

K924231 INTERSPEC APOGEE CLAJun 14, 1993
297 days to decisionK924231 · Product code: **IYO** · Radiology
Source: <https://www.510kdatabase.net/k924231/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Imaging, Pulsed Echo, Ultrasonic (IYO)
Date received	Aug 21, 1992
Decision date	Jun 14, 1993
Days to decision	297 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Interspec, Inc.
Location	Mchenry, IL, US
Contact	ARTHUR J SCHENCK
510(k) history	17 submissions · 16 cleared · 1982-1994

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k924231/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 20, 2026