

K922536 TORQUE DEVICEDec 10, 1992
196 days to decisionK922536 · Product code: **DQX** · CardiovascularSource: <https://www.510kdatabase.net/k922536/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Wire, Guide, Catheter (DQX)
Date received	May 28, 1992
Decision date	Dec 10, 1992
Days to decision	196 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Procedure Products, Inc.
Location	Mchenry, IL, US
Contact	BOB EVERETT
510(k) history	16 submissions · 16 cleared · 1981-2017

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k922536/> 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 June 1, 2026