FDA 510(k) Premarket Notification

K833801 RIGHT CORONARY GUIDING CATHETER

CLEARED

Mar 12, 1984
132 days to decision

K833801 · Product code: **DQY** · Cardiovascular
Source: https://www.510kdatabase.net/k833801/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Percutaneous (DQY)
Date received	Nov 1, 1983
Decision date	Mar 12, 1984
Days to decision	132 days
Third-party review	No

APPLICANT

Company	**Advanced Cardiovascular Systems, Inc.**
Location	Santa Clara, CA, US
510(k) history	103 submissions · 100 cleared · 1982-2002