section has changed since the previous report.

(c) *When to report.* Annual progress reports for postmarketing study commitments entered into by applicants shall be reported to FDA within 60 days of the anniversary date of the U.S. approval of the application for the product.

(d) *Where to report.* Submit two copies of the annual progress report of postmarketing studies to the Food and Drug Administration, Center for Biologics Evaluations and Research, Document Control Center (HFM–99), 1401 Rockville Pike, Rockville, MD 20852–1448.

(e) *Public disclosure of information.* Except for the information described in this paragraph, FDA may publicly disclose any information concerning a postmarketing study, within the meaning of this section, if the agency determines that the information is necessary to identify an applicant or to establish the status of the study including the reasons, if any, for failure to conduct, complete, and report the study. Under this section, FDA will not publicly disclose trade secrets, as defined in § 20.61 of this chapter, or information, described in § 20.63 of this chapter, the disclosure of which would constitute an unwarranted invasion of personal privacy.

Dated: October 13, 2000.

Margaret M. Dotzel,

Associate Commissioner for Policy.

[FR Doc. 00–27731 Filed 10–26–00; 8:45 am]

BILLING CODE 4160–01–F

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 876

[Docket No. 94N–0380]

Gastroenterology and Urology Devices; Effective Date of the Requirement for Premarket Approval of the Implanted Mechanical/Hydraulic Urinary Continence Device; Correction

AGENCY: Food and Drug Administration, HHS.

ACTION: Final rule; correction.

SUMMARY: The Food and Drug Administration (FDA) is correcting a final rule that appeared in the **Federal Register** of September 26, 2000 (65 FR 57726). The final rule requires the filing of a premarket approval application or a notice of completion of a product development protocol for the implanted mechanical/hydraulic urinary continence device, a generic type of medical device intended for the treatment of urinary incontinence. In the final rule, the effective date was stated incorrectly. This document corrects that error.

FOR FURTHER INFORMATION CONTACT: Nicole L. Wolanski, Center for Devices and Radiological Health (HFZ–470), Food and Drug Administration, 9200 Corporate Blvd., Rockville, MD 20850, 301–594–2194.

SUPPLEMENTARY INFORMATION: In FR Doc. 00–24632 appearing on page 57726 in the **Federal Register** of September 26, 2000, the following correction is made:

1. On page 57726, in the second column, under the **EFFECTIVE DATE** caption, the date ''October 26, 2000'' is corrected to read ''September 26, 2000.''

Dated: October 19, 2000.

Linda S. Kahan,

Deputy Director for Regulations Policy, Center for Devices and Radiological Health.

[FR Doc. 00–27736 Filed 10–27–00; 8:45 am]

BILLING CODE 4160–01–F

DEPARTMENT OF DEFENSE

Department of the Air Force

32 CFR Part 811

RIN 0701–AA62

Release, Dissemination, and Sale of Visual Information Materials

AGENCY: Department of the Air Force, DoD.

ACTION: Final rule.

SUMMARY: The Department of the Air Force is revising our rules on the Release, Dissemination, and Sale of Visual Information Materials to reflect current polices. These rules implement Air Force Instruction (AFI) 33–117, Visual Information Management, and apply to all Air Force activities.

EFFECTIVE DATE: December 1, 2000.

ADDRESSES: Mr. Raymond Dabney, HQ AFCIC/ITSM, 1250 Air Force Pentagon, Washington, DC 20330–1250, 703–588–6136.

FOR FURTHER INFORMATION CONTACT: Mr. Raymond Dabney, HQ AFCIC/ITSM, 703–588–6136.

SUPPLEMENTARY INFORMATION: The Department of the Air Force is revising our rules on the Release, Dissemination, and Sale of Visual Information Materials of the Code of Federal Regulations (CFRs) (32 CFR part 811) to reflect current policies. This part implements Air Force Instruction (AFI) 33–117, Visual Information Management, and apply to all Air Force activities. This part was published as a proposed rule in the **Federal Register** on December 28, 1999 (64 FR 72621, FR Doc. 99–33604). Comments were solicited for 60 days ending on February 28, 2000. No comments were received.

List of Subjects in 32 CFR Part 811

Archives and records, Motion pictures.

For the reasons stated in the preamble, the Department of the Air Force is revising 32 CFR Part 811 to read as follows:

PART 811—RELEASE, DISSEMINATION, AND SALE OF VISUAL INFORMATION MATERIALS

Sec.

Authority: 10 U.S.C. 8013.

§ 811.1 Exceptions.

The regulations in this part do not apply to:

(a) Visual information (VI) materials made for the Air Force Office of Special Investigations for use in an investigation or a counterintelligence report. (See Air Force Instruction (AFI) 90–301, The Inspector General Complaints, which describes who may use these materials.)

(b) VI materials made during Air Force investigations of aircraft or missile mishaps according to AFI 91–204, Safety Investigations and Reports. (See AFI 90–301.)

§ 811.2 Release of visual information materials.

(a) Only the Secretary of the Air Force for Public Affairs (SAF/PA) clears and releases Air Force materials for use outside Department of Defense (DoD), according to AFI 35–205, Air Force Security and Policy Review Program.

(b) The Secretary of the Air Force for Legislative Liaison (SAF/LL) arranges the release of VI material through SAF/PA when a member of Congress asks for them for official use.

(c) The International Affairs Division (HQ USAF/CVAII) or, in some cases, the major command (MAJCOM) Foreign Disclosure Office, must authorize release of classified and unclassified