DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Agency Forms Undergoing Paperwork Reduction Act Review

The Centers for Disease Control and Prevention (CDC) publishes a list of information collection requests under review by the Office of Management and Budget (OMB) in compliance with the requirement of Section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995. To request a copy of these requests, call the CDC Reports Clearance Officer at (404) 639–5960 or send an e-mail to omb@cdc.gov. Send written comments to CDC Desk Officer, Office of Management and Budget, Washington, DC or by fax to (202) 395–5680. Written comments should be received within 30 days of this notice.

ANNUALIZED BURDEN TABLE

<table>
<thead>
<tr>
<th>Questionnaire (respondent)</th>
<th>Number of respondents</th>
<th>Number of responses per respondent</th>
<th>Average burden per respondent in hours</th>
</tr>
</thead>
<tbody>
<tr>
<td>Screener Questionnaire</td>
<td>10,000</td>
<td>1</td>
<td>5/60</td>
</tr>
<tr>
<td>Family Core (adult family member)</td>
<td>58,000</td>
<td>1</td>
<td>23/60</td>
</tr>
<tr>
<td>Adult Core (sample adult)</td>
<td>44,250</td>
<td>1</td>
<td>14/60</td>
</tr>
<tr>
<td>Child Core (adult family member)</td>
<td>17,550</td>
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<td>9/60</td>
</tr>
<tr>
<td>Child Record Check (medical provider)</td>
<td>2,120</td>
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<td>5/60</td>
</tr>
<tr>
<td>Child Immunization Provider (adult family member)</td>
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<td>1</td>
<td>4/60</td>
</tr>
<tr>
<td>Supplements (adult family Member)</td>
<td>58,000</td>
<td>1</td>
<td>18/60</td>
</tr>
<tr>
<td>Reinterview Survey</td>
<td>4,000</td>
<td>1</td>
<td>5/60</td>
</tr>
</tbody>
</table>

SUMMARY: The National Institute for Occupational Safety and Health (NIOSH) of the Centers for Disease Control and Prevention (CDC) is conducting a review of the processes of the National Occupational Research Agenda (NORA). In 2006, NORA entered its second decade with an industry sector-based structure. In 2011, as NORA reaches the halfway point of its second decade, NIOSH is conducting a review of NORA processes to learn how adjustments can be made to maximize outcomes through the remainder of the second decade (2012–2016). The goal is to look at NORA processes across the ten NORA industry...
DEPARTMENT OF HEALTH AND HUMAN SERVICES
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[Docket Number NIOSH–245]

Notice of Public Meeting on 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: Notice of public meeting.

SUMMARY: The National Institute for Occupational Safety and Health (NIOSH) of the Centers for Disease Control and Prevention (CDC) will hold a public meeting to discuss and obtain comments on the draft document, “Criteria for a Recommended Standard: Occupational Exposure to Diacetyl and 2,3-pentanedione.” A copy of the draft document will be posted on the Internet at http://www.cdc.gov/niosh/docket/review/docket245/default.html for Docket number NIOSH–245 on August 12, 2011. This notice serves as advance notice of the meeting and public comment period.

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Date and Time
Address
Status
Speaker Registration
Agenda
Supplementary Information
I. Matters to Be Discussed
II. Transcripts
III. Public Comment Period
Contact Person for More Information

DATES AND TIME: August 26, 2011, 8 a.m.–4 p.m., Eastern Time. Please note that public comments may end before the time indicated, following the last call for comments. Members of the public who wish to provide comments should plan to attend the meeting at the start time listed.


Status: The meeting is open to the public limited only by the space available. The meeting room accommodates 150 people. To pre-register for the meeting, interested parties should contact the NIOSH Docket Office at nioshdocket@cdc.gov or by fax at (513) 533–8285. Due to limited space, notification of intent to attend the meeting must be made to the NIOSH Docket Office no later than August 19, 2011. Priority for attendance will be given to those providing oral comments. Other requests to attend the meeting will then be accommodated on a first-come basis.

Speaker Registration: Persons wanting to provide oral comments on the draft document should contact the NIOSH Docket Office at nioshdocket@cdc.gov or by fax at (513) 533–8285. Presenters will be permitted approximately 10 minutes, and will be informed if additional time becomes available. All requests to present should contain the name, address, telephone number, and relevant business affiliations of the presenter, and the topic of the presentation. Oral comments made at the public meeting must also be submitted to the NIOSH Docket Office in writing in order to be considered by the Agency.

Agenda: The meeting will begin with an introduction and presentation by Federal officials, followed by presentations from attendees who wish to speak. Each speaker will be limited to ten minutes. If all pre-registered presentations are complete before the end of the time, there will be an open session to receive comments from anyone who has not signed up on the speaker registration list who may wish to speak. Open session comments will also be limited to 10 minutes per speaker. After the last speaker or at 4 p.m., whichever occurs first, the meeting will be adjourned.

SUPPLEMENTARY INFORMATION:

I. Matters To Be Discussed

At the public meeting, special emphasis will be placed on the following topics:

1. Hazard identification, risk estimation, and discussion of health effects for diacetyl and 2,3-pentanedione;
2. Basis of the Recommended Exposure Limit for diacetyl and 2,3-pentanedione;
3. Workplaces and occupations where exposure to diacetyl and 2,3-pentanedione occur;
4. Current exposure measurement methods;
5. Current strategies for controlling occupational exposure to diacetyl and 2,3-pentanedione: e.g., engineering controls, work practices, medical surveillance, and personal protective equipment;
6. Oral comments provided to NIOSH on the draft criteria document.

II. Transcripts

Transcripts will be prepared and posted to NIOSH Docket number 245 approximately 30 days after the meeting. If a person making a comment gives his or her name, no attempt will be made to redact that name. NIOSH will take reasonable steps to ensure that individuals making public comments are aware of the fact that their comments (including their name, if provided) will appear in a transcript of the meeting posted on a public Web site. Such reasonable steps include: (a) A statement read at the start of the meeting stating that transcripts will be posted and names of speakers will not be redacted; and (b) A printed copy of the statement mentioned in (a) above will be displayed on the table where individuals sign up to make public comments. If individuals in making a statement reveal personal information (e.g., medical information) about