FDA 510(k) Premarket Notification

K924591 SCHNEIDER SOLID-7 SOPTIP GUIDING CATHETER

CLEARED

Dec 8, 1992
89 days to decision

K924591 · Product code: **DQY** · Cardiovascular
Source: https://www.510kdatabase.net/k924591/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Percutaneous (DQY)
Date received	Sep 10, 1992
Decision date	Dec 8, 1992
Days to decision	89 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Schneider Intl., Ltd.**
Location	Minneapolis, MN, US
Contact	ROBERT L ULLEN
510(k) history	22 submissions · 22 cleared · 1989-1995