

**K934122 SCIMED TRANSEND EX STEERABLE GUIDE WIRE
AND ACCESSORIES**Oct 7, 1993
43 days to decisionK934122 · Product code: **DQX** · Cardiovascular
Source: <https://www.510kdatabase.net/k934122/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Wire, Guide, Catheter (DQX)
Date received	Aug 25, 1993
Decision date	Oct 7, 1993
Days to decision	43 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Scimed Peripheral Interventions
Location	Plymouth, MN, US
Contact	DARLENE A THOMETZ
510(k) history	8 submissions · 8 cleared · 1993-1995

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k934122/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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