

**K931577 SCHNEIDER MAGNUM PERIPHERAL  
RECANALIZATION GUIDEWI**Jun 23, 1993  
85 days to decisionK931577 · Product code: **DQX** · Cardiovascular  
Source: <https://www.510kdatabase.net/k931577/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Wire, Guide, Catheter (DQX)
Date received	Mar 30, 1993
Decision date	Jun 23, 1993
Days to decision	85 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Schneider Intl., Ltd.</b>
Location	Minneapolis, MN, US
Contact	AMY PETERSON
510(k) history	22 submissions · 22 cleared · 1989-1995

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k931577/>. Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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