FDA 510(k) Premarket Notification

K871021 GUIDING CATHETERS 9F JL4

CLEARED

Jun 16, 1987
92 days to decision

K871021 · Product code: **DQO** · Cardiovascular
Source: https://www.510kdatabase.net/k871021/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular, Diagnostic (DQO)
Date received	Mar 16, 1987
Decision date	Jun 16, 1987
Days to decision	92 days
Third-party review	No

APPLICANT

Company	**Devices For Vascular Intervention, Inc.**
Location	Redwood City, CA, US
Contact	JAMES F PFEIFFER
510(k) history	14 submissions · 14 cleared · 1987-1994