## MQSA Archived Document

Although some of the information in this document has been modified or no longer applies to MQSA regulatory requirements, this item is presented here for research and historical reference.

U.S. Food and Drug Administration - Center for Devices and Radiological Health)

## **Guidance for Industry and for FDA Staff**

## GUIDANCE FOR REQUEST AND ISSUANCE OF INTERIM NOTICE LETTERS FOR MAMMOGRAPHY FACILITIES UNDER THE MAMMOGRAPHY QUALITY STANDARDS ACT, 42 U.S.C. § 263(b)

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Please note that the contents of this document have been incorporated into the Policy Guidance Help System.

The Policy Guidance Help System can be found at: http://www.fda.gov/cdrh/mammography/browstest.html

Information about the Policy Guidance Help System can be found at <a href="http://www.fda.gov/cdrh/mammography/guidance-rev.html">http://www.fda.gov/cdrh/mammography/guidance-rev.html</a>

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