

K951720 SCIMED EXPO ANGIOGRAPHIC CATHETERJun 22, 1995
69 days to decisionK951720 · Product code: **DQO** · CardiovascularSource: <https://www.510kdatabase.net/k951720/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular, Diagnostic (DQO)
Date received	Apr 14, 1995
Decision date	Jun 22, 1995
Days to decision	69 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k951720/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated June 29, 2026