for broiler and fryer chicken feeds containing 68 grams/ton (g/ton) lasalocid and 10 to 50 g/ton bacitracin methylene disalicylate used for the prevention of coccidiosis, and for increased rate of weight gain and improved feed efficiency; and for broiler chicken feeds containing 68 to 113 g/ton lasalocid and 4 to 50 g/ton bacitracin methylene disalicylate used for the prevention of coccidiosis, and for improved feed efficiency. The NADA is approved as of December 4, 2002, and the regulations are amended in 21 CFR 558.311 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 supplemental application may be seen in the Dockets Management Branch (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)(2) 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.

§ 558.311 [Amended]

■ 2. Section 558.311 *Lasalocid* is amended in the table in paragraph (e)(1)(iv) under the ''Limitations'' column by removing ''withdraw 3 days before slaughter'', and in the table in paragraph (e)(1)(x) under the ''Limitations'' column by removing ''withdraw 3 days before slaughter;''.

Dated: March 21, 2003.

Steven D. Vaughn,

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

[FR Doc. 03–7535 Filed 3–31–03; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 558

New Animal Drugs for Use in Animal Feeds; Monensin; Technical Amendment

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 the approved caution statements that must appear on animal feeds containing monensin. This action is being taken to improve the accuracy of the regulations.

DATES: This rule is effective March 31, 2003.

FOR FURTHER INFORMATION CONTACT: Mohammad I. Sharar, Center for Veterinary Medicine (HFV–2), Food and Drug Administration, 7500 Standish Pl., Rockville, MD 20855, 301–827–0159, e-mail: msharar@cvm.fda.gov.

SUPPLEMENTARY INFORMATION: FDA has found that the animal drug regulations do not reflect the approved caution statements that must appear on animal feeds containing monensin. The regulation in 21 CFR 558.355 is being amended to correct inaccurate references to mature turkeys and guinea fowl that were incorporated into the regulations in the **Federal Register** published on July 26, 2000 (65 FR 45879). This action is being taken to improve the accuracy of the regulations.

Publication of this document constitutes final action on these changes under the Administrative Procedure Act (5 U.S.C. 553). Notice and public procedure are unnecessary because FDA is merely correcting nonsubstantive errors.

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.

§ 558.355 [Amended]

2. Section 558.355 *Monensin* is amended in paragraph (d)(6), in the first sentence, by removing the phrase '', other equines, mature turkeys, or guinea fowl'' and by adding in its place the phrase ''or other equines'' and in the second sentence by removing ''and guinea fowl''.

Dated: March 25, 2003.

Clifford Johnson,

Director, Office of Surveillance and Compliance, Center for Veterinary Medicine.

[FR Doc. 03–7598 Filed 3–28–03; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 558

New Animal Drugs for Use in Animal Feeds; Decoquinate; Correction

AGENCY: Food and Drug Administration, HHS.

ACTION: Final rule; correction.

SUMMARY: The Food and Drug Administration (FDA) is correcting a rule that appeared in the**Federal Register** of December 5, 2002 (67 FR 72370). The rule amended the animal drug regulations to reflect approval of a supplemental new animal drug application (NADA). FDA is correcting the range of approved concentrations of decoquinate Type A medicated article that may be used to make certain combination drug Type C medicated feeds for cattle. This correction is being made so the decoquinate regulations accurately reflect previously approved concentrations. This document corrects those errors.

DATES: This rule is effective March 31, 2003.

FOR FURTHER INFORMATION CONTACT: George K. Haibel, Center for Veterinary Medicine (HFV–6), Food and Drug Administration, 7519 Standish Pl.,