FDA 510(k) Premarket Notification

K903839 INTERSPEC APOGEE

CLEARED

Nov 16, 1990
87 days to decision

K903839 · 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	**Interspec, Inc.**
Location	Mchenry, IL, US
Contact	KAI E THOMENIUS
510(k) history	17 submissions · 16 cleared · 1982-1994