

K930964 ACUSON MODEL #29001 (DIAGNOSTIC ULTRASOUND DEVICE)

Oct 26, 1994
609 days to decision

K930964 · Product code: IYO · Radiology
Source: <https://www.510kdatabase.net/k930964/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Imaging, Pulsed Echo, Ultrasonic (IYO)
Date received	Feb 24, 1993
Decision date	Oct 26, 1994
Days to decision	609 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Acuson Corp.
Location	Mountain View, CA, US
Contact	SHEILA PICKERING
510(k) history	37 submissions · 37 cleared · 1988-2002

510k Database - www.510kdatabase.net
 Device record: <https://www.510kdatabase.net/k930964/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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