

the advertising and sale of products to the public. Attachment A is entitled "Legal Notice" and is a summary of the injunction provisions of the proposed order.

Part XIV of the proposed order requires that the Commission be notified of any change in the corporation that might affect compliance obligations under the order. Part XV of the proposed order requires that for a period of three (3) years, the individual respondent notify the Commission of the discontinuance of his current business or employment or of his affiliation with any new business or employment involving the sale of consumer products and/or services.

Part XVI of the proposed order requires the respondents to file a compliance report with the Commission.

Part XVII of the proposed order states that, absent certain circumstance, the order will terminate twenty (20) years from the date it is issued.

The purpose of this analysis is to facilitate public comment on the proposed consent order. It is not intended to constitute an official interpretation of the agreement and proposed order or to modify their terms in any way.

By direction of the Commission.

**Donald S. Clark,**  
*Secretary.*

[FR Doc. 02-4375 Filed 2-22-02; 8:45 am]  
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## GENERAL SERVICES ADMINISTRATION

### Interagency Committee for Medical Records (ICMR); Automation of Medical Standard Form 519A

**AGENCY:** Office of Communications,  
GSA.

**ACTION:** Guideline on Automating  
Medical Standard Forms.

Background: The Interagency Committee on Medical Records (ICMR) is aware of numerous activities using computer-generated medical forms, many of which are not mirror-like images of the genuine paper Standard/Optional Form. With GSA's approval the ICMR eliminated the requirement that every electronic version of a medical Standard/Optional form be reviewed and granted an exception. The committee proposed to set required fields standards and that activities developing computer-generated versions adhere to the required fields but not necessarily to the image. The ICMR

plans to review medical Standard/Optional forms which are commonly used and/or commonly computer-generated. We will identify those fields which are required, those (if any) which are optional, and the required format (if necessary). Activities may not add or delete data elements that would change the meaning of the form. This would require written approval from the ICMR. Using the process by which overprints are approved for paper Standard/Optional forms, activities may add other data entry elements to those required by the committee. With this decision, activities at the local or headquarters level should be able to develop electronic versions which meet the committee's requirements. This guideline controls the "image" or required fields but not the actual data entered into the field.

**SUMMARY:** With GSA's approval, the Interagency Committee of Medical Records (ICMR) eliminated the requirement that every electronic version of a medical Standard/Optional form be reviewed and granted an exception. The following fields must appear on the electronic version of the following form:

#### ELECTRONIC ELEMENTS FOR SF 519A

| Item                                                                                                                                                                                                                                                                                                                                                                                      | Placement <sup>1</sup>       |
|-------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|------------------------------|
| Radiologic consultation request/report. Standard Form 519A (Rev. 8/1983)(Form ID).                                                                                                                                                                                                                                                                                                        | Top of form.                 |
| 1-Medical Record .....                                                                                                                                                                                                                                                                                                                                                                    | Bottom right corner of form. |
| 2-Physician .....                                                                                                                                                                                                                                                                                                                                                                         | Bottom left corner of form.  |
| 3-Radiology .....                                                                                                                                                                                                                                                                                                                                                                         | Bottom left corner of form.  |
| Data Entry Fields:<br>Patient information (Text)<br>Last name<br>First name<br>Middle name<br>Medical facility<br>Age<br>Sex<br>SSN (Sponsor)<br>Ward/clinic<br>Register No.<br>Examination requested (Use SF 519B for multiple exams)<br>Requested by<br>Telephone number<br>Location of medical records<br>Film number<br>Date requested<br>Pregnant—Yes (Checkbox)<br>Pregnant—No (No) | Above below listed items.    |

#### ELECTRONIC ELEMENTS FOR SF 519A—Continued

| Item                                                                                                                                                                                                                                                     | Placement <sup>1</sup> |
|----------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|------------------------|
| Specific reason(s) for Request (Complaints and findings)<br>Date of examination (Month, day, year)<br>Date of report (Month, day, year)<br>Date of transcription (Month, day, year)<br>Radiologic report<br>Signature<br>Location of radiologic facility |                        |

<sup>1</sup> If no specific placement, data element may be in any order.

**FOR FURTHER INFORMATION CONTACT:** CDR Katherine Ciacco Palatianos, Indian Health Service, Department of Health and Human Services, 5600 Fishers Lane, Room 6A-55, Rockville, MD 20857 or E-Mail at [kciacco@hgs.ihs.gov](mailto:kciacco@hgs.ihs.gov).

**DATES:** Effective February 25, 2002.

Dated: February 12, 2002.

**CDR Katherine Ciacco Palatianos,**  
*Chairperson, Interagency Committee on Medical Records.*

[FR Doc. 02-4452 Filed 2-22-02; 8:45 am]

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## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Centers for Disease Control and Prevention

[60 Day-02-28]

### Proposed Data Collections Submitted for Public Comment and Recommendations

In compliance with the requirement of section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995 for opportunity for public comment on proposed data collection projects, the Centers for Disease Control and Prevention (CDC) will publish periodic summaries of proposed projects. To request more information on the proposed projects or to obtain a copy of the data collection plans and instruments, call the CDC Reports Clearance Officer on (404) 639-7090.

Comments are invited 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)

ways to enhance 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. Send comments to Anne O'Connor, CDC Assistant Reports Clearance Officer, 1600 Clifton Road, MS-D24, Atlanta, GA 30333. Written comments should be received within 60 days of this notice.

*Proposed Project:* National Public Health Performance Standards Program Local Public Health Governance Performance Assessment Instrument—New—Public Health Practice Program Office (PHPPO), Centers for Disease Control and Prevention (CDC).

Since 1998, the CDC National Public Health Performance Standards Program

has convened workgroups with the National Association of County and City Health Officials (NACCHO), the Association of State and Territorial Health Officials (ASTHO), the National Association of Local Boards of Health (NALBOH), the American Public Health Association (APHA), and the Public Health Foundation (PHF) to develop performance standards for public health systems based on the ten Essential Services of Public Health. In the Spring of 2001, CDC conducted field tests with the local public health governance instruments in the state of Massachusetts.

CDC is now proposing to implement a voluntary data collection to assess the capacity of local boards of health to deliver the Essential Public Health Services. This data collection will

provide a framework for local boards of health to evaluate their effectiveness. Electronic data submission will be the method of choice. If computer technology in local jurisdictions does not support electronic submission, hard copy survey instruments will be available. Local jurisdictions using hard copy survey instruments will receive assistance from State or local level field coordinators for web-based data entry.

Local boards of health will respond to the survey. An estimated 33% of approximately 3,200 United States local boards are expected to participate in the National Performance Standards Program per year.

There are no costs to respondents. The burden hours are estimated to be 30,198.

| Respondents                         | Number of respondents | Number of responses/respondent | Average burden/response (in hrs.) | Total burden (in hrs.) |
|-------------------------------------|-----------------------|--------------------------------|-----------------------------------|------------------------|
| Local Boards of Health Year 1 ..... | 1,066                 | 1                              | 10                                | 10,660                 |
| Local Boards of Health Year 2 ..... | 1,066                 | 1                              | 10                                | 10,660                 |
| Local Boards of Health Year 3 ..... | 1,066                 | 1                              | 10                                | 10,660                 |
| Total .....                         |                       |                                |                                   | 30,198                 |

Dated: February 13, 2002.

**John Moore,**

*Acting Associate Director for Policy, Planning and Evaluation, Centers for Disease Control and Prevention.*

[FR Doc. 02-4371 Filed 2-22-02; 8:45 am]

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## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Centers for Medicare and Medicaid Services

[Document Identifier: CMS-10036]

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

**AGENCY:** Centers for Medicare and Medicaid Services.

In compliance with the requirement of section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, the Centers for Medicare and Medicaid Services (CMS) (formerly known as 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.

#### *Type of Information Collection*

*Request:* Revision of a currently approved collection;

#### *Title of Information Collection:*

Inpatient Rehabilitation Assessment Instrument and Data Set for PPS for Inpatient Rehabilitation Facilities and Supporting Regulations in 42 CFR, Parts 412 and 413;

*Form No.:* CMS-10036 (OMB# 0938-0842);

*Use:* This is a request to use the IRF-PAI and its supporting manual for the implementation phase of the inpatient rehabilitation PPS. There have been no revisions or modifications to the instrument; however, this submission includes the current manual/instructions which has been revised. Use of this instrument will enable CMS to implement a classification system and payment system for the Legislatively mandated inpatient rehabilitation hospital and exempt units Prospective Payment System (PPS);

*Frequency:* On occasion;

*Affected Public:* Business or other for-profit, and Not-for-profit institutions;

*Number of Respondents:* 359,000;

*Total Annual Responses:* 359,000;

*Total Annual Hours:* 269,250.

To obtain copies of the supporting statement and any related forms for the proposed paperwork collections referenced above, access CMS's Web site address at <http://www.hcfa.gov/regs/prdact95.htm>, or e-mail your request, including your address, phone number, OMB number, and CMS 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 CMS Paperwork Clearance Officer designated at the following address: CMS, Office of Information Services, Security and Standards Group, Division of CMS Enterprise Standards, Attention: Dawn Willingham, CMS-10036, Room N2-14-26, 7500 Security Boulevard, Baltimore, Maryland 21244-1850.

Dated: February 14, 2002.

**John P. Burke, III,**

*Reports Clearance Officer, Security and Standards Group, Division of CMS Enterprise Standards.*

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