

K983879 ALOKA SSD-900 DIAGNOSTIC ULTRASOUND SYSTEM

Mar 22, 1999
140 days to decision

K983879 · Product code: **IYO** · Radiology
Source: <https://www.510kdatabase.net/k983879/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Imaging, Pulsed Echo, Ultrasonic (IYO)
Date received	Nov 2, 1998
Decision date	Mar 22, 1999
Days to decision	140 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Aloka Co., Ltd.
Location	Tokyo, JP
Contact	PAUL D SMOLENSKI
510(k) history	39 submissions · 39 cleared · 1982-2011

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Device record: <https://www.510kdatabase.net/k983879/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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