(g) Other FAA AD Provisions

The following provisions also apply to this AD:

(1) *Alternative Methods of Compliance (AMOCs):* The Manager, Standards Office, FAA, has the authority to approve AMOCs for this AD, if requested using the procedures found in 14 CFR 39.19. Send information to ATTN: Albert Mercado, Aerospace Engineer, FAA, Small Airplane Directorate, 901 Locust, Room 301, Kansas City, Missouri 64106; telephone: (816) 329–4119; fax: (816) 329–4090; email: *albert.mercado@faa.gov.* Before using any approved AMOC on any airplane to which the AMOC applies, notify your appropriate principal inspector (PI) in the FAA Flight Standards District Office (FSDO), or lacking a PI, your local FSDO.

(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.

(3) *Reporting Requirements:* For any reporting requirement in this AD, a federal agency may not conduct or sponsor, and a person is not required to respond to, nor shall a person be subject to a penalty for failure to comply with a collection of information subject to the requirements of the Paperwork Reduction Act unless that collection of information displays a current valid OMB Control Number. The OMB Control Number for this information collection is 2120–0056. Public reporting for this collection of information is estimated to be approximately 5 minutes per response, including the time for reviewing instructions, completing and reviewing the collection of information. All responses to this collection of information are mandatory. Comments concerning the accuracy of this burden and suggestions for reducing the burden should be directed to the FAA at: 800 Independence Ave. SW., Washington, DC 20591, Attn: Information Collection Clearance Officer, AES–200.

(h) Related Information

Refer to MCAI European Aviation Safety Agency (EASA) AD No. 2012–0202, dated October 1, 2012; and Reims Aviation S.A. Service Bulletin No. F406–74, dated September 26, 2012, for related information. For service information related to this AD, contact Reims Aviation Industries, Aérodrome de Reims Prunay, 51360 Prunay, France; telephone + 33 3 26 48 46 65; fax + 33 3 26 49 18 57; email: *stephan.lapagne@reims-aviation.fr;* Internet: *www.geciaviation.com/en/f406.html.* You may review copies of the referenced service information at the FAA, Small Airplane Directorate, 901 Locust, Kansas City, Missouri 64106. For information on the availability of this material at the FAA, call (816) 329–4148.

Issued in Kansas City, Missouri, on November 29, 2012.

Earl Lawrence,

Manager, Small Airplane Directorate, Aircraft Certification Service.

[FR Doc. 2012–29395 Filed 12–4–12; 8:45 am]

BILLING CODE 4910–13–P

DEPARTMENT OF COMMERCE

Minority Business Development Agency

15 CFR Part 1400

[Docket No. 121130667–2667–01]

Petition for Inclusion of the Arab-American Community in the Groups Eligible for MBDA Services

AGENCY: Minority Business Development Agency, Commerce.

ACTION: Notice of proposed rulemaking and request for comments; amendment.

SUMMARY: The Minority Business Development Agency (MBDA) publishes this notice to extend the date on which it plans to make its decision on a petition from the American-Arab Anti-Discrimination Committee requesting formal designation as a group eligible for MBDA's services from November 30, 2012 to March 1, 2013.

FOR FURTHER INFORMATION CONTACT: For further information about this Notice, contact Josephine Arnold, Minority Business Development Agency, 1401 Constitution Avenue NW., Room 5053, Washington, DC 20230, (202) 482–5461.

SUPPLEMENTARY INFORMATION: On May 30, 2012, the Minority Business Development Agency (MBDA) published a notice of proposed rulemaking and request for comments regarding a petition received on January 11, 2012 from the American-Arab Anti-Discrimination Committee (ADC) requesting formal designation of Arab-Americans as a minority group that is socially or economically disadvantaged pursuant to 15 CFR part 1400. MBDA has published several notices in the **Federal Register** to extend the date for making a decision on the merits of the petition. On September 4, 2012, MBDA published an amendment to extend the deadline for the decision until November 30, 2012. The Agency has determined that an additional ninety (90) day period for consideration of the policy implications associated with the petition is necessary. Therefore, the Agency has determined that the time in which it will make its decision on the petition will be on or before March 1, 2013. This extension will not prejudice the petitioner.

Minority Business Development Agency.

David Hinson,

National Director.

[FR Doc. 2012–29431 Filed 12–4–12; 8:45 am]

BILLING CODE 3510–21–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Parts 500, 520, 522, 524, 529, 556, and 558

[Docket No. FDA–2012–N–1067]

RIN 0910–AG17

New Animal Drugs; Updating Tolerances for Residues of New Animal Drugs in Food

AGENCY: Food and Drug Administration, HHS.

ACTION: Proposed rule.

SUMMARY: The Food and Drug Administration (FDA) is proposing to revise the animal drug regulations regarding tolerances for residues of approved and conditionally approved new animal drugs in food by standardizing, simplifying, and clarifying the determination standards and codification style. In addition, we are proposing to add definitions for key terms. The purpose of the revision is to enhance understanding of tolerance determination and improve the readability of the regulations.

DATES: Submit either electronic or written comments by March 5, 2013. See section VI of this document for the proposed effective date of a final rule based on this proposed rule.

ADDRESSES: You may submit comments, identified by Docket No. FDA–2012–N–1067 and RIN number 0910–AG17, 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 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