

**K950094 SCIMED 6 FRENCH MICRO-MAX(TM) GUIDE
CATHETER**Mar 3, 1995
52 days to decisionK950094 · Product code: **DQY** · Cardiovascular
Source: <https://www.510kdatabase.net/k950094/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Percutaneous (DQY)
Date received	Jan 10, 1995
Decision date	Mar 3, 1995
Days to decision	52 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Scimed Life Systems, Inc.
Location	Mchenry, IL, US
Contact	DIANE M LOWE
510(k) history	109 submissions · 108 cleared · 1977-1998

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k950094/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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