

Dated: April 27, 2004.

**David C. Stalfort,**

*Commander, U.S. Coast Guard, Captain of the Port Memphis.*

[FR Doc. 04-10353 Filed 5-5-04; 8:45 am]

BILLING CODE 4910-15-P

## POSTAL SERVICE

### 39 CFR Part 111

#### Permissible Barcode Symbology for Parcels Eligible for the Barcode Discount

**AGENCY:** Postal Service.

**ACTION:** Final rule with request for comments.

**SUMMARY:** We are amending the information published in the **Federal Register** on July 14, 1998 [63 FR 37946], that announced new requirements for Package Services parcels eligible for the barcode discount. The barcode discount was extended to Standard Mail® machinable parcels in the **Federal Register** on December 15, 2000 [65 FR 78537] that announced the R2000-1 rate case. The standards implementing the new requirements were subsequently published in *Postal Bulletin* 22122 (2-19-04, pages 6-8). The *Postal Bulletin* notice allowed for the optional use of the human-readable presentation of the ZIP Code™. This final rule modifies the standards to now require mailers to include the human-readable equivalent of the ZIP Code with all barcodes.

**DATES:** Effective May 6, 2004. Submit comments on or before May 20, 2004.

**ADDRESSES:** Mail or deliver comments to the Manager, Mailing Standards, Attn: Obataiye B. Akinwale, U.S. Postal Service, 1735 N Lynn Street, Room 3025, Arlington, VA 22209-6038. Written comments may be submitted also by facsimile transmission to (703) 292-4058. Copies of all written comments will be available for inspection and photocopying between 9 a.m. and 4 p.m., Monday through Friday, at Postal Service Headquarters Library, 11th Floor North, 475 L'Enfant Plaza SW., Washington, DC.

**FOR FURTHER INFORMATION CONTACT:** Obataiye B. Akinwale at (703) 292-3643.

**SUPPLEMENTARY INFORMATION:** On July 14, 1998, the Postal Service published in the **Federal Register** [63 FR 37946] a final rule setting forth Domestic Mail Manual (DMM) standards for Package Services barcodes. The DMM standards were subsequently published in *Postal Bulletin* 22122 (2-19-04, pages 6-8). Under the previous rule, the human-

readable equivalent of the ZIP Code information was optional. Under the new rule, mailers are required to include the human-readable equivalent of the ZIP Code information to be eligible for the barcode discount. No other changes are made to the standards in DMM C850.

Using the UCC/EAN Code 128 barcode symbology will benefit mailers in a number of ways:

- Increased accuracy and improved service—reduces manual processing of parcels.
- Variable length—compact, accurate, and reliable.
- Easy data capture capabilities—international availability.

In order to reduce the looping of mail in processing, this rule requires the printing of the applicable AI and ZIP Code or ZIP+4® code whenever a barcode is printed.

#### List of Subjects in 39 CFR Part 111

Administrative practice and procedure, Postal Service.

■ For the reasons noted above, the Postal Service adopts the following changes to the *Domestic Mail Manual*, which is incorporated by reference in the *Code of Federal Regulations* (CFR). See 39 CFR part 111.

#### PART 111—[AMENDED]

■ 1. The authority citation for 39 CFR Part 111 continues to read as follows:

**Authority:** 5 U.S.C. 552(a); 39 U.S.C. 101, 401, 403, 404, 3001-3011, 3201-3219, 3403-3406, 3621, 3626, 5001.

■ 2. Revise the *Domestic Mail Manual* (DMM) as follows:

#### Domestic Mail Manual (DMM)

\* \* \* \* \*

#### C Characteristics and Content

\* \* \* \* \*

#### C800 Automation-Compatible Mail

\* \* \* \* \*

#### C850 Barcoding Standards for Parcels

\* \* \* \* \*

#### 2.0 Barcode Characteristics

\* \* \* \* \*

#### 2.5 Human-Readable Information

The human-readable information on the barcode must conform to one of the following options:

*[Revise item a to read as follows:]*

a. If the barcode is printed on the delivery address label and in close proximity to the address, the human-readable equivalent of the ZIP Code or ZIP+4 code (omitting the AI "420")

encoded in the barcode preceded by the word "ZIP" must be printed between 1/8 inch and 1/2 inch below the barcode in 10-point or larger bold, sans serif type. This standard applies to barcodes printed under 1.1 or 1.2a, 1.2b, and 1.3.

\* \* \* \* \*

*[Revise item c to read as follows:]*

c. For barcodes printed under 1.2 or 1.3, the human-readable presentation of the concatenated barcode must include the AI "91" and the full tracking identification number as text, the AI "420," and the ZIP Code or ZIP+4 code. The AI "420" and ZIP Code information must be parsed separately from the main body of the barcode text (e.g., 420 99999 9101 2345 6789 1234 5678).

\* \* \* \* \*

We will publish an appropriate amendment to 39 CFR part 111 to reflect these changes.

**Neva R. Watson,**

*Attorney, Legislative.*

[FR Doc. 04-10154 Filed 5-5-04; 8:45 am]

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## ENVIRONMENTAL PROTECTION AGENCY

### 40 CFR Part 63

[OAR-2002-0045, FRL-7657-2]

RIN 2060-AK53

#### National Emission Standards for Hazardous Air Pollutants for Chemical Recovery Combustion Sources at Kraft, Soda, Sulfite, and Stand-Alone Semichemical Pulp Mills

**AGENCY:** Environmental Protection Agency (EPA).

**ACTION:** Final rule; technical corrections.

**SUMMARY:** On February 18, 2003, the EPA promulgated amendments to the national emission standards for hazardous air pollutants (NESHAP) for chemical recovery combustion sources at kraft, soda, sulfite, and stand-alone semichemical pulp mills. The technical corrections in the final rule correct several cross-references in order to be consistent with the text shifts made in the February 18, 2003 amendments.

**DATES:** *Effective Date:* The technical corrections are effective May 6, 2004.

**ADDRESSES:** Docket ID No. OAR-2002-0045 and Docket ID No. A-94-67, containing supporting information used in the development of the final rule, are available for public viewing at the EPA Docket Center (Air Docket), EPA West, Room B-102, 1301 Constitution Avenue, NW., Washington, DC. The EPA Docket Center Public Reading

Room is open from 8:30 a.m. to 4:30 p.m., Monday through Friday, excluding legal holidays. The telephone number for the Public Reading Room is (202) 566-1744, and the telephone number for the Air Docket is (202) 566-1742.

**FOR FURTHER INFORMATION CONTACT:** Mr. Jeff Telander, Minerals and Inorganic Chemicals Group, Emission Standards

Division (C504-05), Office of Air Quality Planning and Standards, U.S. EPA, Research Triangle Park, NC 27711, telephone number (919) 541-5427, facsimile number (919) 541-5600, electronic mail (e-mail address [telander.jeff@epa.gov](mailto:telander.jeff@epa.gov)).

**SUPPLEMENTARY INFORMATION:** *Regulated Entities.* Categories and entities

potentially regulated by this action are those kraft, soda, sulfite, and stand-alone semichemical pulp mills with chemical recovery processes that involve the combustion of spent pulping liquor. Categories and entities potentially regulated by this action include:

Category	NAICS code*	Examples of regulated entities
Industry .....	32211 32212 32213	Kraft, soda, sulfite, and stand-alone semichemical pulp mills.
Federal government .....	.....	Not affected.
State/local/tribal government .....	.....	Not affected.

\* North American Industrial Classification System.

This table is not intended to be exhaustive, but rather provides a guide for readers regarding entities likely to be regulated by this action. To determine whether your facility is regulated by this action, you should carefully examine the applicability criteria in § 63.860 of the national emission standards. If you have questions regarding the applicability of this action to a particular entity, consult the person listed in the preceding **FOR FURTHER INFORMATION CONTACT** section of this document. *Docket.* The EPA has established an official public docket for this action including both Docket ID No. OAR-2002-0045 and Docket ID No. A-94-67. The official public docket consists of the documents specifically referenced in the final rule, any public comments received, and other information related to the final rule. All items may not be listed under both docket numbers, so interested parties should inspect both docket numbers to ensure that they have received all materials relevant to the final rule. Although a part of the official docket, the public docket does not include Confidential Business Information or other information whose disclosure is restricted by statute. The official public docket is available for public viewing at the EPA Docket Center (Air Docket), EPA West, Room B-102, 1301 Constitution Avenue, NW., Washington, DC. The EPA Docket Center Public Reading Room is open from 8:30 a.m. to 4:30 p.m., Monday through Friday, excluding legal holidays. The telephone number for the Public Reading Room is (202) 566-1744, and the telephone number for the Air Docket is (202) 566-1742.

*Electronic Access.* You may access this **Federal Register** document electronically through the EPA Internet under the **Federal Register** listings at

<http://www.epa.gov/fedrgstr/>. You may also access a copy of the final rule incorporating the provisions of the **Federal Register** notice through the Technology Transfer Network (TTN) at <http://www.epa.gov/ttn/atw/pulp/pulppg.html>. An electronic version of the public docket is available through EPA's electronic public docket and comment system, EPA Dockets. You may use EPA Dockets at <http://www.epa.gov.edocket/> to view public comments, access the index listing of the contents of the official public docket, and to access those documents in the public docket that are available electronically. Although not all docket materials may be available electronically, you may still access any of the publicly available docket materials through the docket facility identified above. Once in the system, select "search," then key in the appropriate docket identification number.

*Background.* On February 18, 2003, we published a direct final rule (68 FR 7706) and parallel proposal (68 FR 7735) amending the NESHAP for chemical recovery combustion sources at kraft, soda, sulfite, and stand-alone semichemical pulp mills (40 CFR part 63, subpart MM). The amendments clarified and consolidated the monitoring and testing requirements and added a site-specific alternative standard for one pulp mill. The consolidation of the monitoring and testing requirements resulted in significant text shifts within and between the monitoring and testing sections.

The technical corrections in the final rule correct the following cross-references in order to be consistent with the text shifts made in the February 18, 2003 amendments:

- The reference in § 63.866(a)(1) to the procedures in § 63.864(b)(2) for establishing operating ranges is corrected to refer to the procedures in § 63.864(j);
  - The references in §§ 63.866(b) and 63.867(c) to the ongoing compliance provisions in § 63.864(c), (c)(1) and (2) are revised to refer to the provisions in § 63.864(k), (k)(1) and (2), respectively;
  - The reference in § 63.866(c)(4) to the compliance determinations made under § 63.865(a) through (e) is corrected to refer to the compliance determinations made under § 63.865(a) through (d); and
  - The references in the General Provisions table (under §§ 63.7(a)(1) and 63.7(h)) to the performance test exemption in § 63.864(a)(6) are corrected to refer to the exemption in § 63.865(c)(1).
- Section 553(d) of 5 U.S.C. allows an agency, upon a finding of good cause, to make a rule effective immediately. Because today's final rule simply corrects cross-references in order to be consistent with text shifts made in the February 18, 2003 amendments, does not add any requirements necessitating additional time for compliance, and otherwise does not substantively change the requirements of the final rule or otherwise affect sources' ability to comply with the final rule or any compliance obligation a source may have, we find good cause to make the final rule effective immediately.

**Statutory and Executive Order Review**

Under Executive Order 12866 (58 FR 51736, October 4, 1993), this action is not a "significant regulatory action" and is, therefore, not subject to review by the Office of Management and Budget. Because EPA has made a "good cause" finding that this action is not subject to notice and comment requirements

under the Administrative Procedure Act or any other statute, it is not subject to the regulatory flexibility provisions of the Regulatory Flexibility Act (5 U.S.C. 601 *et seq.*), or to sections 202 and 205 of the Unfunded Mandates Reform Act of 1995 (UMRA) (Public Law 104-4). In addition, this action does not significantly or uniquely affect small governments or impose a significant intergovernmental mandate, as described in sections 203 and 204 of the UMRA. This action also does not significantly or uniquely affect the communities of tribal governments, as specified by Executive Order 13084 (63 FR 27655, May 10, 1998). This action will not have substantial direct effects on the States, on the relationship between the national government and the States, or on the distribution of power and responsibilities among the various levels of government, as specified in Executive Order 13132 (64 FR 43255, August 10, 1999). This action also is not subject to Executive Order 13045 (62 FR 19885, April 23, 1997) because it is not economically significant.

Section 12(d) of the National Technology Transfer and Advancement Act (NTTAA) of 1995 (Public Law 104-113; 15 U.S.C. 272 note) directs EPA to use voluntary consensus standards in its regulatory and procurement activities unless to do so would be inconsistent with applicable law or otherwise impractical. The EPA is not proposing/adopting any voluntary consensus standards in this action.

This action does not involve special consideration of environmental justice related issues as required by Executive Order 12898 (59 FR 7629, February 16, 1994). In issuing these technical corrections, EPA has taken the necessary steps to eliminate drafting errors and ambiguity, minimize potential litigation, and provide a clear legal standard for affected conduct, as required by section 3 of Executive Order 12988 (61 FR 4729, February 7, 1996). The EPA has complied with Executive Order 12630 (53 FR 8859, March 15, 1988) by examining the takings implications of these technical corrections in accordance with the "Attorney General's Supplemental Guidelines for the Evaluation of Risk and Avoidance of Unanticipated

Takings" issued under the executive order. These technical corrections do not impose an information collection burden under the provisions of the Paperwork Reduction Act of 1995 (44 U.S.C. 3501 *et seq.*).

The Congressional Review Act (5 U.S.C. 801 *et seq.*), as added by the Small Business Regulatory Enforcement Fairness Act of 1996, generally provides that, before a rule may take effect, the agency promulgating the rule must submit a rule report, which includes a copy of the rule, to each House of the Congress and to the Comptroller General of the United States. Section 808 allows the issuing agency to make a rule effective sooner than otherwise provided by the Congressional Review Act if the agency makes a good cause finding that notice and public procedure is impracticable, unnecessary, or contrary to the public interest. This determination must be supported by a brief statement (5 U.S.C. 808(2)). As stated previously, EPA has made such a good cause finding, including the reasons therefor, and established an effective date of May 6, 2004. The EPA will submit a report containing the rule and other required information to the U.S. Senate, the U.S. House of Representatives, and the Comptroller General of the United States prior to publication of the rule in the **Federal Register**. This action is not a "major rule" as defined by 5 U.S.C. 804(2).

#### List of Subjects in 40 CFR Part 63

Environmental protection, Administrative practice and procedure, Air pollution control, Hazardous substances, Reporting and recordkeeping requirements.

Dated: April 28, 2004.

**Robert Brenner,**

*Acting Assistant Administrator for Air and Radiation.*

■ For the reasons set out in the preamble, title 40, chapter I, part 63 of the Code of Federal Regulations is amended as follows:

#### PART 63—[AMENDED]

■ 1. The authority citation for part 63 continues to read as follows:

**Authority:** 42 U.S.C. 7401 *et seq.*

#### Subpart MM—National Emission Standards for Hazardous Air Pollutants for Chemical Recovery Combustion Sources at Kraft, Soda, Sulfite, and Stand-Alone Semicheical Pulp Mills

■ 2. Section 63.866 is amended by revising paragraphs (a)(1) introductory text, (b), and (c)(4) to read as follows:

##### § 63.866 Recordkeeping requirements.

(a) \* \* \*

(1) Procedures for responding to any process parameter level that is inconsistent with the level(s) established under § 63.864(j), including the procedures in paragraphs (a)(1)(i) and (ii) of this section:

\* \* \* \* \*

(b) The owner or operator of an affected source or process unit must maintain records of any occurrence when corrective action is required under § 63.864(k)(1), and when a violation is noted under § 63.864(k)(2).

\* \* \* \* \*

(c) \* \* \*

(4) Records and documentation of supporting calculations for compliance determinations made under §§ 63.865(a) through (d);

\* \* \* \* \*

■ 3. Section 63.867 is amended by revising paragraph (c) introductory text to read as follows:

##### § 63.867 Reporting requirements.

\* \* \* \* \*

(c) *Excess emissions report.* The owner or operator must report quarterly if measured parameters meet any of the conditions specified in paragraph (k)(1) or (2) of § 63.864. This report must contain the information specified in § 63.10(c) of this part as well as the number and duration of occurrences when the source met or exceeded the conditions in § 63.864(k)(1), and the number and duration of occurrences when the source met or exceeded the conditions in § 63.864(k)(2). Reporting excess emissions below the violation thresholds of § 63.864(k) does not constitute a violation of the applicable standard.

\* \* \* \* \*

■ 4. Table 1 to Subpart MM is amended by revising the entries for §§ 63.7(a)(1) and 63.7(h) to read as follows:

TABLE 1 TO SUBPART MM OF PART 63—GENERAL PROVISIONS APPLICABILITY TO SUBPART MM

General Provisions reference	Summary of requirements	Applies to subpart MM	Explanation
63.7(a)(1)	Performance testing requirements—applicability.	Yes	§ 63.865(c)(1) specifies the only exemption from performance testing allowed under subpart MM.
63.7(h)	Waiver of performance tests	Yes	§ 63.865(c)(1) specifies the only exemption from performance testing allowed under subpart MM.

\* \* \* \* \*  
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**ENVIRONMENTAL PROTECTION AGENCY**

**40 CFR Part 439**

**Pharmaceutical Manufacturing Point Source Category**

*CFR Correction*

In Title 40 of the Code of Federal Regulations, Parts 425 to 699, revised as of July 1, 2003, the duplicated text from pages 401 and 408 is removed and the following text is reinstated.

Text to be reinstated on page 401:

\* \* \* \* \*

Appendix A to Part 439—Tables

**Authority:** 33 U.S.C. 1311, 1314, 1316, 1317, 1318, 1342 and 1361.

**Source:** 48 FR 49821, Oct. 27, 1983, unless otherwise noted.

**General**

**§ 439.0 Applicability.**

(a) This part applies to process wastewater discharges resulting from the research and manufacture of pharmaceutical products, which are generally, but not exclusively, reported under SIC 2833, SIC 2834 and SIC 2836 (1987 Standard Industrial Classification Manual).

(b) Although not reported under SIC 2833, SIC 2834 and SIC 2836, discharges from the manufacture of other pharmaceutical products to which this part applies include (but are not limited to):

- (1) Products manufactured by one or more of the four types of manufacturing processes described in subcategories A, B, C or D of this part, and considered

by the Food and Drug Administration to be pharmaceutical active ingredients;

(2) Multiple end-use products (e.g., components of formulations, chemical intermediates, or final products) derived from pharmaceutical manufacturing operations and intended for use primarily in pharmaceutical applications;

(3) Pharmaceutical products and intermediates not subject to other categorical limitations and standards, provided the manufacturing processes generate process wastewaters that are similar to those derived from the manufacture of pharmaceutical products elsewhere (an example of such a product is citric acid);

(4) Cosmetic preparations that are reported under SIC 2844 and contain pharmaceutical active ingredients, or active ingredients that are intended for the treatment of a skin condition. (These preparations do not include products such as lipsticks or perfumes that serve to enhance appearance, or provide a pleasing odor, but do not enhance skin care. Also excluded are deodorants, manicare preparations, shaving preparations and non-medicated shampoos that do not function primarily as a skin treatment.)

(c) The provisions of this part do not apply to wastewater discharges resulting from the manufacture of the following products, or as a result of providing one or more of the following services:

- (1) Surgical and medical instruments and apparatus reported under SIC 3841;
- (2) Orthopedic, prosthetic, and surgical appliances and supplies reported under SIC 3842;
- (3) Dental equipment and supplies reported under SIC 3843;
- (4) Medical laboratory services reported under SIC 8071;
- (5) Dental laboratory services reported under SIC 8072;
- (6) Outpatient care facility services reported under SIC 8081;

(7) Health and allied services reported under SIC 8091, and not classified elsewhere;

(8) Diagnostic devices other than those reported under SIC 3841;

(9) Animal feed products that include pharmaceutical active ingredients such as vitamins and antibiotics, where the major portion of the product is non-pharmaceutical, and the resulting process wastewater is not characteristic of process wastewater from the manufacture of pharmaceutical products;

(10) Food and beverage products fortified with vitamins or other pharmaceutical active ingredients, where the major portion of the product is non-pharmaceutical, and the resulting process wastewater is not characteristic of process wastewater from the manufacture of pharmaceutical products;

(11) Pharmaceutical products and intermediates subject to the provisions of 40 CFR part 414, provided their manufacture results in less than 50 percent of the total flow of process wastewater that is regulated by 40 CFR part 414 at the facility.

[63 FR 50424, Sept. 21, 1998]

**§ 439.1 General definitions.**

As used in this part:

(a) The general definitions, abbreviations and methods of analysis in 40 CFR part 401 shall apply.

\* \* \* \* \*

Text to be reinstated on page 408:

\* \* \* \* \*

standards specified in §§ 439.23 and 439.24.

[68 FR 12273, Mar. 13, 2003]

**§ 439.26 Pretreatment standards for existing sources (PSES).**

Except as provided in 40 CFR 403.7 and 403.13, any existing source subject to this subpart must achieve the