Compliance: Required as indicated, unless accomplished previously.

To prevent malfunction of the electrical circuits controlling the ''BATT. TEMP.'' red warning light, the ''ENGINE CHIP'' amber caution and the rotor revolutions-per-minute (RPM) signal output to the vehicle engine management display (VEMD), accomplish the following:

(a) Within 15 hours time-in-service (TIS), inspect the ASU No. 2 printed circuit board (PCB), part number SE 03022, to determine if resistor R8 is installed.

(b) If the resistor R8 is not installed, within 50 hours TIS, replace the PCB with an airworthy PCB that has a resistor R8 installed.

Note 2: Eurocopter Service Bulletin No. 77.00.07, dated March 27, 2000, pertains to the subject of this AD.

(c) An alternative method of compliance or adjustment of the compliance time that provides an acceptable level of safety may be used if approved by the Manager, Regulations Group, Rotorcraft Directorate, FAA. Operators shall submit their requests through an FAA Principal Maintenance Inspector, who may concur or comment and then send it to the Manager, Regulations Group.

Note 3: Information concerning the existence of approved alternative methods of compliance with this AD, if any, may be obtained from the Regulations Group.

(d) Special flight permits may be issued in accordance with 14 CFR 21.197 and 21.199 to operate the helicopter to a location where the requirements of this AD can be accomplished.

(e) This amendment becomes effective on May 19, 2003.

Note 4: The subject of this AD is addressed in Direction Generale De L'Aviation Civile (France) AD 2001–319–083(A), dated July 25, 2001.

Issued in Fort Worth, Texas, on April 8, 2003.

Michele M. Owsley,

Acting Manager, Rotorcraft Directorate, Aircraft Certification Service.

[FR Doc. 03–9012 Filed 4–11–03; 8:45 am]

BILLING CODE 4910–13–P

SECURITIES AND EXCHANGE COMMISSION

17 CFR Part 210

[Release Nos. 33–8128A; 34–46464A; FR–63A; File No. S7–08–02]

RIN 3235–AI33

Acceleration of Periodic Report Filing Dates and Disclosure Concerning Web Site Access to Reports; Correction

AGENCY: Securities and Exchange Commission.

ACTION: Final rule; technical amendments.

SUMMARY: This document contains corrections to final rules which were published in the **Federal Register** on Monday, September 16, 2002 (67 FR 58480). The rules relate to the acceleration of the filing of quarterly and annual reports under the Securities Exchange Act of 1934 by certain accelerated filers.

EFFECTIVE DATE: April 14, 2003.

FOR FURTHER INFORMATION CONTACT: Jeffrey J. Minton, Special Counsel, Office of Rulemaking, Division of Corporation Finance, at (202) 942–2910, U.S. Securities and Exchange Commission, 450 Fifth Street, NW., Washington, DC 20549–0312.

SUPPLEMENTARY INFORMATION:

I. Background

On September 5, 2002, the Commission adopted,[1] among other things, changes to rules 3–01[2] and 3–12[3] of Regulation S–X[4] under the Securities Act of 1933 (the ''Securities Act'').[5] These rules relate to the timeliness of financial information in Commission filings, such as Securities Act registration statements and proxy statements and information statements under section 14[6] of the Securities Exchange Act of 1934 (the ''Exchange Act'').[7] The changes were made to conform the timeliness requirements for these filings made by accelerated filers to changes adopted to the deadlines for Forms 10–K[8] and 10–Q[9] for accelerated filers, as defined in Exchange Act rule 12b–2.[10] The new deadlines will be phased-in over three years.

After we adopted the amendments to rules 3–01 and 3–12 of Regulation S–X, questions arose regarding the appropriate phase-in period for an accelerated filer required to update interim financial information in registration statements filed or that become effective 134 days after the end of the filer's fiscal year. This is the period after audited financial statements for the most recently completed fiscal year are already required to be filed on Form 10–K and on or after the date most registrants are required to have filed interim financial statements for the first quarter on Form 10–Q. Concerns arose that the phase-in periods in the conforming amendments to rules 3–01 and 3–12 of Regulation S–X do not match the phase-in periods described in the adopting release.

Accordingly, the amendments set forth in this document clarify that the phase-in periods applicable to accelerated filers who need to update interim information in accordance with amended rules 3–01 and 3–12 of Regulation S–X match the phase-in periods for filing quarterly information on Form 10–Q. The corrections clarify that updated interim information is required within 130 days after the end of the registrant's fiscal year for fiscal years ending on or after December 15, 2003 and before December 15, 2004, and within 125 days after the end of the registrant's fiscal year for fiscal years ending on or after December 15, 2004. The changes are technical corrections to clarify the rules as described in the original adopting release, and do not alter the phase-in periods for these requirements as described in the original adopting release.

II. Need for Correction

As published, the final regulations contain errors which may prove to be misleading and are in need of clarification.

Text of Amendments

List of Subjects in 17 CFR Part 210

Reporting and recordkeeping requirements, Securities.

■ In accordance with the foregoing, the Commission amends Title 17, chapter II of the Code of Federal Regulations as follows:

PART 210—FORM AND CONTENT OF AND REQUIREMENTS FOR FINANCIAL STATEMENTS, SECURITIES ACT OF 1933, SECURITIES EXCHANGE ACT OF 1934, PUBLIC UTILITY HOLDING COMPANY ACT OF 1935, INVESTMENT COMPANY ACT OF 1940, INVESTMENT ADVISERS ACT OF 1940, AND ENERGY POLICY AND CONSERVATION ACT OF 1975

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

Authority: 15 U.S.C. 77f, 77g, 77h, 77j, 77s, 77z–2, 77aa(25), 77aa(26), 78j–1, 78l, 78m, 78n, 78o(d), 78u–5, 78w(a), 78ll(d), 79e(b), 79j(a), 79n, 79t(a), 80a–8, 80a–20, 80a–29, 80a–30, 80a–31, 80a–37(a), unless otherwise noted.

■ 2. Section 210.3–01 is amended by revising paragraphs (e)(1) and (i)(2) to read as follows:

§ 210.3–01 Consolidatedbalance sheets.

* * * * *

(e) * * *

(1) For accelerated filers (as defined in § 240.12b–2 of this chapter):

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[1] *See* Release No. 33–8128 (Sept. 5, 2002) [67 FR 58480].

[2] 17 CFR 210.3–01.

[3] 17 CFR 210.3–12.

[4] 17 CFR 210.1–01 *et seq.*

[5] 15 U.S.C. 77a *et seq.*

[6] 15 U.S.C. 78n

[7] 15 U.S.C. 78a *et seq.*

[8] 17 CFR 249.310.

[9] 17 CFR 308a.

[10] 17 CFR 240.12b–2.

(i) 135 days for fiscal years ending on or after December 15, 2002 and before December 15, 2003;

(ii) 130 days for fiscal years ending on or after December 15, 2003 and before December 15, 2004; and

(iii) 125 days for fiscal years ending on or after December 15, 2004; and

* * * * *

(i) * * *

(2) For purposes of paragraph (e) of this section, the number of days shall be:

(i) For accelerated filers (as defined in § 240.12b–2 of this chapter):

(A) 134 days subsequent to the end of the registrant's most recent fiscal year for fiscal years ending on or after December 15, 2002 and before December 15, 2003;

(B) 129 days subsequent to the end of the registrant's most recent fiscal year for fiscal years ending on or after December 15, 2003 and before December 15, 2004; and

(C) 124 days subsequent to the end of the registrant's most recent fiscal year for fiscal years ending on or after December 15, 2004; and

(ii) 134 days subsequent to the end of the registrant's most recent fiscal year for all other registrants.

■ 3. Section 210.3–12 is amended by revising paragraph (g)(1) to read as follows:

§ 210.3–12 Age of financial statements at effective date of registration statement or at mailing date of proxy statement.

* * * * *

(g)(1) For purposes of paragraph (a) of this section, the number of days shall be:

(i) For accelerated filers (as defined in § 240.12b–2 of this chapter):

(A) 135 days for fiscal years ending on or after December 15, 2002 and before December 15, 2003;

(B) 130 days for fiscal years ending on or after December 15, 2003 and before December 15, 2004; and

(C) 125 days for fiscal years ending on or after December 15, 2004; and

(ii) 135 days for all other registrants.

* * * * *

Dated: April 8, 2003.

Margaret H. McFarland,

Deputy Secretary.

[FR Doc. 03–8998 Filed 4–11–03; 8:45 am]

BILLING CODE 8010–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 341

[Docket No. 76N–052G]

RIN 0910–AA01

Cold, Cough, Allergy, Bronchodilator, and Antiasthmatic Drug Products for Over-the-Counter Human Use; Final Monograph for Combination Drug Products; Correction

AGENCY: Food and Drug Administration, HHS.

ACTION: Final rule; correction.

SUMMARY: The Food and Drug Administration (FDA) is correcting a final rule that appeared in the **Federal Register** of December 23, 2002 (67 FR 78158). The document issued a final monograph that established conditions under which over-the-counter (OTC) cold, cough, allergy, bronchodilator, and antiasthmatic (cough-cold) combination drug products are generally recognized as safe and effective and not misbranded as part of its ongoing review of OTC drug products.

DATES: The regulation is effective December 23, 2004.

FOR FURTHER INFORMATION CONTACT: Cazemiro R. Martin or Gerald M. Rachanow, Center for Drug Evaluation and Research (HFD–560), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–2222.

SUPPLEMENTARY INFORMATION: In FR Doc. 02–32158 appearing on page 78158 in the **Federal Register** of Monday, December 23, 2002, the following corrections are made:

§ 341.40 [Corrected]

1. On page 78168, in the second column, in Part 341 *Cold, Cough, Allergy, Bronchodilator, and Antiasthmatic Drug Products for Over-the-Counter Human Use*, under the authority citation, in amendment 2, ''Section 341.40 is added to subpart C to read as follows:'' is corrected to read ''Section 341.40 is added to subpart B to read as follows:''

§ 341.70 [Corrected]

2. On page 78170, in the second column, in § 341.70 *Labeling of OTC drug products containing ingredients that are used for treating concurrent symptoms (in either a single-ingredient or combination drug product)*, in paragraph (b), ''Repeat every hour as needed or as directed by a doctor.'' is corrected to read ''Repeat every 2 hours as needed or as directed by a doctor.''

Dated: April 8, 2003.

Jeffrey Shuren,

Assistant Commissioner for Policy.

[FR Doc. 03–9067 Filed 4–11–03; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Parts 510 and 558

New Animal Drugs; Change of Sponsor

AGENCY: Food and Drug Administration, HHS.

ACTION: Final rule.

SUMMARY: The Food and Drug Administration (FDA) is amending the animal drug regulations to reflect a change of sponsor for seven approved new animal drug applications (NADAs) for clopidol Type A medicated articles and combination drug medicated chicken and turkey feeds from Aventis Animal Nutrition, Inc., to Merial Ltd.

DATES: This rule is effective April 14, 2003.

FOR FURTHER INFORMATION CONTACT: Lonnie W. Luther, Center for Veterinary Medicine (HFV–102), Food and Drug Administration, 7500 Standish Pl., Rockville, MD 20855, 301–827–0209.

SUPPLEMENTARY INFORMATION: Aventis Animal Nutrition, Inc., 3480 Preston Ridge Rd., suite 650, Alpharetta, GA 30005–8891, has informed FDA that it has transferred ownership of, and all rights and interest in, NADA 34–393, 40–264, 41–541, 44–016, 46–209, 49–934, and 99–150 for clopidol Type A medicated articles and certain combination clearances for use in medicated feeds for chickens and turkeys to Merial Ltd., 3239 Satellite Blvd., Bldg. 500, Duluth, GA 30096–4640. Accordingly, the agency is amending the regulations in § 558.175 (21 CFR 558.175) to reflect the transfer of ownership. Section 558.175 is also being changed to a table format.

Following the change of sponsor of these NADAs, Aventis Animal Nutrition, Inc., is no longer the sponsor of any approved applications. Therefore, 21 CFR 510.600(c) is being amended to remove the entries for this sponsor.

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.