

approved the information collection requirements contained in this AD and has assigned OMB Control Number 2120-0056.

(1) If the inspection is accomplished after the effective date of this AD: Submit the report within 10 days after the inspection.

(2) If the inspection was accomplished before the effective date of this AD: Submit the report within 10 days after the effective date of this AD.

#### Parts Installation

(h) As of the effective date of this AD, no person may install a CFRP rudder, any series of P/N A55471500, on any airplane, unless the CFRP rudder has been inspected and any applicable corrective action has been accomplished in accordance with paragraphs (f)(2) and (f)(3) of this AD.

#### Alternative Methods of Compliance (AMOCs)

(i)(1) The Manager, International Branch, ANM-116, FAA, has the authority to approve AMOCs for this AD, if requested in accordance with the procedures found in 14 CFR 39.19.

(2) Before using any AMOC approved in accordance with § 39.19 on any airplane to which the AMOC applies, notify the appropriate principal inspector in the FAA Flight Standards Certificate Holding District Office.

#### Related Information

(j) The European Aviation Safety Agency's airworthiness directive 2006-0066, dated March 24, 2006, also addresses the subject of this AD.

#### Material Incorporated by Reference

(k) You must use Airbus All Operators Telex A310-55A2043, dated March 2, 2006, or Airbus All Operators Telex A300-55A6042, dated March 2, 2006, as applicable; and Airbus Technical Disposition 943.0046/06, dated March 2, 2006; to perform the actions that are required by this AD, unless the AD specifies otherwise. (Only page 1 of Airbus All Operators Telex A310-55A2043 and Airbus All Operators Telex A300-55A6042 contains the document number and date of the document; no other page of the document contains this information.) The Director of the Federal Register approved the incorporation by reference of these documents in accordance with 5 U.S.C. 552(a) and 1 CFR part 51. Contact Airbus, 1 Rond Point Maurice Bellonte, 31707 Blagnac Cedex, France, for a copy of this service information. You may review copies at the Docket Management Facility, U.S. Department of Transportation, 400 Seventh Street, SW., room PL-401, Nassif Building, Washington, DC; on the Internet at <http://dms.dot.gov>; or at the National Archives and Records Administration (NARA).

For information on the availability of this material at the NARA, call (202) 741-6030, or go to [http://www.archives.gov/federal\\_register/code\\_of\\_federal\\_regulations/ibr\\_locations.html](http://www.archives.gov/federal_register/code_of_federal_regulations/ibr_locations.html).

Issued in Renton, Washington, on March 24, 2006.

**Ali Bahrami,**

*Manager, Transport Airplane Directorate, Aircraft Certification Service.*

[FR Doc. 06-3119 Filed 3-28-06; 12:45 pm]

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## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Food and Drug Administration

#### 21 CFR Part 3

#### Change of Telephone Number; Technical Amendment

**AGENCY:** Food and Drug Administration, HHS.

**ACTION:** Final rule; technical amendment.

**SUMMARY:** The Food and Drug Administration (FDA) is amending its regulations to reflect a change in telephone number for the Office of Combination Products (OCP). This action is editorial in nature and is intended to improve the accuracy of the agency's regulations.

**DATES:** March 30, 2006.

**FOR FURTHER INFORMATION CONTACT:** Leigh Hayes, Office of Combination Products (HFG-3), Food and Drug Administration, 15800 Crabbs Branch Way, suite 200, Rockville, MD 20855, 301-427-1934.

**SUPPLEMENTARY INFORMATION:** FDA is amending its regulations in 21 CFR part 3 to reflect a change in the telephone number for the OCP.

Publication of this document constitutes final action on this change under the Administrative Procedure Act (5 U.S.C. 553). Notice and public procedures are unnecessary because FDA is merely correcting a nonsubstantive error.

#### List of Subjects in 21 CFR Part 3

Administrative practice and procedure, Biologics, Drugs, Medical devices.

■ Therefore, under the Federal Food, Drug, and Cosmetic Act and under authority delegated to the Commissioner of Food and Drugs, 21 CFR Part 3 is amended as follows:

#### PART 3—PRODUCT JURISDICTION

■ 1. The authority citation for 21 CFR part 17 continues to read as follows:

**Authority:** 21 U.S.C. 321, 351, 353, 355, 360, 360c-360f, 360h-360j, 360gg-360ss, 360bbb-2, 371(a), 379e, 381, 394; 42 U.S.C. 216, 262, 264.

#### § 3.6 [Amended]

■ 2. Section 3.6 is amended by removing “301-827-9229” and by adding in its place “301-427-1934”.

Dated: March 23, 2006.

**Jeffrey Shuren,**

*Assistant Commissioner for Policy.*

[FR Doc. 06-3046 Filed 3-29-06; 8:45 am]

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## DEPARTMENT OF THE INTERIOR

### Minerals Management Service

#### 30 CFR Parts 250 and 251

RIN 1010-AC81

#### Oil and Gas and Sulphur Operations in the Outer Continental Shelf (OCS)—Geological and Geophysical (G&G) Explorations of the OCS—Proprietary Terms and Data Disclosure

**AGENCY:** Minerals Management Service (MMS), Interior.

**ACTION:** Final rule.

**SUMMARY:** This rule expands the circumstances under which MMS allows inspection of G&G data and information. The rule also modifies the start dates of proprietary terms for geophysical data and information and any derivatives of these data and information that MMS acquires. In addition, the rule clarifies the proprietary terms of geological data and information MMS acquires pursuant to a permit.

**DATES:** *Effective Date:* May 1, 2006.

**FOR FURTHER INFORMATION CONTACT:** George Dellagiarino or David Zinzer at (703) 787-1628.

**SUPPLEMENTARY INFORMATION:** This final rule implements changes put forward by our notice of proposed rulemaking (NPR) published July 17, 2002 (67 FR 46942). The comment period ended September 16, 2002. MMS received 10 sets of written comments and recommendations in response to the NPR. Two sets of comments and recommendations were from industry associations, and eight were from permittees and third party users of G&G data and information collected on the OCS. We have carefully considered each of these comments and recommendations. We did not adopt recommendations that did not appear to be in the public's best interest.

#### Discussion and Analysis of Comments

MMS has decided to proceed with the final rule after carefully considering all written comments on the proposed