



DEPARTMENT OF HEALTH & HUMAN SERVICES

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Public Health Service

Central Region

Telephone (973) 526-6007

Food and Drug Administration
Waterview Corporate Center
10 Waterview Blvd., 3rd Floor
Parsippany, NJ 07054

April 22, 2002

WARNING LETTER

CERTIFIED MAIL
RETURN RECEIPT REQUESTED

Valentin L. Fagel, M.D.
Owner
Pennsville Radiology
248 South Broadway
Pennsville, New Jersey 08070

FILE NO.: 02-NWJ-21
Inspection ID NO.: 1304270011

Dear Dr. Fagel:

A representative from the State of New Jersey under contract to the Food and Drug Administration (FDA) inspected your facility on March 22 2002. This inspection revealed a serious regulatory problem involving mammography performed at your facility.

Under a United States Federal law, the Mammography Quality Standards Act of 1992 (MQSA) 42 U.S.C. 263b, your facility must meet specific requirements for mammography. These requirements help protect the health of women by assuring that a facility can perform quality mammography. The inspection revealed the following violation(s) of Section 354(f) of the Act [42 U.S.C. 263b(f)] at your facility (identified on your inspection report):

- Your facility has not implemented a system that provides the results of mammographic examinations to its patients. [See 21 CFR § 900.12(c)(2)].
- Five mammography reports were reviewed and it was found that five of the five random reports did not contain an acceptable assessment category. [See 21 CFR § 900.12(c)(1)(iv)]

The specific problems noted above appeared on your MQSA Facility Inspection Report, which was issued to your facility at the close of the inspection.

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Because these conditions may be symptomatic of serious underlying problems that could compromise the quality of mammography at your facility, they represent serious violations of the law which may result in FDA taking regulatory action without further notice to you. These actions include, but are not limited to, placing your facility under a Directed Plan of Correction, charging your facility for the cost of on-site monitoring, assessing civil money penalties up to \$10,000 for each failure to substantially comply with, or each day of failure to substantially comply with, the Standards, suspension or revocation of your facility's FDA certificate, or obtaining a court injunction against further mammography (see Sections 354(h) through (j) of the Act, 42 U.S.C. 263b(h) through (j)).

It is necessary for you to act on this matter immediately. Please explain to this office in writing within 15 working days from the date you received this letter:

- the specific steps you have taken to correct the violations noted in this letter;
- each step your facility is taking to prevent the recurrence of similar violations;
- equipment settings (including technique factors), raw test data, and calculated final results, where appropriate; and
- sample records that demonstrate proper record keeping procedures.

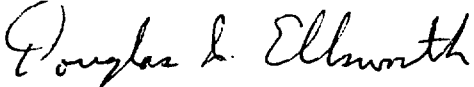
Please submit your response to Rosa L. Brown, Compliance Technician, Food and Drug Administration, New Jersey District, 10 Waterview Blvd, 3rd Floor, Parsippany, New Jersey 07054.

Finally, you should understand that there are many FDA requirements pertaining to mammography. This letter pertains only to findings disclosed during the inspection of your facility and does not necessarily address other obligations you have under the law. You may obtain general information about all of FDA's requirements for mammography facilities by contacting the Mammography Quality Assurance Program, Food and Drug Administration, P.O. Box 6057, Columbia, MD 21045-6057 (1-800-838-7715) or through the Internet at <http://www.fda.gov>.

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If you have technical questions about mammography facility requirements, or about the content of this letter, please feel free to contact Elizabeth A. Laudig at (410) 779-5441.

Sincerely,



DOUGLAS I. ELLSWORTH
District Director
New Jersey District Office

cc: Priscilla F. Butler, M.S.
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