

products identified in this rulemaking action.

**Regulatory Findings**

We have determined that this proposed AD would not have federalism implications under Executive Order 13132. This proposed AD would not have a substantial direct effect on the States, on the relationship between the national Government and the States, or on the distribution of power and responsibilities among the various levels of government.

For the reasons discussed above, I certify that the proposed regulation:

1. Is not a "significant regulatory action" under Executive Order 12866;
2. Is not a "significant rule" under the DOT Regulatory Policies and Procedures (44 FR 11034, February 26, 1979); and
3. Will not have a significant economic impact, positive or negative, on a substantial number of small entities under the criteria of the Regulatory Flexibility Act.

We prepared a regulatory evaluation of the estimated costs to comply with this proposed AD and placed it in the AD docket. See the **ADDRESSES** section for a location to examine the regulatory evaluation.

**List of Subjects in 14 CFR Part 39**

Air transportation, Aircraft, Aviation safety, Safety.

**The Proposed Amendment**

Accordingly, under the authority delegated to me by the Administrator, the FAA proposes to amend 14 CFR part 39 as follows:

**PART 39—AIRWORTHINESS DIRECTIVES**

1. The authority citation for part 39 continues to read as follows:

**Authority:** 49 U.S.C. 106(g), 40113, 44701.

**§ 39.13 [Amended]**

2. The Federal Aviation Administration (FAA) amends § 39.13 by adding the following new airworthiness directive (AD):

**Fokker Services B.V.:** Docket No. FAA–2005–22748; Directorate Identifier 2005–NM–127–AD.

**Comments Due Date**

(a) The FAA must receive comments on this AD action by November 21, 2005.

**Affected ADs**

(b) None.

**Applicability**

(c) This AD applies to Fokker Model F.28 Mark 0070 and 0100 airplanes, certificated in

any category, as identified in Fokker Service Bulletin SBF100–52–069, Revision 3, dated December 18, 2002.

**Unsafe Condition**

(d) This AD results from reports of the airstairs-type passenger door opening during flight. We are issuing this AD to prevent rapid decompression of the airplane, or ejection of a passenger or crew member out the door during flight.

**Compliance**

(e) You are responsible for having the actions required by this AD performed within the compliance times specified, unless the actions have already been done.

**Installation**

(f) Within 30 months after the effective date of this AD, modify the passenger door and install new placards, in accordance with the Accomplishment Instructions of Fokker Service Bulletin SBF100–52–069, Revision 3, dated December 18, 2002; including Manual Change Notification—Operational Documentation MCNO–F100–031, dated December 3, 2001, and Manual Change Notification—Maintenance Documentation MCNO–F100–064, Revision 2, dated December 18, 2002; and including the drawings listed in Table 1 of this AD. To conform to certain Office of the Federal Register requirements for incorporating these materials by reference, the table identifies the date of the service bulletin for undated drawings.

TABLE 1.—DRAWINGS INCLUDED IN SERVICE BULLETIN SBF100–52–069

Fokker drawing	Sheet	Issue	Date
W41074 .....	065	DB .....	December 18, 2002.
W41418 .....	003	L .....	December 18, 2002.
W41418 .....	005	E .....	December 18, 2002.
W41418 .....	006	E .....	December 18, 2002.
W41418 .....	007	E .....	December 18, 2002.
W41418 .....	008	M .....	December 18, 2002.
W42310 .....	001	D .....	August 14, 2000.
W42310 .....	002	B .....	August 14, 2000.
W42310 .....	003	F .....	June 11, 2001.
W59243 .....	024	AU .....	June 12, 2001.
W59261 .....	017	W .....	August 9, 2002.
W59261 .....	025	S .....	July 3, 2001.

**Alternative Methods of Compliance (AMOCs)**

(g)(1) The Manager, International Branch, ANM–116, Transport Airplane Directorate, 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 14 CFR 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**

(h) Dutch airworthiness directive 2002–057, dated April 29, 2002, also addresses the subject of this AD.

Issued in Renton, Washington, on October 14, 2005.

**Kalene C. Yanamura,**

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

[FR Doc. 05–21054 Filed 10–20–05; 8:45 am]

**BILLING CODE 4910–13–P**

**DEPARTMENT OF TRANSPORTATION**

**Office of the Secretary**

**14 CFR Part 382**

**[Docket No. OST–2005–22298]**

**RIN 2105–AC29**

**Nondiscrimination on the Basis of Disability in Air Travel—Medical Oxygen and Portable Respiration Assistive Devices**

**AGENCY:** Office of the Secretary (OST), U.S. Department of Transportation (DOT).

**ACTION:** Extension of comment period on proposed rule.

**SUMMARY:** The Department is extending through January 30, 2006, the period for interested persons to submit comments to its proposed rule on medical oxygen and portable respiration assistive devices.

**DATES:** Comments must be received by January 30, 2006. Comments received after this date will be considered to the extent practicable.

**ADDRESSES:** You may submit comments identified by the docket number [OST-2005-22298] by any of the following methods: (1) Federal eRulemaking Portal: <http://www.regulations.gov> (follow the instructions for submitting comments); (2) Web site: <http://dms.dot.gov> (follow the instructions for submitting comments on the DOT electronic docket site); (3) Fax: 1-202-493-2251; (4) Mail: Docket Management System; U.S. Department of Transportation, 400 Seventh Street, SW., Nassif Building, Room PL-401, Washington, DC 20590-001; or (5) Hand Delivery: To the Docket Management System; Room PL-401 on the plaza level of the Nassif Building, 400 Seventh Street, SW., Washington, DC between 9 a.m. and 5 p.m., Monday through Friday, except Federal Holidays.

You must include the agency name and docket number [OST-2005-22298] or the Regulatory Identification Number (RIN) for this notice at the beginning of your comment. Note that all comments received will be posted without change to <http://dms.dot.gov> including any personal information provided. Please see the Privacy Act section of this document. You may view the public docket through the Internet at <http://dms.dot.gov> or in person at the Docket Management System office at the above address.

**FOR FURTHER INFORMATION CONTACT:** Blane A. Workie, Office of Assistant General Counsel for Aviation Enforcement and Proceedings, 400 7th Street, SW., Room 4116, Washington DC 29590. Phone: 202-366-9342. TTY: 202-366-0511. Fax: 202-366-7152. E-mail: [blane.workie@dot.gov](mailto:blane.workie@dot.gov).

**SUPPLEMENTARY INFORMATION:** On September 7, 2005, the Department of Transportation (DOT or Department) issued a notice of proposed rulemaking (NPRM) that proposed to require airlines to provide in-flight medical oxygen without charge, to test certain respiratory assistive devices and to permit their use if safe. See 70 FR 53108. The NPRM would apply to certain U.S. and foreign air carriers operating to and from the U.S. The

original comment closing date is November 7, 2005.

The Air Carrier Association of America (ACAA), the Air Transport Association (ATA), the National Air Carrier Association (NACA), and the Regional Airline Association (RAA) jointly requested an extension of the comment period to consider "the enormous technical, operational and cost issues raised by the multiple actions required by the NPRM." They requested an extension of more than sixty days to January 30, 2006, at least partially because an extension of sixty days in this rulemaking would place the close of the comment period in the holiday season. This request was supported by comments from the International Air Transport Association (IATA), which further explained that IATA has begun the process of gathering comments from its in-flight, dangerous goods, passenger services, operations, medical and regulatory contacts but that gathering and collating such feedback is a significant task that requires time.

The Department concurs that an extension of the comment period is necessary to allow members of industry sufficient time to analyze the impact of the proposed rule and believes that this extension would result in more thorough comments to the docket than might otherwise be possible without delaying final action in the rulemaking proceeding. We do not anticipate the need for any further extensions. Accordingly, the Department finds that good cause exists to extend the comment period on the proposed rule from November 7, 2005, to January 30, 2006.

Issued in Washington, DC this 17th day of October, 2005, under authority assigned to me by 14 CFR 385.17(c).

**Neil Eisner,**

*Assistant General Counsel for Regulation and Enforcement, U.S. Department of Transportation.*

[FR Doc. 05-21078 Filed 10-20-05; 8:45 am]

**BILLING CODE 4910-62-P**

## ENVIRONMENTAL PROTECTION AGENCY

### 40 CFR Part 158

[OPP-2005-0415; FRL-7734-2]

RIN 2070-AD51

### Pesticides; Data Requirements for Biochemical and Microbial Pesticides; Notification to the Secretary of Agriculture

**AGENCY:** Environmental Protection Agency (EPA).

**ACTION:** Notification to the Secretary of Agriculture.

**SUMMARY:** This document notifies the public that the Administrator of EPA has forwarded to the Secretary of Agriculture a draft proposed rule as required by section 25(a) of the Federal Insecticide, Fungicide, and Rodenticide Act (FIFRA). As described in the Agency's semi-annual Regulatory Agenda, the draft proposed rule updates and revises data requirements for the registration of microbial and biochemical pesticide products to reflect current scientific knowledge and understanding. These data requirements and those already codified in part 158 of title 40 of the Code of Federal Regulations (CFR), are intended to provide EPA with data and other information necessary for the registration of biochemical and microbial pesticide products.

**ADDRESSES:** EPA has established a docket for this action under Docket identification (ID) number OPP-2005-0415. All documents in the docket are listed in the EDOCKET index at <http://www.epa.gov/edocket>. Although listed in the index, some information is not publicly available, i.e., CBI or other information whose disclosure is restricted by statute. Certain other material, such as copyrighted material, is not placed on the Internet and will be publicly available only in hard copy form. Publicly available docket materials are available either electronically in EDOCKET or in hard copy at the Public Information and Records Integrity Branch (PIRIB), Rm. 119, Crystal Mall #2, 1801 S. Bell St., Arlington, VA, Monday through Friday, excluding legal holidays. The Docket telephone number is (703) 305-5805.

**FOR FURTHER INFORMATION CONTACT:** Candace Brassard, Field and External Affairs Division (7506C), Office of Pesticide Programs, Environmental Protection Agency, 1200 Pennsylvania Ave., NW., Washington DC 20460-0001; telephone number: 703-305-6598; e-mail address:

[brassard.candace@epa.gov](mailto:brassard.candace@epa.gov).

### SUPPLEMENTARY INFORMATION:

#### I. General Information

##### A. Does This Action Apply to Me?

This action is directed to the public in general. It simply announces the submission of a draft proposed rule to USDA and does not otherwise affect any specific entities. This action may, however, be of particular interest to producers or registrants of a biochemical or microbial pesticide product. This proposal also may affect