FDA 510(k) Premarket Notification

K933517 AUTOMATIC PULL BACK DEVICE

CLEARED

Oct 15, 1993
87 days to decision

K933517 · Product code: **DQX** · Cardiovascular
Source: https://www.510kdatabase.net/k933517/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Wire, Guide, Catheter (DQX)
Date received	Jul 20, 1993
Decision date	Oct 15, 1993
Days to decision	87 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Cardiovascular Imaging Systems, Inc.**
Location	Sunnyvale, CA, US
Contact	YUE-TEH JANG
510(k) history	19 submissions · 19 cleared · 1989-1997