Dated: October 31, 1996.

Joseph R. Carter,

Acting Associate Director for Management and Operations, Centers for Disease Control and Prevention (CDC).

[FR Doc. 96–28484 Filed 11–5–96; 8:45 am]

BILLING CODE 4163–18–P

National Institute for Occupational Safety and Health Draft Document ''Engineering Control Guidelines for Hot Mix Asphalt Pavers''; Correction

AGENCY: National Institute for Occupational Safety and Health (NIOSH), Centers for Disease Control and Prevention (CDC), Department of Health and Human Services.

ACTION: Notice; corrections.

SUMMARY: This notice makes corrections in the request for comments on the draft document ''Engineering Control Guidelines for Hot Mix Asphalt Pavers'' published in the Federal Register on Thursday, October 3, 1996 [61 FR 51708].

FOR FURTHER INFORMATION CONTACT: Technical information may be obtained from Joann Wess or Ralph Zumwalde, NIOSH, CDC, 4676 Columbia Parkway, M/S C–32, Cincinnati, Ohio 45226, telephone (513) 533–8319.

SUPPLEMENTARY INFORMATION: In the notice document beginning on page 51708 in the issue of Thursday, October 3, 1996, make the following corrections:

On page 51711, in the first column, the following equation should be inserted in the last sentence of paragraph 5 after ''* * * the following equation:''

$$Q_{(exh)} = \frac{Q_{(SF_6)}}{C^*_{(SF_6)}} \times 10^6$$

On page 51711, in the second column, the paragraph beginning, ''To quantify capture efficiency * * *'' line 11 should read, ''test and should be ±3% or better. The''.

On page 51711, in the second column, the paragraph beginning, ''At least five consecutive measurements * * *'' the following equation should be inserted after ''* * * the following equation:''

$$\eta = \frac{C_{(SF_6)}}{C^*_{(SF_6)}} \times 100$$

On page 51711, in the second column, the paragraph beginning ''If the SF_6 volumetric * * *'' the following equation should be inserted after ''* * * using the following:''

$$\eta = \frac{C_{(SF_6)} \times Q_{(exh)}}{Q_{(SF_6)} \times 10^6} \times 100$$

On page 51712, first column, the paragraph beginning ''At least five consecutive measurements * * *'', disregard the equation shown after ''* * * volumetric flow rate from Equation 1.'' and insert the following equation instead.

$$Q_{(exh)} = \frac{\frac{0.903}{28.3}}{21.85 - 0.0057} \times 10^6 = 1460 \ cfm$$

On page 51712, second column, under the heading ''Statistics,'' after the sentence ''Calculate the estimated standard deviation:'' disregard the equation shown and insert the following equation instead.

$s=\{(87.9-87.5)^2+(92.1-87.5)^2+(83.3-87.5)^2+(86.7-87.5)^2)/(4-1)\}^{0.5}$

$=\{(0.16+21.16+17.64+0.64)/3\}^{0.5}=3.63$

On page 51712, third column, after the sentence ''Calculate a test statistic (T):'', disregard the test statistic shown and insert the following:

$$T = \frac{m - (t \times s)}{n^{0.5}}$$

For this example:

$$T = \frac{87.5 - (2.35 \times 3.63)}{4^{0.5}} = 83.2$$

Dated: October 29, 1996.

Linda Rosenstock,

Director, National Institute for Occupational Safety and Health, Centers for Disease Control and Prevention (CDC).

[FR Doc. 96–28499 Filed 11–5–96; 8:45 am]

BILLING CODE 4163–19–P

Board of Scientific Counselors, National Center for Infectious Diseases: Meeting

In accordance with section 10(a)(2) of the Federal Advisory Committee Act (Pub. L. 92–463), the Centers for Disease Control and Prevention (CDC) announces the following committee meeting.

Name: Board of Scientific Counselors, National Center for Infectious Diseases (NCID).

Times and Dates: 11:00 a.m.–5:30 p.m., December 5, 1996; 8:00 a.m.–2:30 p.m., December 6, 1996.

Place: CDC, Auditorium B, 1600 Clifton Road, NE, Atlanta, Georgia 30333.

Status: Open to the public, limited only by the space available.

Purpose: The Board of Scientific Counselors, NCID, provides advice and guidance to the Director, CDC, and Director, NCID, in the following areas: program goals and objectives; strategies; program organization and resources for infectious disease prevention and control; and program priorities.

Matters To Be Discussed: The agenda will focus on:

1. NCID Update.
2. Minority and Women's Health.
3. Global Emerging Infectious Diseases.
4. Current Scientific Issues.
5. Workgroup Sessions: Emerging Infectious Disease FY 1997 and FY 1998 Planning:

a. Surveillance and Response Capacity
b. Research
c. Prevention and Control
d. Laboratory Infrastructure

6. Workgroup Reports.
7. Recommendations.

Other agenda items include announcements/introductions; follow-up on actions recommended by the Board (May 1996); and consideration of future directions, goals, and recommendations.

Agenda items are subject to change as priorities dictate.

Written comments are welcome and should be received by the contact person listed below prior to the opening of the meeting.

Contact Person for More Information: Diane S. Holley, Office of the Director, NCID, CDC, Mailstop C–20, 1600 Clifton Road, NE, Atlanta, Georgia 30333, telephone 404/639–0078.

Dated: October 25, 1996.

Carolyn J. Russell,

Director, Management Analysis and Services Office, Centers for Disease Control and Prevention (CDC).

[FR Doc. 96–28498 Filed 11–5–96; 8:45 am]

BILLING CODE 4163–18–P

Food and Drug Administration

[Docket No. 96N–0403]

Agency Information Collection Activities: Proposed Collection; Comment Request; Extension

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing an opportunity for public comment on the proposed collection of certain information by the agency. Under the Paperwork Reduction Act of 1995, Federal agencies are required to publish notice in the Federal Register concerning each proposed collection of information, including each proposed extension of an existing collection of information, and to allow 60 days for public comment in response to the notice. This notice solicits comments on the recordkeeping and labeling requirements for food irradiation processors.

DATES: Submit written comments on the collection of information by January 6, 1997.

ADDRESSES: Submit written comments on the collection of information to the Dockets Management Branch (HFA–305), Food and Drug Administration, 12420 Parklawn Dr., rm. 1–23, Rockville, MD 20857. All comments should be identified with the docket number found in brackets in the heading of this document.

FOR FURTHER INFORMATION CONTACT: Kim A. Sanders, Office of Information Resources Management (HFA–250), Food and Drug Administration, 5600 Fishers Lane, rm. 16B–19, Rockville, MD 20857, 301–827–1473.

SUPPLEMENTARY INFORMATION: Under the Paperwork Reduction Act of 1995 (the PRA) (44 U.S.C. 3501–3520), Federal agencies must obtain approval from the Office of Management and Budget (OMB) for each collection of information they conduct or sponsor. ''Collection of information'' is defined in 44 U.S.C. 3502(3) and 5 CFR 1320.3(c) and includes agency requests or requirements that members of the public submit reports, keep records, or provide information to a third party. Section 3506(c)(2)(A) of the PRA (44 U.S.C. 3506(c)(2)(A)) requires Federal agencies to provide a 60-day notice in the Federal Register concerning each proposed collection of information, including each proposed extension of an existing collection of information, before submitting the collection to OMB for approval. To comply with this requirement, FDA is publishing notice of the proposed collection of information listed below.

With respect to the following collection of information, FDA invites comments on: (1) Whether the proposed collection of information is necessary for the proper performance of FDA's functions, including whether the information will have practical utility; (2) the accuracy of FDA's estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques, when appropriate, and other forms of information technology.

Irradiation in the Production, Processing, and Handling of Food (21 CFR Part 179)—(OMB Control Number 0910–0186)—Extension

Under sections 201(s) and 409 of the Federal Food, Drug, and Cosmetic Act (the act), food irradiation is subject to regulation as a food additive (21 U.S.C. 321(s) and 348). The regulations providing for uses of irradiation in the production, processing, and handling of food are found in part 179 (21 CFR part 179).

Section 179.25(e) requires that food processors who treat food with radiation make and retain, for 1 year past the expected shelf life of the products up to a maximum of 3 years, specified records relating to the irradiation process (e.g., the food treated, lot identification, scheduled process, etc.).

Section 179.26(c) requires that food processors label retail packages of irradiated foods with an FDA prescribed logo and statement, ''Treated with radiation'' or ''Treated by irradiation.'' To assure safe use of radiation sources, § 179.21(b)(1) requires that the label of sources bear appropriate and accurate information identifying the source of radiation (§ 179.21(b)(1)(i)) and the maximum energy of radiation emitted by X-ray tube sources (§ 179.21(b)(1)(ii)). Section 179.21(b)(2) requires that the label or accompanying labeling bear adequate directions for installation and use (§ 179.21(b)(2)(i)), a statement that