Dated: December 19, 2013.

John Howard,

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

[FR Doc. 2013–30905 Filed 12–24–13; 8:45 am]

BILLING CODE 4163–19–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[CDC–2013–0021; NIOSH 245–A]

Notice of Request for Comments on Chapters 6 and 8 of the NIOSH document titled: ''Criteria for a Recommended Standard: Occupational Exposure to Diacetyl and 2,3-pentanedione''

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

ACTION: Request for Comments.

SUMMARY: The National Institute for Occupational Safety and Health (NIOSH) of the Centers for Disease Control and Prevention (CDC) is inviting comments on Chapter 6 and a new section of Chapter 8 of the draft document, ''*Criteria for a Recommended Standard: Occupational Exposure to Diacetyl and 2,3-pentanedione.''* To view the notice and related materials, visit *http://www.regulations.gov* and enter CDC–2013–0021 in the search field and click ''Search.'' *Public Comment Period: Comments must be received by* February 10, 2014.

Status: Comments are being sought from individuals including scientists and representatives from various government agencies, industry, labor, and other stakeholders, and also the public.

ADDRESSES: You may submit comments, identified by CDC–2013–0021 and Docket Number NIOSH 245–A by either of the following two methods:

Federal rulemaking portal: http://www.regulations.gov. Follow the instructions for submitting comments.

Mail: NIOSH Docket Office, Robert A. Taft Laboratories, 4676 Columbia Parkway, MS–C34, Cincinnati, Ohio 45226.

Instructions: All information received in response to this notice must include the agency name and docket number [CDC–2013–0021; NIOSH 245–A]. All relevant comments received, including any personal information provided, will be posted without change to *http://www.regulations.gov.* All electronic comments should be formatted as Microsoft Word. Please make reference to CDC–2013–0021 and Docket Number NIOSH 245–A. To access the docket, read background documents or read comments, go to *http://www.regulations.gov.* To access any prior background documents or previous comments received please go to NIOSH Docket 245 (*http://www.cdc.gov/niosh/docket/archive/docket245.html*). All information received in response to this notice will be available for public examination and copying at the NIOSH Docket Office, 4676 Columbia Parkway, Cincinnati, Ohio 45226.

FOR MORE INFORMATION CONTACT: Lauralynn Taylor McKernan, ScD CIH NIOSH, 4676 Columbia Parkway C–14, Cincinnati, OH 45226, telephone (513) 533–8542, Fax (513) 533–8588, email *LMcKernan@cdc.gov.*

Dated: December 19, 2013.

John Howard,

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

[FR Doc. 2013–30900 Filed 12–24–13; 8:45 am]

BILLING CODE 4163–19–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Proposed Information Collection Activity; Comment Request

Proposed Projects: To establish a systematic method of reporting suicides and suicide attempts by refugees.

Title: Refugee Suicide Report Form (RSR).

OMB No.: 0970–NEW.

Description

Pursuant to section 412(b)(4) of the Immigration and Nationality Act, the Administration for Children and Families' Office of Refugee Resettlement (ORR), as the designee for the Secretary of Health and Human Services, is authorized to identify and monitor refugees with certain medical conditions that affect the public health and require treatment.

The intent of this collection activity is to allow ORR to systematically gather information on suicides and suicide attempts among refugee populations resettled in the U.S. Data will be collected on individuals who have made suicide attempts or completed a suicide. The data will be analyzed to identify trends and factors related to suicidal behavior. In addition, the data will be used to plan, implement, and evaluate suicide prevention and intervention activities, in collaboration with local, state, and national government agencies and organizations serving the refugee population.

Respondents: State Governments.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
Refugee Suicide Report Form (RSR)	100 or more	1	0.5	50

Estimated Total Annual Burden Hours:

In compliance with the requirements of Section 506(c)(2)(A) of the Paperwork Reduction Act of 1995, the Administration for Children and Families is soliciting public comment on the specific aspects of the information collection described above. Copies of the proposed collection of information can be obtained and comments may be forwarded by writing to the Administration for Children and Families, Office of Planning, Research and Evaluation, 370 L'Enfant Promenade SW., Washington, DC 20447, Attn: ACF Reports Clearance Officer. Email address: *infocollection@acf.hhs.gov.* All requests should be identified by the title of the information collection.

The Department specifically requests comments on: (a) Whether the proposed collection of information is necessary

for the proper performance of the functions of the agency, including whether the information shall have practical utility; (b) the accuracy of the agency's estimate of the burden of the proposed collection of information; (c) the quality, utility, and clarity of the information to be collected; and (d) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques or other forms of information technology. Consideration will be given to comments and suggestions submitted within 60 days of this publication.

Robert Sargis,

Reports Clearance Officer.

[FR Doc. 2013–30809 Filed 12–24–13; 8:45 am]

BILLING CODE 4184–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2013–N–1620]

Agency Information Collection Activities; Proposed Collection; Comment Request; Information From United States Firms and Processors That Export to the European Community

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA or we) is announcing an opportunity for public comment on the proposed collection of certain information by the Agency. Under the Paperwork Reduction Act of 1995 (the PRA), 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 reporting requirements in implementing the lists of United States (U.S.) firms/processors exporting shell eggs, dairy products, game meat, game meat products, animal casings, gelatin, and collagen to the European Community (the EC).

DATES: Submit either electronic or written comments on the collection of information by February 24, 2014.

ADDRESSES: Submit electronic comments on the collection of information to *http://www.regulations.gov*. Submit written comments on the collection of information to the Division of Dockets Management (HFA–305), Food and Drug Administration, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852. All comments should be identified with the docket number found in brackets in the heading of this document.

FOR FURTHER INFORMATION CONTACT: FDA PRA Staff, Office of Operations, Food and Drug Administration, 1350 Piccard Dr., PI50–400B, Rockville, MD 20850, *PRAStaff@fda.hhs.gov*.

SUPPLEMENTARY INFORMATION: Under 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 set forth in this document.

With respect to the following collection of information, FDA invites comments on these topics: (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.

Information From United States Firms and Processors That Export to the European Community (OMB Control Number 0910–0320)—Extension

The EC is a group of 27 European countries that have agreed to harmonize their commodity requirements to facilitate commerce among member States. EC legislation for intra-EC trade has been extended to trade with non-EC countries, including the United States. For certain food products, including those listed in this document, EC legislation requires assurances from the responsible authority of the country of origin that the processor of the food is in compliance with applicable regulatory requirements. The European Commission, the executive branch of the EC, requires countries trading with any of the EC member countries to provide lists of firms and processors approved to export certain animal-derived commodities to the EC. As stated in the notice published in the **Federal Register** of April 4, 1996 (61 FR 15077), we established a list of U.S. firms and processors that intended to export shell eggs, dairy products, and game meat and game meat products to the EC.

Although our 1996 **Federal Register** notice did not include on the list firms and processors exporting gelatin and raw, bulk collagen intended for human consumption, EC directives require that shipments of gelatin and raw, bulk collagen products be accompanied by certification stating that the product, derived from ruminant bones, bovine hides, and pigskins, has been produced in compliance with EC Council Directive 2003/863/EC. The directive contains the requirements for sourcing, manufacture, transport, and storage of raw materials and manufacture of finished products and requires lists identifying non-EC firms and processors that meet EC requirements and have the appropriate animal and public health certificates. Therefore, we revised this information collection in order to facilitate exports of gelatin and raw, bulk collagen originating from the United States into the EC. We announced OMB approval of the revised information collection in the **Federal Register** of May 10, 2011 (76 FR 27061).

We request the following information from each firm or processor seeking to be included on the lists for shell eggs, dairy products, game meat, game meat products, and animal casings:

- Business name and address;
- Name and telephone number of person designated as business contact;
- Lists of products presently being shipped to the EC and those intended to be shipped in the next 6 months;
- Name and address of manufacturing plants for each product; and
- Names and affiliations of any Federal, State, or local governmental Agencies that inspect the plant, government-assigned plant identifier such as plant number, and last date of inspection.

We use the information to maintain lists of firms and processors that have demonstrated current compliance with