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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

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21 CFR Part 5

Delegation of Authority and Organization; Center for Veterinary Medicine; Technical Amendment

AGENCY: Food and Drug Administration, HHS.

ACTION: Final rule; technical amendment.

SUMMARY: The Food and Drug Administration (FDA) is amending the regulation to reflect a redelegation of authority with respect to approval of supplemental new animal drug applications to the Director, Division of Manufacturing Technology, Office of New Animal Drug Evaluation, Center for Veterinary Medicine (CVM). This action is necessary to ensure the continued accuracy of the regulation.

EFFECTIVE DATE: December 22, 1998.

FOR FURTHER INFORMATION CONTACT:

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SUPPLEMENTARY INFORMATION: In the **Federal Register** of December 22, 1998 (63 FR 70650), FDA amended the regulation for redelegations of authority to incorporate provisions for feed mill licensing and to update positions and component titles associated with the authorities under § 5.83 (21 CFR 5.83). The regulation in § 5.83(c)(1) reflected a redelegation with respect to approval of certain supplemental new animal drug applications to the position title "Director, Division of

Human Food Safety, Office of New Animal Drug Evaluation, CVM.” The correct position title to which these functions should have been redelegated is the “Director, Division of Manufacturing Technologies, Office of New Animal Drug Evaluation, CVM,” because that office is responsible for supplemental applications for manufacturing. Therefore, the redelegation to the “Director, Division of Human Safety Office of New Animal Drug Evaluation, CVM” has been amended and a new redelegation made to the “Director, Division of Manufacturing Technologies, Office of New Animal Drug Evaluation, CVM.” The regulation in § 5.83(c)(1) has been revised to reflect this redelegation. Further redelegation of authorities is not authorized at this time. Authority delegated to a position may be exercised by a person officially designated to serve in such position in an acting capacity or on a temporary basis.

List of Subjects in 21 CFR Part 5

Authority delegations (Government agencies), Imports, Organization and functions (Government agencies).

Therefore, under the Federal Food, Drug, and Cosmetic Act and under authority delegated to the Commissioner of Food and Drugs, 21 CFR part 5 is amended to read as follows:

PART 5—DELEGATIONS OF AUTHORITY AND ORGANIZATION

1. The authority citation for 21 CFR part 5 continues to read as follows:

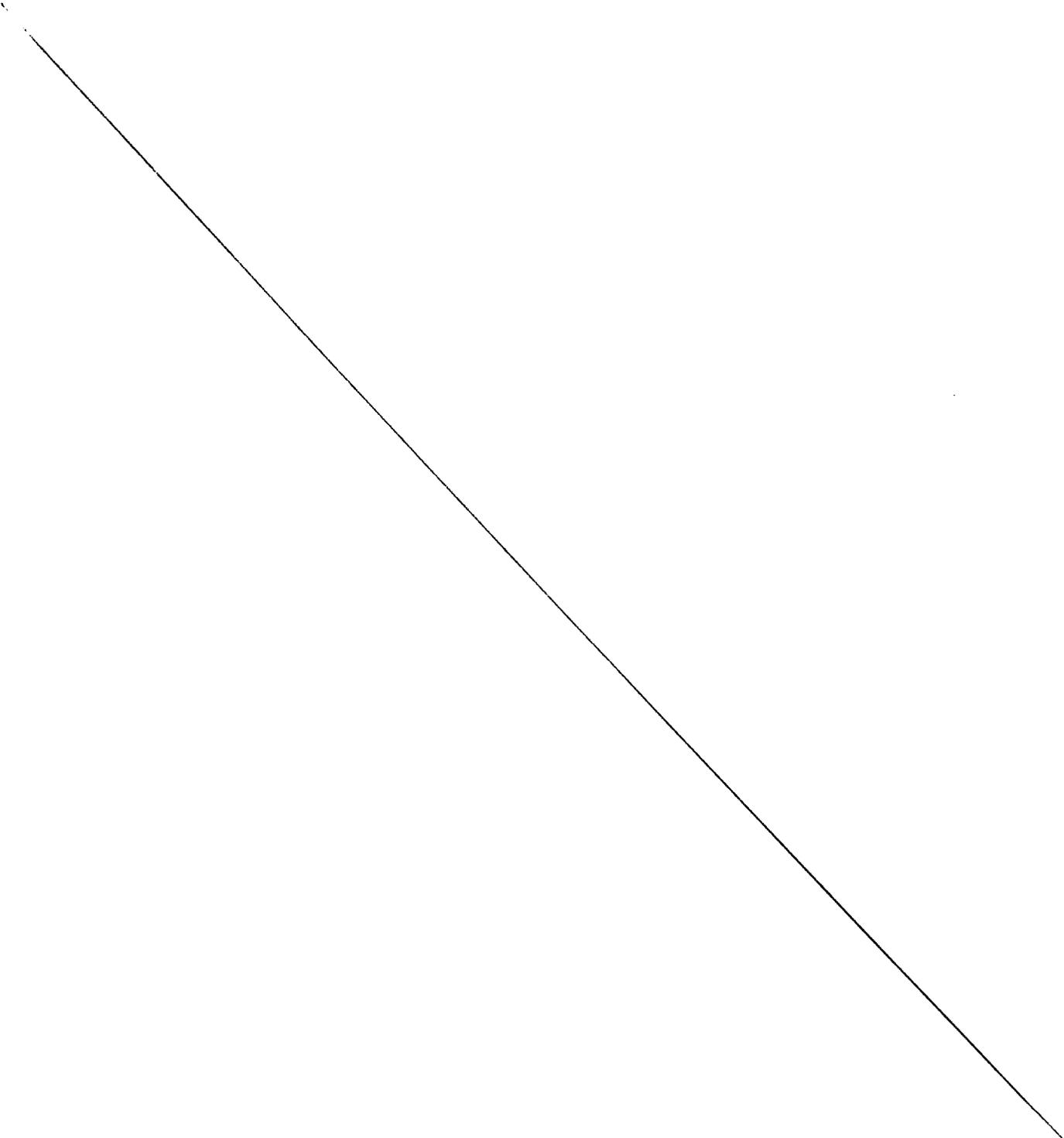
Authority: 5 U.S.C. 504, 552, App. 2; 7 U.S.C. 138a, 2271; 15 U.S.C. 638, 1261–1282, 3701–3711a; 15 U.S.C. 1451–1461; 21 U.S.C. 41–50, 61–63, 141–149, 321–394, 467f, 679(b), 801–886, 1031–1309; 35 U.S.C. 156; 42 U.S.C. 241, 242, 242a, 242l, 242n, 243, 262, 263, 264, 265, 300u–300u–5, 300aa–1; 1395y, 3246b, 4332, 4831(a), 10007–10008; E.O. 11921, 41 FR 24294, 3 CFR 1977 Comp., p. 124–131; E.O. 12591, 52 FR 13414, 3 CFR, 1988 Comp., p. 220–223.

2. Section 5.83 is amended by revising paragraph (c)(1) to read as follows:

§ 5.83 Approval of new animal drug applications, medicated feed mill license applications, and their supplements.

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(c) * * *



(1) The Director, Division of Manufacturing Technologies, Office of New Animal Drug Evaluation, CVM.

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Dated: April 23, 1999
April 23, 1999



William K. Hubbard
Acting Deputy Commissioner for
Policy

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