

**K913783 CORDIS PTCA GUIDING CATHETER, MODIFICATION**Oct 18, 1991  
65 days to decisionK913783 · Product code: **DQY** · CardiovascularSource: <https://www.510kdatabase.net/k913783/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Percutaneous (DQY)
Date received	Aug 14, 1991
Decision date	Oct 18, 1991
Days to decision	65 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Dow Corning Wright</b>
Location	Mchenry, IL, US
Contact	ERIN MCGURK-BURLESON
510(k) history	74 submissions · 52 cleared · 1979-1994

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k913783/> 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 May 22, 2026