

**K914128 CONNECTING SET**Nov 27, 1991  
75 days to decisionK914128 · Product code: **DQO** · Cardiovascular  
Source: <https://www.510kdatabase.net/k914128/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular, Diagnostic (DQO)
Date received	Sep 13, 1991
Decision date	Nov 27, 1991
Days to decision	75 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

---

Company	<b>Procedure Products, Inc.</b>
Location	Mchenry, IL, US
Contact	ROBERT B EVERETT
510(k) history	16 submissions · 16 cleared · 1981-2017

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k914128/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated June 1, 2026