

Substances	Limitations
<p style="text-align: center;">* * *</p> <p><i>N,N</i>-Bis(2-hydroxyethyl)dodecanamide produced when diethanolamine is made to react with methyl laurate such that the finished product: Has a minimum melting point of 36 °C; has a minimum amide assay of 90 percent; contains no more than 2 percent by weight of free diethanolamine; and contains no more than 0.5 percent by weight of <i>N,N</i>, bis(2-hydroxyethyl)piperazine, as determined by paper chromatography method.</p> <p style="text-align: center;">* * *</p>	<p style="text-align: center;">* * *</p> <p>For use only:</p> <ol style="list-style-type: none"> 1. As an antistatic agent at levels not to exceed 0.5 percent by weight of molded or extruded polyethylene containers intended for contact with honey, chocolate syrup, liquid sweeteners, condiments, flavor extracts and liquid flavor concentrates, grated cheese, light and heavy cream, yogurt, and foods of Type VIII as described in Table 1 of § 176.170(c) of this chapter. 2. As an antistatic agent at levels not to exceed 0.2 percent by weight in polypropylene films complying with § 177.1520 of this chapter, and used in contact with food of Types I, II, III, IV, V, VI-B, VII, VIII, and IX described in Table 1 of § 176.170(c) of this chapter, and under conditions of use B through H described in Table 2 of § 176.170(c) of this chapter. The average thickness of such polypropylene film shall not exceed 0.001 inches (30 micrometers). <p style="text-align: center;">* * *</p>

Dated: May 23, 1997.
Fred R. Shank,
Director, Center for Food Safety and Applied Nutrition.
 [FR Doc. 97-15011 Filed 6-9-97; 8:45 am]
 BILLING CODE 4160-01-F

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 872

[Docket No. 95N-0033]

Dental Devices; Endodontic Dry Heat Sterilizer; Corrections and Technical Amendment

AGENCY: Food and Drug Administration, HHS.

ACTION: Final rule; corrections and technical amendment.

SUMMARY: The Food and Drug Administration (FDA) is correcting a final rule that appeared in the **Federal Register** of January 21, 1997 (62 FR 2900). The document issued a final rule to require the filing of a premarket approval application or a notice of completion of a product development protocol for the endodontic dry heat sterilizer, a medical device. The document was published with some errors. This document corrects those errors.

EFFECTIVE DATE: January 21, 1997.

FOR FURTHER INFORMATION CONTACT: Joseph M. Sheehan, Center for Devices and Radiological Health (HFZ-215), Food and Drug Administration, 1350 Piccard Dr., Rockville, MD 20850, 301-827-2974.

The Corrections

In FR Doc. 97-1336, beginning on page 2900 in the **Federal Register** of

Tuesday, January 21, 1997, the following corrections are made:

1. On page 2900, in the third column, in the second full paragraph, in the thirty-first line, "September 5, 1995" is corrected to read "June 22, 1995" and on that same page, in the third column, in the second full paragraph, in the thirty-second line, "August 7, 1995" is corrected to read "September 5, 1995".

2. On page 2902, in the second column, in the second paragraph, in the fourth line, and on that same page, in the second column, in the third paragraph, in the twenty-second line, "September 5, 1995" is corrected to read "April 21, 1997".

3. On page 2902, in the second column, in the third paragraph, in the twenty-eighth line, "August 7, 1995" is corrected to read "March 21, 1997."

The Technical Amendment

§ 872.6730 [Amended]

4. Section 872.6730 *Endodontic dry heat sterilizer* is amended in paragraph (c) by removing "September 5, 1995" each time it appears and adding in its place "April 21, 1997".

Dated: May 27, 1997.

Joseph A. Levitt,
Deputy Director for Regulations Policy, Center for Devices and Radiologica Health.

[FR Doc. 97-15013 Filed 6-9-97; 8:45 am]

BILLING CODE 4160-01-F

POSTAL SERVICE

39 CFR Part 111

Special Services Reform; Implementation Standards

AGENCY: Postal Service.

ACTION: Supplementary final rule.

SUMMARY: This supplementary final rule sets forth the remaining Domestic Mail

Manual (DMM) standards adopted by the Postal Service to implement the Decision of the Governors of the United States Postal Service on the Recommended Decision of the Postal Rate Commission on Special Services and Fees, Docket No. MC96-3. These standards constitute only minor changes or refinements to internal operational procedures that have been made since publication of the final rule in the **Federal Register** on May 12, 1997 (62 FR 26086-26099).

The standards in this supplementary final rule do not, in any way, affect the fees or attributes of the special services as they were published in the final rule for post office box service and caller service, certified mail, insurance (insured mail and Express Mail), parcel airlift, registered mail, return receipt service, return receipt for merchandise service, and stamped cards (formerly named postal cards). Although no substantive changes have been made to the final rule, this supplementary final rule does respond to comments that the Postal Service had sought with publication of the final rule.

EFFECTIVE DATE: June 8, 1997.

FOR FURTHER INFORMATION CONTACT: Neil Berger, (202) 268-2859.

SUPPLEMENTARY INFORMATION: On June 7, 1996, pursuant to its authority under 39 U.S.C. 3621, et seq., the Postal Service filed with the Postal Rate Commission (PRC) a request for a recommended decision on several special service reform proposals. The PRC designated the filing as Docket No. MC96-3. The PRC published a notice of the filing, with a description of the Postal Service's proposals, on June 21, 1996, in the **Federal Register** (61 FR 31968-31979).

Pursuant to 39 U.S.C. 3624, on April 2, 1997, the PRC issued its