

longer responsible for the day-to-day operations of the facility.

(Approved by the Office of Management and Budget under control number 0579-0258)

[62 FR 27934, May 22, 1997, as amended at 62 FR 54758, Oct. 22, 1997; 63 FR 32119, June 12, 1998; 68 FR 62226, Nov. 3, 2003; 74 FR 14709, Apr. 1, 2009]

§ 71.21 Tissue and blood testing at slaughter.

(a) Any person moving livestock or poultry interstate for slaughter or rendering may only move the animals to a slaughtering establishment or a rendering establishment that has been listed by the Administrator⁸ for the purposes of this part. Livestock or poultry may not be removed from the premises of a slaughtering establishment or a rendering establishment listed by the Administrator except under a permit issued by APHIS, and in accordance with applicable FSIS regulations in this title. A slaughtering establishment or rendering establishment may receive livestock or poultry in interstate commerce only if the establishment has been listed by the Administrator. The Administrator may list a slaughtering establishment or a rendering establishment after determining that collecting samples for testing from the establishment is not currently necessary for the purposes of APHIS disease surveillance programs and the establishment has agreed to allow testing and to provide the access and facilities required by this section upon future APHIS notification that testing is required at the establishment. The Administrator will list a slaughtering or rendering establishment after determining that it meets the following facility and access requirements:

(1) The establishment provides space and equipment in accordance with paragraph (b) of this section⁹ within

⁸A list of these slaughtering or rendering establishments may be obtained by writing to National Center for Animal Health Programs, VS, APHIS, 4700 River Road Unit 43, Riverdale, MD 20737-1231.

⁹FSIS also has equipment and space requirements for official establishments at § 307.2(c) of this title.

their facility for blood and tissue sample collection;

(2) The establishment allows APHIS, FSIS, or APHIS contractors to take blood and tissue samples from all livestock or poultry at the facility without cost to the United States, and specifically allows these personnel access to the processing line to collect samples; and

(3) The establishment allows APHIS, FSIS, or APHIS contractors to record the identification of individual animals and retain any external or internal identification devices.

(b) The establishment must provide office and sample collection space, including necessary furnishings, light, heat, and janitor service, rent free, for the use by APHIS, FSIS, or APHIS contractors collecting samples for blood and tissue testing under this section. The Administrator will inform each establishment of the exact amount and type of space required, taking into account whether APHIS will be conducting complete tests at the facility, or only collecting samples and sending them elsewhere for testing. At the discretion of the Administrator, small plants need not furnish facilities as prescribed in this section if adequate facilities exist in a nearby convenient location. In granting or denying listing of an establishment, the Administrator will consider whether the space at the facility:

(1) Is conveniently located, properly ventilated, and provided with lockers suitable for the protection and storage of supplies;

(2) Has sufficient light to be adequate for proper conduct of sample collection and processing;

(3) Includes racks, receptacles, or other suitable devices for retaining such parts as the head, glands, and viscera, and all parts and blood to be collected, until after the post-mortem examination is completed;

(4) Includes tables, benches, and other equipment on which sample collection and processing are to be performed, of such design, material, and construction as to enable sample collection and processing in a safe, ready, efficient, and clean manner;

(5) Has adequate arrangements, including liquid soap and cleansers, for

cleansing and disinfecting hands, dissection tools, floors, and other articles and places that may be contaminated by diseased carcasses or otherwise; and

(6) Has adequate facilities, including denaturing materials, for the proper disposal in accordance with this chapter of tissue, blood, and other waste generated during test sample collection.

(c) The Administrator will give the operator of the establishment actual notice that APHIS, FSIS, or an APHIS contractor will be taking blood and/or tissue samples at the establishment. The Administrator may give the operator of the establishment notice in any form or by any means that the Administrator reasonably believes will reach the operator of the establishment prior to the start of sample collection.

(1) The notice will include the anticipated date and time sample collection will begin. The notice will also include the anticipated ending date and time.

(2) The Administrator will give the operator of the establishment as much advance notice as possible. However, the actual amount of notice will depend on the specific situation.

(d) *Denial and withdrawal of listing.* The Administrator may deny or withdraw the listing of an establishment upon a determination that the establishment is not in compliance with the requirements of this section.

(1) In the case of a denial, the operator of the establishment will be informed of the reasons for the denial and may appeal the decision in writing to the Administrator within 10 days after receiving notification of the denial. The appeal must include all of the facts and reasons upon which the person relies to show that the establishment was wrongfully denied listing. The Administrator will grant or deny the appeal in writing as promptly as circumstances permit, stating the reason for his or her decision. If there is a conflict as to any material fact, a hearing will be held to resolve the conflict. Rules of practice concerning the hearing will be adopted by the Administrator.

(2) In the case of withdrawal, before such action is taken, the operator of the establishment will be informed of the reasons for the proposed with-

drawal. The operator of the establishment may appeal the proposed withdrawal in writing to the Administrator within 10 days after being informed of the reasons for the proposed withdrawal. The appeal must include all of the facts and reasons upon which the person relies to show that the reasons for the proposed withdrawal are incorrect or do not support the withdrawal of the listing. The Administrator will grant or deny the appeal in writing as promptly as circumstances permit, stating the reason for his or her decision. If there is a conflict as to any material fact, a hearing will be held to resolve the conflict. Rules of practice concerning the hearing will be adopted by the Administrator. However, withdrawal shall become effective pending final determination in the proceeding when the Administrator determines that such action is necessary to protect the public health, interest, or safety. Such withdrawal shall be effective upon oral or written notification, whichever is earlier, to the operator of the establishment. In the event of oral notification, written confirmation shall be given as promptly as circumstances allow. This withdrawal shall continue in effect pending the completion of the proceeding, and any judicial review thereof, unless otherwise ordered by the Administrator.

(Approved by the Office of Management and Budget under control number 0579–0212)

[69 FR 10150, Mar. 4, 2004]

§71.22 Removal and loss of official identification devices.

Official identification devices are intended to provide permanent identification of livestock and to ensure the ability to find the source of animal disease outbreaks. Removal of these devices, including devices applied to imported animals in their countries of origin and recognized by the Administrator as official, is prohibited except at the time of slaughter. If an official identification device is lost and it is necessary to retag an animal with a new official number, every effort should be made to correlate the new official number with the previous official number of the animal. If an official identification device applied to an imported animal in its country of origin