anthropometry that are to be used in computer simulation?

(4) What are the essential cab accommodation models to be developed once updated anthropometric and workspace data become available?

The public is invited to attend and will have the opportunity to provide comments. NIOSH will use this information to assess the scientific basis for the current pilot project and the possible large-scale project on U.S. truck driver anthropometric and workspace data.

Status: The consortium meeting will include scientists and representatives from various government agencies, industry and other stakeholders and is open to the public, limited only by the space available. The meeting room accommodates 40 people. Due to limited space, notification of intent to attend the meeting must be made to Jinhua Guan, PhD, not later than April 14, 2006. Dr. Guan can be reached by telephone at (304) 599–4676 or by e-mail at *ezg6@cdc.gov*. Requests to attend the meeting will be accommodated on a first-come basis.

Non-U.S. Citizens: Because of CDC Security Regulations, non-U.S. citizens wishing to attend this meeting must provide the following information in writing to Barbara Phillips (telephone: 304–285–6325; fax: (304) 285–6039; e-mail: *djp2@cdc.gov*) no later than April 14, 2006:

1. Visitor's Full Name:
2. Gender:
3. Date of Birth:
4. Place of Birth (city, province, state, country):
5. Citizenship:
6. Passport Number:
7. Date of Passport Issue:
8. Date of Passport Expiration:
9. Type of Visa:
10. Visitor's Organization:
11. Organization Address:
12. Organization Telephone Number:
13. Visitor's Position/Title within the Organization:

This information will be transmitted to the CDC Security Office for approval. Visitors will be notified as soon as approval has been obtained.

A copy of the research protocol titled ''NIOSH Pilot Study of Truck Driver Anthropometric and Workspace Dimensions'' can be obtained from the CDC Internet at *http://www.cdc.gov/niosh/docs* or a hard copy may be requested from the Docket Officer, Karen Dragon (see contact information below).

ADDRESSES: Comments should be submitted to the NIOSH Docket Office, ATTN: Karen Dragon, Robert A. Taft Laboratories, 4676 Columbia Parkway, M/S C–34, Cincinnati, Ohio 45226, telephone 513/533–8303, fax 513/533–8285.

Comments may also be submitted directly through the Web site (*http://www.cdc.gov/niosh/docs/*) or by e-mail to: *niocindocket@cdc.gov*. E-mail attachments should be formatted in Microsoft Word. Comments should be submitted to NIOSH no later than June 30, 2006, and should reference docket number NIOSH–068 in the subject heading.

Oral comments made at the public meeting must also be submitted to the docket office in writing in order to be considered by the Agency.

All information received in response to this notice will be available for public examination and copying at the NIOSH Docket Office, Room 111, 4676 Columbia Parkway, Cincinnati, Ohio 45226.

Contact Person for Technical Information: Jinhua Guan, PhD, telephone (304) 285–6333, Division of Safety Research, NIOSH, 1095 Willowdale Road, Morgantown, WV 26505.

Dated: April 3, 2006.

James D. Seligman,

Chief Information Officer, Centers for Disease Control and Prevention.

[FR Doc. E6–5168 Filed 4–7–06; 8:45 am]

BILLING CODE 4163–19–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 2005D–0274]

Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Voluntary Hazard Analysis and Critical Control Point Manuals for Operators and Regulators of Retail and Food Service Establishments

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a collection of information entitled ''Voluntary Hazard Analysis and Critical Control Point Manuals for Operators and Regulators of Retail and Food Service Establishments'' has been approved by the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995.

FOR FURTHER INFORMATION CONTACT: Jonna Capezzuto, Office of Management Programs (HFA–250), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–4659.

SUPPLEMENTARY INFORMATION: In the **Federal Register** of December 22, 2005 (70 FR 76056), the agency announced that the proposed information collection had been submitted to OMB for review and clearance under 44 U.S.C. 3507. An agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a currently valid OMB control number. OMB has now approved the information collection and has assigned OMB control number 0910–0578. The approval expires on March 31, 2009. A copy of the supporting statement for this information collection is available on the Internet at *http://www.fda.gov/ohrms/dockets*.

Dated: April 3, 2006.

Jeffrey Shuren,

Assistant Commissioner for Policy.

[FR Doc. E6–5142 Filed 4–7–06; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 2005N–0327]

Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Blood Establishment Registration and Product Listing, Form FDA 2830

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a collection of information entitled ''Blood Establishment Registration and Product Listing, Form FDA 2830'' has been approved by the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995.

FOR FURTHER INFORMATION CONTACT: Jonna Capezzuto, Office of Management Programs (HFA–250), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–4659.

SUPPLEMENTARY INFORMATION: In the **Federal Register** of January 25, 2006 (71 FR 4145), the agency announced that the proposed information collection had been submitted to OMB for review and clearance under 44 U.S.C. 3507. An agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a currently valid OMB control number. OMB has now approved the

information collection and has assigned OMB control number 0910–0052. The approval expires on March 31, 2009. A copy of the supporting statement for this information collection is available on the Internet at *http://www.fda.gov/ohrms/dockets*.

Dated: April 3, 2006.

Jeffrey Shuren,

Assistant Commissioner for Policy.

[FR Doc. E6–5146 Filed 4–7–06; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 2005N–0190]

Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Export Certificates for Food and Drug Administration-Regulated Products

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a collection of information entitled ''Export Certificates for FDA-Regulated Products'' has been approved by the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995.

FOR FURTHER INFORMATION CONTACT: Jonna Capezzuto, Office of Management Programs (HFA–250), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–4659.

SUPPLEMENTARY INFORMATION: In the **Federal Register** of January 25, 2006 (71 FR 4147), the agency announced that the proposed information collection had been submitted to OMB for review and clearance under 44 U.S.C. 3507. An agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a currently valid OMB control number. OMB has now approved the information collection and has assigned OMB control number 0910–0498. The approval expires on March 31, 2009. A copy of the supporting statement for this information collection is available on the Internet at *http://www.fda.gov/ohrms/dockets*.

Dated: April 3, 2006.

Jeffrey Shuren,

Assistant Commissioner for Policy.

[FR Doc. E6–5148 Filed 4–7–06; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 2005N–0389]

Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Reprocessed Single-Use Device Labeling

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a collection of information entitled ''Reprocessed Single-Use Device Labeling'' has been approved by the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995.

FOR FURTHER INFORMATION CONTACT: Karen Nelson, Office of Management Programs (HFA–250), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–1482.

SUPPLEMENTARY INFORMATION: In the **Federal Register** of December 15, 2005 (70 FR 74324), the agency announced that the proposed information collection had been submitted to OMB for review and clearance under 44 U.S.C. 3507. An agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a currently valid OMB control number. OMB has now approved the information collection and has assigned OMB control number 0910–0577. The approval expires on January 31, 2009. A copy of the supporting statement for this information collection is available on the Internet at *http://www.fda.gov/ohrms/dockets*.

Dated: April 3, 2006.

Jeffrey Shuren,

Assistant Commissioner for Policy.

[FR Doc. E6–5150 Filed 4–7–06; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 2005N–0343]

Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Guidance for Requesting an Extension to Use Existing Label Stock After the Trans Fat Labeling Effective Date of January 1, 2006

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a proposed collection of information has been submitted to the Office of Management and Budget (OMB) for review and clearance under the Paperwork Reduction Act of 1995.

DATES: Fax written comments on the collection of information by May 10, 2006.

ADDRESSES: OMB is still experiencing significant delays in the regular mail, including first class and express mail, and messenger deliveries are not being accepted. To ensure that comments on the information collection are received, OMB recommends that written comments be faxed to the Office of Information and Regulatory Affairs, OMB, Attn: Fumie Yokota, Desk Officer for FDA, FAX: 202–395–6974.

FOR FURTHER INFORMATION CONTACT: Jonna Capezzuto, Office of Management Programs (HFA–250), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–4659.

SUPPLEMENTARY INFORMATION: In compliance with 44 U.S.C. 3507, FDA has submitted the following proposed collection of information to OMB for review and clearance.

Guidance for Requesting an Extension to Use Existing Label Stock After the Trans Fat Labeling Effective Date of January 1, 2006—(OMB Control Number 0910–0571)—Extension

FDA issued a final rule (the *trans* fat final rule) on July 11, 2003, (68 FR 41434) to require food labels to bear the gram (g) amount of *trans fat* without a percent Daily Value (% DV) directly under the saturated fat line on the Nutrition Facts panel (*http://www.cfsan.fda.gov/~acrobat/fr03711a.pdf).* The *trans* fat final rule affects almost all manufacturers of packaged, labeled food sold in the United States. FDA believes that most businesses, including small