

**K962641 MAXIMA FORTE HOLLOW FIBER OXYGENATOR WITH PLASMA RESISTANT FIBER**Dec 16, 1996  
164 days to decisionK962641 · Product code: DTZ · Cardiovascular  
Source: <https://www.510kdatabase.net/k962641/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Oxygenator, Cardiopulmonary Bypass (DTZ)
Date received	Jul 5, 1996
Decision date	Dec 16, 1996
Days to decision	164 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Medtronic Vascular</b>
Location	Walker, MI, US
Contact	DEBRA KRIDNER
510(k) history	475 submissions · 453 cleared · 1977-2023

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k962641/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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