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Director, Office of Scientific Integrity (OSI), Office of the Associate Director for Science (OADS), Office of the Director, Centers for Disease Control and Prevention.

The Centers for Medicare & Medicaid Services

Agency Information Collection Activities: Proposed Collection; Comment Request

AGENCY: Centers for Medicare & Medicaid Services, HHS.

In compliance with the requirement of section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, the Centers for Medicare & Medicaid Services (CMS) is publishing the following summary of proposed collections for public comment. Interested persons are invited to send comments regarding this burden estimate or any other aspect of this collection of information, including any of the following subjects: (1) The necessity and utility of the proposed information collection for the proper performance of the agency’s functions; (2) the accuracy of the estimated burden; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) the use of automated collection techniques or other forms of information technology to minimize the information collection burden.

1. Type of Information Collection Request: Extension. Title of Information Collection: Detailed Notice of Discharge (DND). Use: When a Medicare beneficiary requests a Quality Improvement Organization review of his/her inpatient hospital discharge, hospitals and Medicare plans have used the DND to provide the beneficiary with a detailed explanation regarding the reason for discharge. Form Number: CMS–10066 (OCN 0938–1019). Frequency: Yearly. Affected Public: Private Sector (business or other for-profit and not-for-profit institutions). Number of Respondents: 6,169. Total Annual Responses: 12,852. Total Annual Hours: 12,852. (For policy questions regarding this collection contact Evelyn Blaemire at 410–786–1803. For all other issues call 410–786–1326.)

2. Type of Information Collection Request: Extension. Title of Information Collection: Important Message from Medicare (IM). Use: Hospitals have used the IM to inform original Medicare, Medicare Advantage, and other Medicare plan beneficiaries who are hospital inpatients about their hospital rights and discharge rights. In particular, the IM provides information about when a beneficiary will and will not be liable for charges for a continued stay in a hospital and offers a detailed description of the Quality Improvement Organization review process. Form Number: CMS–R–193 (OCN 0938–0692). Frequency: Yearly. Affected Public: Private Sector (business or other for-profit and not-for-profit institutions). Number of Respondents: 6,169. Total Annual Responses: 19,840,000. Total Annual Hours: 2,976,000. (For policy questions regarding this collection contact Evelyn Blaemire at 410–786–1803. For all other issues call 410–786–1326.)

To obtain copies of the supporting statement and any related forms for the proposed paperwork collections referenced above, access CMS’ Web Site address at http://www.cms.hhs.gov/PaperworkReductionActof1995, or Email your request, including your address, phone number, OMB number, and CMS document identifier, to Paperwork@cms.hhs.gov, or call the Reports Clearance Office on (410) 786–1326.

In commenting on the proposed information collections please reference the document identifier or OMB control number. To be assured consideration, comments and recommendations must be submitted in one of the following ways by May 6, 2013:

1. Electronically. You may submit your comments electronically to http://www.regulations.gov. Follow the instructions for “Comment or Submission” or “More Search Options” to find the information collection document(s) accepting comments.

2. By regular mail. You may mail written comments to the following address: CMS, Office of Strategic Operations and Regulatory Affairs, Division of Regulations Development, Attention: Document Identifier/OMB Control Number ___, Room C4–26–05, 7500 Security Boulevard, Baltimore, Maryland 21244–1850.
DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Proposed Information Collection Activity; Comment Request

Proposed Projects

Title: ADP & Services Conditions for FFP for ACF.
OMB No.: 0992–0005.
Description: The Advance Planning Document (ADP) process, established in the rules at 45 CFR Part 95, Subpart F, is the procedure by which States request and obtain approval for Federal financial participation in their cost of acquiring Automatic Data Processing (ADP) equipment and services. State agencies that submit ADP requests provide the Department of Health and Human Services (HHS) with the following information necessary to determine the States’ needs to acquire the requested ADP equipment and/or services:
(1) A statement of need;
(2) A requirements analysis and feasibility study;
(3) A proposed activity schedule; and,
(4) A proposed budget.
HHS’ determination of a State Agency’s need to acquire requested ADP equipment or services is authorized at sections 402(a)(5), 452(a)(1), 1902(a)(4) and 1102 of the Social Security Act.
Respondents: States.

ANNUAL BURDEN ESTIMATES

<table>
<thead>
<tr>
<th>Instrument</th>
<th>Number of respondents</th>
<th>Number of responses per respondent</th>
<th>Average burden hours per response</th>
<th>Total burden hours</th>
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<tbody>
<tr>
<td>RFP and Contract</td>
<td>4</td>
<td>1.5</td>
<td>4</td>
<td>324</td>
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<tr>
<td>Emergency Funding Request</td>
<td>5</td>
<td>1</td>
<td>2</td>
<td>1</td>
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<tr>
<td>Biennial Reports</td>
<td>54</td>
<td>20</td>
<td>1.2</td>
<td>4,896</td>
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<td>Advance Planning Document</td>
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<td>1</td>
<td>120</td>
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<tr>
<td>Operational Advance Planning Document</td>
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<td>600</td>
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<td><strong>Estimated Total Annual Burden Hours</strong></td>
<td></td>
<td></td>
<td></td>
<td><strong>5,902</strong></td>
</tr>
</tbody>
</table>

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.

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

Cheng Yi Liang: Debarment Order

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is issuing an order under the Federal Food, Drug, and Cosmetic Act (the FD&C Act) permanently debarring Cheng Yi Liang, from providing services in any capacity to a person that has an approved or pending drug product application. We base this order on a finding that Mr. Liang was convicted of a felony under Federal law for conduct relating to the development or approval, including the process for the development or approval, of a drug product. Mr. Liang was given notice of the proposed permanent debarment and an opportunity to request a hearing within the timeframe prescribed by regulation. Mr. Liang failed to respond. Mr. Liang’s failure to respond constitutes a waiver of his right to a hearing concerning this action.

DATES: This order is effective March 6, 2013.

ADDRESSES: Submit applications for special termination of debarment to the Division of Dockets Management (HFA–305), Food and Drug Administration, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852.

FOR FURTHER INFORMATION CONTACT: Kenny Shade, Office of Regulatory Affairs, Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–796–4640.

SUPPLEMENTARY INFORMATION:

I. Background

Section 306(a)(2)(A) of the FD&C Act (21 U.S.C. 335a(a)(2)(A)) requires debarment of an individual if FDA finds that the individual has been convicted of a felony under Federal law for