

**K875325 MODIFIED OCCLUDER/FLUSHING CATHETER
STOPCOCK**Feb 18, 1988
51 days to decisionK875325 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k875325/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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