FDA 510(k) Premarket Notification

K901592 MODEL MD-5 DOPPLER

CLEARED

Oct 22, 1990
200 days to decision

K901592 · Product code: **DXK** · Cardiovascular
Source: https://www.510kdatabase.net/k901592/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Echocardiograph (DXK)
Date received	Apr 5, 1990
Decision date	Oct 22, 1990
Days to decision	200 days
Third-party review	No

APPLICANT

Company	**D. E. Hokanson, Inc.**
Location	Mchenry, IL, US
Contact	EUGENE HOKANSON
510(k) history	19 submissions · 19 cleared · 1979-2004