

K941710 SCIMED TRANSEND STEERABLE GUIDE WIRE AND ACCESSORIESSep 1, 1994
148 days to decisionK941710 · Product code: **DQX** · Cardiovascular
Source: <https://www.510kdatabase.net/k941710/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Wire, Guide, Catheter (DQX)
Date received	Apr 6, 1994
Decision date	Sep 1, 1994
Days to decision	148 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/k941710/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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