Food and Drug Administration, HHS

§ 14.122

Food, Drug, and Cosmetic Act and, as required, any other product for which the Food and Drug Administration has regulatory responsibility, and makes appropriate recommendations to the Commissioner of Food and Drugs.

(d) Center for Devices and Radiological Health—

(1) Medical Devices Advisory Committee.

(i) Date established: October 27, 1990.

(ii) Function: Reviews and evaluates data on the safety and effectiveness of marketed and investigational devices and makes recommendations for their regulation.

(2) Device Good Manufacturing Practice Advisory Committee.

(i) Date established: May 17, 1987.

(ii) Function: Reviews proposed regulations for good manufacturing practices governing the methods used in, and the facilities and controls used for, the manufacture, packing, storage, and installation of devices, and makes recommendations on the feasibility and reasonableness of the proposed regulations.

(3) Technical Electronic Product Radiation Safety Standards Committee.

(i) Date established: October 18, 1968.


(4) National Mammography Quality Assurance Advisory Committee.

(i) Date established: July 6, 1993.

(ii) Function: Advises on developing appropriate quality standards and regulations for the use of mammography facilities.

(e) National Center for Toxilogical Research—Science Advisory Board.

(i) Date established: June 2, 1973.

(2) Function: Advises on establishment and implementation of a research program that will assist the Commissioner of Food and Drugs to fulfill regulatory responsibilities.

(f) Center for Food Safety and Applied Nutrition—Food Advisory Committee.

(i) Date established: December 15, 1991.

(ii) Function: The committee provides advice on emerging food safety, food science, and nutrition issues that FDA considers of primary importance in the next decade.

[54 FR 9036, Mar. 3, 1989]

EDITORIAL NOTE: For FEDERAL REGISTER citations affecting §14.100, see the List of CFR Sections Affected, which appears in the Finding Aids section of the printed volume and at www.fdsys.gov.

EFFECTIVE DATE NOTE: At 79 FR 20095, Apr. 11, 2014, §14.100 was amended by revising the paragraph (c)(9) heading and paragraph (c)(9)(ii), effective Apr. 11, 2014. For the convenience of the user, the revised text is set forth as follows:

§ 14.100 List of standing advisory committees.

* * * * *

(c) * * *

(9) Bone, Reproductive and Urologic Drugs Advisory Committee.

(i) * * *

(ii) Function: Advises the Commissioner or designee in discharging responsibilities as they relate to helping to ensure safe and effective drugs for human use and, as required, any other product for which the Food and Drug Administration has regulatory responsibility.

* * * * *

Subpart G—Technical Electronic Products Radiation Safety Standards Committee

§ 14.120 Establishment of the Technical Electronic Product Radiation Safety Standards Committee (TEPRSSC).

The Technical Electronic Product Radiation Safety Standards Committee (TEPRSSC), consisting of 15 members, is established in accordance with the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 360kk(f)(1)(A)) to provide consultation before the Commissioner prescribes any performance standard for an electronic product.

[44 FR 22351, Apr. 13, 1979, as amended at 78 FR 17087, Mar. 20, 2013]

§ 14.122 Functions of TEPRSSC.

(a) In performing its function of advising the Commissioner, TEPRSSC—

(1) May propose electronic product radiation safety standards to the Commissioner for consideration;