presentations from the public will be scheduled between approximately 8:35 a.m. and 9 a.m., 11 a.m. and 11:30 a.m., 2 p.m. and 2:30 p.m., and 4:30 p.m. and 5 p.m. on July 22, 2004; and between approximately 10:15 a.m. and 11:15 a.m. and 2 p.m. and 2:30 p.m. on July 23, 2004. Time allotted for each presentation may be limited. Those desiring to make formal oral presentations should notify the contact person before July 12, 2004, and submit a brief statement of the general nature of the evidence or arguments they wish to present, the names and addresses of proposed participants, and an indication of the approximate time requested to make their presentation.

Persons attending FDA’s advisory committee meetings are advised that the agency is not responsible for providing access to electrical outlets.

FDA welcomes the attendance of the public at its advisory committee meetings and will make every effort to accommodate persons with physical disabilities or special needs. If you require special accommodations due to a disability, please contact AnnMarie Williams, Conference Management Staff, 301–594–1283, ext. 113, at least 7 days in advance of the meeting.

Notice of this meeting is given under the Federal Advisory Committee Act (5 U.S.C. app. 2).


Peter J. Pitts,
Associate Commissioner for External Relations.

[FR Doc. 04–13727 Filed 6–17–04; 8:45 am]
BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

Dental Products Panel of the Medical Devices Advisory Committee; Notice of Meeting

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

This notice announces a forthcoming meeting of a public advisory committee of the Food and Drug Administration (FDA). At least one portion of the meeting will be closed to the public.

Name of Committee: Dental Products Panel of the Medical Devices Advisory Committee.

General Function of the Committee: To provide advice and recommendations to the agency on FDA’s regulatory issues.

Date and Time: The meeting will be held on July 13, 2004, from 8 a.m. to 5 p.m.

Location: Hilton Washington DC North/Gaithersburg, Ballroom Salons A and B, 620 Perry Pkwy., Gaithersburg, MD.

Contact Person: Michael E. Adjodha, Center for Devices and Radiological Health (HFZ–480), Food and Drug Administration, 9200 Corporate Blvd., Rockville, MD 20850, 301–827–5283, ext. 123, e-mail: mea@cdrh.fda.gov, or FDA Advisory Committee Information Line 800–741–8138 (301–443–0572 in the Washington, DC area), code 3014512518. Please call the Information Line for up-to-date information on this meeting.

Agenda: The committee will discuss, make recommendations, and vote on a premarket approval application for a bone grafting material, which contains a wound-healing and revascularization agent, for treatment of dental osseous defects. Background information, including the agenda and questions for the committee, will be available to the public 1 business day before the meeting on the Internet at http://www.fda.gov/cdrh/panel/index.html. Material will be posted on July 12, 2004.

Procedure: On July 13, 2004, from 8:30 a.m. to 5 p.m., the meeting is open to the public. Interested persons may present data, information, or views, orally or in writing, on issues pending before the committee. Written submissions may be made to the contact person by July 1, 2004. Oral presentations from the public will be scheduled for approximately 30 minutes at the beginning of committee deliberations and for approximately 30 minutes near the end of the deliberations. Time allotted for each presentation may be limited. Those desiring to make formal oral presentations should notify the contact person before July 1, 2004, and submit a brief statement of the general nature of the evidence or arguments they wish to present, the names and addresses of proposed participants, and an indication of the approximate time requested to make their presentation.

Closed Committee Deliberations: On July 13, 2004, from 8 a.m. to 8:30 a.m., the meeting will be closed to permit FDA to present to the committee trade secret and/or confidential commercial information regarding pending and future agency issues (5 U.S.C. 552b(c)(4)).

Persons attending FDA’s advisory committee meetings are advised that the agency is not responsible for providing access to electrical outlets.

FDA welcomes the attendance of the public at its advisory committee meetings and will make every effort to accommodate persons with physical disabilities or special needs. If you require special accommodations due to a disability, please contact AnnMarie Williams, Conference Management Staff, 301–594–1283, ext. 113, at least 7 days in advance of the meeting.

Notice of this meeting is given under the Federal Advisory Committee Act (5 U.S.C. app. 2).


Peter J. Pitts,
Associate Commissioner for External Relations.

[FR Doc. 04–13727 Filed 6–17–04; 8:45 am]
BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 2004N–0226]

Food and Drug Administration Modernization Act of 1997: Modifications to the List of Recognized Standards, Recognition List Number: 010

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing a publication containing modifications the agency is making to the list of standards FDA recognizes for use in premarket reviews (FDA Recognized Consensus Standards). This publication, entitled “Modifications of the List of Recognized Standards, Recognition List Number: 010” (Recognition List Number: 010), will assist manufacturers who elect to declare conformity with consensus standards to meet certain requirements for medical devices.

DATES: Submit written or electronic comments concerning this document at any time. See section VII of this document for the effective date of the recognition of standards announced in this document.

ADDRESSES: Submit written requests for single copies on a 3.5” diskette of “Modification to the List of Recognized Standards, Recognition List Number: 010” to the Division of Small Manufacturers Assistance, Center for Devices and Radiological Health (HFZ–220), Food and Drug Administration, 1350 Piccard Dr., Rockville, MD 20850. Send two self-addressed adhesive labels to assist that office in processing your
requests, or fax your request to 301–443–8818. Submit written comments concerning this document or to recommend additional standards for recognition to the contact person (see FOR FURTHER INFORMATION CONTACT). Submit electronic comments by e-mail: standards@cdrh.fda.gov. This document may also be accessed on FDA’s Internet site at http://www.fda.gov/cdrh/fedregin.html. See section VI of this document for electronic access to the searchable database for the current list of “FDA Recognized Consensus Standards,” including Recognition List Number: 010 modifications and other standards related information.

FOR FURTHER INFORMATION CONTACT:
Carol L. Herman, Center for Devices and Radiological Health (HFZ–84), Food and Drug Administration, 2094 Gaither Rd., Rockville, MD 20850, 301–594–4766, ext. 156.

SUPPLEMENTARY INFORMATION:

I. Background

Section 204 of the Food and Drug Administration Modernization Act of 1997 (FDAMA) (Public Law 105–115) amended section 514 of the Federal Food, Drug, and Cosmetic Act (the act) (21 U.S.C. 360d). Amended section 514 allows FDA to recognize consensus standards, developed by international and national organizations, for use in satisfying portions of device premarket review submissions or other requirements.

In a notice published in the Federal Register of February 25, 1998 (63 FR 9561), FDA announced the availability of a guidance document entitled “Recognition and Use of Consensus Standards.” This notice described how FDA will implement its standard recognition program and provided the initial list of recognized standards.

In Federal Register notices published on October 16, 1998 (63 FR 55617), July 12, 1999 (64 FR 37546), November 15, 2000 (65 FR 69022), May 7, 2001 (66 FR 23032), January 14, 2002 (67 FR 1774), October 2, 2002 (67 FR 61893), April 28, 2003 (68 FR 22391), and March 8, 2004 (69 FR 10712), FDA modified its initial list of recognized standards. These notices described the addition, withdrawal, and revision of certain standards recognized by FDA. The agency maintains hypertext markup language (HTML) and portable document format (PDF) versions of the list of FDA Recognized Consensus Standards. Both versions are publicly accessible at the agency’s Internet site. See section VI of this document for electronic access information. Interested persons should review the supplementary information sheet for the standard to understand fully the extent to which FDA recognizes the standard.

II. Modifications to the List of Recognized Standards, Recognition List Number: 010

FDA is announcing the addition, withdrawal, correction, and revision of certain consensus standards the agency will recognize for use in satisfying premarket reviews and other requirements for devices. FDA will incorporate these modifications in the list of FDA Recognized Consensus Standards in the agency’s searchable database. FDA will use the term “Recognition List Number: 010” to identify these current modifications.

In the following table, FDA describes modifications that involve: (1) The withdrawal of standards and their replacement by others, (2) the correction of errors made by FDA in listing previously recognized standards, and (3) the changes to the supplementary information sheets of recognized standards that describe revisions to the applicability of the standards.

In section III of this document, FDA lists modifications the agency is making that involve the initial addition of standards not previously recognized by FDA.

A. Anesthesia

<table>
<thead>
<tr>
<th>Old Item No.</th>
<th>Standard</th>
<th>Change</th>
<th>Replacement Item No.</th>
</tr>
</thead>
<tbody>
<tr>
<td>19</td>
<td>ISO 8382:1988, Resuscitators Intended for Use with Humans</td>
<td>Processes impacted, extent of recognition, relevant guidance</td>
<td>19</td>
</tr>
<tr>
<td>42</td>
<td>ISO 5360:1993, Anaesthetic vaporizers—Agent-specific filling systems</td>
<td>Devices affected, processes impacted, extent of recognition</td>
<td>42</td>
</tr>
</tbody>
</table>

B. General

<table>
<thead>
<tr>
<th>Old Item No.</th>
<th>Standard</th>
<th>Change</th>
<th>Replacement Item No.</th>
</tr>
</thead>
<tbody>
<tr>
<td>2</td>
<td>IEC 60601–1, Medical Electrical Equipment—Part 1: General Requirements for Safety</td>
<td>Contact person</td>
<td>2</td>
</tr>
</tbody>
</table>

C. General Hospital/General Plastic Surgery
<table>
<thead>
<tr>
<th>Old Item No.</th>
<th>Standard</th>
<th>Change</th>
<th>Replacement Item No.</th>
</tr>
</thead>
<tbody>
<tr>
<td>10</td>
<td>IEC 60601–2–38, Medical Electrical Equipment—Part 2: Particular Requirements for the Safety of Electrically Operated Hospital Beds</td>
<td>Withdrawn and replaced with newer version</td>
<td>111</td>
</tr>
</tbody>
</table>

**D. In Vitro Diagnostic**

<table>
<thead>
<tr>
<th>Old Item No.</th>
<th>Standard</th>
<th>Change</th>
<th>Replacement Item No.</th>
</tr>
</thead>
<tbody>
<tr>
<td>47</td>
<td>NCCLS MM2–A2 Immunoglobulin and T-Cell Receptor Gene Rearrangement Assays; Approved Guideline—Second Edition</td>
<td>Withdrawn and replaced with newer version</td>
<td>98</td>
</tr>
<tr>
<td>84</td>
<td>CEN 13640, Stability Testing of In Vitro Diagnostic Reagents</td>
<td>Correction to date of standard</td>
<td>84</td>
</tr>
</tbody>
</table>

**E. Materials**

<table>
<thead>
<tr>
<th>Old Item No.</th>
<th>Title of Standard</th>
<th>Change</th>
<th>Replacement Item No.</th>
</tr>
</thead>
<tbody>
<tr>
<td>39</td>
<td>ASTM F2052–02, Standard Test Method for Measurement of Magnetically Induced Displacement Force on medical Devices in the Magnetic Resonance Environment</td>
<td>Recognizing a newer version with a revised title</td>
<td>70</td>
</tr>
<tr>
<td>Old Item No.</td>
<td>Title of Standard</td>
<td>Change</td>
<td>Replacement Item No.</td>
</tr>
<tr>
<td>-------------</td>
<td>-----------------------------------------------------------------------------------</td>
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<td>----------------------</td>
</tr>
<tr>
<td>76</td>
<td>ISO 6474–94, Implants for surgery—Ceramic materials based on high purity alumina</td>
<td>Transferred from Orthopedics 76 to Materials 66</td>
<td>66</td>
</tr>
</tbody>
</table>

**F. Radiology**

<table>
<thead>
<tr>
<th>Old Item No.</th>
<th>Standard</th>
<th>Change</th>
<th>Replacement Item No.</th>
</tr>
</thead>
<tbody>
<tr>
<td>5</td>
<td>ANSI Ph 2.50–1983, Photography—Direct-Exposing Medical and Dental Radiographic Film/Process Systems—Determination of ISO Speed and Average Gradient</td>
<td>Title correction</td>
<td>5</td>
</tr>
<tr>
<td>7</td>
<td>ISO/IEC 10918–1:1994, Information Technology—Digital Compression and Coding of Continuous—Tone Still Images: Requirements and Guidelines</td>
<td>Title correction</td>
<td>7</td>
</tr>
<tr>
<td>8</td>
<td>IEC 60336 (R1993), X-ray Tube Assemblies for Medical Diagnosis Characteristics of Focal Spots</td>
<td>Title correction</td>
<td>8</td>
</tr>
<tr>
<td>17</td>
<td>NEMA MS 8–1993 (2000), Characterization of the Specific Absorption Rate for Magnetic Resonance Imaging Systems</td>
<td>Reaffirmation</td>
<td>17</td>
</tr>
<tr>
<td>23</td>
<td>NEMA XR 10–1986 (R1992, R1998), Measurement of the Maximum Symmetrical Radiation Field from a Rotating Node X-ray Tube used for Medical Diagnosis</td>
<td>Reaffirmation</td>
<td>23</td>
</tr>
<tr>
<td>24</td>
<td>NEMA XR 11–1993 (R1999), Test Standard for Determination of the Limiting Spatial Resolution of X-ray Image Intensifier Systems</td>
<td>Title correction</td>
<td>24</td>
</tr>
<tr>
<td>29</td>
<td>NEMA XR 19–1993 (R1999), Thermal and Loading Characteristics of X-ray Tubes used for Medical Diagnosis</td>
<td>Reaffirmation</td>
<td>29</td>
</tr>
<tr>
<td>Old Item No.</td>
<td>Standard</td>
<td>Change</td>
<td>Replacement Item No.</td>
</tr>
<tr>
<td>-------------</td>
<td>--------------------------------------------------------------------------</td>
<td>---------------------------------------------</td>
<td>----------------------</td>
</tr>
<tr>
<td>44</td>
<td>AIUM AOMS—Acoustic Output Measurement Standard for Diagnostic Ultrasound Equipment</td>
<td>Title correction and reaffirmation</td>
<td>44</td>
</tr>
<tr>
<td>46</td>
<td>AIUM RTD1—Standard for Real-Time Display of Thermal and Mechanical Acoustic Output Indices on Diagnostic Ultrasound Equipment Revision 1</td>
<td>Title correction and reaffirmation</td>
<td>46</td>
</tr>
<tr>
<td>48</td>
<td>AIUM AOL—Acoustic Output Labeling Standard for Diagnostic Ultrasound Equipment: A Standard for How Manufacturers Should Specify Acoustic Output Data</td>
<td>Title correction</td>
<td>48</td>
</tr>
<tr>
<td>66</td>
<td>AIUM MUS—Medical Ultrasound Safety</td>
<td>Title correction and reaffirmation</td>
<td>66</td>
</tr>
<tr>
<td>72</td>
<td>NEMA UD 3–1998, Revision 1, Standard for Real Time Display of Thermal and Mechanical Acoustic Output Indices on Diagnostic Ultrasound Equipment</td>
<td>Title correction</td>
<td>72</td>
</tr>
<tr>
<td>11</td>
<td>NEMA MS 2–2003, Determination of Two-Dimensional Geometric Distortion in Diagnostic Magnetic Resonance Images</td>
<td>Withdrawn and replaced with newer version</td>
<td>95</td>
</tr>
<tr>
<td>12</td>
<td>NEMA MS 3–2003, Determination of Image Uniformity in Diagnostic Magnetic Resonance Images</td>
<td>Withdrawn and replaced with newer version</td>
<td>96</td>
</tr>
<tr>
<td>77</td>
<td>NEMA MS–1–2001, Determination of Signal to Noise Ratio (SNR) in Diagnostic Magnetic Resonance Images</td>
<td>Withdrawn and replaced with newer version</td>
<td>97</td>
</tr>
<tr>
<td>69</td>
<td>NEMA MS 6–1991 (R2000), Characterization of Special Purpose Coils for Diagnostic Magnetic Resonance Images</td>
<td>Reaffirmation</td>
<td>69</td>
</tr>
<tr>
<td>3</td>
<td>ANSI IT1.49–1995, Photography (Films)—Medical Radiographic Cassettes/Screens/Films-Dimensions</td>
<td>Withdrawn and replaced with Item 98</td>
<td>98</td>
</tr>
<tr>
<td>14</td>
<td>NEMA MS 5–2003, Determination of Slice Thickness in Diagnostic Magnetic Resonance Imaging</td>
<td>Withdrawn and replaced with newer version</td>
<td>99</td>
</tr>
</tbody>
</table>

### G. Sterility

<table>
<thead>
<tr>
<th>Old Item No.</th>
<th>Standard</th>
<th>Change</th>
<th>Replacement Item No.</th>
</tr>
</thead>
</table>

### III. Listing of New Entries

The listing of new entries and consensus standards added as “Modifications to the List of Recognized Standards,” under Recognition List Number: 010, is as follows:

#### A. Anesthesia

<table>
<thead>
<tr>
<th>Item No.</th>
<th>Title of Standard</th>
<th>Reference No. and Date</th>
</tr>
</thead>
<tbody>
<tr>
<td>47</td>
<td>Ancillary devices for expired air resuscitation</td>
<td>AS 4259–1995</td>
</tr>
</tbody>
</table>

#### B. General
### IV. List of Recognized Standards

FDA maintains the agency’s current list of FDA Recognized Consensus Standards in a searchable database that may be accessed directly at FDA’s Internet site at http://www.accessdata.fda.gov/scripts/cdrh/cfdocs/cfStandards/search.cfm. FDA will incorporate the modifications and minor revisions described in this notice into the database, and upon publication in the Federal Register, this recognition of consensus standards will be effective. FDA will announce additional modifications and minor revisions to the list of recognized consensus standards, as needed, in the Federal Register once a year, or more often, if necessary.

#### VI. Electronic Access

In order to receive “Guidance on the Recognition and Use of Consensus Standards” via your fax machine, call the CDRH Facts-On-Demand system at 800–899–0381 or 301–827–0111 from a touch-tone telephone. Press 1 to enter the system. At the second voice prompt press 1 to order a document. Enter the document number 321 followed by the pound sign (#). Follow the remaining voice prompts to complete your request.

You may obtain a copy of “Guidance on the Recognition and Use of Consensus Standards” by using the Internet. CDRH maintains a site on the Internet for easy access to information including text, graphics, and files that you may download to a personal computer with access to the Internet. Updated on a regular basis, the CDRH home page includes the guidance as well as the current list of recognized standards and other standards related documents. After publication in the Federal Register, this notice announcing “Modification to the List of Recognized Standards, Recognition List Number: 010” will be available on the CDRH home page. You may access the CDRH home page at http://www.fda.gov/cdrh.


#### VII. Submission of Comments and Effective Date

Interested persons may submit to the contact person (see FOR FURTHER INFORMATION CONTACT) written or electronic comments regarding this document. Two copies of any mailed comments are to be submitted, except that individuals may submit only one paper copy. Comments are to be identified

<table>
<thead>
<tr>
<th>Item No.</th>
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</tr>
</thead>
<tbody>
<tr>
<td>33</td>
<td>Medical Electrical Equipment—Parts 1 to 8: General requirements for safety—Collateral Standard: Alarm systems—Requirements, tests, and guidelines—General requirements and guidelines for alarm systems in medical equipment</td>
<td>IEC 60601–1–8:2003</td>
</tr>
</tbody>
</table>

#### C. In Vitro Diagnostic

<table>
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<tr>
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#### D. Materials

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</thead>
</table>

#### E. Radiology

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<tr>
<th>Item No.</th>
<th>Title of Standard</th>
<th>Reference No. and Date</th>
</tr>
</thead>
<tbody>
<tr>
<td>98</td>
<td>Medical Electrical Equipment—Dosimeters with Ionization Chambers as Used in Radiotherapy</td>
<td>IEC 60731—Amendment 1 2002–06</td>
</tr>
</tbody>
</table>
with the docket number found in brackets in the heading of this
document. FDA will consider any
comments received in determining
whether to amend the current listing of
“Modifications to the List of Recognized
Standards, Recognition List Number:
010.” These modifications to the list or
recognized standards are effective upon
publication of this notice in the Federal
Register.

Linda S. Kahan,
Deputy Director, Center for Devices and
Radiological Health.

DEPARTMENT OF HEALTH AND
HUMAN SERVICES
Substance Abuse and Mental Health
Services Administration

Notice of a Meeting

Pursuant to Public Law 92–463, notice is hereby given of a meeting of
the Substance Abuse and Mental Health Services Administration (SAMHSA)
National Advisory Council on June 30
and July 1, 2004.

The SAMHSA National Advisory
Council meeting will be open to the
public. The meeting will include the
SAMHSA Administrator’s Report, and
discussions on Mental Health Systems
Transformation, the Co-occurring
Report, SAMHSA’s Strategic Prevention
Framework Initiative, suicide
prevention, and SAMHSA’s Access to
Recovery Initiative. The meeting will
also include a discussion of the
Agency’s current legislative highlights,
and an update on the Interagency
Coordinating Committee on the
Prevention of Underage Drinking.

Attendance by the public will be
limited to space available. Public
comments are welcome. Please
communicate with the individual listed
as contact below to make arrangements
to comment or to request special
accommodations for persons with
disabilities.

Substantive program information, a
summary of the meeting, and a roster of
Council members may be obtained
either by accessing the SAMHSA
www.samhsa.gov/council/council or by
communicating with the contact whose
name and telephone number is listed
below. The transcript for the meeting
will also be available on the SAMHSA
Council Web site.

Committee Name: SAMHSA National
Advisory Council.

Date/Time: Wednesday, June 30,
2004, 9 a.m. to 4:45 p.m. (Open),
Thursday, July 1, 2004, 9 a.m. to 12:15
p.m. (Open).

Place: Hilton Washington Embassy
Row Hotel, Ambassador Room, 2015
Massachusetts Avenue, NW.,
Washington, DC 20036.

Contact: Toian Vaughn, Executive
Secretary, 5600 Fishers Lane, Parklawn
Building, Room 12C–05, Rockville, MD
20857. Telephone: (301) 443–7016;
FAX: (301) 443–1450 and E-mail:
tvaughn@samhsa.gov.

Toian Vaughn,
Committee Management Officer, SAMHSA.

DEPARTMENT OF HOMELAND
SECURITY
Federal Emergency Management
Agency

Agency Information Collection
Activities: Submission for OMB
Review; Comment Request

AGENCY: Federal Emergency
Management Agency, Emergency
Preparedness and Response Directorate,

ACTION: Notice and request for
comments.

SUMMARY: The Federal Emergency
Management Agency has submitted the
following proposed information
collection to the Office of Management
and Budget for review and clearance in
accordance with the requirements of the
Paperwork Reduction Act of 1995 (44

Title: Federal Emergency Management
Agency (FEMA) Mitigation Success
Story Database.

Type of Information Collection:
Existing collection in use without an
OMB Control Number.

OMB Number: OMB No. 1660–NEW6.

Abstract: This Web-based database
serves a dual purpose in providing a
centralized and user-friendly venue for
gaining and disseminating knowledge
about effective and efficient mitigation
strategies implemented in communities
nationwide. By sharing information,
communities and individuals can learn
about available Federal programs to
support implementation of mitigation
projects relevant to individual
conditions and characteristics.

Affected Public: State, local and tribal
governments, individuals, business or
other for-profit organizations, not-for
profit institutions, and Federal
government.

Number of Respondents: 150.

Estimated Time per Respondent: The
electronic submission takes
approximately 30 minutes for filling in
all fields in the submission form, and
approximately 1 hour to conceptualize
the narrative description for a total of
1.5 hours. Respondents choosing to
supply the information directly to
FEMA Regional or HQ staff or to a
Disaster Field Office (DFO) staff may
spend up to 4 hours, which includes
initial interview and follow-up sessions
(when needed and agreed upon by the
respondent on a voluntary basis).

Estimated Total Annual Burden
Hours: 563 hours.

Frequency of Response: Time.

Comments: Interested persons are
invited to submit written comments on
the proposed information collection to
the Office of Information and Regulatory
Affairs at OMB, Attention: Desk Officer
for the Emergency Preparedness and
Response Directorate/Federal
Emergency Management Agency, U.S.
Department of Homeland Security, 725
17th Street, NW., Docket Library Room
10102, Washington, DC 20503.

Comments must be submitted on or
before July 19, 2004. In addition,
interested persons may also send
comments to FEMA (see contact
information below).

FOR FURTHER INFORMATION CONTACT:
Requests for additional information or
copies of the information collection
should be made to Muriel B. Anderson,
Chief, Records Management, FEMA at
500 C Street, SW., Room 316,
Washington, DC 20472, facsimile
number (202) 646–3347, or e-mail
address FEMA–Information-
Collections@dhs.gov.

Edward W. Kernan,
Branch Chief, Information Resources
Management Branch, Information
Technology Services Division.

DEPARTMENT OF HOMELAND
SECURITY
Federal Emergency Management
Agency

FEMA–1518–DR

Iowa; Amendment No. 3 to Notice of a
Major Disaster Declaration

AGENCY: Federal Emergency
Management Agency, Emergency
Preparedness and Response Directorate,
Department of Homeland Security.