

**K884183 BURRON TEFLON-COATED GUIDEWIRE**Aug 3, 1989  
303 days to decisionK884183 · Product code: **DQX** · Cardiovascular  
Source: <https://www.510kdatabase.net/k884183/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Wire, Guide, Catheter (DQX)
Date received	Oct 4, 1988
Decision date	Aug 3, 1989
Days to decision	303 days
Third-party review	No

**APPLICANT**

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Company	<b>Burron Medical, Inc.</b>
Location	Bethlehem, PA, US
Contact	PEGGY KEIFFER
510(k) history	9 submissions · 9 cleared · 1988-1991

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k884183/>; 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 27, 2026