TABLE 1.—ESTIMATED ANNUAL REPORTING BURDEN[1]

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

[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[1]

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

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

Dated: January 14, 2003.

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)''.

Dated: January 14, 2003.

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

[Docket No. FR–4814–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