

**K820887 MITRAL VALVE ORIFICE SIZER**Jul 8, 1982  
100 days to decisionK820887 · Product code: **DTI** · Cardiovascular  
Source: <https://www.510kdatabase.net/k820887/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Sizer, Heart-valve, Prosthesis (DTI)
Date received	Mar 30, 1982
Decision date	Jul 8, 1982
Days to decision	100 days
Third-party review	No

**APPLICANT**

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

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Device record: <https://www.510kdatabase.net/k820887/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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