Lawrenceburg, TN—Lawrenceburg Muni, NDB RWY 16, Orig.
Abingdon, VA—Virginia Highlands, NDB RWY 24, Amdt. 1
Shell Lake, WI—Shell Lake Muni, NDB RWY 31, Amdt. 1

* * * *Effective April 11, 1985*

Battle Creek, MI—W. K. Kellogg Regional, NDB RWY 23, Amdt. 15

* * * *Effective March 1, 1985*

Jackson, TN—McKellar Field, NDB RWY 2, Amdt. 6

* * * *Effective February 28, 1985*

Gainesville, GA—Lee Gilmer Memorial, NDB RWY 4, Amdt. 3
Gastonia, NC—Gastonia Muni, NDB RWY 3, Amdt. 3

* * * *Effective February 27, 1985*

Glasgow, MT—Glasgow Intl, NDB RWY 30, Amdt. 1

* * * *Effective February 26, 1985*

Soldotna, AK—Soldotna, NDB RWY 25, Amdt. 1

4. By amending § 97.29 ILS, ILS/DME, ISMLS, MLS, MLS/DME and MLS/RNAV SIAPs identified as follows:

* * * *Effective May 9, 1985*

Seattle, WA—Seattle-Tacoma Intl, ILS RWY 34R, Amdt. 9

* * * *Effective April 25, 1985*

Albany, GA—Albany-Dougherty County, ILS RWY 4, Amdt. 8
Atlanta, GA—The William B Hartsfield Atlanta Intl, ILS RWY 26R, Amdt. 1
Chicago, IL—Chicago-O'Hare Intl, ILS RWY 9L, Amdt. 5
Detroit, MI—Willow Run, ILS RWY 23L, Amdt. 2
Greensboro, NC—Greensboro-High Point-Winston Salem Regnl, ILS RWY 5, Amdt. 2
Greensboro, NC—Greensboro-High Point-Winston Salem Regnl, ILS RWY 14, Amdt. 16
Greensboro, NC—Greensboro-High Point-Winston Salem Regnl, ILS RWY 23, Amdt. 4
Providence, RI—Theodore Francis Green State, ILS/DME RWY 34, Amdt. 4
Greenville, SC—Donaldson Center, ILS RWY 4, Orig.
Pierre, SD—Pierre Muni, ILS RWY 31, Amdt. 8

* * * *Effective April 11, 1985*

Bridgeport, CT—Igor I. Sikorsky Memorial, ILS RWY 6, Amdt. 5
Wilmington, DE—Greater Wilmington-New Castle County, ILS RWY 1, Amdt. 18
Battle Creek, MI—W. K. Kellogg Regional, ILS RWY 23, Amdt. 15

* * * *Effective March 1, 1985*

Jackson, TN—McKellar Field, ILS RWY 2, Amdt. 7

5. By amending § 97.31 RADAR SIAPs identified as follows:

* * * *Effective April 25, 1985*

Detroit, MI—Willow Run, RADAR-1, Amdt. 5
Greensboro, NC—Greensboro-High Point-Winston Salem Regnl, RADAR-1, Amdt. 6

* * * *Effective April 11, 1985*

Reno, NV—Reno Cannon Intl, RADAR-1, Orig.

6. By amending § 97.33 RNAV SIAPS identified as follows:

* * * *Effective May 9, 1985*

Half Moon Bay, CA—Half Moon Bay, RNAV-A, Amdt. 1
St. Louis, MO—Lambert-St. Louis Intl, RNAV RWY 12R, Amdt. 1
St. Louis, MO—Lambert-St. Louis Intl, RNAV RWY 30L, Amdt. 11

* * * *Effective April 25, 1985*

Hampton, IA—Hampton Muni, RNAV RWY 17, Amdt. 1
Greensboro, NC—Greensboro-High Point-Winston Salem Regnl, RNAV RWY 23, Amdt. 3, Cancelled

* * * *Effective April 11, 1985*

Wilmington, DE—Greater Wilmington-New Castle County, RNAV RWY 9, Amdt. 3

(Secs. 307, 313(a), 601, and 1110, Federal Aviation Act of 1958 (49 U.S.C. 1348, 1354(a), 1421, and 1510); 49 U.S.C. 106(g) (Revised, Pub. L. 97-449, January 12, 1983); and 14 CFR 11.49(b)(3).)

Note.—The FAA has determined that this regulation only involves an established body of technical regulations for which frequent and routine amendments are necessary to keep them operationally current. It, therefore—(1) is not a "major rule" under Executive Order 12291; (2) is not a "significant rule" under DOT Regulatory Policies and Procedures (44 FR 11034; February 26, 1979); and (3) does not warrant preparation of a regulatory evaluation as the anticipated impact is so minimal. For the same reason, the FAA certifies that this amendment will not have a significant economic impact on a substantial number of small entities under the criteria of the Regulatory Flexibility Act.

Note.—The incorporation by reference in the preceding document was approved by the Director of the Federal Register on December 31, 1980, and reapproved as of January 1, 1982.

Issued in Washington, D.C. on March 8, 1985.

John S. Kern,

Acting Director of Flight Operations.

[FR Doc. 85-6447 Filed 3-18-85; 8:45 am]

BILLING CODE 4910-13-M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 610

General Biological Products Standards; OMB Approval of Requirements

AGENCY: Food and Drug Administration.

ACTION: Final rule.

SUMMARY: The Food and Drug Administration (FDA) announces that the Office of Management and Budget (OMB) has approved the collection of information requirements in the biologics regulation on official release of samples and protocols (21 CFR 610.2). The agency is amending that regulation to reflect OMB's approval under OMB control number 0910-0206.

EFFECTIVE DATE: March 19, 1985.

FOR FURTHER INFORMATION CONTACT: Michael Hooton, Center for Drugs and Biologics (HFN-368), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301-443-1306.

SUPPLEMENTARY INFORMATION: Because this amendment merely reflects OMB's approval of a collection of information requirement, notice and public procedure and delayed effective date are unnecessary (5 U.S.C. 553(b)(B) and (d)).

List of Subjects in 21 CFR Part 610

Biologics, Labeling.

PART 610—GENERAL BIOLOGICAL PRODUCTS STANDARDS

§ 610.2 [Amended]

Therefore, under the Public Health Service Act (sec. 351, 58 Stat. 702 as amended (42 U.S.C. 262)) and under authority delegated to the Commissioner of Food and Drugs (21 CFR 5.10), Part 610 is amended in § 610.2 *Requests for samples and protocols: official release* by adding at the end of the section the parenthetical statement "(Information collection requirements approved by the Office of Management and Budget under control number 0910-0206.)".

Dated: March 13, 1985.

Mervin H. Shumate,

Acting Associate Commissioner for Regulatory Affairs.

[FR Doc. 85-6463 Filed 3-14-85; 10:31 am]

BILLING CODE 4160-01-M

21 CFR Part 610

[Docket No. 81N-0417]

Additional Standards for Anti-Human Globulin; Correction

AGENCY: Food and Drug Administration.

ACTION: Final rule; correction.

SUMMARY: The Food and Drug Administration is correcting the final rule that amended the biologics regulation to establish additional standards for Anti-Human Globulin (50

FR 5574; February 11, 1985). Tests precluded or not required were incorrectly listed. This document corrects that error.

EFFECTIVE DATE: May 13, 1985.

FOR FURTHER INFORMATION CONTACT: Joseph Wilczek, Center for Drugs and Biologics (HFN-368), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301-443-1306.

SUPPLEMENTARY INFORMATION: In FR Doc. 85-3305, appearing on page 5574 in the **Federal Register** of Monday, February 11, 1985, the following correction is made:

§ 610.12 [Corrected]

On page 5579, in the first column under § 610.12 *Sterility*, paragraph (g)(4)(i) is corrected to read "(4) *Test precluded or not required*. (i) The tests prescribed in this section need not be performed for Whole Blood, Cryoprecipitated AHF, Platelets, Red Blood Cells, Plasma, Source Plasma, Smallpox Vaccine, Reagent Red Blood Cells, or Anti-Human Globulin."

Dated: March 13, 1985.

Mervin H. Shumate,

Acting Associate Commissioner for Regulatory Affairs.

[FR Doc. 85-6464 Filed 3-14-85; 10:31 am]

BILLING CODE 4160-01-M

DEPARTMENT OF HOUSING AND URBAN DEVELOPMENT

Office of Assistant Secretary for Housing—Federal Housing Commissioner

24 CFR Parts 1710 and 1720

[Docket No. R-85-1230; FR-2080]

Interstate Land Sales Registration and Exemption Guidelines (Interpretative Rule); Technical Amendments and Corrections

AGENCY: Office of Assistant Secretary for Housing—Federal Housing Commissioner, HUD.

ACTION: Final rule; technical amendments and corrections.

SUMMARY: This rule amends and corrects the Department's regulations and Final Guidelines that implement the Interstate Land Sales Full Disclosure Act.

EFFECTIVE DATE: April 29, 1985.

FOR FURTHER INFORMATION CONTACT: John L. Brady, Director, Interstate Land Sales Registration Division, Department of Housing and Urban Development, Room 6278, 451 7th Street, SW., Washington, DC 20410 (202) 755-0502. (This is not a toll-free number.)

SUPPLEMENTARY INFORMATION: This rule makes technical amendments to Title 24 of the Code of Federal Regulations and corrects a final rule and final guidelines published in the **Federal Register** on August 6, 1984 (49 FR 31366, 31372) relating to the Department's Interstate Land Sales Registration Program.

The Department has determined that this document need not be published as a proposed rule, as generally required by the Administrative Procedure Act (APA), since this rulemaking merely makes technical amendments to existing HUD regulations.

A Finding of No Significant Impact with respect to the environment required by the National Environmental Policy Act (42 U.S.C. 4321-4347) is unnecessary, since these technical amendments are categorically excluded under HUD regulations at 24 CFR 50.20(k).

This rule does not constitute a "major rule" as that term is defined in Section 1(b) of Executive Order 12291 on Federal Regulation issued on February 17, 1981. Analysis of this rule indicates that it does not: (1) Have an annual effect on the economy of $100 million or more; (2) cause a major increase in cost or prices for consumers, individual industries, Federal, State or local government agencies, or geographic regions; or (3) have a significant adverse effect on competition, employment, investment, productivity, innovation, or on the ability of United States-based enterprises to compete with foreign-based enterprises in domestic or export markets.

As required by section 605(b) of the Regulatory Flexibility Act (5 U.S.C. 601), the Undersigned hereby certifies that this rule does not have a significant economic impact on a substantial number of small entities because it merely makes technical amendments to the Department's regulations.

This rule was not listed in the Department's Semiannual Agenda of Regulations published on October 22, 1984 (49 FR 41684).

List of Subjects

24 CFR Part 1710

Consumer protection; Land sales; Reporting and recordkeeping requirements.

24 CFR Part 1720

Administrative practice and procedure.

Accordingly, the Department amends 24 CFR Chapter X as follows:

PART 1710—LAND REGISTRATION

§ 1710.1 [Amended]

1. In § 1710.1, the term "Blanked encumbrance" appearing alphabetically in the list of definitions is removed and the term "Blanket encumbrance" is added in its place.

2. In § 1710.20, the section heading and paragraph (c) are revised to read as follows:

§ 1710.20 Requirements for registering a subdivision—Statement of Record—filing and form.

* * * * *

(c) *State filings.* A Statement of Record submitted under the provisions of 24 CFR Part 1710, Subpart C—Certification of Substantially Equivalent State Law, shall consist of the materials designated by the Certification Agreement between the Secretary and the certified State in which the subdivision is located.

§ 1710.114 [Amended]

3. Section 1710.114(a) is amended by removing the reference to number one, "(1)" which appears after the paragraph heading and before the word "Unless".

4. In § 1710.212, paragraph (d)(3) is revised to read as follows:

§ 1710.212 Financial information.

* * * * *

(d) * * *

(3) If the developer no longer has an active sales program on the date this report is due, the information set forth in § 1710.310(c)(7)(iii) may be furnished in lieu of this report.

* * * * *

PART 1720—FORMAL PROCEDURES AND RULES OF PRACTICE

§ 1720.505 [Amended]

5. Section 1720.505 currently consists of two undesignated paragraphs. The two paragraphs are designated paragraph (a) and paragraph (b), respectively. In addition, new paragraph (a) contains references to designated paragraphs (a) and (b) which are removed and references to paragraphs (1) and (2) are added respectively in their place.

§ 1710.21 [Amended]

6. In FR Doc. 84-20695, appearing on page 31366 in the August 6, 1984 issue of the **Federal Register**, in the amendatory language for amendment number 25, appearing in the first column on page 31370, the reference to "Section 1710.12" is corrected to read "Section 1710.21".