is found installed and the S/N of the FDU is listed in table 1 of this AD: Before further flight, replace the FDU with a serviceable FDU, in accordance with the instructions of Airbus AOT A330–26A3052, dated April 19, 2011 (for Model A330–200 and –300 series airplanes); Airbus AOT A340–200/300–26A4044, dated April 19, 2011 (for Model A340–200 and –300 series airplanes); or Airbus AOT A340–500/600–26A5024, dated April 19, 2011 (for Model A340–500 and –600 series airplanes).

TABLE 1—AFFECTED P/N 3711–00 FIRE DETECTION UNITS

Serial Nos.
ZL0683.
ZL0718.
ZL0721 through ZL0725 inclusive.
ZL0727.
ZL0729 through ZL0731 inclusive.
ZL0736.
ZL0738.
ZL0740.
ZL0742.
ZL0743.
ZL0745.

TABLE 1—AFFECTED P/N 3711–00 FIRE DETECTION UNITS—Continued

Serial Nos.
ZL0747.
ZL0770.
ZL0772.
ZL0775.
ZL0788.
ZL0804.

Note 1: Some of the affected P/N 3711–00 FDUs have been installed in production on certain airplanes, as indicated in table 2 of this AD.

TABLE 2—FDU INSTALLED IN PRODUCTION

Model A330–200 and –300 airplanes manufacturer serial numbers	Position	S/N
1177	ENG2 FDU (1WD2)	ZL0683
1191	ENG2 FDU (1WD2)	ZL0723
1192	ENG1 FDU (1WD1)	ZL0721
	ENG2 FDU (1WD2)	ZL0722
1193	APU FDU (13WG)	ZL0718
1195	ENG1 FDU (1WD1)	ZL0740
1196	ENG1 FDU (1WD1)	ZL0742
	ENG2 FDU (1WD2)	ZL0736
	APU FDU (13WG)	ZL0743
1198	ENG2 FDU (1WD2)	ZL0738
1199	APU FDU (13WG)	ZL0731
1200	ENG1 FDU (1WD1)	ZL0747
1206	ENG2 FDU (1WD2)	ZL0770

Parts Installation

(i) As of the effective date of this AD, no person may install on any airplane, any P/N 3711–00 FDU with a serial number listed in table 1 of this AD, unless the FDU has been reworked and re-identified by L'Hotellier as specified in the instructions in Airbus AOT A330–26A3052, dated April 19, 2011 (for Model A330–200 and –300 series airplanes); Airbus AOT A340–200/300–26A4044, dated April 19, 2011 (for Model A340–200 and –300 series airplanes); or Airbus AOT A340–500/600–26A5024, dated April 19, 2011 (for Model A340–500 and –600 series airplanes).

FAA AD Differences

Note 2: This AD differs from the MCAI and/or service information as follows:

No differences.

Other FAA AD Provisions

(j) The following provisions also apply to this AD:

(1) *Alternative Methods of Compliance (AMOCs):* The Manager, International Branch, ANM–116, Transport Airplane Directorate, FAA, has the authority to approve AMOCs for this AD, if requested using the procedures found in 14 CFR 39.19. In accordance with 14 CFR 39.19, send your request to your principal inspector or local Flight Standards District Office, as appropriate. If sending information directly to the International Branch, send it to *Attn:* Vladimir Ulyanov, Aerospace Engineer, International Branch, ANM–116, Transport Airplane Directorate, FAA, 1601 Lind Avenue, SW., Renton, Washington 98057–3356; telephone (425) 227–1138; fax (425) 227–1149. Information may be e-mailed to: *9-ANM-116-AMOC-REQUESTS@faa.gov.* Before using any approved AMOC, notify your appropriate principal inspector, or lacking a principal inspector, the manager of the local flight standards district office/certificate holding district office. The AMOC approval letter must specifically reference this AD.

(2) *Airworthy Product:* For any requirement in this AD to obtain corrective actions from a manufacturer or other source, use these actions if they are FAA-approved. Corrective actions are considered FAA-approved if they are approved by the State of Design Authority (or their delegated agent). You are required to assure the product is airworthy before it is returned to service.

Related Information

(k) Refer to MCAI EASA Airworthiness Directive 2011–0073, dated April 20, 2011; Airbus AOT A330–26A3052, dated April 19, 2011; Airbus AOT A340–200/300–26A4044, dated April 19, 2011; and Airbus AOT A340–500/600–26A5024, dated April 19, 2011; for related information.

Issued in Renton, Washington, on September 7, 2011.

Jeffrey E. Duven,

Acting Manager, Transport Airplane Directorate, Aircraft Certification Service.

[FR Doc. 2011–23470 Filed 9–13–11; 8:45 am]

BILLING CODE 4910–13–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 352

[Docket No. FDA–1978–N–0018] (formerly Docket No. 1978N–0038)

RIN 0910–ZA40

Sunscreen Drug Products for Over-the-Counter Human Use; Request for Data and Information Regarding Dosage Forms; Extension of Comment Period

AGENCY: Food and Drug Administration, HHS.

ACTION: Advance notice of proposed rulemaking; request for data and information; extension of comment period.

SUMMARY: The Food and Drug Administration (FDA) is extending the comment period for the advance notice of proposed rulemaking (ANPRM) that published on June 17, 2011. The ANPRM is requesting data and information on certain dosage forms of over-the-counter (OTC) sunscreen drug products marketed without approved applications. The comment period for that ANPRM will end on September 15, 2011. This document extends the comment period to October 17, 2011.

DATES: Submit either electronic or written data and information by October 17, 2011.

ADDRESSES: You may submit comments, identified by Docket No. FDA–1978–N–0018 (formerly Docket No. 1978N–0038) and/or RIN number 0910–ZA40, by any of the following methods:

Electronic Submissions

Submit electronic comments in the following way:

- *Federal eRulemaking Portal: http://www.regulations.gov*. Follow the instructions for submitting comments.

Written Submissions

Submit written submissions in the following ways:

- *Fax:* 301–827–6870.
- *Mail/Hand delivery/Courier (for paper, disk, or CD–ROM submissions):* Division of Dockets Management (HFA–305), Food and Drug Administration, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852.

Instructions: All submissions received must include the Agency name, Docket No. FDA–1978–N–0018, and RIN 0910–ZA40 for this rulemaking. All comments received may be posted without change to *http://www.regulations.gov*, including any personal information provided if not marked as confidential. For additional information on submitting comments, see the ''Request for Comments'' heading of the **SUPPLEMENTARY INFORMATION** section of this document.

Docket: For access to the docket to read background documents or comments received, go to *http://www.regulations.gov*, insert the docket number, found in brackets in the heading of this document, into the ''Search'' box and follow the prompts and/or go to the Division of Dockets Management, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852.

FOR FURTHER INFORMATION CONTACT: Reynold Tan, Center for Drug Evaluation and Research, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 22, Rm. 5411, Silver Spring, MD 20993–0002, 301–796–2090.

SUPPLEMENTARY INFORMATION:

I. Background

In the **Federal Register** of June 17, 2011 (76 FR 35669) (the June 17, 2011, ANPRM), FDA published an ANPRM that requested data and information on OTC sunscreen products marketed without approved applications that are formulated in certain dosage forms. FDA requested these data to help establish OTC monograph conditions, including dosage form specifications, for OTC sunscreen drug products. Among the data requested is data necessary to resolve specific questions about the effectiveness and safety of OTC sunscreens in spray dosage forms.

II. Extension of the Comment Period

In response to the June 17, 2011, ANPRM, three submissions (Refs. 1, 2, and 3) requested an extension of the comment period, which will end on September 15, 2011. Two of the submissions requested that FDA extend the comment period by 30 days so that the comment period totals 4 months (Refs. 1 and 2). The other submission requested that FDA extend the comment period by 90 to 180 days so that the comment period totals 6 to 9 months (Ref. 3). The submissions cited the need for additional time to evaluate their available data and to organize and submit the data and information that best addresses FDA's request while simultaneously implementing the new requirements for their sunscreen products imposed by the Labeling and Effectiveness Testing final rule that published in the **Federal Register** of June 17, 2011 (76 FR 35620).

FDA is extending the comment period to end on October 17, 2011. A total comment period of 4 months is sufficient for the public to submit comments to the ANPRM.

III. Request for Comments

Interested persons may submit to the Division of Dockets Management (see **ADDRESSES**) either electronic or written comments, data, and information by October 17, 2011. It is only necessary to submit one set of comments, data, and information. It is no longer necessary to two copies of mailed comments, data, and information. Identify submissions with the docket number found in brackets in the heading of this document, and may be accompanied by supporting information. Received submissions may be seen in the Division of Dockets Management between 9 a.m. and 4 p.m., Monday through Friday. Information submitted after the closing date will not be considered except by petition under 21 CFR 10.30.

IV. References

The following references are on display in the Division of Dockets Management (see **ADDRESSES**) under Docket No. FDA–1978–N–0018 and may be seen by interested persons between 9 a.m. and 4 p.m., Monday through Friday.

1. Comment No. FDA–1978–N–0018–DRAFT–5225.

2. Comment No. FDA–1978–N–0018–DRAFT–5227.

3. Comment No. FDA–1978–N–0018–DRAFT–5228.

Dated: September 9, 2011.

Leslie Kux,

Acting Assistant Commissioner for Policy.

[FR Doc. 2011–23479 Filed 9–13–11; 8:45 am]

BILLING CODE 4160–01–P

DEPARTMENT OF THE INTERIOR

Bureau of Ocean Energy Management, Regulation and Enforcement

30 CFR Part 250

[Docket ID BOEM–2011–0003]

RIN 1010–AD73

Oil and Gas and Sulphur Operations in the Outer Continental Shelf—Revisions to Safety and Environmental Management Systems

AGENCY: Bureau of Ocean Energy Management, Regulation and Enforcement (BOEMRE), Interior.

ACTION: Proposed rule.

SUMMARY: This rulemaking proposes to amend BOEMRE regulations to require operators to develop and implement additional provisions in their Safety and Environmental Management Systems (SEMS) programs for oil, gas, and sulphur operations in the Outer Continental Shelf (OCS). These revisions pertain to developing and implementing stop work authority and ultimate work authority, requiring employee participation in the development and implementation of SEMS programs, and establishing requirements for reporting unsafe working conditions. In addition, this proposed rule requires independent third parties to conduct audits of operators' SEMS programs and establishes further requirements relating to conducting job safety analysis (JSA) for activities identified in an operator's SEMS program. We believe that these new requirements will further reduce the likelihood of accidents, injuries, and spills in connection with OCS activities that are regulated under BOEMRE jurisdiction, by requiring OCS operators to specifically address issues associated with human behavior as it applies to their SEMS program.

DATES: Submit comments by November 14, 2011. BOEMRE may not fully consider comments received after this date. Submit comments to the Office of Management and Budget on the information collection burden in this