Abstract: BLM proposes to collect information from Alaska Native permittees under its Reindeer Grazing Program to assess the compatibility of grazing on the land with multiple-use objectives for the area.

Bureau Form Number: 4201–1, Grazing Lease or Permit Application; 4132–2, Grazing Permit.

Frequency: Once.

Description of Respondents: Alaska Natives, groups of Alaska Natives, or associations or corporations of Alaska Natives who want to graze reindeer on public lands in Alaska that are vacant and unappropriated.

Annual Responses: 6.

Annual Burden Hours: 7.5.

Bureau Clearance Officer: Carole Smith, (202) 452–0367.

Dated: August 18, 1998

Carole J. Smith,

Bureau of Land Management, Information Clearance Officer.

[FR Doc. 98–23598 Filed 9–1–98; 8:45 am]

BILLING CODE 4310–84–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 98N–0363]

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

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice; correction.

SUMMARY: The Food and Drug Administration (FDA) is correcting a notice that appeared in the **Federal Register** of July 8, 1998 (63 FR 36921). The document announced an opportunity for public comment on the proposed collection of certain information by the agency. The document published with an incorrect address. This document corrects that error.

FOR FURTHER INFORMATION CONTACT: Carolyn C. Harris, Office of Policy (HF–27), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–443–2994.

SUPPLEMENTARY INFORMATION: In FR Doc. 98–18145, appearing on page 36921, in the **Federal Register** of July 8, 1998, the following correction is made: On page 36921, in the third column, under the ''**ADDRESSES**'' caption, beginning in the fifth line ''12420 Parklawn Dr., rm. 1–23, Rockville, MD 20857'' is corrected to read ''5630 Fishers Lane, rm. 1061, Rockville, MD 20852''.

Dated: August 26, 1998.

William K. Hubbard,

Associate Commissioner for Policy Coordination.

[FR Doc. 98–23584 Filed 9–1–98; 8:45 am]

BILLING CODE 4160–01–F

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

Technical Electronic Products Radiation Safety Standards 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). The meeting will be open to the public.

Name of Committee: Technical Electronic Products Radiation Safety Standards Committee.

General Function of the Committee: To provide advice on technical feasibility, reasonableness, and practicality of performance standards for electronic products to control the emission of radiation under 42 U.S.C. 263f(f)(1)(A).

Date and Time: The meeting will be held on September 23, 1998, 8:30 a.m. to 6 p.m., and September 24, 1998, 8:30 a.m. to 3:45 p.m.

Location: Hilton Hotel, Salons A and B, 620 Perry Pkwy., Gaithersburg, MD.

Contact Person: Orhan H. Suleiman, Center for Devices and Radiological Health (HFZ–240), Food and Drug Administration, 1350 Piccard Dr., Rockville, MD 20850, 301–594–3332, or FDA Advisory Committee Information Line, 1–800–741–8138 (301–443–0572 in the Washington, DC area), code 12399. Please call the Information Line for up-to-date information on this meeting.

Agenda: On September 23, 1998, the committee will: (1) Discuss possible proposed amendments to performance standards for fluoroscopic imaging systems (21 CFR 1020), and (2) hear presentations on x-ray scanning security devices and the indoor tanning industry. On September 24, 1998, the committee will: (1) Discuss electronic article surveillance systems and metal detectors, and the potential for electromagnetic interference with the operation of medical devices, and (2) hear a presentation on medical telemetry systems and the impact of changes in communications standards.

Procedure: 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 September 8, 1998. On September 23, 1998, oral presentations from the public will be scheduled between approximately 11:15 a.m. and 12 m., and between approximately 4:30 p.m. and 5:15 p.m., and on September 24, 1998, oral presentations from the public will be scheduled between approximately 1 p.m. and 2 p.m. Time allotted for each presentation may be limited. Those desiring to make formal oral presentations should notify the contact person before September 15, 1998, 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.

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

Dated: August 25, 1998.

Randolph Wykoff,

Acting Deputy Commissioner for Operations.

[FR Doc. 98–23585 Filed 9–1–98; 8:45 am]

BILLING CODE 4160–01–F

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 96N–0373]

Agency Information Collection Activities; Announcement of OMB Approval; Request for Information From U.S. Processors that Export to the European Community

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a collection of information entitled ''Request for Information From U.S. Processors that Export to the European Community'' has been approved by the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995 (the PRA).

FOR FURTHER INFORMATION CONTACT: Margaret R. Schlosburg, Office of Information Resources Management (HFA–250), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–1223.

SUPPLEMENTARY INFORMATION: In the **Federal Register** of June 1, 1998 (63 FR 29738), the agency announced that the proposed information collection had been submitted to OMB for review and