the imposition of a control in the form of a licensing requirement for exports and reexports of all items subject to the EAR destined to the nine additional entities related to Mayrow General Trading that were added to General Order No. 3 with the September 6, 2006, final rule.

Since August 21, 2001, the Act has been in lapse and the President, through Executive Order 13222 of August 17, 2001 (3 CFR, 2001 Comp. 783 (2002)), as extended most recently by the Notice of August 3, 2006 (71 FR 44551 (August 7, 2006)), has continued the EAR in effect under the International Emergency Economic Powers Act (50 U.S.C. 1701–1706 (2000)) (''IEEPA''). BIS continues to carry out the provisions of the Act, as appropriate and to the extent permitted by law, pursuant to Executive Order 13222.

Rulemaking Requirements

1. This rule has been determined to be not significant for purposes of Executive Order 12866.

2. Notwithstanding any other provision of law, no person is required to respond to nor be subject to a penalty for failure to comply with a collection of information, subject to the requirements of the Paperwork Reduction Act of 1995 (44 U.S.C. 3501 et seq.) (PRA), unless that collection of information displays a currently valid Office of Management and Budget (OMB) Control Number. This regulation involves collections previously approved by the OMB under control numbers 0694–0088, ''Multi-Purpose Application,'' which carries a burden hour estimate of 58 minutes to prepare and submit form BIS–748. Miscellaneous and recordkeeping activities account for 12 minutes per submission. Total burden hours associated with the Paperwork Reduction Act and Office and Management and Budget control number 0694–0088 are expected to increase slightly as a result of this rule. Send comments regarding these burden estimates or any other aspect of this collection of information, including suggestions for reducing the burden, to David Rostker, OMB Desk Officer, by e-mail at *david_rostker@omb.eop.gov* or by fax to (202) 395–7285; and to the Regulatory Policy Division, Bureau of Industry and Security, Department of Commerce, P.O. Box 273, Washington, DC 20044, e-mail: *publiccomments@bis.doc.gov.*

3. This rule does not contain policies with Federalism implications as that term is defined in Executive Order 13132.

4. The provisions of the Administrative Procedure Act (5 U.S.C. 553) requiring notice of proposed rulemaking, the opportunity for public participation, and a delay in effective date, are inapplicable because this regulation involves a military or foreign affairs function of the United States. (*See* 5 U.S.C. 553(a)(1)) Further, no other law requires that a notice of proposed rulemaking and an opportunity for public comment be given for this rule. Because a notice of proposed rulemaking and an opportunity for public comment are not required to be given for this rule by 5 U.S.C. 553, or by any other law, the analytical requirements of the Regulatory Flexibility Act, 5 U.S.C. 601 *et seq.*, are not applicable.

List of Subjects in 15 CFR Part 736

Exports, Foreign trade.

■ Accordingly, part 736 of the Export Administration Regulations (15 CFR part 736) is corrected by making the following correcting amendment:

PART 736—[CORRECTED]

■ 1. The authority citation for 15 CFR part 736 is revised to read as follows:

Authority: 50 U.S.C. app. 2401 *et seq.*; 50 U.S.C. 1701 *et seq.*; 22 U.S.C. 2151 (note), Public Law 108–175; E.O. 12938, 59 FR 59099, 3 CFR, 1994 Comp., p. 950; E.O. 13020, 61 FR 54079, 3 CFR, 1996 Comp. p. 219; E.O. 13026, 61 FR 58767, 3 CFR, 1996 Comp., p. 228; E.O. 13222, 66 FR 44025, 3 CFR, 2001 Comp., p. 783; E.O. 13338, 69 FR 26751, May 13, 2004; Notice of October 25, 2005, 70 FR 62027 (October 27, 2005); Notice of August 3, 2006, 71 FR 44551 (August 7, 2006).

■ 2. General Order 3 to Supplement No. 1 to part 736, paragraph (a)(1) is correctly revised to read as follows:

Supplement No. 1 to Part 736—General Orders

* * * * *

(a) *License requirements.* (1) Effective June 5, 2006, a license is required to export or reexport any item subject to the EAR to Mayrow General Trading or entities related, as follows: Micatic General Trading; Majidco Micro Electronics; Atlinx Electronics; Micro Middle East Electronics; Narinco; Farrokh Nia Yaghmaei, a.k.a., Farrokh Nia Yaghmayi; and H. Ghasir. Mayrow General Trading and all entities related described in paragraph (a)(1) are located in Dubai, United Arab Emirates. This license requirement applies with respect to any transaction in which any of the above-named entities will act as purchaser, intermediate consignee, ultimate consignee, or end-user of the items.

* * * * *

Eileen Albanese,

Director, Office of Exporter Services.

[FR Doc. E6–15135 Filed 9–12–06; 8:45 am]

BILLING CODE 3510–33–P

DEPARTMENT OF ENERGY

Federal Energy Regulatory Commission

18 CFR Part 35

[Docket No. RM02–12–002]

Standardization of Small Generator Interconnection Agreements and Procedures

AGENCY: Federal Energy Regulatory Commission, DOE.

ACTION: Order on clarification; correction.

SUMMARY: This document corrects an error in an Order on Clarification that the Federal Energy Regulatory Commission published in the **Federal Register** on July 27, 2006. The Order on Clarification erroneously omitted text from two sections within the appendices of the document.

DATES: *Effective Date:* August 28, 2006.

FOR FURTHER INFORMATION CONTACT: Michael G. Henry (Legal Information), Office of the General Counsel, Federal Energy Regulatory Commission at (202) 502–8532.

SUPPLEMENTARY INFORMATION: In FR Document E6–11989, published July 27, 2006 (71 FR 42587) make the following correction to appendices 2 and 3 of the document:

Appendix 2: Revised System Impact Study Agreement and Appendix 3: Revised Facilities Study Agreement [Corrected]

On page 42591, column 2, section ''21.0 *Reservation of Rights* and on page 42592, column 3, section ''19.0 *Reservation of Rights,* the language is corrected to read as follows for both of these sections:

''*Reservation of Rights:* The Transmission Provider shall have the right to make a unilateral filing with FERC to modify this Agreement with respect to any rates, terms and conditions, charges, classifications of service, rule or regulation under section 205 of any other applicable provision of the Federal Energy Power Act and FERC rules and regulations thereunder, and the Interconnection Customer shall have

the right to make a unilateral filing with FERC to modify this Agreement under any applicable provision of the Federal Power Act and FERC's rules and regulations; provided that each party shall have the right to protest any such filing by the other Party and to participate fully in any proceeding before FERC in which such modifications may be considered. Nothing in this Agreement shall limit the rights of the parties or of FERC under sections 205 or 206 of the Federal Power Act and FERC's rules and regulations, except to the extent that the parties otherwise agree as provided herein.''

Magalie R. Salas,

Secretary.

[FR Doc. E6–15126 Filed 9–12–06; 8:45 am]

BILLING CODE 6717–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 558

New Animal Drugs for Use in Animal Feeds; Chlortetracycline

AGENCY: Food and Drug Administration, HHS.

ACTION: Final rule, technical amendment.

SUMMARY: The Food and Drug Administration (FDA) is amending the animal drug regulations to reflect approval of a supplemental new animal drug application (NADA) filed by Alpharma Inc. The supplemental NADA provides for use of an approved Type A medicated article containing chlortetracycline to formulate a free-choice loose mineral Type C medicated feed for beef and nonlactating dairy cattle.

DATES: This rule is effective September 13, 2006.

FOR FURTHER INFORMATION CONTACT: Eric S. Dubbin, Center for Veterinary Medicine (HFV–126), Food and Drug Administration, 7500 Standish Pl., Rockville, MD 20855, 301–827–0232, e-mail: *eric.dubbin@fda.hhs.gov*.

SUPPLEMENTARY INFORMATION: Alpharma Inc., One Executive Dr., Fort Lee, NJ 07024, filed NADA 48–761 for use of AUREOMYCIN 90 Granular (chlortetracycline) Type A medicated article to formulate a free-choice loose mineral Type C medicated feed for beef and nonlactating dairy cattle as an aid in the control of active infection of anaplasmosis caused by *Anaplasma marginale* susceptible to chlortetracycline. The supplemental NADA is approved as of July 28, 2006, and the regulations are amended in 21 CFR 558.128 to reflect the approval. The basis of approval is discussed in the freedom of information summary.

In accordance with the freedom of information provisions of 21 CFR part 20 and 21 CFR 514.11(e)(2)(ii), a summary of safety and effectiveness data and information submitted to support approval of this application may be seen in the Division of Dockets Management (HFA–305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852, between 9 a.m. and 4 p.m., Monday through Friday.

The agency has determined under 21 CFR 25.33(a)(1) that this action is of a type that does not individually or cumulatively have a significant effect on the human environment. Therefore, neither an environmental assessment nor an environmental impact statement is required.

This rule does not meet the definition of ''rule'' in 5 U.S.C. 804(3)(A) because it is a rule of ''particular applicability.'' Therefore, it is not subject to the congressional review requirements in 5 U.S.C. 801–808.

List of Subjects in 21 CFR Part 558

Animal drugs, Animal feeds.

■ Therefore, under the Federal Food, Drug, and Cosmetic Act and under authority delegated to the Commissioner of Food and Drugs and redelegated to the Center for Veterinary Medicine, 21 CFR part 558 is amended as follows:

PART 558—NEW ANIMAL DRUGS FOR USE IN ANIMAL FEEDS

■ 1. The authority citation for 21 CFR part 558 continues to read as follows:

Authority: 21 U.S.C. 360b, 371.

■ 2. In § 558.128, redesignate paragraph (e)(6) as paragraph (e)(7); and add new paragraph (e)(6) to read as follows:

§ 558.128 Chlortetracycline.

* * * * *

(e) * * *

(6) It is used as a free-choice, loose mineral Type C feed as follows:

(i) *Specifications*.

Ingredient	Percent	International Feed No.
Dicalcium Phosphate	46.20	6–26–335
Sodium Chloride (Salt)	15.00	6–04–152
Magnesium Oxide	10.67	6–02–756
Cottonseed Meal	10.00	5–01–625
Trace Mineral/Vitamin Premix[1]	3.80	
Calcium Carbonate	3.50	6–01–069
Dried Cane Molasses	3.00	4–04–695
Potassium Chloride	2.00	6–03–755
Mineral Oil	2.00	8–03–123
Iron Oxide	0.50	6–02–431
Chlortetracycline Type A medicated article (90 gram/lb)	3.33	

[1]Content of vitamin and trace mineral premixes may be varied. However, they should be comparable to those used for other free-choice feeds. Formulation modifications require FDA approval prior to marketing. Selenium must comply with 21 CFR 573.920. Ethylenediamine dihydroiodide (EDDI) should comply with FDA Compliance Policy Guides Sec. 651.100 (CPG 7125.18).

(ii) *Amount*. 6,000 grams per ton.

(iii) *Indications for use*. Beef and nonlactating dairy cattle: As an aid in the control of active infection of anaplasmosis caused by *Anaplasma marginale* susceptible to chlortetracycline.

(iv) *Limitations*. Feed continuously on a free-choice basis at a rate of 0.5 to 2.0 mg chlortetracycline per head per day.

(v) *Sponsor*. See No. 046573 in § 510.600(c) of this chapter.

* * * * *

Dated: August 30, 2006.

Steven D. Vaughn,

Director, Office of New Animal Drug Evaluation, Center for Veterinary Medicine.

[FR Doc. E6–15103 Filed 9–12–06; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF THE TREASURY

Internal Revenue Service

26 CFR Part 54

[TD 9277]

RIN 1545–BE30

Employer Comparable Contributions to Health Savings Accounts Under Section 4980G; Correction

AGENCY: Internal Revenue Service (IRS), Treasury.

ACTION: Correcting amendment.

SUMMARY: This document contains a correction to final regulations (TD 9277) that were published in the **Federal**