

**K880776 7 FR. TRIPLE LUMEN CENTRAL VEIN CATHETERI.  
SYSTEM**May 2, 1988  
66 days to decisionK880776 · Product code: **DQR** · Cardiovascular  
Source: <https://www.510kdatabase.net/k880776/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Cannula, Catheter (DQR)
Date received	Feb 26, 1988
Decision date	May 2, 1988
Days to decision	66 days
Third-party review	No

**APPLICANT**

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Company	<b>Medsurg Industries, Inc.</b>
Location	Mchenry, IL, US
Contact	TOM BONNER
510(k) history	41 submissions · 26 cleared · 1982-1994

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**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k880776/>; 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