(1) When preparing an EA, EIS, or FONSI, USMS personnel in charge of preparing the document will invite comment from affected Federal, tribal, State, local agencies, and other interested persons, as early as the scoping process;

(2) The USMS will disseminate information to potentially interested or affected parties, such as local communities and Indian tribes, through such means as news releases to various local media, announcements to local citizens groups, public hearings, and posted signs near the affected area;

(3) The USMS will mail notice to those individuals or groups who have requested one on a specific action or similar actions;

(4) For matters of national concern, the USMS will publish notification in the **Federal Register**, and will send notification by mail to national organizations reasonably expected to be interested;

(5) If a decision is made to develop an EIS, the USMS will publish a NOI in the **Federal Register** as soon as possible;

(6) The personnel in charge of preparing the NEPA analysis and documentation will invite public comment and maintain two-way communication channels throughout the NEPA process, provide explanations of where interested parties can obtain information on status reports of the NEPA process and other relevant documents, and keep all public affairs officers informed;

(7) The USMS will establish a Web site to keep the public informed; and

(8) During the NEPA process, responsible personnel will consult with local government and tribal officials, leaders of citizen groups, and members of identifiable population segments within the potentially affected environment, such as farmers and ranchers, homeowners, small business owners, minority and disadvantaged communities, and tribal members.

10. Scoping

Prior to starting the NEPA analysis, USMS personnel responsible for preparing either an EA or EIS, shall engage in an early scoping process to identify the significant issues to be examined in depth, and to identify and eliminate from detailed study those issues which are not significant or which have been adequately addressed by prior environmental review. The scoping process should identify any other environmental analyses being conducted relevant to the proposed action, address timing and set time limits with respect to the NEPA process, set page limits, designate respective responsibilities among the lead and cooperating agencies, identify any other environmental review and consultation requirements to allow for integration with the NEPA analysis, and hold an early scoping meeting that may be integrated with other initial planning meetings.

11. Mitigation and Monitoring

USMS personnel, who are responsible for preparing NEPA analyses and documents, will consider mitigation measures to avoid or minimize environmental harm. EAs and EISs will consider reasonable mitigation measures relevant to the proposed action and alternatives. Paragraph 5(b) of this Appendix describes the requirements for documenting mitigation measures in a ROD.

12. Supplementing an EA or EIS

When substantial changes are made to a proposed action that is relevant to environmental concerns, a supplement will be prepared for an EA or a draft or a final EIS. A supplement will also be prepared when significant new circumstances arise or new relevant information surfaces concerning and bearing upon the proposed action or its impacts. Any necessary supplement shall be processed in the same way as an original EA or EIS, with the exception that new scoping is not required. Any supplement shall be added to the formal administrative record, if such record exists.

13. Compliance With Other Environmental Statutes

To the extent practicable, a NEPA document shall include information necessary to assure compliance with all applicable environmental statutes.

Dated: November 8, 2006.

John F. Clark,

Director, United States Marshals Service.

[FR Doc. E6–20940 Filed 12–7–06; 8:45 am]

BILLING CODE 4410–04–P

DEPARTMENT OF DEFENSE

Department of the Army

32 CFR Parts 626 and 627

The Biological Defense Safety Program and Technical Safety Requirements

AGENCY: Department of the Army, DOD.

ACTION: Final rule; removals.

SUMMARY: The Department of the Army is removing its regulations concerning the biological Defense Safety Program and its requirements because it is now superseded through consolidation with other Army safety regulations into Army Regulation (AR) 385–10, Army Safety Program and does not affect the general public.

EFFECTIVE DATE: December 8, 2006.

FOR FURTHER INFORMATION CONTACT: Mr. Kenneth W. Proper, (703) 601–2408.

SUPPLEMENTARY INFORMATION: The Office of the Director of Army Safety (DACS–SF), is the proponent for the regulation represented by 32 CFR Part 626 and the DA PAM represented by 32 CFR 627. The Office of the Director of Army Safety has consolidated the Army regulation, represented by 32 CFR Part 626 into AR 385–10, Army Safety Program. This regulation was extensively revised during the consolidation process, and the new consolidated regulation does not affect the general public.

The Office of the Director of Army Safety has extensively revised the DA PAM, represented by 32 CFR 627 to reflect the consolidation effect and to update it to address new biological safety techniques and requirements and determined that the revised DA PAM does not affect the general public.

List of Subjects in 32 CFR Parts 626 and 627

Biologics, Government contracts, Hazardous substances, National defense, Occupational safety and health, Research.

PART 626 AND 627—[REMOVED]

■ Accordingly, for reasons stated in the preamble, under the authority of 5 U.S.C. 102, 10 U.S.C. 21, 111, 151–158, 42 U.S.C. 216; sec. 361, 50 U.S.C. 1431, Pub. L. 101–510, 104 Stat. 1516, 58 Stat. 703 and 264; 49 U.S.C. App 1803, 1804, 1807, and 1808, 29 CFR 1910. 1450(e), 32 CFR Part 626, Biological Defense Safety Program and 32 CFR Part 627, The Biological Defense Safety Program, Technical Safety Requirements (DA Pamphlet 385–69), are removed in their entirety.

Brenda S. Bowen,

Army Federal Register Liaison Officer.

[FR Doc. 06–9598 Filed 12–7–06; 8:45 am]

BILLING CODE 3710–08–M

DEPARTMENT OF DEFENSE

Department of the Army

32 CFR Part 656

Installations, Use of Off-Road Vehicles on Army Land

AGENCY: Department of the Army, DOD.

ACTION: Final rule; removal.

SUMMARY: The Department of the Army is resending AR 385–55, Prevention of Motor Vehicle Accidents, and has consolidated its requirements into AR 385–10, Army Safety Program. During consolidation, the section concerning the use of non-tactical off-road vehicles on Army land was removed.

EFFECTIVE DATE: December 8, 2006.

FOR FURTHER INFORMATION CONTACT: Mr. Kenneth W. Proper, (703) 601–2408.

SUPPLEMENTARY INFORMATION: The Office of the Director of Army Safety (DACS–SF), is the proponent for the regulation represented by 32 CFR Part 656. The Office of the Director of Army Safety has consolidated the Army regulation, represented by 32 CFR Part 656 into AR 385–10, Army Safety Program. This regulation was extensively revised

during the consolidation process, and the new consolidated regulation does not affect the general public. Similar requirements concerning the use of off-road vehicles on Army land are now provided by 32 CFR 650, Environmental Protection and Enhancement (AR 200–1) and 32 CFR 651, Environmental Analysis Of Army Actions (AR 200–2) which when taken into combination provided greater and wider protection on installation than did 32 CFR Part 656 or AR 385–55.

List of Subjects in 32 CFR Part 656

Environmental protection, Federal buildings and facilities, Traffic regulations.

PART 656—[REMOVED]

■ Accordingly, for reasons stated in the preamble, under the authority 10 U.S.C. 3012, 32 CFR Part 656, Installations, Use of Off-Road Vehicles on Army Land, is removed in its entirety.

Brenda S. Bowen,

Army Federal Register Liaison Officer.

[FR Doc. 06–9599 Filed 12–8–06; 8:45 am]

BILLING CODE 3710–08–M

ENVIRONMENTAL PROTECTION AGENCY

40 CFR Part 180

[EPA–HQ–OPP–2006–0654; FRL–8093–4]

Cyproconazole; Pesticide Tolerances for Emergency Exemptions

AGENCY: Environmental Protection Agency (EPA).

ACTION: Final rule.

SUMMARY: This regulation establishes a time-limited tolerance for residues of cyproconazole ((2RS,3RS)-2-(4-chlorophenyl)-3-cyclopropyl-1-(1H -1,2,4- triazole-1-yl)butan-2-ol) in or on soybean seed. This action is associated with EPA's granting of an emergency exemption under section 18 of the Federal Insecticide, Fungicide, and Rodenticide Act (FIFRA) authorizing use of the pesticide on soybeans. This regulation establishes a maximum permissible level for residues of cyproconazole in this food commodity. The tolerance will expire and be revoked on December 31, 2009.

DATES: This regulation is effective December 8, 2006. Objections and requests for hearings must be received on or before February 6, 2007, and must be filed in accordance with the instructions provided in 40 CFR part 178 (see also Unit I.C. of the **SUPPLEMENTARY INFORMATION**).

ADDRESSES: EPA has established a docket for this action under docket identification (ID) number EPA–HQ–OPP–2006–0654. All documents in the docket are listed on the regulations.gov website. Although listed in the index, some information is not publicly available, e.g., Confidential Business Information (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 in the electronic docket at *http://www.regulations.gov*, or, if only available in hard copy, at the Office of Pesticide Programs (OPP) Regulatory Public Docket in Room S-4400, One Potomac Yard (South Building), 2777 South Crystal Drive Arlington, VA 22202–3553. The hours of operation of this Docket Facility are from 8:30 a.m. to 4 p.m., Monday through Friday, excluding legal holidays. The Docket telephone number is (703) 305–5805.

FOR FURTHER INFORMATION CONTACT: Carmen Rodia, Registration Division (7505P), Office of Pesticide Programs, Environmental Protection Agency, 1200 Pennsylvania Ave., NW., Washington, DC 20460–0001; telephone number: (703) 306–0327; fax: (703) 308–8041; e-mail address: *rodia.carmen@epa.gov*.

SUPPLEMENTARY INFORMATION:

I. General Information

A. Does this Action Apply to Me?

You may be potentially affected by this action if you are an agricultural producer, food manufacturer, or pesticide manufacturer. Potentially affected entities may include, but are not limited to:

- Crop production (NAICS code 111).
- Animal production (NAICS code 112).
- Food manufacturing (NAICS code 311).
- Pesticide manufacturing (NAICS code 32532).

This listing is not intended to be exhaustive, but rather provides a guide for readers regarding entities likely to be affected by this action. Other types of entities not listed in this unit could also be affected. The North American Industrial Classification System (NAICS) codes have been provided to assist you and others in determining whether this action might apply to certain entities. If you have any questions regarding the applicability of this action to a particular entity, consult the person listed under **FOR FURTHER INFORMATION CONTACT**.

B. How Can I Access Electronic Copies of this Document?

In addition to accessing an electronic copy of this **Federal Register** document through the electronic docket at *http://www.regulations.gov*, you may access this **Federal Register** document electronically through the EPA Internet under the ''**Federal Register**'' listings at *http://www.epa.gov/fedrgstr*. You may also access a frequently updated electronic version of 40 CFR part 180 through the Government Printing Office's pilot e-CFR site at *http://www.gpoaccess.gov/ecfr*.

C. Can I File an Objection or Hearing Request?

Under section 408(g) of the FFDCA, as amended by the FQPA, any person may file an objection to any aspect of this regulation and may also request a hearing on those objections. The EPA procedural regulations which govern the submission of objections and requests for hearings appear in 40 CFR part 178. You must file your objection or request a hearing on this regulation in accordance with the instructions provided in 40 CFR part 178. To ensure proper receipt by EPA, you must identify docket ID number EPA–HQ–OPP–2006–0654 in the subject line on the first page of your submission. All requests must be in writing, and must be mailed or delivered to the Hearing Clerk on or before February 6, 2007.

In addition to filing an objection or hearing request with the Hearing Clerk as described in 40 CFR part 178, please submit a copy of the filing that does not contain any CBI for inclusion in the public docket that is described in **ADDRESSES**. Information not marked confidential pursuant to 40 CFR part 2 may be disclosed publicly by EPA without prior notice. Submit your copies, identified by docket ID number EPA–HQ–OPP–2006–0654, by one of the following methods:

- Federal eRulemaking Portal: *http://www.regulations.gov*. Follow the on-line instructions for submitting comments.
- *Mail*: Office of Pesticide Programs (OPP), Regulatory Public Docket (7502P), Environmental Protection Agency, 1200 Pennsylvania Avenue, NW., Washington, DC 20460–0001.
- *Delivery*: OPP Regulatory Public Docket (7502P), Environmental Protection Agency, Room S-4400, One Potomac Yard (South Building), 2777 South Crystal Drive, Arlington, VA 22202–3553. Deliveries are only accepted during the Docket's normal hours of operation (8:30 a.m. to 4 p.m., Monday through Friday, excluding legal holidays). Special arrangements should