TABLE 1.—ESTIMATED ANNUAL REPORTING BURDEN

<table>
<thead>
<tr>
<th>21 CFR Section</th>
<th>No. of Respondents</th>
<th>Annual Frequency per Response</th>
<th>Total Annual Responses</th>
<th>Hours per Response</th>
<th>Total Hours</th>
</tr>
</thead>
<tbody>
<tr>
<td>600.80(c)(1) and (e)</td>
<td>95</td>
<td>146.72</td>
<td>13,938</td>
<td>1</td>
<td>13,938</td>
</tr>
<tr>
<td>600.80(c)(2)</td>
<td>95</td>
<td>106.34</td>
<td>10,102</td>
<td>28</td>
<td>282,856</td>
</tr>
<tr>
<td>600.81</td>
<td>95</td>
<td>3.57</td>
<td>339</td>
<td>1</td>
<td>339</td>
</tr>
<tr>
<td>600.90</td>
<td>12</td>
<td>1</td>
<td>12</td>
<td>1</td>
<td>12</td>
</tr>
<tr>
<td><strong>Total</strong></td>
<td></td>
<td></td>
<td><strong>297,145</strong></td>
<td></td>
<td><strong>297,145</strong></td>
</tr>
</tbody>
</table>

1 There are no capital costs or operating and maintenance costs associated with this collection of information.

Under table 2 of this document, the number of respondents is based on the number of manufacturers subject to those regulations. Based on information obtained from CBER’s database system, there were approximately 329 licensed manufacturers of biological products. However, the number of recordkeepers listed for § 600.12(a) through (e), excluding paragraph (b)(2), is estimated to be 111. This number excludes manufacturers of blood and blood components because their burden hours for recordkeeping have been reported under § 606.160 in OMB control number 0910–0116. The total annual records is based on the annual average of lots released (6,747), number of recalls made (1,646) and total number of AER reports received (24,040) in the years 2000 and 2001. The hours per record are based on FDA’s experience. FDA estimates the burden of this recordkeeping as follows:

TABLE 2.—ESTIMATED ANNUAL RECORDKEEPING BURDEN

<table>
<thead>
<tr>
<th>21 CFR Section</th>
<th>No. of Respondents</th>
<th>Annual Frequency per Response</th>
<th>Total Annual Responses</th>
<th>Hours per Response</th>
<th>Total Hours</th>
</tr>
</thead>
<tbody>
<tr>
<td>600.12</td>
<td>111</td>
<td>60.78</td>
<td>6,747</td>
<td>32</td>
<td>215,904</td>
</tr>
<tr>
<td>600.12(b)(2)</td>
<td>329</td>
<td>5.00</td>
<td>1,646</td>
<td>24</td>
<td>39,504</td>
</tr>
<tr>
<td>600.80(i)</td>
<td>95</td>
<td>253.05</td>
<td>24,040</td>
<td>1</td>
<td>24,040</td>
</tr>
<tr>
<td><strong>Total</strong></td>
<td></td>
<td></td>
<td><strong>279,448</strong></td>
<td></td>
<td><strong>279,448</strong></td>
</tr>
</tbody>
</table>

1 There are no capital costs or operating and maintenance costs associated with this collection of information.

Margaret M. Dotzel,
Assistant Commissioner for Policy.
[FR Doc. 03–1406 Filed 1–22–03; 8:45 am]
BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES
Food and Drug Administration
[Docket No. 00N–1529]
Elaine Yee-Ling Lai; Debarment Order; Correction
AGENCY: Food and Drug Administration, HHS.
ACTION: Notice; correction.
SUMMARY: The Food and Drug Administration (FDA) is correcting a notice that appeared in the Federal Register of November 13, 2002 (67 FR 68877). The document announced the issuance of an order under the Federal Food, Drug, and Cosmetic Act debarring Ms. Elaine Yee-Ling Lai for 5 years from providing services in any capacity to a person that has an approved or pending drug product application. The document was published with an inadvertent error. This document corrects that error.

FOR FURTHER INFORMATION CONTACT:
Joyce Strong, Office of Policy (HF–27), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–7010.

SUPPLEMENTARY INFORMATION: In FR Doc. 02–28715, appearing on page 68877 in the Federal Register of Wednesday, November 13, 2002, the following correction is made:

1. On page 68877, in the third column, under section II, in the fourth line “(21 CFR 5.99)” is corrected to read “(21 CFR 5.34)”.

Margaret M. Dotzel,
Assistant Commissioner for Policy.
[FR Doc. 03–1404 Filed 1–22–03; 8:45 am]
BILLING CODE 4160–01–S

DEPARTMENT OF HOUSING AND URBAN DEVELOPMENT
[FR Doc. 03–1404–N–01]
Notice of Proposed Information Collection: Comment Request Annual Progress Report (APR) for Competitive Homeless Assistance Programs
AGENCY: Office of the Assistant Secretary for Community Planning and Development, HUD.
ACTION: Notice.
SUMMARY: The proposed information collection requirement described below will be submitted to the Office of Management and Budget (OMB) for review, as required by the Paperwork Reduction Act. The Department is soliciting public comments on the subject proposal.
DATES: Comments Due Date: March 24, 2003.
ADDRESSES: Interested persons are invited to submit comments regarding this proposal. Comments should refer to the proposal by name and/or OMB...