

**K820736 CENTRAL VENOUS ACCESS CATHETER**Apr 15, 1982  
28 days to decisionK820736 · Product code: **DQY** · CardiovascularSource: <https://www.510kdatabase.net/k820736/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Percutaneous (DQY)
Date received	Mar 18, 1982
Decision date	Apr 15, 1982
Days to decision	28 days
Third-party review	No

**APPLICANT**

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Company	<b>Hemed, Inc.</b>
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1982-1982

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k820736/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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