

the Affordable Care Act and 45 CFR 155.220(a)(1) expands the role of agents/brokers by permitting them to enroll qualified individuals or small employers/employees in qualified health plans (QHPs) through the Exchanges, and assist individuals in applying for Advance Premium Tax Credits (APTCs) and Cost Sharing Reductions (CSRs). To participate as facilitators to enrollment, agents/brokers must register with the FFE, complete a training course covering eligibility and enrollment criteria for assisting in QHP enrollment, and sign agreements that formalize their understanding and commitment to adhere to the rules of the program. This requirement is specific to the FFE and does not automatically apply to State-based Exchanges (SBEs). This ICR serves as the formal request for renewal of the existing data collection. *Form Number:* CMS-10464 (OMB control number: 0938-1204); *Frequency:* Annually; *Affected Public:* Private Sector (Business or other for-profits) *Number of Respondents:* 64,000; *Number of Responses:* 64,000; *Total Annual Hours:* 15,360. (For questions regarding this collection contact Madeline Pellish at 301-492-4390).

Dated: July 26, 2022.

**William N. Parham, III,**

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

[FR Doc. 2022-16331 Filed 7-28-22; 8:45 am]

**BILLING CODE 4120-01-P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Food and Drug Administration

[Docket No. FDA-2022-N-0766]

#### Hospira, Inc., et al.; Withdrawal of Approval of 21 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** on May 20, 2022. The document announced the withdrawal of approval (as of June 21, 2022) of 21 abbreviated new drug applications (ANDAs) from multiple applicants. The document indicated that FDA was withdrawing approval of the following ANDAs after receiving withdrawal requests from Bionpharma Inc., 600 Alexander Rd., Suite 2-4B, Princeton, NJ 08540: ANDA 065301, Cefadroxil

Tablets, Equivalent to (EQ) 1 gram (g) base; ANDA 065307, Cefadroxil Oral Suspension, EQ 250 milligrams (mg) base/5 milliliters (mL) and EQ 500 mg base/5 mL; ANDA 065309, Cefadroxil Capsules, EQ 500 mg base; ANDA 065326, Cephalexin Oral Suspension, EQ 125 mg base/5 mL and EQ 250 mg base/5 mL; from Sunny Pharmtech Inc., 175 SW 166th Ave., Pembroke Pines, FL 33027; ANDA 203581, Glyburide Tablets, 1.25 mg, 2.5 mg, and 5 mg; and from Unicorn Pharmaceuticals, 5 Links Circle, Durham, NC, 27707; ANDA 204137, Omeprazole and Sodium Bicarbonate Capsules, 20 mg; 1.1 g. Before FDA withdrew the approval of these ANDAs, Bionpharma Inc., Sunny Pharmtech Inc., and Unicorn Pharmaceuticals informed FDA that they did not want the approval of the ANDAs withdrawn. Because Bionpharma Inc. timely requested that approval of ANDAs 065301, 065307, 065309, and 065326 not be withdrawn, the approvals are still in effect. Because Sunny Pharmtech Inc. timely requested that ANDA 203581 not be withdrawn, the approval is still in effect. Because Unicorn Pharmaceuticals timely requested that ANDA 204137 not be withdrawn, the approval is still in effect.

#### FOR FURTHER INFORMATION CONTACT:

Martha Nguyen, Center for Drug Evaluation and Research, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 75, Rm. 1676, Silver Spring, MD 20993-0002, 240-402-6980, [Martha.Nguyen@fda.hhs.gov](mailto:Martha.Nguyen@fda.hhs.gov).

**SUPPLEMENTARY INFORMATION:** In the **Federal Register** of Friday, May 20, 2022 (87 FR 30962), in FR Doc. 2022-10924, the following correction is made:

On page 30963, in the table, the entries for ANDAs 065301, 065307, 065309, 065326, 203581, and 204137 are removed.

Dated: July 25, 2022.

**Lauren K. Roth,**

*Associate Commissioner for Policy.*

[FR Doc. 2022-16281 Filed 7-28-22; 8:45 am]

**BILLING CODE 4164-01-P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Indian Health Service

#### Special Diabetes Program for Indians

*Announcement Type:* New.

*Funding Announcement Number:* HHS-2023-IHS-SDPI-0001.

*Assistance Listing (Catalog of Federal Domestic Assistance or CFDA) Number:* 93.237.

#### Key Dates

*Application Deadline Date:* October 7, 2022.

*Earliest Anticipated Start Date:* January 1, 2023.

#### I. Funding Opportunity Description

##### Statutory Authority

The Indian Health Service (IHS) is accepting applications for the Special Diabetes Program for Indians (SDPI—formerly Community-Directed SDPI). This program is authorized under the Snyder Act, 25 U.S.C. 13; the Transfer Act, 42 U.S.C. 2001(a); and section 330C of the Public Health Service Act, codified at 42 U.S.C. 254c-3. This program is described in the Assistance Listings located at <https://sam.gov/content/home/> (formerly known as the CFDA) under 93.237.

##### Background

Diabetes is a complex and costly chronic disease that requires tremendous long-term efforts to prevent and treat. Although diabetes is a nationwide public health problem, American Indian/Alaska Native (AI/AN) people are disproportionately affected. In 2019, 14.5 percent of AI/AN people aged 18 years or older had diagnosed diabetes, compared to 7.4 percent of non-Hispanic white people [CDC, 2021. <https://www.cdc.gov/diabetes/data/statistics-report/diagnosed-diabetes.html>]. In addition, AI/AN people have higher rates of diabetes-related morbidity and mortality than the general U.S. population [O'Connell, 2010 (<https://diabetesjournals.org/care/article/33/7/1463/39326/Racial-Disparities-in-Health-Status-A-comparison-of>); Cho, 2014 (<http://ajph.aphapublications.org/doi/full/10.2105/AJPH.2014.301968>)]. Strategies to address the prevention and treatment of diabetes in AI/AN communities are urgently needed.

In response to the burgeoning diabetes epidemic among AI/AN people, Congress established the SDPI through the Balanced Budget Act of 1997. SDPI is a \$150 million per year program that provides awards for diabetes treatment and prevention services. The IHS administers SDPI, with programmatic oversight provided by the IHS Division of Diabetes Treatment and Prevention (DDTP).

##### Purpose

The purpose of this program is to provide diabetes treatment and/or prevention activities and/or services (also referred to as “activities/services”) for AI/AN communities. Awardees will implement one SDPI Diabetes Best