"significant rule" under DOT Regulatory Policies and Procedures (44 FR 11034, February 26, 1979); and (3) will not have a significant economic impact, positive or negative, on a substantial number of small entities under the criteria of the Regulatory Flexibility Act. A final evaluation has been prepared for this action and it is contained in the Rules Docket. A copy of it may be obtained from the Rules Docket at the location provided under the caption **ADDRESSES**.

List of Subjects in 14 CFR Part 39

Air transportation, Aircraft, Aviation safety, Incorporation by reference, Safety.

Adoption of the Amendment

Accordingly, pursuant to the authority delegated to me by the Administrator, the Federal Aviation Administration amends part 39 of the Federal Aviation Regulations (14 CFR part 39) as follows:

PART 39—AIRWORTHINESS DIRECTIVES

1. The authority citation for part 39 continues to read as follows:

Authority: 49 U.S.C. 106(g), 40113, 44701.

§39.13 [Amended]

2. Section 39.13 is amended by adding a new airworthiness directive to read as follows:

2000–16–06 Bell Helicopter Textron

Canada: Amendment 39–11860. Docket No. 99–SW–84–AD.

Applicability: Model 430 helicopters, serial numbers 49001 through 49018, 49020 through 49043, and 49045 through 49051, certificated in any category.

Note 1: This AD applies to each helicopter identified in the preceding applicability provision, regardless of whether it has been otherwise modified, altered, or repaired in the area subject to the requirements of this AD. For helicopters that have been modified, altered, or repaired so that the performance of the requirements of this AD is affected, the owner/operator must request approval for an alternative method of compliance in accordance with paragraph (c) of this AD. The request should include an assessment of the effect of the modification, alteration, or repair on the unsafe condition addressed by this AD; and if the unsafe condition has not been eliminated, the request should include specific proposed actions to address it.

Compliance: Required within 150 hours time-in-service after the effective date of this AD, unless accomplished previously.

To prevent a jammed flight control and subsequent loss of control of the helicopter, accomplish the following:

(a) Remove the arm clamp screws (screws) in the yaw, roll, pitch, and collective syncro resolvers and replace them with airworthy screws in accordance with the Accomplishment Instructions in Alert Service Bulletin 430–99–11, dated May 7, 1999 (ASB).

(b) Install a guard bracket on the yaw, roll, pitch, and collective syncro resolvers in accordance with the Accomplishment Instructions in the ASB.

(c) An alternative method of compliance or adjustment of the compliance time that provides an acceptable level of safety may be used if approved by the Manager, Regulations Group, Rotorcraft Directorate, FAA. Operators shall submit their requests through an FAA Principal Maintenance Inspector, who may concur or comment and then send it to the Manager, Regulations Group.

Note 2: Information concerning the existence of approved alternative methods of compliance with this AD, if any, may be obtained from the Regulations Group.

(d) Special flight permits may be issued in accordance with sections 21.197 and 21.199 of the Federal Aviation Regulations (14 CFR 21.197 and 21.199) to operate the helicopter to a location where the requirements of this AD can be accomplished.

(e) The modifications shall be done in accordance with the Accomplishment Instructions in Bell Helicopter Textron Alert Service Bulletin 430-99-11, dated May 7, 1999. This incorporation by reference was approved by the Director of the Federal Register in accordance with 5 U.S.C. 552(a) and 1 CFR part 51. Copies may be obtained from Bell Helicopter Textron Canada, 12,800 Rue de l'Avenir, Mirabel, Quebec JON1LO. telephone (800) 463-3036, fax (514) 433-0272. Copies may be inspected at the FAA, Office of the Regional Counsel, Southwest Region, 2601 Meacham Blvd., Room 663, Fort Worth, Texas; or at the Office of the Federal Register, 800 North Capitol Street, NW., suite 700, Washington, DC.

(f) This amendment becomes effective on September 20, 2000.

Note 3: The subject of this AD is addressed in Transport Canada (Canada) AD No. CF– 99–26, dated September 28, 1999.

Issued in Fort Worth, Texas, on August 2, 2000.

Henry A. Armstrong,

Manager, Rotorcraft Directorate, Aircraft Certification Service.

[FR Doc. 00–20404 Filed 8–15–00; 8:45 am] BILLING CODE 4910–13–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

42 CFR Part 70

Food and Drug Administration

21 CFR Part 1240

[Docket No. 00N-1317]

Control of Communicable Diseases; Apprehension and Detention of Persons With Specific Diseases; Transfer of Regulations

AGENCIES: Food and Drug Administration and Centers for Disease Control and Prevention, HHS. **ACTION:** Final rule.

SUMMARY: The Secretary of Health and Human Service (the Secretary) is transferring a portion of the Food and Drug Administration (FDA) "Control of Communicable Diseases" regulations to the Centers for Disease Control and Prevention (CDC). In general, these regulations provide the Secretary with the authority to apprehend, detain, or conditionally release individuals to prevent the spread of specified communicable diseases. The regulations implement the provisions of the Public Health Service Act (PHS Act) to prevent the introduction, transmission, or spread of communicable diseases from one State or possession into any other State or possession. CDC will have authority for interstate quarantine over persons, while FDA will retain regulatory authority over animals and other products that may transmit or spread communicable diseases. The Secretary is taking this action to consolidate regulations designed to control the spread of communicable diseases, thereby increasing the agencies' efficiency and effectiveness. **DATES:** This rule is effective September 15, 2000.

ADDRESSES: Submit written comments on the information collection requirements to the Office of Information and Regulatory Affairs, Office of Management and Budget (OMB), New Executive Office Bldg., 725 17th St. NW., rm. 10235, Washington, DC 20503, Attn: Desk Officer for CDC.

FOR FURTHER INFORMATION CONTACT:

James E. Barrow, National Center for Infectious Diseases (E–03), Centers for Disease Control and Prevention, 1600 Clifton Rd. NE., Atlanta, GA 30333, 404–639–8107; or

Captain Lawrence C. Edwards, Retail Food and Interstate Travel Team (HFS–627), Food and Drug Administration, 200 "C" St. SW., Washington DC 20204, 202–205– 8280.

SUPPLEMENTARY INFORMATION:

I. Background

Under sections 361 and 369 of the PHS Act, as amended, the Secretary issues and enforces regulations necessary to prevent the introduction. transmission, or spread of communicable diseases in the United States. FDA and CDC have been delegated regulatory responsibility under these provisions. The regulations contained in part 1240 (21 CFR part 1240), pertain to interstate control of communicable diseases and are currently administered by FDA. Regulations to prevent the introduction, transmission, or spread of communicable diseases from foreign countries into the United States are currently administered by CDC in 42 CFR part 71. The Secretary is taking this action to consolidate regulations designed to control the spread of communicable diseases, thereby increasing the agencies' efficiency and effectiveness.

The Secretary is transferring regulatory authority contained in the following sections of 21 CFR part 1240 to CDC: Section 1240.3 introductory text and paragraphs (b), (c), (e), (g), (h), (l), (n), and (p); § 1240.30; and Subpart C-**Restrictions on Travel of Persons** (consisting of §§ 1240.40, 1240.45, 1240.50, 1240.54, 1240.55, and 1240.57). The transferred regulations will be sequentially renumbered in 42 CFR part 70. In addition, "Director of the Centers for Disease Control and Prevention" has been inserted in new 42 CFR 70.2 in place of "Commissioner of Food and Drugs," currently in 21 CFR 1240.30.

Although regulatory authority with respect to interstate quarantine over persons in §§ 1240.3, 1240.30, and

1240.45 is being transferred to CDC, FDA is retaining its regulatory authority in §§ 1240.3, 1240.30, and 1240.45 with respect to animals and other products that may transmit or spread communicable diseases, to ensure that FDA's Center for Food Safety and Applied Nutrition has the necessary authorities to prevent the spread of disease from any conveyance engaged in interstate travel or in the event of inadequate local control. Current §1240.45 is also being moved to 21 CFR part 1240, subpart B (Administrative Procedures), and subpart C is being removed and reserved. The Secretary is issuing this rule without publishing a general notice of proposed rulemaking because such a notice is not required for this rule of agency organization, procedure, or practice under 5 U.S.C. 553(b)(A).

Table 1 reflects the actions the Secretary is taking:

TABLE 1

Current 21 CFR Sections (FDA)	New 42 CFR Sections (CDC)
1240.3 introductory text, and paragraphs (b), (c), (e), (g), (h), (l), (n), and (p) transferring and retaining.	70.1
1240.30 transferring and retaining	70.2
1240.40 transferred to	70.3
1240.45 transferring and retaining	70.4
1240.50 transferred to	70.5
1240.54 transferred to	70.6
1240.55 transferred to	70.7
1240.57 transferred to	70.8

This transfer of authority does not affect other authority exercised by FDA under sections 361 and 369, or any other sections, of the PHS Act.

II. Environmental Impact

FDA has determined under 21 CFR 25.30(h) that this action 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 the absence of an applicable categorical exclusion, CDC conducted an environmental assessment of this transfer of authority in accordance with the Department of Health and Human Services administrative guidance and determined that the transfer presented no significant impact on the human environment.

III. Federalism

Executive Order 13132 applies when agencies formulate or implement policies or regulations that preempt State law or that have federalism implications. Executive Order 13132

provides that agencies are to examine the constitutional and statutory authority supporting any action that would limit the policymaking discretion of the States and assess carefully the need for such actions. FDA and CDC have examined this rule and have determined that it does not preempt State law and it does not have a substantial direct effect on the States, on the relationship between the National Government and the States, or on the distribution of power and responsibilities among the various levels of government. Rather, the Secretary is taking this action to consolidate regulations to control the spread of communicable diseases, thereby increasing the agencies' efficiency and effectiveness. Therefore, no further action is required by Executive Order 13132.

IV. Analysis of Impacts

FDA and CDC have examined the impacts of this rule under Executive Order 12866. Executive Order 12866 directs agencies to assess all costs and benefits of available regulatory

alternatives and, when regulation is necessary, to select regulatory approaches that maximize net benefits (including potential economic, environmental, public health and safety, and other advantages; distributive impacts; and equity). Executive Order 12866 classifies a rule as significant if it meets any one of a number of specified conditions, including having an annual effect on the economy of \$100 million or more, or adversely affecting in a material way a sector of the economy, competition, or jobs, or if it raises novel legal or policy issues. The agencies find that this final rule, which transfers existing regulatory authority from one agency to the other, is not a significant rule as defined by Executive Order 12866. No analysis is required under the Regulatory Flexibility Act (5 U.S.C. 601–612) because the Secretary is issuing it without publishing a general notice of proposed rulemaking, as explained previously in this document.

V. Paperwork Reduction Act of 1995

This final rule contains information collection provisions that are subject to

review by OMB under the Paperwork Reduction Act of 1995 (44 U.S.C. 3501-3520). In the Federal Register of April 12, 2000 (65 FR 19772), CDC published a document entitled "Proposed Data Collections Submitted for Public Comment and Recommendations" to collect information under those regulations, and sought public comment for 60 days. The comment period closed on June 12, 2000. CDC will now prepare an information collection request for submission to OMB, and will publish another document in the Federal Register announcing submission of the request to OMB and soliciting that comments be submitted to OMB. CDC will publish an additional document in the Federal Register announcing OMB's decision to approve, modify, or disapprove the information collection provisions of this final rule. An agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a currently valid OMB control number.

List of Subjects

21 CFR Part 1240

Communicable diseases, Public health, Travel restrictions, Water supply.

42 CFR Part 70

Communicable diseases, Public health, Quarantine, Reporting and recordkeeping requirements, Travel restrictions.

Therefore, under the Public Health Service Act, 21 CFR Chapter I and 42 CFR Chapter I are amended as follows:

21 CFR Chapter I

PART 1240—CONTROL OF COMMUNICABLE DISEASES

1. The authority citation for 21 CFR part 1240 continues to read as follows:

Authority: 42 U.S.C. 216, 243, 264, 271.

§ 1240.45 [Transferred from Subpart C to Subpart B]

2. Section 1240.45 *Report of disease* is transferred from subpart C to subpart B.

Subpart C [Removed and Reserved]

3. Subpart C is removed and reserved.

42 CFR Chapter I

4. Part 70 is added to subchapter F of Chapter I to read as follows:

PART 70—INTERSTATE QUARANTINE

Secs. 70.1 General definitions.

- 70.2 Measures in the event of inadequate local control.
- 70.3 All communicable diseases.
- 70.4 Report of disease.
- 70.5 Certain communicable diseases; special requirements.
- 70.6 Apprehension and detention of persons with specific diseases.
- 70.7 Responsibility with respect to minors, wards, and patients.
- 70.8 Members of military and naval forces.

Authority: 42 U.S.C. 216, 243, 264, 271.

§70.1 General definitions.

As used in this part, terms shall have the following meaning:

(a) Communicable diseases means illnesses due to infectious agents or their toxic products, which may be transmitted from a reservoir to a susceptible host either directly as from an infected person or animal or indirectly through the agency of an intermediate plant or animal host, vector, or the inanimate environment.

(b) *Communicable period* means the period or periods during which the etiologic agent may be transferred directly or indirectly from the body of the infected person or animal to the body of another.

(c) *Conveyance* means any land or air carrier, or any vessel as defined in paragraph (h) of this section.

(d) *Incubation period* means the period between the implanting of disease organisms in a susceptible person and the appearance of clinical manifestation of the disease.

(e) Interstate traffic means:

(1) The movement of any conveyance or the transportation of persons or property, including any portion of such movement or transportation that is entirely within a State or possession—

(i) From a point of origin in any State or possession to a point of destination in any other State or possession; or

(ii) Between a point of origin and a point of destination in the same State or possession but through any other State, possession, or contiguous foreign country.

(2) Interstate traffic does not include the following:

(i) The movement of any conveyance which is solely for the purpose of unloading persons or property transported from a foreign country, or loading persons or property for transportation to a foreign country.

(ii) The movement of any conveyance which is solely for the purpose of effecting its repair, reconstruction, rehabilitation, or storage.

(f) *Possession* means any of the possessions of the United States, including Puerto Rico and the Virgin Islands. (g) *State* means any State, the District of Columbia, Puerto Rico, and the Virgin Islands.

(h) *Vessel* means any passengercarrying, cargo, or towing vessel exclusive of:

(1) Fishing boats including those used for shell-fishing;

(2) Tugs which operate only locally in specific harbors and adjacent waters;

(3) Barges without means of selfpropulsion;

(4) Construction-equipment boats and dredges; and

(5) Sand and gravel dredging and handling boats.

§70.2 Measures in the event of inadequate local control.

Whenever the Director of the Centers for Disease Control and Prevention determines that the measures taken by health authorities of any State or possession (including political subdivisions thereof) are insufficient to prevent the spread of any of the communicable diseases from such State or possession to any other State or possession, he/she may take such measures to prevent such spread of the diseases as he/she deems reasonably necessary, including inspection, fumigation, disinfection, sanitation, pest extermination, and destruction of animals or articles believed to be sources of infection.

§70.3 All communicable diseases.

A person who has a communicable disease in the communicable period shall not travel from one State or possession to another without a permit from the health officer of the State, possession, or locality of destination, if such permit is required under the law applicable to the place of destination. Stop-overs other than those necessary for transportation connections shall be considered as places of destination.

§70.4 Report of disease.

The master of any vessel or person in charge of any conveyance engaged in interstate traffic, on which a case or suspected case of a communicable disease develops shall, as soon as practicable, notify the local health authority at the next port of call, station, or stop, and shall take such measures to prevent the spread of the disease as the local health authority directs.

§ 70.5 Certain communicable diseases; special requirements.

The following provisions are applicable with respect to any person who is in the communicable period of cholera, plague, smallpox, typhus or yellow fever, or who, having been exposed to any such disease, is in the incubation period thereof:

(a) *Requirements relating to travelers.* (1) No such person shall travel from one State or possession to another, or on a conveyance engaged in interstate traffic, without a written permit of the Surgeon General or his/her authorized representative.

(2) Application for a permit may be made directly to the Surgeon General or to his/her representative authorized to issue permits.

(3) Upon receipt of an application, the Surgeon General or his/her authorized representative shall, taking into consideration the risk of introduction, transmission, or spread of the disease from one State or possession to another, reject it, or issue a permit that may be conditioned upon compliance with such precautionary measures as he/she shall prescribe.

(4) A person to whom a permit has been issued shall retain it in his/her possession throughout the course of his/ her authorized travel and comply with all conditions prescribed therein, including presentation of the permit to the operators of conveyances as required by its terms.

(b) *Requirements relating to operation of conveyances.*(1) The operator of any conveyance engaged in interstate traffic shall not knowingly:

(i) Accept for transportation any person who fails to present a permit as required by paragraph (a) of this section; or

(ii) Transport any person in violation of conditions prescribed in his/her permit.

(2) Whenever a person subject to the provisions of this section is transported on a conveyance engaged in interstate traffic, the operator thereof shall take such measures to prevent the spread of the disease, including submission of the conveyance to inspection, disinfection and the like, as an officer of the Public Health Service designated by the Surgeon General for such purposes deems reasonably necessary and directs.

§70.6 Apprehension and detention of persons with specific diseases.

Regulations prescribed in this part are not applicable to the apprehension, detention, or conditional release of individuals except for the purpose of preventing the introduction, transmission, or spread of the following diseases: Anthrax, chancroid, cholera, dengue, diphtheria, granuloma inguinale, infectious encephalitis, favus, gonorrhea, leprosy, lymphogranuloma venereum, meningococcus meningitis, plague, poliomyelitis, psittacosis, relapsing fever, ringworm of the scalp, scarlet fever, streptococcic sore throat, smallpox, syphilis, trachoma, tuberculosis, typhoid fever, typhus, and yellow fever.

§70.7 Responsibility with respect to minors, wards, and patients.

A parent, guardian, physician, nurse, or other such person shall not transport, or procure or furnish transportation for any minor child or ward, patient or other such person who is in the communicable period of a communicable disease, except in accordance with provisions of this part.

§70.8 Members of military and naval forces.

The provisions of §§ 70.3, 70.4, 70.5, 70.7, and this section shall not apply to members of the military or naval forces, and medical care or hospital beneficiaries of the Army, Navy, Veterans' Administration, or Public Health Service, when traveling under competent orders: *Provided*, That in the case of persons otherwise subject to the provisions of § 70.5 the authority authorizing the travel requires precautions to prevent the possible transmission of infection to others during the travel period.

Dated: August 8, 2000.

Donna E. Shalala,

Secretary of Health and Human Services. [FR Doc. 00–20719 Filed 8–15–00; 8:45 am] BILLING CODE 4160–01–F

DEPARTMENT OF THE TREASURY

Internal Revenue Service

26 CFR Parts 1 and 301

[TD 8896]

RIN 1545-AY37

Modification of Tax Shelter Rules

AGENCY: Internal Revenue Service (IRS), Treasury.

ACTION: Temporary regulations.

SUMMARY: These temporary regulations modify the rules relating to the filing by certain corporate taxpayers of a statement with their Federal corporate income tax returns under section 6011(a), the registration of confidential corporate tax shelters under section 6111(d), and the maintenance of lists of investors in potentially abusive tax shelters under section 6112. These regulations provide the public with additional guidance needed to comply with the disclosure rules, the registration requirement, and the list maintenance requirement applicable to

tax shelters. The temporary regulations affect corporations participating in certain reportable transactions, persons responsible for registering confidential corporate tax shelters, and organizers of potentially abusive tax shelters. The text of these temporary regulations also serves as the text of the proposed regulations set forth in the notice of proposed rulemaking on this subject in the Proposed Rules section of this issue of the **Federal Register**.

DATES: *Effective Date:* These temporary regulations are effective August 11, 2000.

Applicability Date: For dates of applicability, see §§ 1.6011–4T(g), 301.6111–2T(h), and 301.6112–1T, A–22.

FOR FURTHER INFORMATION CONTACT:

Catherine Moore, (202) 622–3080, (not a toll-free number).

SUPPLEMENTARY INFORMATION:

Paperwork Reduction Act

The collections of information contained in these regulations previously have been reviewed and approved by the Office of Management and Budget under control numbers 1545–1685 and 1545–1686. No material changes to these collections of information are made by these regulations.

An agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless the collection of information displays a valid OMB control number.

Books or records relating to a collection of information must be retained as long as their contents may become material in the administration of any internal revenue law. Generally, tax returns and tax return information are confidential, as required by 26 U.S.C. 6103.

Background

This document amends 26 CFR parts 1 and 301 to provide modified rules relating to the disclosure of certain tax shelters by corporate investors on their Federal corporate income tax returns under section 6011, the registration of confidential corporate tax shelters under section 6111, and the maintenance of lists of investors in potentially abusive tax shelters under section 6112.

On February 28, 2000, the IRS issued temporary and proposed regulations regarding section 6011 (TD 8877, REG– 103735–00), section 6111 (TD 8876, REG–110311–98), and section 6112 (TD 8875, REG–103736–00). The regulations were published in the **Federal Register** (65 FR 11205, 65 FR 11215, 65 FR 11211) on March 2, 2000.