

**K993365 GE LOGIQ 700 WITH HARMONIC IMAGING  
MODIFICATION**Nov 5, 1999  
30 days to decisionK993365 · Product code: **IYO** · Radiology  
Source: <https://www.510kdatabase.net/k993365/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	System, Imaging, Pulsed Echo, Ultrasonic (IYO)
Date received	Oct 6, 1999
Decision date	Nov 5, 1999
Days to decision	30 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>Ge Medical Systems, Inc.</b>
Location	Milwaukee, WI, US
Contact	ALLEN SCHUH
510(k) history	54 submissions · 54 cleared · 1997-2009

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k993365/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 27, 2026