

**K834237 GRADUATED GUIDE WIRE BOWL 8**Feb 4, 1984  
60 days to decisionK834237 · Product code: **DQX** · CardiovascularSource: <https://www.510kdatabase.net/k834237/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Wire, Guide, Catheter (DQX)
Date received	Dec 6, 1983
Decision date	Feb 4, 1984
Days to decision	60 days
Third-party review	No

**APPLICANT**

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Company	<b>Sterile Design, Inc.</b>
Location	Mchenry, IL, US
510(k) history	10 submissions · 9 cleared · 1979-1994

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**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k834237/> 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 June 29, 2026