the award of a supplemental award of approximately $330,000 per year to the Education Development Center (EDC) for the Home Visiting Collaborative Improvement and Innovation Network 2.0 (HV CoIIN 2.0) for fiscal years (FY) 2020, 2021 and 2022. The supplement will allow the recipient to build a continuous quality improvement (CQI) health equity framework for the Maternal, Infant and Early Childhood Home Visiting Program (MIECHV).

FOR FURTHER INFORMATION CONTACT: Monique Fountain Hanna, Chief Medical Officer, Division Home Visiting and Early Childhood Systems, HRSA, 5600 Fishers Lane, Room 18N180,