FDA 510(k) Premarket Notification

K875325 MODIFIED OCCLUDER/FLUSHING CATHETER STOPCOCK

CLEARED

Feb 18, 1988
51 days to decision

K875325 · Product code: **DQO** · Cardiovascular
Source: https://www.510kdatabase.net/k875325/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular, Diagnostic (DQO)
Date received	Dec 29, 1987
Decision date	Feb 18, 1988
Days to decision	51 days
Third-party review	No

APPLICANT

Company	**Schneider-Shilley (Usa)**
Location	Minneapolis, MN, US
Contact	ANN MORRISSEY
510(k) history	9 submissions · 9 cleared · 1988-1989