Transcripts: Transcripts of the public workshop may be requested in writing from the Freedom of Information Office (HFI–35), Food and Drug Administration, 5600 Fishers Lane, rm. 6–30, Rockville, MD 20857, approximately 15 working days after the public workshop at a cost of 10 cents per page. A transcript of the public workshop will be available on the Internet at *http://www.fda.gov/cber/minutes/workshop-min.htm*.

Dated: June 11, 2007.

Jeffrey Shuren,

Assistant Commissioner for Policy.

[FR Doc. E7–11613 Filed 6–15–07; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 2007N–0217]

Licensure of Apheresis Blood Products; Public Workshop

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice of public workshop.

The Food and Drug Administration (FDA) is announcing a public workshop entitled: Licensure of Apheresis Blood Products. The purpose of the public workshop is to educate industry on the licensure requirements and license application procedures for Platelets, Pheresis; Red Blood Cells; and Plasma collected by automated blood cell separator devices.

Date and Time: The public workshop will be held on August 15, 2007, from 8 a.m. to 5 p.m.

Location: The public workshop will be held at the Lister Hill Center Auditorium, Bldg. 38A, National Institutes of Health, 8800 Rockville Pike, Bethesda, MD 20894.

Contact Person: Rhonda Dawson, Center for Biologics Evaluation and Research(HFM–302), Food and Drug Administration, 1401 Rockville Pike, suite 200N, Rockville, MD 20852–1448, 301–827–6129, FAX: 301–827–2843, e-mail: *rhonda.dawson@fda.hhs.gov*.

Registration: Mail, fax, or e-mail your registration information (including name, title, firm name, address, telephone and fax numbers) to the contact person by July 31, 2007. There is no registration fee for the public workshop. Early registration is recommended because seating is limited. Registration on the day of the public workshop will be provided on a space available basis beginning at 7:30 a.m.

If you need special accommodations due to a disability, please contact Rhonda Dawson at least 7 days in advance.

SUPPLEMENTARY INFORMATION: The public workshop will feature presentations by experts from government and industry. The workshop will include presentations by FDA on: (1) Requirements for licensure, and applicable regulations and guidances, for Platelets, Pheresis; Red Blood Cells; and Plasma (intended for transfusion) collected by apheresis instruments; (2) the FDA managed review process; and (3) failure investigations of apheresis products. Device manufacturers will present an overview of their devices and review validation procedures and quality control processes. Representatives from blood establishments will present case studies of licensing applications. FDA will lead a question and answer session with workshop participants.

Comments: All individuals wishing to submit questions to be addressed at the public workshop should submit written or electronic comments by July 31, 2007, to the Division of Dockets Management (HFA–305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852. Submit electronic comments to *http://www.fda.gov/dockets/ecomments*. Comments are to be identified with the docket number found in brackets in the heading of this document. Received comments may be seen in the Division of Dockets Management between 9 a.m. and 4 p.m., Monday through Friday.

Transcripts: Transcripts of the public workshop may be requested in writing from the Freedom of Information Office (HFI–35), Food and Drug Administration, 5600 Fishers Lane, rm.6–30, Rockville, MD 20857, approximately 15 working days after the public workshop at a cost of 10 cents per page. A transcript of the public workshop will be available on the Internet at *http://www.fda.gov/cber/minutes/workshop-min.htm*.

Dated: June 11, 2007.

Jeffrey Shuren,

Assistant Commissioner for Policy.

[FR Doc. E7–11615 Filed 6–15–07; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HOMELAND SECURITY

Federal Emergency Management Agency

Agency Information Collection Activities: Proposed Collection; Comment Request

AGENCY: Federal Emergency Management Agency, DHS.

ACTION: Notice and request for comments.

SUMMARY: The Federal Emergency Management Agency (FEMA), as part of its continuing effort to reduce paperwork and respondent burden, invites the general public and other Federal agencies to take this opportunity to comment on a proposed revised information collection. In accordance with the Paperwork Reduction Act of 1995, this notice seeks comments concerning FEMA's National Emergency Training Center (NETC) to approve and coordinate the use of the NETC facility for extracurricular training activities.

SUPPLEMENTARY INFORMATION: The National Emergency Training Center (NETC) is a FEMA facility that houses the Emergency Management Institute (EMI) and the National Fire Academy (NFA). NETC provides training and educational programs for Federal, State, and local personnel in hazard mitigation, emergency response and preparedness, fire prevention and control, disaster response, and long-term disaster recovery. Special groups sponsored by EMI, NFA or other FEMA organizations may use NETC facilities to conduct activities closely related to and in direct support of their activities. Such groups include other Federal departments and agencies, groups charted by Congress such as the American Red Cross, State and local governments, volunteer groups, and national and international associations representing State and local governments.

Collection of Information

Title: Approval and Coordination of requirements to use NETC for Extracurricular Training Activities.

Type of Information Collection: Revision of an existing collection.

OMB Number: 1660–0029.

Form Numbers: FEMA Form 75–10, Request for Housing Accommodations, and FEMA Form 75–11, Request for Use of NETC Facility.

Abstract: Data will be obtained from special groups that request to use NETC facilities for extracurricular training