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]

BILLING CODE 6750–01–P

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.

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:

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.

ELECTRONIC ELEMENTS FOR SF 519A

Item	Placement [1]
Radiologic consultation request/report.	Top of form.
Standard Form 519A (Rev. 8/1983)(Form ID).	Bottom right corner of form.
1-Medical Record	Bottom left corner of form.
2-Physician	Bottom left corner of form.
3-Radiology	Bottom left corner of form.
Data Entry Fields:	
Patient information (Text)	Above below listed items.
Last name	
First name	
Middle name	
Medical facility	
Age	
Sex	
SSN (Sponsor)	
Ward/clinic	
Register No.	
Examination requested (Use SF 519B for multiple exams)	
Requested by	
Telephone number	
Location of medical records	
Film number	
Date requested	
Pregnant—Yes (Checkbox)	
Pregnant—No (No)	
Specific reason(s) for Request (Complaints and findings)	
Date of examination (Month, day, year)	
Date of report (Month, day, year)	
Date of transcription (Month, day, year)	
Radiologic report	
Signature	
Location of radiologic facility	

[1] 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@hge.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)