Nuclear materials, Radiation protection, Reporting and recordkeeping requirements.

10 CFR Part 51

Administrative practice and procedure, Environmental impact statement, Nuclear materials, Nuclear power plants and reactors, Reporting and recordkeeping requirements.

10 CFR Part 52

Administrative practice and procedure, Antitrust, Backfitting, Combined license, Early site permit, Emergency planning, Fees, Incorporation by reference, Inspection, Limited work authorization, Nuclear power plants and reactors, Probabilistic risk assessment, Prototype, Reactor siting criteria, Redress of site, Reporting and recordkeeping requirements, Standard design, Standard design certification.

10 CFR Part 55

Criminal penalties, Manpower training programs, Nuclear power plants and reactors, Reporting and recordkeeping requirements.

For the reasons set out in the preamble and under the authority of the Atomic Energy Act of 1954, as amended; the Energy Reorganization Act of 1974, as amended; and 5 U.S.C. 552 and 553; the NRC is adopting the following amendments to 10 CFR Parts 19, 30, 51, 52, and 55.

PART 19—NOTICES, INSTRUCTIONS AND REPORTS TO WORKERS: INSPECTION AND INVESTIGATIONS

1. The authority citation for Part 19 continues to read in part as follows:

Authority: Secs. 53, 63, 81, 103, 104, 161, 186, 68 Stat. 930, 933, 935, 936, 937, 948, 955, as amended, sec. 234, 83 Stat. 444, as amended, sec. 1701, 106 Stat. 2951, 2952, 2953 (42 U.S.C. 2073, 2093, 2111, 2133, 2134, 2201, 2236, 2282, 2297f) * * *.

§ 19.2 [Amended]

2. In § 19.2, in the first sentence, add the numeral ''70,'' between the numeral ''61,'' and the word ''or''.

PART 30—RULES OF GENERAL APPLICABILITY TO DOMESTIC LICENSING OF BYPRODUCT MATERIAL

3. The authority citation for Part 30 continues to read in part as follows:

Authority: Secs. 81, 82, 161, 182, 183, 186, 68 Stat. 935, 948, 953, 954, 955, as amended, sec. 234, 83 Stat. 444, as amended (42 U.S.C. 2111, 2112, 2201, 2232, 2233, 2236, 2282) * * *.

§ 30.72 [Amended]

4. In § 30.72, Schedule C, in the ''Radioactive material'' column, the entry for Carbon-14 is revised to read, ''Carbon-14 (non-carbon dioxide).''

PART 51—ENVIRONMENTAL PROTECTION REGULATIONS FOR DOMESTIC LICENSING AND RELATED REGULATORY FUNCTIONS

5. The authority citation for Part 51 continues to read as follows:

Authority: Secs. 161, 68 Stat. 948, as amended, sec. 1701, 106 Stat. 2951, 2952, 2953, as amended (42 U.S.C. 2201, 2297f); secs. 201, as amended, 202, 88 Stat. 1242, as amended, 1244 (42 U.S.C. 5841, 5842) * * *.

§ 51.22 [Amended]

6. In § 51.22, in paragraph (c)(14)(ii), remove the words ''10 CFR 35.14 and 35.100'' and add ''10 CFR 35.18.''

§ 51.123 [Amended]

7. In § 51.123, in paragraphs (a) and (b), remove the words ''§ 9.14 of this chapter,'' and add the words ''§ 9.35 of this chapter.''

PART 52—EARLY SITE PERMITS; STANDARD DESIGN CERTIFICATIONS; AND COMBINED LICENSES FOR NUCLEAR POWER PLANTS

8. The authority citation for Part 52 continues to read as follows:

Authority: Secs. 103, 104, 161, 182, 183, 186, 189, 68 Stat. 936, 948, 953, 954, 955, 956, as amended, sec. 234, 83 Stat. 1244, as amended (42 U.S.C. 2133, 2201, 2232, 2233, 2236, 2239, 2282); secs. 201, 202, 206, 88 Stat. 1242, 1244, 1246, 1248, as amended 42 U.S.C. 5841, 5842, 5846).

9. In Appendix O to Part 52, paragraph 1. is revised to read as follows:

Appendix O To Part 52—Standardization of Design: Staff Review of Standard Designs

* * * * *

1. Any person may submit a proposed preliminary or final standard design for a nuclear power reactor of the type described in § 50.22 to the regulatory staff for its review. Such a submittal may consist of either the preliminary or final design for the entire reactor facility or the preliminary or final design of major portions thereof.

* * * * *

PART 55—OPERATORS' LICENSES

10. The authority citation for Part 55 continues to read as follows:

Authority: Secs. 107, 161, 182, 68 Stat. 939, 948, 953, as amended, sec. 234, 83 Stat. 444, as amended (42 U.S.C. 2137, 2201, 2232, 2282); secs. 201, as amended, 202, 88 Stat. 1242, as amended, 1244 (42 U.S.C. 5841, 5842).

§ 55.5 [Amended]

11. In § 55.5, paragraph (b)(2)(iii), remove the words ''799 Roosevelt Road, Glyn Ellyn, IL 60137,'' and add the words ''801 Warrenville Road, Lisle, IL 60532–4351.''

Dated at Rockville, Maryland, this 28th day of February 1996.

For the Nuclear Regulatory Commission.

James M. Taylor,

Executive Director for Operations.

[FR Doc. 96–5815 Filed 3–11–96; 8:45 am]

BILLING CODE 7590–01–P

DEPARTMENT OF TRANSPORTATION

Federal Aviation Administration

14 CFR Part 71

[Airspace Docket No. 95–ANE–31]

Alteration of V–423

AGENCY: Federal Aviation Administration (FAA), DOT.

ACTION: Final rule.

SUMMARY: This rule alters Federal Airway V–423 due to the decommissioning of the Uplands Nondirectional Beacon (NDB).

EFFECTIVE DATE: 0901 UTC, June 20, 1996.

FOR FURTHER INFORMATION CONTACT: Patricia P. Crawford, Airspace and Obstruction Evaluation Branch (ATP–240), Airspace-Rules and Aeronautical Information Division, Air Traffic Rules and Procedures Service, Federal Aviation Administration, 800 Independence Avenue, SW., Washington, DC 20591; telephone: (202) 267–3075.

SUPPLEMENTARY INFORMATION:

The Rule

This amendment to part 71 of the Federal Aviation Regulations alters Federal Airway V–423 due to the decommissioning of the Uplands NDB located in Ottawa, Canada. Specifically, this action eliminates a segment of Federal Airway V–423 due to the decommissioning of that NDB. Because this action is a minor amendment in which the public would not be particularly interested, I find that notice and public procedure under 5 U.S.C. 553(b) are unnecessary. Domestic VOR Federal airways are published in paragraph 6010(a) of FAA Order 7400.9C dated August 17, 1995, and

effective September 16, 1995, which is incorporated by reference in 14 CFR 71.1. The airway listed in this document will be published subsequently in the Order.

The FAA has determined that this regulation only involves an established body of technical regulations for which frequent and routine amendments are necessary to keep them operationally current. It, therefore—(1) Is not a ''significant regulatory action'' under Executive Order 12866; (2) is not a ''significant rule'' under DOT Regulatory Policies and Procedures (44 FR 11034; February 26, 1979); and (3) does not warrant preparation of a regulatory evaluation as the anticipated impact is so minimal. Since this is a routine matter that will only affect air traffic procedures and air navigation, it is certified that this rule will not have a significant economic impact on a substantial number of small entities under the criteria of the Regulatory Flexibility Act.

List of Subjects in 14 CFR Part 71

Airspace, Incorporation by reference, Navigation (air).

Adoption of the Agreement

In consideration of the foregoing, the Federal Aviation Administration amends 14 CFR part 71 as follows:

PART 71—[AMENDED]

1. The authority citation for part 71 is revised to read as follows:

Authority: 49 U.S.C. 106(g), 40103, 40113, 40120; E.O. 10854, 24 FR 9565, 3 CFR, 1959–1963 Comp., p. 389.

§ 71.1 [Amended]

2. The incorporation by reference in 14 CFR 71.1 of the Federal Aviation Administration Order 7400.9C, Airspace Designations and Reporting Points, dated August 17, 1995, and effective September 16, 1995, is amended as follows:

Paragraph 6010(a)—Domestic VOR Federal Airways

* * * * *

V–423 [Revised]

From Williamsport, PA; Binghamton, NY; Ithaca, NY; Syracuse, NY; Watertown, NY; INT Watertown 018° radial and the United States/Canadian Border.

* * * * *

Issued in Washington, DC, on March 4, 1996.

Nancy B. Kalinowski,

Acting Manager, Airspace—Rules and Aeronautical Information Division.

[FR Doc. 96–5834 Filed 3–11–96; 8:45 am]

BILLING CODE 4910–13–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 175

[Docket No. 93F–0358]

Indirect Food Additives: Adhesives and Components of Coatings

AGENCY: Food and Drug Administration, HHS.

ACTION: Final rule.

SUMMARY: The Food and Drug Administration (FDA) is amending the food additive regulations to provide for the safe use of *meta*-tetramethylxylene diisocyanate for reaction with one or more of the polyols and polyesters listed in the adhesive regulations and with dimethylolpropionic acid and trimethylamine, *N*-methyldiethanolamine, 2-dimethylaminoethanol, 2-dimethylamino-2-methyl-1-propanol, and/or 2-amino-2-methyl-1-propanol in the production of polyurethane resins intended for use as components of adhesive formulations used in food packaging applications. This action is in response to a petition filed by Cytec Industries.

DATES: Effective March 12, 1996; written objections and requests for a hearing by April 11, 1996.

ADDRESSES: Submit written objections to the Dockets Management Branch (HFA–305), Food and Drug Administration, 12420 Parklawn Dr., rm. 1–23, Rockville, MD 20857.

FOR FURTHER INFORMATION CONTACT: Vir D. Anand, Center for Food Safety and Applied Nutrition (HFS–216), Food and Drug Administration, 200 C St. SW., Washington, DC 20204, 202–418–3081.

SUPPLEMENTARY INFORMATION: In a notice published in the Federal Register of October 26, 1993 (58 FR 57613), FDA announced that a food additive petition (FAP 3B4401) had been filed by Cytec Industries, c/o Keller and Heckman, 1001 G St. NW., suite 500 West, Washington, DC 20001, proposing that § 175.105 *Adhesives* (21 CFR 175.105) be amended to provide for the safe use of *meta*-tetramethylxylene diisocyanate for reaction with one or more of the polyols and polyesters listed in § 175.105 and with dimethylolpropionic acid and trimethyamine, *N*-methyldiethanolamine, 2-dimethylaminoethanol, 2-dimethylamino-2-methyl-1-propanol, and/or 2-amino-2-methyl-1-propanol in the production of polyurethane resins intended for use as components of adhesive formulations used in food packaging applications. This document is also amending § 175.105(c)(5) to correct inconsistencies in the spelling of the Chemical Abstract Service Registry Numbers.

FDA has evaluated data in the petition and other relevant material. The agency concludes that the proposed use of the additive is safe and that 21 CFR 175.105 should be amended as set forth below.

In accordance with § 171.1(h) (21 CFR 171.1(h)), the petition and the documents that FDA considered and relied upon in reaching its decision to approve the petition are available for inspection at the Center for Food Safety and Applied Nutrition by appointment with the information contact person listed above. As provided in 21 CFR 171.1(h), the agency will delete from the documents any materials that are not available for public disclosure before making the documents available for inspection.

The agency has carefully considered the potential environmental effects of this action. FDA has concluded that the action will not have a significant impact on the human environment, and that an environmental impact statement is not required. The agency's finding of no significant impact and the evidence supporting that finding, contained in an environmental assessment, may be seen in the Dockets Management Branch (address above) between 9 a.m. and 4 p.m., Monday through Friday.

Any person who will be adversely affected by this regulation may at any time on or before April 11, 1996, file with the Dockets Management Branch (address above) written objections thereto. Each objection shall be separately numbered, and each numbered objection shall specify with particularity the provisions of the regulation to which objection is made and the grounds for the objection. Each numbered objection on which a hearing is requested shall specifically so state. Failure to request a hearing for any particular objection shall constitute a waiver of the right to a hearing on that objection. Each numbered objection for which a hearing is requested shall include a detailed description and analysis of the specific factual information intended to be presented in support of the objection in the event that a hearing is held. Failure to include such a description and analysis for any particular objection shall constitute a waiver of the right to a hearing on the objection. Three copies of all documents shall be submitted and shall be identified with the docket number found in brackets in the heading of this