

K991671 HDI 5000 DIAGNOSTIC ULTRASOUND SYSTEM

Jun 18, 1999
 32 days to decision

K991671 · Product code: **IYO** · Radiology
 Source: <https://www.510kdatabase.net/k991671/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Imaging, Pulsed Echo, Ultrasonic (IYO)
Date received	May 17, 1999
Decision date	Jun 18, 1999
Days to decision	32 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Advanced Technology Laboratories, Inc.
Location	Mchenry, IL, US
Contact	TERRENCE J SWEENEY
510(k) history	61 submissions · 59 cleared · 1980-2002

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Device record: <https://www.510kdatabase.net/k991671/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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