

**K903839 INTERSPEC APOGEE**Nov 16, 1990  
87 days to decisionK903839 · Product code: **DXK** · Cardiovascular  
Source: <https://www.510kdatabase.net/k903839/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Echocardiograph (DXK)
Date received	Aug 21, 1990
Decision date	Nov 16, 1990
Days to decision	87 days
Third-party review	No

**APPLICANT**

---

Company	<b>Interspec, Inc.</b>
Location	Mchenry, IL, US
Contact	KAI E THOMENIUS
510(k) history	17 submissions · 16 cleared · 1982-1994

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k903839/>; Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated May 20, 2026