upgrade UI Benefits and Tax Systems by SWAs using Federal funds.

(b) The standard designated in paragraph (a) of this section is effective March 21, 2014.

Eric M. Seleznow,

Acting Assistant Secretary, Employment and Training Administration.

[FR Doc. 2014–03496 Filed 2–18–14; 8:45 am]

BILLING CODE 4510–FN–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Parts 106 and 107

[Docket No. FDA–1995–N–0063 (formerly 95N–0309)]

RIN 0910–AF27

Current Good Manufacturing Practices, Quality Control Procedures, Quality Factors, Notification Requirements, and Records and Reports, for Infant Formula; Correction

AGENCY: Food and Drug Administration, HHS.

ACTION: Interim final rule; request for comments; correction.

SUMMARY: The Food and Drug Administration (FDA) is correcting a document that appeared in the **Federal Register** of February 10, 2014. The document revised our infant formula regulations to establish requirements for current good manufacturing practices, including audits; to establish requirements for quality factors; and to amend FDA's quality control procedures, notification, and record and reporting requirements for infant formula. FDA took the action to improve the protection of infants who consume infant formula products. The document was published with an incorrect docket number. This document corrects that error.

DATES: *Effective Date:* This correction is effective February 19, 2014.

FOR FURTHER INFORMATION CONTACT: Joyce Strong, Office of Policy, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 32, rm. 3208, Silver Spring, MD 20993, 301–796–9148.

SUPPLEMENTARY INFORMATION: In FR Doc. 2014–02148, appearing on page 7934 in the **Federal Register** of February 10, 2014 (79 FR 7934), the following corrections are made:

1. On page 7934, ''FDA–1995–N–0036'' is corrected to read ''FDA–1995–N–0063'' each time it appears.

2. On page 8055, in the second column, ''FDA–1995–N–0036'' is corrected to read ''FDA–1995–N–0063''.

3. On page 8058, in the third column, ''FDA–1995–N–0036'' is corrected to read ''FDA–1995–N–0063''.

Dated: February 13, 2014.

Leslie Kux,

Assistant Commissioner for Policy.

[FR Doc. 2014–03588 Filed 2–18–14; 8:45 am]

BILLING CODE 4160–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 800

[Docket No. FDA–1977–N–0222]

Administrative Detention; Corrections

AGENCY: Food and Drug Administration, HHS.

ACTION: Final rule; correcting amendment.

SUMMARY: The Food and Drug Administration (FDA) published a document in the **Federal Register** on Friday, March 9, 1979 (44 FR 13239). The document established administrative detention procedures for devices intended for human use believed to be adulterated or misbranded. The document was published with a citation in the first column on page 13240 that subsequently was changed by the Nutrition Labeling and Education Act Amendments of 1993. In addition, the document was published with one typographical error in the first column on page 13241. This document corrects these errors.

DATES: This correction is effective February 19, 2014.

FOR FURTHER INFORMATION CONTACT: Jan B. Welch, Center for Devices and Radiological Health, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 66, Rm. 3412, 301–796–5776, FAX: 301–847–8136, *jan.welch@fda.hhs.gov.*

SUPPLEMENTARY INFORMATION: FDA is correcting a final rule that appeared in the **Federal Register** on Friday, March 9, 1979 (44 FR 13239). The final rule established administrative detention procedures for devices intended for human use believed to be adulterated or misbranded. The document was published with a citation to section 201(y) of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 321(y)) (the FD&C Act) in the first column on page 13240 (§ 800.55(g)(1) (21 CFR 800.55(g)(1)) that subsequently was changed to section 201(x) of the FD&C Act by section 3(b) of the Nutrition Labeling and Education Act Amendments of 1993 (Pub. L. 103–80). In addition, the document was published with one typographical error in the first column on page 13241 (§ 800.55(k)(1)) in which the word ''is'' should have been the word ''in''. This document updates the statutory reference in § 800.55(g)(1) and corrects the typographical error in § 800.55(k)(1).

Publication of this rule constitutes final action under the Administrative Procedure Act (5 U.S.C. 553). This amendment to the regulations provides only a technical change and corrects a nonsubstantive error. FDA therefore, for good cause, finds under 5 U.S.C. 553(b)(3)(B) that notice and public comment are unnecessary, and under 5 U.S.C. 553(d)(3) that the rule can become effective upon publication.

FDA has determined under 21 CFR 25.30(i) that this final rule is of a type that, as a class, does not individually or cumulatively have a significant effect on the human environment. Therefore, neither an environmental assessment nor an environmental impact statement is required.

This final rule contains information collection provisions that are subject to review by the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995 (44 U.S.C. 3501–3520). The collections of information in § 800.55 have been approved under OMB control number 0910–0114, which expires April 30, 2016.

List of Subjects in 21 CFR Part 800

Administrative practice and procedure; Medical devices; Ophthalmic goods and services; Packaging and containers; Reporting and recordkeeping requirements.

Therefore, under the Federal Food, Drug, and Cosmetic Act and under authority delegated to the Commissioner of Food and Drugs, 21 CFR Part 800 is amended as follows:

PART 800—GENERAL

■ 1. The authority citation for 21 CFR Part 800 continues to read as follows:

Authority: 21 U.S.C. 321, 334, 351, 352, 355, 360e, 360i, 360k, 361, 362, 371.

■ 2. In § 800.55, revise paragraph (g)(1) and the first sentence of paragraph (k) to read as follows:

§ 800.55 Administrative detention.

* * * * *

(g) *Appeal of a detention order.* (1) A person who would be entitled to claim

the devices, if seized, may appeal a detention order. Any appeal shall be submitted in writing to the FDA District Director in whose district the devices are located within 5 working days of receipt of a detention order. If the appeal includes a request for an informal hearing, as defined in section 201(x) of the act, the appellant shall request either that a hearing be held within 5 working days after the appeal is filed or that the hearing be held at a later date, which shall not be later than 20 calendar days after receipt of a detention order.

* * * * *

(k) *Recordkeeping requirements.* (1) After issuance of a detention order under paragraph (d) of this section, the owner, operator, or agent in charge of any factory, warehouse, other establishment, or consulting laboratory where detained devices are manufactured, processed, packed, or held shall have, or establish, and maintain adequate records relating to how the detained devices may have become adulterated or misbranded, records on any distribution of the devices before and after the detention period, records on the correlation of any in-process detained devices that are put in final form under paragraph (h) of this section to the completed devices, records of any changes in, or processing of, the devices permitted under the detention order, and records of any other movement under paragraph (h) of this section. * * *

* * * * *

Dated: February 12, 2014.

Leslie Kux,

Assistant Commissioner for Policy.

[FR Doc. 2014–03582 Filed 2–18–14; 8:45 am]

BILLING CODE 4160–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 806

[Docket No. FDA–2014–N–0011]

Medical Devices; Reports of Corrections and Removals; Technical Amendment

AGENCY: Food and Drug Administration, HHS.

ACTION: Final rule; technical amendment.

SUMMARY: The Food and Drug Administration (FDA) is amending its regulation regarding reports of corrections to and removals of medical devices to address a minor change as a result of the enactment of the Food and Drug Administration Amendments Act of 2007 (FDAAA). This action is technical in nature and is intended to provide accuracy to the Agency's regulation.

DATES: This rule is effective February 19, 2014.

FOR FURTHER INFORMATION CONTACT: Deborah Yoder, Office of Compliance, Center for Devices and Radiological Health, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 66, Rm. 2676, Silver Spring, MD 20993–0002, 301–796–6109, *Deborah.Yoder@fda.hhs.gov*.

SUPPLEMENTARY INFORMATION: Section 806.1(a) (21 CFR 806.1(a)) refers to a subsection of the Federal Food, Drug, and Cosmetic Act that was redesignated as a result of FDAAA (Pub. L. 110–85). FDA is amending § 806.1(a) to update the obsolete reference.

FDA is publishing the document as a final rule under the Administrative Procedures Act (5 U.S.C. 551, *et seq.*). FDA has determined that good cause exists to dispense with prior notice and public comment under 5 U.S.C. 553(b)(3)(B) and 21 CFR 10.40(e)(1) since such notice and comment are unnecessary because this amendment to the regulation provides only a technical change to update an obsolete citation. In addition, FDA finds good cause to provide for this regulation to be effective immediately upon publication under 5 U.S.C. 553(d)(3).

FDA has determined under 21 CFR 25.30(i) that this final rule is of a type that does not individually or cumulatively have a significant effect on the human environment. Therefore, neither an environmental assessment nor an environmental impact statement is required.

This final rule contains information collection provisions that are subject to review by the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995 (44 U.S.C. 3501–3520). The collections of information in 21 CFR part 806 have been approved under OMB control number 0910–0359, which expires May 31, 2014.

List of Subjects in 21 CFR Part 806

Imports, Medical devices, Reporting and recordkeeping requirements.

Therefore, under the Federal Food, Drug, and Cosmetic Act and under authority delegated to the Commissioner of Food and Drugs, 21 CFR part 806 is amended as follows:

PART 806—MEDICAL DEVICES; REPORTS OF CORRECTIONS AND REMOVALS

■ 1. The authority citation for 21 CFR part 806 continues to read as follows:

Authority: 21 U.S.C. 352, 360, 360i, 360j, 371, 374.

§ 806.1 [Amended]

■ 2. Amend § 806.1(a) by removing ''section 519(f)'' and adding in its place ''section 519(g)''.

Dated: February 12, 2014.

Leslie Kux,

Assistant Commissioner for Policy.

[FR Doc. 2014–03581 Filed 2–18–14; 8:45 am]

BILLING CODE 4160–01–P

NATIONAL FOUNDATION ON THE ARTS AND THE HUMANITIES

National Endowment for the Humanities

45 CFR Part 1171

RIN 3136–AA32

Public Access to NEH Records Under the Freedom of Information Act

AGENCY: National Endowment for the Humanities.

ACTION: Final rule.

SUMMARY: The National Endowment for the Humanities (NEH) is unilaterally rescinding its joint Freedom of Information Act (FOIA) regulations with the National Endowment for the Arts (NEA) and the Institute of Museum and Library Services (IMLS), and issuing its own FOIA regulations. This final rule provides the NEH's procedures for disclosure of its records, as required by the FOIA, 5 U.S.C. 552, as amended. These regulations also provide the procedures for disclosing records of the Federal Council on the Arts and the Humanities (FCAH), an agency for which NEH provides legal counsel.

DATES: The final rule will be effective March 21, 2014.

FOR FURTHER INFORMATION CONTACT: Mara Campbell, Office of the General Counsel, National Endowment for the Humanities, at 202–606–8322, or *mcampbell@neh.gov*.

SUPPLEMENTARY INFORMATION: The NEH along with the NEA, the IMLS, and the FCAH make up the National Foundation on the Arts and Humanities (Foundation). The Foundation was established by the National Foundation on the Arts and Humanities Act of 1965, 20 U.S.C. 951 *et seq*. The NEH along