from engaging in similar acts and practices in the future.

Part I of the order prohibits claims that LASIK surgery services or any other refractive surgery services: (1) Eliminate the need for glasses and contacts for life; (2) pose significantly less risk to patients' eye health than wearing glasses or contacts; or (3) eliminate the risk of glare and haloing, unless the claims are substantiated by competent and reliable scientific evidence. "Refractive surgery services" are defined as any surgical procedure designed to improve the focusing power of the eye by permanently changing the shape of the cornea.

Part II of the order requires that future claims about the benefits, performance, efficacy, or safety of any refractive surgery service be substantiated by competent and reliable scientific evidence.

Part III of the order permits device claims approved by the FDA under any new medical device application.

Parts IV, V, VI, and VII of the order require LCA to keep copies of relevant advertisements and materials substantiating claims made in the advertisements, to provide copies of the order to certain of its personnel, to notify the Commission of changes in corporate structure, and to file compliance reports with the Commission. Part VIII provides that the

a. Head, Face, Neck and Scalp-Normal (Checkbox)

order will terminate after twenty (20) years under certain circumstances.

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

By direction of the Commission.

Donald S. Clark,

Secretary.

[FR Doc. 03–7930 Filed 4–1–03; 8:45 am]

BILLING CODE 6750-01-P

GENERAL SERVICES ADMINISTRATION

Interagency Committee for Medical Records (ICMR); Automation of Medical Standard Form 88

AGENCY: Office of Communications,

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 to ICMR eliminated the requirement that every electronic version of a medical Standard/Optional form be reviewed

and granted an exception. The committee proposes 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 computergenerated. 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 any exception. The following fields must appear on the electronic version of the following form:

ELECTRONIC ELEMENTS FOR SF 88

Item	Placement*
Report of Medical Examination	Top of form. Bottom right corner of form.
Data Entry Fields:	
1. Date of Exam	
2. Last Name	
2. First Name	
2. Middle Name	
3. Identification Number	
4. Grade of Position	
4. Component of Position	
5. Home Address (Number, street or RDFD, city or town, state and ZIP code)	
6. Emergency Contact (Name)	
Emergency Contact (address) Date of Birth	
8. Age 9. Sex—Female (Checkbox)	
9. Sex—Male (Checkbox)	
10. Relationship of Contact	
11. Place of Birth	
12. Agency	
13. Organization Unit	
14a. Total Years Government Service—Military	
14b. Total Years Government Service—Civilian	
15. Name of Examining Facility or Examiner	
15. Address of Examining Facility or Examiner	
16. Rating or Specialty of Examiner	
17. Purpose of Examination	
18. Clinical Evaluation—Check each item in appropriate columns; enter "NE" if not evaluated	Above below listed items

ELECTRONIC ELEMENTS FOR SF 88—Continued

ELECTRONIC ELEMENTS FOR SF 88—Continued	
Item	Placement*
a. Head, Face, Neck and Scalp—Abnormal (Checkbox) b. Ears-General (Internal Canals) (auditory acuity under item 39)—Normal (Checkbox) b. Ears-General (Internal Canals) (auditory acuity under item 281)—Abnormal (Checkbox) c. Drums (Perforations)—Abnormal (Checkbox) d. Nose—Normal (Checkbox) d. Nose—Normal (Checkbox) e. Sinuses—Abnormal (Checkbox) e. Sinuses—Abnormal (Checkbox) f. Mouth and Throat—Abnormal (Checkbox) f. Mouth and Throat—Abnormal (Checkbox) g. Eyes—General (Visual accuity and refraction under item 281–28s)—Normal (Checkbox) g. Eyes—General (Visual accuity and refraction under item 281–28s)—Abnormal (Checkbox) h. Ophtalmoscopic—Normal (Checkbox) h. Ophtalmoscopic—Abnormal (Checkbox) h. Usugs and Chest—Normal (Checkbox) h. Usugs and Chest—Abnormal (Checkbox) h. Lungs and Chest—Abnormal (Checkbox) h. Lungs and Chest—Abnormal (Checkbox) h. Heart (Thrust, size, rhythm, sounds)—Normal (Checkbox) h. Vascular System—Abnormal (Checkbox) h. Abdomen and Viscera (Include hernia)—Abnormal (Checkbox) h. Abdomen and Viscera (Include hernia)—Abnormal (Checkbox) h. Prostate (Over 40 or clinical) indicated)—Abnormal (Checkbox) h. Foetale (Over 40 or clinical) indicated)—Abnormal (Checkbox) h. Foetale (Over 40 or clinical) indicated)—Abnormal (Checkbox) h. Foetale (Checkbox) h. Chauss and Rectum (Hernorrhods, Fistulae) (Hernocult Results)—Normal (Checkbox) h. Foetale Abnormal (Checkbox) h. Foetale Abnormal (Checkbox) h. Foetale Abnormal (Checkbox) h. Feet—N	Placement*
 Notes (Describe every abnormality in detail. Enter pertinent item number before each comment. Continue in item 29 and use additional sheets if necessary) Dental—Acceptable (Checkbox) Dental—Not Acceptable (Checkbox) 	
20. Dental—Not Acceptable (if checked, explain) 20. Dental—Dental Examination not done by Dental Officer 21. Remarks and Additional Dental Defects and Diseases	Ahava halaw Katad
22. Test Results (Copies of results are preferred as attachments) 22a. Urinalysis—Specific Gravity	Above below listed items.
22a. Urine Albumin 22a. Urine Sugar 22b. Syphilis Serology (Specify test used and results)	
22c. EKG 22d. Blood Type and RH Factor	

ELECTRONIC ELEMENTS FOR SF 88—Continued

ltem	Placement*
22e. Chest X-Ray or PPD (Place, date, film number and result)	
22f. Other Tests	
23. Relationship to Sponsor 24a. Sponsor's Name—Last	
24b. Sponsor's Name—First	
24c. Sponsor's Name—MI	
24c. Sponsor's ID Number (SSN or Other)	
25. Depart./Service	
26. Hospital or Medical Facility	
27. Records Maintained At Last Name—First Name—Middle Name	Tan of book none
Last Name—First Name—Middle Name	
Number of Sheets Attached	, , , ,
28. Measurements and Other Findings	
-	items.
28a. Height	
28b. Weight	
28d. Color Hair	
28d. Color Eyes 28e. Build—Slender (Checkbox)	
28e. Build—Medium (Checkbox)	
28e. Build—Heavy (Checkbox)	
28e. Build—Obese (Checkbox)	
28f. Temperature	
28g(1). Blood Pressure (Arm at heart level)—Sitting—Sys.	
28g(1). Blood Pressure (Arm at heart level)—Sitting—Dias. 28g(2). Blood Pressure (Arm at heart level)—Recumbent—Sys.	
28g(2). Blood Pressure (Arm at heart level)—Recumbent—Dias.	
28g(3). Blood Pressure (Arm at heart level)—Standing (5 minutes)—Sys.	
28g(3). Blood Pressure (Arm at heart level)—Standing (5 minutes)—Dias.	
28h(1). Pulse (Arm at heart level)—Sitting	
28h(2). Pulse (Arm at heart level)—Recumbent	
28h(3). Pulse (Arm at heart level)—Standing—3 minutes 28h(4). Pulse (Arm at heart level)—After Exercise	
28h(5). Pulse (Arm at heart level)—2 minutes after exercise	
28i(1). Distant Vision—Right 20/ (number)	
28i(1). Distant Vision—Right—Corrected to 20/ (number)	
28i(2). Distant Vision—Left 20/ (number)	
28i(2). Distant Vision—Left Corrected to 20/ (number)	
28j(1). Refraction—Right—By 28j(1). Refraction—Right—S	
28j(1). Refraction—Right—CX	
28j(2). Refraction—Left—By	
28j̇́(2)́. Refraction—Left—S´	
28j(2). Refraction—Left—CX	
28k(1). Near Vision—Right (Number)	
28k(1). Near Vision—Right—Corrected To (Number) 28k(1). Near Vision—Right—By (Number)	
28k(2). Near Vision—Left (Number)	
28k(2). Near Vision—Left—Corrected To (Number)	
28k(2). Near Vision—Left—By (Number)	
28I(1). Heterophoria (Specify Distance)—ESO	
28I(2). Heterophoria (Specify Distance)—EXO	
28I(3). Heterophoria (Specify Distance)—RH	
28I(4). Heterophoria (Specify Distance)—LH 28I(5). Heterophoria (Specify Distance)—Prism Division	
28I(6). Heterophoria (Specify Distance)—Prism Division 28I(6). Heterophoria (Specify Distance)—Prism Conv. Ct.	
28I(7). Heterophoria (Specify Distance)—PC	
28I(8). Heterophoria (Specify Distance)—PD	
28m(1). Accommodation—Right	
28m(2). Accommodation—Left	
28n(1). Field of Vision—Right	
28n(2). Field of Vision—Left 28o. Color Vision (Test used and result)	
28p. Night Vision (Test used and result)	
28g(1). Depth Perception (Test used and score)—Uncorrected	
28q(2). Depth Perception (Test used and score)—Corrected	
28r. Red Lens Test	
28s(1). Intraocular Tension—Right	
28s(2). Intraocular Tension—Left	
28t. Audiometer—Right Ear—500–512	1

ELECTRONIC ELEMENTS FOR SF 88—Continued

Item	Placement*
28t. Audiometer—Right Ear—2000–2048	
28t. Audiometer—Right Ear—3000–3096	
28t. Audiometer—Right Ear—4000–4096	
28t. Audiometer—Right Ear—6000–6144	
28t. Audiometer—Left Ear—500–512	
28t. Audiometer—Left Ear—100–1024	
28t. Audiometer—Left Ear—2000–2048	
28t. Audiometer—Left Ear—3000–3096	
28t. Audiometer—Left Ear—4000–4096	
28t. Audiometer—Left Ear—6000–6144	
28u. Psychological and Psychomotor (Tests used and score)	
29. Notes (Continued) and Significant or Interval History	
30. Summary of Defects and Diagnoses (List diagnoses with item numbers)	
31. Recommendations—Further Specialist Examinations Indicated (Specify)	
32. Physical Profile—P	
32. Physical Profile—U	
32. Physical Profile—L	
32. Physical Profile—H	
32. Physical Profile—E	
32. Physical Profile—S	
33. Examinee—Is Qualified for (Checkbox)	
33. Examinee—Is Qualified for Explanation	
33. Examinee—Is Not Qualified for (Checkbox)	
33. Examinee—Is Not Qualified for Explanation	
34. Physical Category—A	
34. Physical Category—B	
34. Physical Category—C	
34. Physical Category—E	
35. If Not Qualified, List Disqualifying Defects by Item Number	
36. Typed or Printed Name of Physician	
36. Signature of Physician	
37. Typed or Printed Name of Physician	
37. Signature of Physician	
38. Typed or Printed Name of Dentist or Physician (Indicate which)	
38. Signature of Dentist or Physician	
39. Typed or Printed Name of Reviewing Officer or Approving Authority	
39. Signature of Reviewing Officer or Approving Authority	

^{*}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, Rockville, MD 20857 or e-mail at kciacco@hqe.ihs.gov.

Dated: March 21, 2003.

Katherine Ciacco Palatianos,

Chairperson, Interagency Committee on Medical Records.

[FR Doc. 03-7927 Filed 4-1-03; 8:45 am]

BILLING CODE 6820-34-M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration [Docket No. 02N-0354]

Agency Information Collection Activities; Submission for OMB Review; Comment Request; The Evaluation of Long-Term Antibiotic Drug Therapy for Persons Involved in Anthrax Remediation Activities

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that the proposed collection of information listed below has been submitted to the Office of Management and Budget (OMB) for review and clearance under the Paperwork Reduction Act of 1995.

DATES: Fax written comments on the information collection provisions by May 2, 2003.

ADDRESSES: OMB is still experiencing significant delays in the regular mail, including first class and express mail, and messenger deliveries are not being accepted. To ensure that comments on the information collection are received, OMB recommends that written comments be electronically mailed to sshapiro@omb.eop.gov or faxed to the Office of Information and Regulatory Affairs, OMB, Attn: Stuart Shapiro, Desk Officer for FDA, FAX: 202–395–6974.

FOR FURTHER INFORMATION CONTACT:

Karen L. Nelson, Office of Information Resources Management (HFA–250), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–1482.

SUPPLEMENTARY INFORMATION: In compliance with 44 U.S.C. 3507, FDA has submitted the following proposed collection of information to OMB for review and clearance.