FDA 510(k) Premarket Notification

K900560 ACUSON HEARTSOUND/PULSE/RESPIRATION MODULE

CLEARED

Dec 3, 1990
299 days to decision

K900560 · Product code: **DQC** · Cardiovascular
Source: https://www.510kdatabase.net/k900560/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Phonocardiograph (DQC)
Date received	Feb 7, 1990
Decision date	Dec 3, 1990
Days to decision	299 days
Third-party review	No

APPLICANT

Company	**Acuson Corp.**
Location	Mountain View, CA, US
Contact	T JOHNSON
510(k) history	37 submissions · 37 cleared · 1988-2002