FDA 510(k) Premarket Notification

K993365 GE LOGIQ 700 WITH HARMONIC IMAGING MODIFICATION

CLEARED

Nov 5, 1999
30 days to decision

K993365 · 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	**Ge Medical Systems, Inc.**
Location	Milwaukee, WI, US
Contact	ALLEN SCHUH
510(k) history	54 submissions · 54 cleared · 1997-2009