Powers Act of 1977, as amended, section 505 of the International Security and Development Cooperation Act of 1985, and Executive Orders 12957 and 12959 of March 15, 1995 and May 6, 1995, respectively. This embargo includes prohibitions on export and certain reexport transactions involving Iran, including transactions dealing with items subject to the EAR. (See OFAC's Iranian Transactions Regulations, 31 CFR part 560.)

(1) The controls on exports and reexports to Iran, as specified in the CCL and in paragraph (d) of this section, continue to apply. To avoid duplication, exporters or reexporters are not required to seek separate authorization from BXA for an export or reexport subject both to the EAR and to OFAC's Iranian Transactions Regulations. Therefore, if OFAC authorizes an export or reexport, no separate authorization from BXA is necessary.

(2) Section 3 of the Executive Order directs all agencies of the United States Government to take all appropriate measures within their jurisdiction to carry out the order. Accordingly, no validated license, general license or other authorization constitutes authority for any export or reexport prohibited by the Iranian Transactions Regulations unless authorized by OFAC, and no person may export or reexport items subject to both the EAR and OFAC's Iranian Transactions Regulations without prior OFAC authorization. Any export or reexport prohibited both by the EAR and by the Executive Order and not authorized by OFAC is a violation of the EAR.

(3) Exporters should consult with OFAC (Office of Foreign Assets Control, U.S. Department of the Treasury, 1500 Pennsylvania Avenue, N.W., Annex, 2nd Floor, Washington, D.C. 20220. Telephone (202) 622–2480) for authorization for:

(i) Exports from the United States involving Iran;

(ii) Exports or reexports to Iran from a third country, when the exporter or reexporter is a United States person (as defined in OFAC's Iranian Transactions Regulations, 31 CFR part 560); or

(iii) Reexports to Iran of U.S.-origin items that were subject to any export license application requirements prior to Executive Order 12959 of May 6, 1995.

(Office of Foreign Assets Control, U.S. Department of the Treasury, 1500 Pennsylvania Avenue, N.W., Annex, 2nd Floor, Washington, D.C. 20220. Telephone (202) 622–2480.)

* * * * *

Dated: February 29, 1996.

Sue E. Eckert,

Assistant Secretary for Export Administration.

[FR Doc. 96–5103 Filed 3–4–96; 8:45 am]

BILLING CODE 3510–DT–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 5

Delegations of Authority and Organization; Technical Amendment

AGENCY: Food and Drug Administration, HHS.

ACTION: Final rule; technical amendment.

SUMMARY: The Food and Drug Administration (FDA) is amending the regulations for delegations of authority to set forth the current organizational structure of the agency as well as the current addresses for headquarters and field offices. This action is necessary to ensure accuracy of the regulations.

EFFECTIVE DATE: March 5, 1996.

FOR FURTHER INFORMATION CONTACT: Ellen Rawlings, Division of Management Systems and Policy (HFA–340), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–443–4976.

SUPPLEMENTARY INFORMATION: The regulations are being amended in 21 CFR 5.100 to reflect the current addresses for headquarters and for field and district offices.

Notice and comment on these amendments are not necessary under the Administrative Procedure Act because this is a rule of Agency organization (5 U.S.C. 553(b)).

List of Subjects in 21 CFR Part 5

Authority delegations (Government agencies), Imports, Organization and functions (Government agencies).

Therefore, under the Federal Food, Drug, and Cosmetic Act and under authority delegated to the Commissioner of Food and Drugs, 21 CFR part 5 is amended as follows:

PART 5—DELEGATIONS OF AUTHORITY AND ORGANIZATION

1. The authority citation for 21 CFR part 5 continues to read as follows:

Authority: 5 U.S.C. 504, 552, App. 2; 7 U.S.C. 138a, 2271; 15 U.S.C. 638, 1261–1282, 3701–3711a; secs. 2–12 of the Fair Packaging and Labeling Act (15 U.S.C. 1451–1461); 21 U.S.C. 41–50, 61–63, 141–149, 467f, 679(b), 801–886, 1031–1309; secs. 201–903 of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 321–394); 35 U.S.C. 156; secs. 301, 302, 303, 307, 310, 311, 351, 352, 354, 361, 362, 1701–1706; 2101, 2125, 2127, 2128 of the Public Health Service Act (42 U.S.C. 241, 242, 242a, 242l, 242n, 243, 262, 263, 263b, 264, 265, 300u–300u–5, 300aa–1, 300aa–25, 300aa–27, 300aa–28); 42 U.S.C. 1395y, 3246b, 4332, 4831(a), 10007–10008; E.O. 11490, 11921, and 12591; secs. 312, 313, 314 of the National Childhood Vaccine Injury Act of 1986, Pub. L. 99–660 (42 U.S.C. 300aa–1 note).

§ 5.100 [Amended]

2. Section 5.100 is amended by revising footnotes 9 and 12, and by adding new footnote 17 to the entry for ''Division of Clinical Laboratory Devices.''To read as follows:

§ 5.100 Headquarters.

* * * * *

Center for Biologics Evaluation and Research[9]

* * * * *

Office of Device Evaluation[12]

* * * * *

Division of Clinical Laboratory Devices[17]

* * * * *

Dated: February 26, 1996.

William K. Hubbard,

Associate Commissioner for Policy Coordination.

[FR Doc. 96–4977 Filed 3–4–96; 8:45 am]

BILLING CODE 4160–01–F

DEPARTMENT OF JUSTICE

28 CFR Part 52

[AG ORDER No. 2012–96]

RIN 1105–AA43

Revision of Policy Concerning Consent To Try Civil Matters Before Magistrate Judges

AGENCY: Department of Justice.

ACTION: Final rule.

SUMMARY: The United States Department of Justice is publishing a final rule to revise and clarify Department policy concerning consent to try civil matters before magistrate judges.

EFFECTIVE DATE: This final rule is effective March 5, 1996.

FOR FURTHER INFORMATION CONTACT: Mary C. Morgan, Deputy Assistant Attorney General, Office of Policy

[9] Mailing address: 1401 Rockville Pike, suite 200N, Rockville, MD 20852–1448.

[12] Mailing address: 9200 Corporate Blvd., Rockville, MD 20850.

[17] See footnote 13.