FDA 510(k) Premarket Notification

CLEARED

K060542 GE VIVID 7 MODEL FC008XX, GE VIVID 7 MODEL FC009XX

Mar 31, 2006
30 days to decision

K060542 · Product code: **IYN** · Radiology
Source: https://www.510kdatabase.net/k060542/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	System, Imaging, Pulsed Doppler, Ultrasonic (IYN)
Date received	Mar 1, 2006
Decision date	Mar 31, 2006
Days to decision	30 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**General Electric Co.**
Location	Mchenry, IL, US
Contact	ALLEN SCHUH
510(k) history	254 submissions · 254 cleared · 1976-2011