FDA 510(k) Premarket Notification

K063085 MODIFICATION TO ACUSON SEQUOIA ULTRASOUND SYSTEM

CLEARED

Nov 14, 2006
35 days to decision

K063085 · Product code: **IYO** · Radiology
Source: https://www.510kdatabase.net/k063085/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	System, Imaging, Pulsed Echo, Ultrasonic (IYO)
Date received	Oct 10, 2006
Decision date	Nov 14, 2006
Days to decision	35 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Siemens Medical Solutions USA, Inc.**
Location	Hoffman Estates, IL, US
Contact	SHEILA W PICKERING
510(k) history	778 submissions · 778 cleared · 1980-2026