FDA 510(k) Premarket Notification

K914009 SCHNEIDER SOFTIP(R) A+(TM) ANGIOGRAPHIC CATHETER

CLEARED

Dec 6, 1991
88 days to decision

K914009 · Product code: **DQO** · Cardiovascular
Source: https://www.510kdatabase.net/k914009/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular, Diagnostic (DQO)
Date received	Sep 9, 1991
Decision date	Dec 6, 1991
Days to decision	88 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

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