

K000571 GE LOGIQ 700Apr 18, 2000
56 days to decisionK000571 · Product code: **IYO** · Radiology
Source: <https://www.510kdatabase.net/k000571/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	System, Imaging, Pulsed Echo, Ultrasonic (IYO)
Date received	Feb 22, 2000
Decision date	Apr 18, 2000
Days to decision	56 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	GE Medical Systems
Location	Milwaukee, WI, US
Contact	ALLEN SCHUH
510(k) history	169 submissions · 166 cleared · 1989-2022

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k000571/> 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