

**K800555 PARTNER SEVEN DISP. BUBBLE OXYGENATOR**Apr 4, 1980  
23 days to decisionK800555 · Product code: **DTZ** · CardiovascularSource: <https://www.510kdatabase.net/k800555/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Oxygenator, Cardiopulmonary Bypass (DTZ)
Date received	Mar 12, 1980
Decision date	Apr 4, 1980
Days to decision	23 days
Third-party review	No

**APPLICANT**

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Company	<b>Cardiovascular Research, Inc.</b>
Location	Mchenry, IL, US
510(k) history	2 submissions · 2 cleared · 1980-1984

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**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k800555/> 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 July 8, 2026