

**K881884 HARBOR CENTRAL VENOUS CATHETER**Jul 28, 1988  
85 days to decisionK881884 · Product code: **DQR** · CardiovascularSource: <https://www.510kdatabase.net/k881884/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Cannula, Catheter (DQR)
Date received	May 4, 1988
Decision date	Jul 28, 1988
Days to decision	85 days
Third-party review	No

**APPLICANT**

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Company	<b>Harbor Medical Devices, Inc.</b>
Location	Boston, MA, US
Contact	JOSHUA TOLKOFF
510(k) history	7 submissions · 7 cleared · 1988-1989

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k881884/>; 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