Request for Referendum will be conducted in accordance with the Act and applicable regulations. Pursuant to 5 U.S.C. 553, good cause is found for not postponing the effective date of the action until 30 days after publication in the **Federal Register** in order to conduct the Request for Referendum within the timeframes that appear in the Act.

List of Subjects in 7 CFR Part 1220

Administrative practice and procedure, Advertising, Agricultural research, Marketing agreements, Reporting and recordkeeping requirements, Soybeans and soybean products.

■ For the reasons set forth in the preamble, 7 CFR part 1220 is amended as follows:

PART 1220—SOYBEAN PROMOTION, RESEARCH, AND CONSUMER INFORMATION

■ 1. The authority citation for part 1220 continues to read as follows:

Authority: 7 U.S.C. 6301–6311 and 7 U.S.C. 7401.

Subpart F—Procedures to Request a Referendum

■ 2. In § 1220.616, paragraph (d) is revised to read as follows:

§ 1220.616 General.

* * * * *

(d) For purposes of paragraphs (b) and (c) of this section, the number of soybean producers in the United States is determined to be 589,182.

§ 1220.622 [Amended]

■ 3. In § 1220.622, paragraph (b) the Web site ''*http://www.ams.usda.gov/lsg/mpb/rp-soy.htm*'' is removed and a new Web site ''*http://www.ams.usda.gov/lsmarketing programs*'' is added in its place.

■ 4. In § 1220.628, paragraph (a) is revised to read as follows:

§ 1220.628 Results of the request for referendum.

(a) The Administrator, FSA, shall submit to the Administrator, AMS, the reports from all State FSA offices. The Administrator, AMS shall tabulate the results of the Request for Referendum. USDA will issue an official press release announcing the results of the Request for Referendum and publish the same results in the **Federal Register**. In addition, USDA will post the official results at the following Web site: ''*http://www.ams.usda.gov/lsmarketingprograms*''.

Subsequently, State reports and related papers shall be available for public inspection upon request during normal business hours in the Marketing Programs Branch office, Livestock and Seed Program, AMS, USDA, Room 2628–S, STOP 0251, 1400 Independence Avenue, SW., Washington, DC.

* * * * *

Dated: February 24, 2009.

Robert C. Keeney,

Acting Associate Administrator.

[FR Doc. E9–4292 Filed 2–27–09; 8:45 am]

BILLING CODE 3410–02–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 522

[Docket No. FDA–2009–N–0665]

Implantation or Injectable Dosage Form New Animal Drugs; Ivermectin

AGENCY: Food and Drug Administration, HHS.

ACTION: Final rule.

SUMMARY: The Food and Drug Administration (FDA) is amending the animal drug regulations to reflect approval of a supplemental abbreviated new animal drug application (ANADA) filed by IVX Animal Health, Inc. The supplemental ANADA adds claims for persistent effectiveness against various species of external and internal parasites when cattle are treated with a 1-percent ivermectin solution by subcutaneous injection.

DATES: This rule is effective March 2, 2009.

FOR FURTHER INFORMATION CONTACT: John K. Harshman, Center for Veterinary Medicine (HFV–104), Food and Drug Administration, 7500 Standish Pl., Rockville, MD 20855, 240–276–8197, e-mail: *john.harshman@fda.hhs.gov*.

SUPPLEMENTARY INFORMATION: IVX Animal Health, Inc., 3915 South 48th Street Ter., St. Joseph, MO 64503, filed a supplement to ANADA 200–228 that provides for use of PHOENECTIN (ivermectin) Injection 1% for the treatment and control of parasites in cattle. The supplemental ANADA adds claims for persistent effectiveness against various species of external and internal parasites of cattle. The supplemental ANADA is approved as of January 23, 2009, and the regulations are amended in 21 CFR 522.1192 to reflect the approval.

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 522

Animal drugs.

■ 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 522 is amended as follows:

PART 522—IMPLANTATION OR INJECTABLE DOSAGE FORM NEW ANIMAL DRUGS

■ 1. The authority citation for 21 CFR part 522 continues to read as follows:

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

§ 522.1192 [Amended]

■ 2. In § 522.1192, in paragraph (b)(2), remove ''No. 055529'' and in its place add ''Nos. 055529 and 059130''; and remove paragraph (b)(3).

Dated: February 18, 2009.

Steven D. Vaughn,

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

[FR Doc. E9–4304 Filed 2–27–09; 8:45 am]

BILLING CODE 4160–01–S