effective September 15, 2012, is amended as follows:

Paragraph 6010 VOR Federal Airways.

(a) Domestic VOR Federal airways.

* * * * *

V–233

From Spinner, IL; INT Spinner 061° and Roberts, IL, 233° radials; Roberts; Knox, IN; Goshen, IN; Litchfield, MI; Lansing, MI; Mount Pleasant, MI; INT Mount Pleasant 351° and Gaylord, MI, 207° radials; Gaylord; to Pellston, MI.

* * * * *

Issued in Washington, DC, March 13, 2013.

Gary A. Norek,

Manager, Airspace Policy and ATC Procedures Group.

[FR Doc. 2013–06365 Filed 3–25–13; 8:45 am]

BILLING CODE 4910–13–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Parts 814, 822, 862, 864, 866, 868, 870, 872, 874, 876, 878, 880, 882, 884, 886, 888, 890, and 892

[Docket No. FDA–2013–N–0011]

Medical Devices; Technical Amendment

AGENCY: Food and Drug Administration, HHS.

ACTION: Final rule; technical amendment.

SUMMARY: The Food and Drug Administration (FDA) is amending certain medical device regulations to correct minor errors in the Code of Federal Regulations (CFR). This action is editorial in nature and is intended to provide accuracy and clarity to the Agency's regulations.

DATES: This rule is effective March 26, 2013.

FOR FURTHER INFORMATION CONTACT: Abigail Corbin, Center for Devices and Radiological Health, Food and Drug Administration, 10903 New Hampshire Ave. Bldg. 66, rm. 4430, Silver Spring, MD 20993–0002, 301–796–9142.

SUPPLEMENTARY INFORMATION: FDA is amending certain regulations in 21 CFR parts 814, 822, 862, 864, 866, 868, 870, 872, 874, 876, 878, 880, 882, 884, 886, 888, 890, and 892. This action corrects minor spelling errors and outdated Web site addresses affecting certain regulations regarding medical devices.

Publication of this document constitutes final action of these changes under the Administrative Procedure Act (5 U.S.C. 553). These amendments are merely correcting nonsubstantive errors. FDA therefore, for good cause, finds under 5 U.S.C. 553(b)(3)(B) and (d)(3) that notice and public comment are unnecessary.

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. In addition, FDA has determined that this final rule contains no collection of information. Therefore, clearance by the Office of Management and Budget under the Paperwork Reduction Act of 1995 is not required.

List of Subjects

21 CFR Part 814

Administrative practice and procedure, Confidential business information, Medical devices, Medical research, Reporting and recordkeeping requirements.

21 CFR Part 822

Medical devices, Reporting and recordkeeping requirements.

21 CFR Part 862

Medical devices.

21 CFR Part 864

Blood, Medical devices, Packaging and containers.

21 CFR Part 866

Biologics, Laboratories, Medical devices.

21 CFR Part 868, 870, 872, 874, 876, 878, and 880

Medical devices.

21 CFR Part 882

Medical devices, Neurological devices.

21 CFR Part 884

Medical devices.

21 CFR Part 886

Medical devices, Ophthalmic goods and services.

21 CFR Part 888

Medical devices.

21 CFR Part 890

Medical devices, Physical medicine devices.

21 CFR Part 892

Medical devices, Radiation protection, X-rays.

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

§§ 814.20, 822.7, 822.15, 862.1, 864.1, 866.1, 868.1, 870.1, 872.1, 874.1, 876.1, 878.1, 880.1, 882.1, 884.1, 886.1, 888.1, 890.1, and 892.1 [Amended]

1. In the table below, for each section indicated in the left column, remove the Web address indicated in the middle column from wherever it appears in the section, and add the Web address indicated in the right column:

Section	Remove	Add
814.20	*http://www.fda.gov/cdrh/devadvice/pma/*	*http://www.fda.gov/MedicalDevices/DeviceRegulationandGuidance/HowtoMarketYourDevice/PremarketSubmissions/PremarketApprovalPMA/default.htm.*
822.7	*http://www.fda.gov/cdrh/ombudsman/dispute.html.*	*http://www.fda.gov/AboutFDA/CentersOffices/OfficeofMedicalProductsandTobacco/CDRH/CDRHOmbudsman/default.htm.*
822.15	www.fda.gov/cdrh/ombudsman/	*http://www.fda.gov/AboutFDA/CentersOffices/OfficeofMedicalProductsandTobacco/CDRH/CDRHOmbudsman/default.htm.*
862.1, 864.1, 866.1, 868.1, 870.1, 872.1, 874.1, 876.1, 878.1, 880.1, 882.1, 884.1, 886.1, 888.1, 890.1, and 892.1.	*http://www.fda.gov/cdrh/guidance.html*	*http://www.fda.gov/MedicalDevices/DeviceRegulationandGuidance/GuidanceDocuments/default.htm.*

PART 870—CARDIOVASCULAR DEVICES

■ 2. The authority citation for 21 CFR part 870 continues to read as follows:

Authority: 21 U.S.C. 351, 360, 360c, 360e, 360j, 371.

■ 3. Amend § 870.3600 by revising the second sentence in paragraph (a) to read as follows:

§ 870.3600 External pacemaker pulse generator.

(a) *Identification.* * * * This device, which is used outside the body, is used as a temporary substitute for the heart's intrinsic pacing system until a permanent pacemaker can be implanted, or to control irregular heartbeats in patients following cardiac surgery or a myocardial infarction. * * *

* * * * *

■ 4. Amend § 870.5300 by revising the section heading to read as follows:

§ 870.5300 DC-defibrillator (including paddles).

* * * * *

PART 882—NEUROLOGICAL DEVICES

■ 5. The authority citation for 21 CFR part 882 continues to read as follows:

Authority: 21 U.S.C. 351, 360, 360c, 360e, 360j, 371.

■ 6. Amend § 882.5870 by revising the second sentence in paragraph (a) to read as follows:

§ 882.5870 Implanted peripheral nerve stimulator for pain relief.

(a) *Identification.* * * * The stimulator consists of an implanted receiver with electrodes that are placed around a peripheral nerve and an external transmitter for transmitting the stimulating pulses across the patient's skin to the implanted receiver.

* * * * *

PART 886—OPHTHALMIC DEVICES

■ 7. The authority citation for 21 CFR part 886 continues to read as follows:

Authority: 21 U.S.C. 351, 360, 360c, 360e, 360j, 371.

■ 8. Amend § 886.1120 by revising the section heading to read as follows:

§ 886.1120 Ophthalmic camera.

* * * * *

Dated: March 20, 2013.

Leslie Kux,

Assistant Commissioner for Policy.

[FR Doc. 2013–06826 Filed 3–25–13; 8:45 am]

BILLING CODE 4160–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 1005

[Docket No. FDA–2007–N–0091; (formerly 2007N–0104)]

Service of Process on Manufacturers; Manufacturers Importing Electronic Products Into the United States; Agent Designation; Change of Address

AGENCY: Food and Drug Administration, HHS.

ACTION: Final rule; technical amendment.

SUMMARY: The Food and Drug Administration (FDA) is amending a final rule that appeared in the **Federal Register** of April 9, 2007 (72 FR 17397 at 17401) to reflect changes to the Center for Devices and Radiological Health's address. This action is editorial in nature and is intended to improve the accuracy of the Agency's regulations.

DATES: This rule is effective March 26, 2013

FOR FURTHER INFORMATION CONTACT: Prince P. Kangoma, Center for Devices and Radiological Health, Food and Drug Administration, 10903 New Hampshire Ave. Bldg. 66, rm. G628B, 301–796–6627, FAX: 301–595–7850, email: *Prince.Kangoma@fda.hhs.gov.*

SUPPLEMENTARY INFORMATION: FDA is making technical amendments in the Agency's regulations under 21 CFR 1005.25(b) as a result of an office relocation. The former address was 9200 Corporate Blvd., Rockville, MD 20850. The new address is 10903 New Hampshire Ave., Silver Spring, MD 20993–0002. Designation of agent by manufacturers of electronic products offering such products for importation into the United States must be addressed to the Center for Devices and Radiological Health, Document Mail Center—WO66–G609. Publication of this document constitutes final action of these changes under the Administrative Procedures Act (5 U.S.C. 553). Notice and public procedures are unnecessary because FDA is merely updating nonsubstantive content.

List of Subjects in 21 CFR Part 1005

Administrative practice and procedure; Electronic product; Imports; Radiation protection; Surety bonds.

Therefore, under the Federal Food, Drug, and Cosmetic Act and under authority delegated to the Commissioner of Food and Drugs, 21 CFR part 1005 is amended as follows:

PART 1005—IMPORTATION OF ELECTRONIC PRODUCTS

■ 1. The authority citation for 21 CFR part 1005 continues to read as follows:

Authority: 21 U.S.C. 360ii, 360mm.

§ 1005.25 [Amended]

■ 2. Section 1005.25(b) is amended by removing ''Center for Devices and Radiological Health (HFZ–240), 9200 Corporate Blvd., Rockville, MD 20850'' and by adding in its place ''Center for Devices and Radiological Health, 10903 New Hampshire Ave., Document Mail Center—WO66–G609, Silver Spring, MD 20993–0002''.

Dated: March 20, 2013.

Leslie Kux,

Assistant Commissioner for Policy.

[FR Doc. 2013–06864 Filed 3–25–13; 8:45 am]

BILLING CODE 4160–01–P

DEPARTMENT OF HOUSING AND URBAN DEVELOPMENT

24 CFR Part 970

Demolition or Disposition of Public Housing Projects

CFR Correction

■ In Title 24 of the Code of Federal Regulations, Parts 700 to 1699, revised as of April 1, 2012, on page 490, in § 970.27, redesignate paragraph (c) introductory text as paragraph (b).

[FR Doc. 2013–07091 Filed 3–25–13; 8:45 am]

BILLING CODE 1505–01–D

DEPARTMENT OF THE TREASURY

Internal Revenue Service

26 CFR Part 1

Section 482: Methods To Determine Taxable Income in Connection With a Cost Sharing Arrangement

CFR Correction

■ In Title 26 of the Code of Federal Regulations, Part 1 (§§ 1.441 to 1.500), revised as of April 1, 2012, on page 604, in § 1.482–1, in paragraph (c)(1), before the last sentence, reinstate the following sentence:

§ 1.482–1 Allocation of income and deductions among taxpayers.

* * * * *