

otherwise not permitted by the Environmental Protection Agency.

**(b) Timing of report**

A laboratory shall submit the report required under subsection (a) to the Secretary, the Secretary of Health and Human Services, and the owner of such food as soon as practicable after the completion of the analysis of such food.

**(c) Guidelines**

The Secretary shall adopt standardized reporting guidelines to be applied to laboratories under this section and shall provide such guidelines to laboratories accredited under this chapter, as well as other sources of information regarding applicable pesticide chemical tolerances.

(Pub. L. 101-624, title XIII, §1326, Nov. 28, 1990, 104 Stat. 3565.)

**§ 138f. Fees**

**(a) In general**

At the time that an application for accreditation is received by the Secretary and annually thereafter, a laboratory seeking accreditation by the Secretary under the authority of this chapter, the Federal Meat Inspection Act (21 U.S.C. 601 et seq.), or the Poultry Products Inspection Act (21 U.S.C. 451 et seq.) shall pay to the Secretary a nonrefundable accreditation fee. All fees collected by the Secretary shall be credited to the account from which the expenses of the laboratory accreditation program are paid and, subject to subsection (e), shall be available immediately and remain available until expended to pay the expenses of the laboratory accreditation program.

**(b) Amount of fee**

The fee required under this section shall be established by the Secretary in an amount that will offset the cost of the laboratory accreditation programs administered by the Secretary under the statutory authorities set forth in subsection (a).

**(c) Reimbursement of expenses**

Each laboratory that is accredited under a statutory authority set forth in subsection (a) or that has applied for accreditation under such authority shall reimburse the Secretary for reasonable travel and other expenses necessary to perform onsite inspections of the laboratory.

**(d) Adjustment of fees**

The Secretary may, on an annual basis, adjust the fees imposed under this section as necessary to support the full costs of the laboratory accreditation programs carried out under the statutory authorities set forth in subsection (a).

**(e) Appropriations prerequisite**

No fees collected under this section may be used to offset the cost of laboratory accreditation without appropriations made under subsection (f).

**(f) Authorization of appropriations**

There are authorized to be appropriated each fiscal year such sums as may be necessary for laboratory accreditation services under this section.

(Pub. L. 101-624, title XIII, §1327, Nov. 28, 1990, 104 Stat. 3565; Pub. L. 102-237, title X, §1017, Dec. 13, 1991, 105 Stat. 1904.)

REFERENCES IN TEXT

The Federal Meat Inspection Act, referred to in subsec. (a), is titles I to V of act Mar. 4, 1907, ch. 2907, as added Pub. L. 90-201, Dec. 15, 1967, 81 Stat. 584, and Pub. L. 110-246, title XI, §11015(a), June 18, 2008, 122 Stat. 2124, which are classified generally to subchapters I to IV-A (§601 et seq.) of chapter 12 of Title 21, Food and Drugs. For complete classification of this Act to the Code, see Short Title note set out under section 601 of Title 21 and Tables.

The Poultry Products Inspection Act, referred to in subsec. (a), is Pub. L. 85-172, Aug. 28, 1957, 71 Stat. 441, as amended, which is classified generally to chapter 10 (§451 et seq.) of Title 21. For complete classification of this Act to the Code, see Short Title note set out under section 451 of Title 21 and Tables.

AMENDMENTS

1991—Pub. L. 102-237 amended section generally, in subsec. (a), inserting provisions relating to Federal Meat Inspection Act and Poultry Products Inspection Act and provisions relating to crediting and availability of fees, in subsec. (b), substituting provisions relating to fee under this section for provisions relating to fee under subsec. (a) of this section, and provisions relating to laboratory accreditation programs administered by Secretary under statutory authorities set forth in subsec. (a) of this section for provisions relating to program established under this chapter, in subsec. (c), substituting provisions relating to statutory authority set forth in subsec. (a) of this section for provisions relating to this chapter, in subsec. (d), substituting provisions relating to laboratory accreditation programs under statutory authority set forth in subsec. (a) of this section for provisions relating to program established under this chapter, and adding subsecs. (e) and (f).

**§ 138g. Public disclosure**

The results of the evaluations of laboratories conducted by the Secretary under this chapter shall be made available to the Secretary of Health and Human Services and to the public on request.

(Pub. L. 101-624, title XIII, §1328, Nov. 28, 1990, 104 Stat. 3565.)

**§ 138h. Regulations**

The Secretary shall promulgate regulations to carry out this chapter.

(Pub. L. 101-624, title XIII, §1329, Nov. 28, 1990, 104 Stat. 3565.)

**§ 138i. Effect of other laws**

Nothing in this chapter shall alter the authority of the Secretary of Health and Human Services under the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 301 et seq.).

(Pub. L. 101-624, title XIII, §1330, Nov. 28, 1990, 104 Stat. 3565.)

REFERENCES IN TEXT

The Federal Food, Drug, and Cosmetic Act, referred to in text, is act June 25, 1938, ch. 675, 52 Stat. 1040, as amended, which is classified generally to chapter 9 (§301 et seq.) of Title 21, Food and Drugs. For complete classification of this Act to the Code, see section 301 of Title 21 and Tables.