

**K914009 SCHNEIDER SOFTIP(R) A+(TM) ANGIOGRAPHIC CATHETER**

Dec 6, 1991  
88 days to decision

K914009 · Product code: **DQO** · Cardiovascular  
Source: <https://www.510kdatabase.net/k914009/>

**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular, Diagnostic (DQO)
Date received	Sep 9, 1991
Decision date	Dec 6, 1991
Days to decision	88 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

---

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

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k914009/> 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 July 2, 2026