

K833801 RIGHT CORONARY GUIDING CATHETERMar 12, 1984
132 days to decisionK833801 · Product code: **DQY** · CardiovascularSource: <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

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k833801/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated July 1, 2026