

EXHIBIT G-9



DEPARTMENT OF HEALTH & HUMAN SERVICES

Public Health Service

Food and Drug Administration
Rockville MD 20857

July 26, 2002

Facsimile

FDA MQSA Facility ID#: 110957

Amile Korangy

Baltimore Imaging Center

724 Maiden Choice Lane, Suite 102

Baltimore, MD 21228

Dear Amile Korangy:

The enclosed Food and Drug Administration (FDA) Interim Notice is being sent to your facility because: (1) your facility has applied for accreditation from an FDA-approved Accreditation Body (AB), and (2) FDA has received information from the Accreditation Body indicating your facility's application is sufficiently complete for review purposes. Until you receive your Provisional Certificate, while your facility's application is in progress, this Interim Notice certifies your facility as a lawful provider of mammography services under the requirements of the Mammography Quality Standards Act of 1992 (MQSA). This Interim Notice expires on September 8, 2002.

This document means that FDA acknowledges that your facility is certified by FDA to lawfully provide mammography services during the interim period while you are awaiting receipt of your FDA Mammography Facility Provisional Reinstatement Certificate, including: the operation of x-ray equipment to produce mammograms; the processing of films, interpretation of mammograms; medical record keeping; reporting and patient notification; and outcomes data collection.

Furthermore, as an FDA certified mammography facility, you must:

treat the notice as you would the certificate and uphold the requirements to the Mammography Quality Standards Act (MQSA) of 1992; and

- prominently post this notice in the mammography facility where it can be viewed by mammography patients until it can be replaced with an FDA Mammography Facility Provisional Reinstatement Certificate.

If accreditation is denied, this Provisional Reinstatement Interim Notice is **no longer valid** and you must cease performing mammography services immediately and no longer display this notice. This Interim Notice also should no longer be displayed upon receipt of a certificate or provisional certificate from FDA.

Baltimore Imaging Center
Baltimore, Maryland
FEI#1000512417 CFN 1124028
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8/8, 8/12, 8/14, 8/21, 8/22, 9/3, 9/5, 9/6/2002
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If you have any questions about this Interim Notice or questions or comments regarding your certificate, the certification process, inspections, or general MQSA information, you may use our 800 hotline number, fax number or address provided below.

If you wish us to direct future correspondence to someone other than yourself, please notify your Accreditation Body to change the official contact designated for your facility.

FDA Contact Information:

Hotline: 1-800-838-7715
Fax Contact: Food and Drug Administration
1-410-290-6351
Address: Mammography Quality Assurance Program
Food and Drug Administration
P.O. Box 6057
Columbia, MD 21045-6057

Sincerely yours,



CAPT John L. McCrohan, USPHIS
Director
Division of Mammography Quality
and Radiation Programs
Office of Health and Industry Programs
Center for Devices and Radiological Health

Enclosure - Interim Notice (Provisional Reinstatement)