

**K915351 SHERPA GUIDING CATHETER WITH CATHETER INTRODUCER**

Jun 29, 1992  
216 days to decision

K915351 · Product code: **DQO** · Cardiovascular  
Source: <https://www.510kdatabase.net/k915351/>

**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular, Diagnostic (DQO)
Date received	Nov 26, 1991
Decision date	Jun 29, 1992
Days to decision	216 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Medtronic Interventional Vascular</b>
Location	Danvers, MA, US
Contact	KIRK DALY
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/k915351/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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