

**K920889 MINIMAX(R)/MAXIMA(R) HOLLOW FIBER
OXYGENATORS**Sep 15, 1992
216 days to decisionK920889 · Product code: **DTZ** · Cardiovascular
Source: <https://www.510kdatabase.net/k920889/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Oxygenator, Cardiopulmonary Bypass (DTZ)
Date received	Feb 12, 1992
Decision date	Sep 15, 1992
Days to decision	216 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Medtronic Vascular
Location	Walker, MI, US
Contact	NORMA L LOWE
510(k) history	475 submissions · 453 cleared · 1977-2023

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k920889/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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