

**K891230 MAXIMA* HOLLOW FIBER OXYGENATOR
W/FILTERED RESERV.**May 19, 1989
73 days to decisionK891230 · Product code: **DTZ** · Cardiovascular
Source: <https://www.510kdatabase.net/k891230/>**SUBMISSION DETAILS**

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|-----------------------|--|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Oxygenator, Cardiopulmonary Bypass (DTZ) |
| Date received | Mar 7, 1989 |
| Decision date | May 19, 1989 |
| Days to decision | 73 days |
| Third-party review | No |

APPLICANT

| | |
|----------------|---|
| Company | Medtronic Blood Systems, Inc. |
| Location | Anaheim, CA, US |
| Contact | SHARON ROCKWELL |
| 510(k) history | 15 submissions · 15 cleared · 1988-1995 |

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k891230/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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