FDA 510(k) Premarket Notification

K081121 MODIFICATION TO ACUSON X150 ULTRASOUND SYSTEM

CLEARED

Jun 4, 2008
44 days to decision

K081121 · Product code: **IYO** · Radiology
Source: https://www.510kdatabase.net/k081121/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Imaging, Pulsed Echo, Ultrasonic (IYO)
Date received	Apr 21, 2008
Decision date	Jun 4, 2008
Days to decision	44 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Siemens Medical Solutions USA, Inc., Ultrasound DI**
Location	Mountain View, CA, US
Contact	MARTINA VOGT
510(k) history	11 submissions · 11 cleared · 2007-2017