or Principal Avionics Inspector, as appropriate, or lacking a principal inspector, your local Flight Standards District Office.

**Related Information**

(j) For more information about this AD, contact Georgios Roussos, Aerospace Engineer, Systems and Equipment Branch, ANM–130S, FAA, Seattle Aircraft Certification Office, 1601 Lind Avenue, SW., Renton, Washington 98057–3356; telephone (425) 917–6482; fax (425) 917–6590.

(k) For service information identified in this AD, contact Boeing Commercial Airplanes, Attention: Data & Services Management, P.O. Box 3707, MC 2H–65, Seattle, Washington 98124–2207; telephone 206–544–5000, extension 1; fax 206–766–5680; *e-mail me.boecom@boeing.com;* Internet *https://www.myboeingfleet.com.* You may review copies of the referenced service information at the FAA, Transport Airplane Directorate, 1601 Lind Avenue, SW., Renton, Washington. For information on the availability of this material at the FAA, call 425–227–1221.

Issued in Renton, Washington, on December 10, 2010.

**Ali Bahrami,**

*Manager, Transport Airplane Directorate, Aircraft Certification Service.*

[FR Doc. 2010–31828 Filed 12–17–10; 8:45 am]

**BILLING CODE 4910–13–P**

## SECURITIES AND EXCHANGE COMMISSION

### 17 CFR Parts 240 and 249

**[Release No. 34–63347; File No. S7–35–10]**

**RIN 3235–AK79**

## Security-Based Swap Data Repository Registration, Duties, and Core Principles

*Correction*

In proposed rule document 2010–29719 beginning on page 77306 in the issue of December 10, 2010, make the following corrections:

1. On page 77320, in the third column, footnote 74, in the fourth line, "recordkeeping" should read "record keeping".

2. On page 77321, in the second column, below the heading *Request for Comment,* in the fifth bulleted paragraph, in the tenth line, "requiring" should read "require".

3. On page 77324, in the third column, footnote 90, in the fifth line, "recordkeeping" should read "record keeping".

4. On page 77338, the last line of text in the third column, prior to footnote 164 on the page, should read "information maintained by the SDR,[165]".

5. On the same page, in the same column, after footnote 164, add footnote 165 to read as follows:

[165] *See* Public Law 111–203 (adding Exchange Act Section 12(n)(5)(D)(i)).

6. On page 77347, in the second column, in the tenth line from the bottom of the page, "conflict" should read "conflicts".

7. On page 77356, in the third column, in thirty-first line, "systematically" should read "systemically".

8. On the same page, in the same line of the same column, "Therefor" should read "Therefore".

**§ 249.1500 [Corrected]**

9. On page 77375, in § 249.1500, before the first line in the first column, insert the following text:

**EXHIBITS—BUSINESS ORGANIZATION**

13. List as Exhibit A any person as defined in Section 3(a)(9) of the

10. On the same page, in the second column, in the fifth, eleventh, and fifteenth lines from the bottom of the page, "l5" should read "15".

[FR Doc. C1–2010–29719 Filed 12–17–10; 8:45 am]

**BILLING CODE 1505–01–D**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Food and Drug Administration

### 21 CFR Part 500

**[Docket No. FDA–2010–N–0612]**

## Animal Drugs, Feeds, and Related Products; Regulation of Carcinogenic Compounds in Food-Producing Animals

**AGENCY:** Food and Drug Administration, HHS.

**ACTION:** Proposed rule.

**SUMMARY:** The Food and Drug Administration (FDA) is proposing to amend its regulations regarding compounds of carcinogenic concern used in food-producing animals. Specifically, the Agency is clarifying the definition of "$S_o$" and revising the definition of "$S_m$" so that it conforms to the clarified definition of $S_o$. Other clarifying and conforming changes are also being made.

**DATES:** Submit either electronic or written comments on the proposed rule by March 7, 2011. Submit comments on information collection issues under the Paperwork Reduction Act of 1995 by January 19, 2011 (*see* the "Paperwork Reduction Act of 1995" section of this document).

**ADDRESSES:** You may submit comments, identified by Docket No. FDA–2010–N–0612, by any of the following methods, except that comments on information collection issues under the Paperwork Reduction Act of 1995 must be submitted to the Office of Regulatory Affairs, Office of Management and Budget (OMB) (*see* the "Paperwork Reduction Act of 1995" section of this document).

**Electronic Submissions**

Submit electronic comments in the following way:

- *Federal eRulemaking Portal: http://www.regulations.gov.* Follow the instructions for submitting comments.

**Written Submissions**

Submit written submissions in the following ways:

- *Fax:* 301–827–6870.
- *Mail/Hand delivery/Courier (for paper, disk, or CD–ROM submissions):* Division of Dockets Management (HFA–305), Food and Drug Administration, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852.

*Instructions:* All submissions received must include the Agency name and Docket No. and Regulatory Information Number (RIN) (if a RIN number has been assigned) for this rulemaking. All comments received may be posted without change to *http://www.regulations.gov,* including any personal information provided. For additional information on submitting comments, *see* the "Comments" heading of the **SUPPLEMENTARY INFORMATION** section of this document.

*Docket:* For access to the docket to read background documents or comments received, go to *http://www.regulations.gov* and insert the docket number, found in brackets in the heading of this document, into the "Search" box and follow the prompts and/or go to the Division of Dockets Management, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852.

**FOR FURTHER INFORMATION CONTACT:** Kevin Greenlees, Center for Veterinary Medicine (HFV–100), Food and Drug Administration, 7500 Standish Pl., Rockville, MD 20855, 301–827–6975. e-mail: *kevin.greenlees@fda.hhs.gov.*

**SUPPLEMENTARY INFORMATION:**

**I. Background**

The Federal Food, Drug, and Cosmetic Act (the FD&C Act) contains three anticancer, or Delaney, clauses: Sections 409(c)(3)(A), 512(d)(1)(I), and 721(b)(5)(B)(i) (21 U.S.C. 348(c)(3)(A), 360b(d)(1)(I), and 379e(b)(5)(B)(i)), pertaining to food additives, new animal drugs, and color additives, respectively.