

Boulevard, Baltimore, Maryland 21244-1850.

Dated: April 28, 1998.

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**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**Health Care Financing Administration**

[Document Identifier: HCFA-116, HCFA-R-148, and HCFA-R-231]

**Agency Information Collection Activities: Proposed Collection; Comment Request**

**AGENCY:** Health Care Financing Administration, HHS.

In compliance with the requirement of section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, the Health Care Financing Administration (HCFA), Department of Health and Human Services, 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:* Revision of a currently approved collection; *Title of Information Collection:* Clinical Laboratory Improvement Amendments (CLIA) Application Form and Supporting Regulations in 42 CFR 493.1-2001; *Form No.:* HCFA-116 (OMB# 0938-0581); *Use:* These certification requirements have been established for any entity that performs testing on human beings for diagnostic or treatment purposes. If a laboratory conducts relatively simple tests that are categorized as waived or provider performed microscopy test procedures (PPMP), it must obtain a certificate of waiver or certificate of PPMP. If the laboratory conducts any tests outside of these two categories, it must apply for

a certificate of compliance or certificate of accreditation and initially obtain a registration certificate. These certificates ensure that laboratories are in compliance with CLIA.; *Frequency:* Biennially; *Affected Public:* Business or other for profit, not for profit institutions, Federal Government, and State, local or tribal government; *Number of Respondents:* 16,000; *Total Annual Responses:* 16,000; *Total Annual Hours:* 20,000.

2. *Type of Information Collection Request:* Extension of a currently approved collection; *Title of Information Collection:* Limitation on Provider-Related Donations and Health Care-Related Taxes; Limitations on Payments to Disproportionate Share Hospitals; Medicaid and Supporting Regulations in 42 CFR 433.68, 433.74, 447.74 and 447.272; *Form No.:* HCFA-R-148 (OMB# 0938-0618); *Use:* These information collection requirements specify limitations on the amount of Federal financial participation available for medical assistance expenditures in a fiscal year. States receive donated funds from providers and revenues are generated by health care related taxes. These donations and revenues are used to fund medical assistance programs.; *Frequency:* Quarterly; *Affected Public:* State, Local, or Tribal Government; *Number of Respondents:* 51; *Total Annual Responses:* 51; *Total Annual Hours:* 3,892.

3. *Type of Information Request:* Revision of a currently approved collection; *Title of Information Collection:* Medicare+Choice (M+C) Providers Sponsored Organization (PSO) Waiver Request Form and Supporting Regulations in 42 CFR 422.374; *Form Number:* HCFA-R-231; *Use:* The PSO waiver request form is for use by PSO's that do not have a State risk-bearing entity license and that wish to enter into a M+C contract with HCFA to provide prepaid health care services to eligible Medicare beneficiaries. HCFA will use the information requested on this form to determine whether the applicant is eligible for a waiver of the state licensure requirement for M+C organizations as allowed under section 1855(a)(2) of the Social Security Act.; *Frequency:* One-time.; *Affected Public:* Business or other for-profit, not-for-profit institutions, and Federal Government.; *Annual Number of Respondents:* 30.; *Total Annual Responses:* 30.; *Total Annual Hours Requested:* 300.

To obtain copies of the supporting statement and any related forms for the proposed paperwork collections referenced above, access HCFA's Web Site address at <http://www.hcfa.gov/>

[regs/prdact95.htm](http://www.hcfa.gov/regs/prdact95.htm), or E-mail your request, including your address, phone number, OMB number, and HCFA document identifier, to [Paperwork@hcfa.gov](mailto:Paperwork@hcfa.gov), or call the Reports Clearance Office on (410) 786-1326. Written comments and recommendations for the proposed information collections must be mailed within 60 days of this notice directly to the HCFA Paperwork Clearance Officer designated at the following address: HCFA, Office of Information Services, Information Technology Investment Management Group, Division of HCFA Enterprise Standards, Attention: Louis Blank, Room C2-26-17, 7500 Security Boulevard, Baltimore, Maryland 21244-1850.

Dated: April 24, 1998.

**John P. Burke III,**

*HCFA Reports Clearance Officer, HCFA Office of Information Services, Information Technology Investment Management Group, Division of HCFA Enterprise Standards.*

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**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**Health Care Financing Administration**

[Document Identifier: HCFA-R-235]

**Agency Information Collection Activities: Proposed Collection; Comment Request**

In compliance with the requirement of section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, the Health Care Financing Administration (HCFA), Department of Health and Human Services, is publishing the following summary of proposed collections for public comment. Interested persons are invited to send comments regarding the 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.

*Type of Information Collection Request:* New Collection; *Title of Information Collection:* Data Use Agreement Information Collection Requirements, model agreement, and Supporting regulations; *Form No.:*