

**K950026 MEDTRONIC(R) 8.2 FRENCH SHERPA(R) SUPER PEAK FLOW(TM) CORONARY GUIDING CATHETER**Apr 5, 1995  
91 days to decisionK950026 · Product code: **DQY** · Cardiovascular  
Source: <https://www.510kdatabase.net/k950026/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Percutaneous (DQY)
Date received	Jan 4, 1995
Decision date	Apr 5, 1995
Days to decision	91 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Medtronic Interventional Vascular</b>
Location	Danvers, MA, US
Contact	JOSEPH O MAGLIOZZI
510(k) history	21 submissions · 21 cleared · 1992-1999

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k950026/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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