FDA 510(k) Premarket Notification

K071134 SONOSITE MAXX SERIES DIAGNOSTIC ULTRASOUND SYSTEM

CLEARED

May 8, 2007
15 days to decision

K071134 · Product code: **IYO** · Radiology
Source: https://www.510kdatabase.net/k071134/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Imaging, Pulsed Echo, Ultrasonic (IYO)
Date received	Apr 23, 2007
Decision date	May 8, 2007
Days to decision	15 days
Third-party review	Yes
Summary / Statement	Summary

APPLICANT

Company	**Sonosite,Inc.**
Location	Bothell, WA, US
Contact	DAINA L GRAHAM
510(k) history	23 submissions · 23 cleared · 1998-2011