(iii) *Free-choice feeds*—(A) *Amount.* 5 mg/kg body weight (2.27 mg/lb), including the following formulations:

Ingredient[1]	Percent	International Feed No.
(*1*) Free-choice, dry Type C feed:		
Salt (sodium chloride)	59.00	6–04–152
Monosodium phosphate	31.16	6–04–288
Dried cane molasses	3.12	4–04–695
Zinc sulfate	0.76	6–05–556
Copper sulfate	0.45	6–01–720
Fenbendazole 20% Type A article	5.51	n/a
(*2*) Free-choice, dry Type C feed:		
Salt (sodium chloride)	35.93	6–04–152
Dicalcium phosphate (18.5% P)	32.44	6–00–080
Calcium carbonate (38% Ca)	15.93	6–01–069
Magnesium oxide (56% Mg)	10.14	6–02–756
Zinc sulfate	1.47	6–05–556
Mineral oil	1.00	8–03–123
Dried cane molasses (46% sugars)	0.98	4–04–695
Potassium iodide	0.01	6–03–759
Fenbendazole 20% Type A article	2.10	n/a
(*3*) Free-choice, liquid Type C feed:		
Cane molasses[2]	80.902	4–13–251
Water	9.36	n/a
Urea solution, 55%	7.05	5–05–707
Phosphoric acid 75% (feed grade)	2.00	6–03–707
Xantham gum	0.20	8–15–818
Trace minerals	0.20	n/a
Vitamin premix	0.01	n/a
Fenbendazole 20% Type A article	0.278	n/a

[1]The content of any added vitamin and trace mineral may be varied; however, they should be comparable to those used by the manufacturer for other free-choice cattle feeds. Formulation modifications require FDA approval prior to marketing. Selenium is not approved for the free-choice formulations described in paragraph (e)(3)(iii) of this section. Free-choice cattle feeds containing selenium must comply with published regulations (*see* 21 CFR 573.920).

[2]The percentage of cane molasses and water in the formulation may be adjusted as needed in order to bring the brix value of the molasses to the industry standard of 79.5 brix.

(B) *Indications for use.* As in paragraph (e)(3)(i) of this section.

(C) *Limitations.* Feed a total of 5 mg of fenbendazole per kg (2.27 mg/lb) of body weight to cattle over a 3- to 6-day period. Retreatment may be needed after 4 to 6 weeks. Cattle must not be slaughtered within 13 days following last treatment. For dairy cattle the milk discard time is zero hours. A withdrawal period has not been established for this product in pre-ruminating calves. Do not use in calves to be processed for veal.

* * * * *

Dated: September 29, 2008.

Bernadette Dunham,

Director, Center for Veterinary Medicine.

[FR Doc. E8–23845 Filed 10–7–08; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 801

[Docket No. FDA–2008–N–0148]

Medical Devices; Hearing Aids; Technical Data Amendments; Confirmation of Effective Date

AGENCY: Food and Drug Administration, HHS.

ACTION: Direct final rule; confirmation of effective date.

SUMMARY: The Food and Drug Administration (FDA) is confirming the effective date of October 15, 2008, for the final rule that appeared in the **Federal Register** of June 2, 2008 (73 FR 31358). The direct final rule amends the hearing aid labeling to reference the most recent version of the consensus standard used to determine the technical data to be included in labeling for hearing aids. This document confirms the effective date of the direct final rule.

DATES: Effective date confirmed: October 15, 2008.

FOR FURTHER INFORMATION CONTACT: Eric A. Mann, Center for Devices and Radiological Health (HFZ–460), Food and Drug Administration,9200 Corporate Blvd., Rockville, MD 20850, 240–276–4242.

SUPPLEMENTARY INFORMATION: In the **Federal Register** of June 2, 2008 (73 FR 31358), FDA solicited comments concerning the direct final rule for a 75-day period ending August 18, 2008. FDA stated that the effective date of the direct final rule would be on October 15, 2008, 60 days after the end of the comment period, unless any significant adverse comment was submitted to FDA during the comment period. FDA received one letter of comment on the direct final rule. However, this comment does not constitute a significant adverse comment. Therefore, FDA is confirming the effective date of the direct final rule. The comment received and FDA's response to the comment are discussed as follows:

The only comment on the direct final rule requested clarification regarding the applicability of the proposed change in the standard of the American National Standards Institute to hearing aid models that were tested and characterized prior to the effective date

of the direct final rule on October 15, 2008. The comment interpreted the proposed change as being only prospectively applied to new models undergoing test procedures on or after the effective date of the proposed change. FDA agrees that the proposed change applies only to new hearing aid models undergoing characterization on or after the effective date of October 15, 2008; hearing aid models tested prior to this date are subject only to the characterization standard cited in the regulation at the time they were tested.

Authority: Therefore, under the Federal Food, Drug, and Cosmetic Act and the Public Health Service Act, and under authority delegated to the Commissioner of Food and Drugs, the amendments issued thereby become effective on October 15, 2008.

Dated: October 2, 2008.

Jeffrey Shuren,

Associate Commissioner for Policy and Planning.

[FR Doc. E8–23717 Filed 10–7–08; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF THE INTERIOR

Minerals Management Service

[Docket No. MMS–2008–MRM–0021]

30 CFR Part 210

RIN 1010–AD20

Reporting Amendments

AGENCY: Minerals Management Service (MMS), Interior.

ACTION: Correcting amendment.

SUMMARY: The MMS published a final rule in the **Federal Register** on Wednesday, March 26, 2008 (73 FR 15885), announcing amendments to existing regulations for reporting production and royalties on oil, gas, coal and other solid minerals, and geothermal resources produced from Federal and Indian leases. This docutment corrects the final rule, which contained a clerical error in the tables identifying OMB-approved information collections and their corresponding forms.

DATES: *Effective Date:* Effective on October 8, 2008.

FOR FURTHER INFORMATION CONTACT: Hyla Hurst, Regulatory Specialist, Minerals Management Service, Minerals Revenue Management, P.O. Box 25165, MS 302B2, Denver, Colorado 80225; telephone (303) 231–3495; or e-mail *Hyla.Hurst@mms.gov.*

SUPPLEMENTARY INFORMATION: A final rule was published in the **Federal Register** on March 26, 2008 (73 FR 15885) containing a clerical error in the preamble and the regulatory text in the tables listing OMB-approved information collections. The forms approved under OMB Control Number 1010–0139 were incorrectly identified on page 15889 in the preamble and page 15893 in the regulatory text. Both tables contain the same error. Form MMS–4054 (Parts A, B, and C) and Form MMS–4058 are correctly identified as shown below in the table at § 210.10.

List of Subjects in 30 CFR Part 210

Coal, Solid minerals, Continental Shelf, Electronic funds transfers, Geothermal energy, Government contracts, Indian lands, Mineral royalties, Natural gas, Penalties, Petroleum, Oil and gas, Public lands—mineral resources, Reporting and recordkeeping requirements.

■ Accordingly, 30 CFR Part 210 is corrected by making the following amendments:

PART 210—FORMS AND REPORTS

■ 1. The authority citation for part 210 continues to read as follows:

Authority: 5 U.S.C. 301 et. seq. ; 25 U.S.C. 396, 2107; 30 U.S.C. 189, 190, 359, 1023, 1751(a); 31 U.S.C. 3716, 9701; 43 U.S.C. 1334, 1801 et. seq. ; and 44 U.S.C. 3506(a).

■ 2. In § 210.10, the table is amended by revising the entry for OMB number 1010–0139 to read as follows:

§ 210.10 What are the OMB-approved information collections?

* * * * *

OMB Control No. and short title	Form or information collected
* *	* * * * * * *
1010–0139, 30 CFR Parts 210 and 216, Production Accounting.	Form MMS–4054 (Parts A, B, and C), Oil and Gas Operations Report. Form MMS–4058, Production Allocation Schedule Report.
* *	* * * * * * *

Dated: September 30, 2008.

C. Stephen Allred,

Assistant Secretary for Land and Minerals Management.

[FR Doc. E8–23788 Filed 10–7–08; 8:45 am]

BILLING CODE 4310–MR–P

DEPARTMENT OF VETERANS AFFAIRS

38 CFR Part 17

RIN 2900–AM95

Dental Care—Provision of One-Time Outpatient Dental Care for Certain Veterans

AGENCY: Department of Veterans Affairs.

ACTION: Final rule.

SUMMARY: The Department of Veterans Affairs (VA) is amending its regulations regarding the authority to provide one-time outpatient dental treatment to eligible veterans following discharge or release from active duty. In section 1709 of Public Law 110–181, the National Defense Authorization Act for Fiscal Year 2008, Congress amended the eligibility criteria for the one-time dental treatment benefit. This rule is necessary to incorporate the statutory amendments into VA regulations.

DATES: *Effective Date:* October 8, 2008.

FOR FURTHER INFORMATION CONTACT: Tony Guagliardo, Director, Business Policy, Chief Business Office (163), Veterans Health Administration, Department of Veterans Affairs, 810