

**K896671 SARNs MEMBRANE OXYGENATOR W/INTEGRAL RESER. #9461**Feb 14, 1990  
79 days to decisionK896671 · Product code: **DTZ** · Cardiovascular  
Source: <https://www.510kdatabase.net/k896671/>**SUBMISSION DETAILS**

---

|                       |  |
|-----------------------|--|
| Decision              | Substantially Equivalent (Cleared)       |
| Submission type       | Traditional                              |
| Device classification | Oxygenator, Cardiopulmonary Bypass (DTZ) |
| Date received         | Nov 27, 1989                             |
| Decision date         | Feb 14, 1990                             |
| Days to decision      | 79 days                                  |
| Third-party review    | No                                       |
| Combination product   | No                                       |
| PCCP authorized       | No                                       |

**APPLICANT**

---

|                |   |
|----------------|---|
| Company        | <b>3M Company</b>                                   |
| Location       | White City, OR, US                                  |
| Contact        | D SHIMOKOCHI  |
| Website        | <a href="http://www.3m.com/">http://www.3m.com/</a> |
| 510(k) history | 331 submissions · 322 cleared · 1976-2025           |

---

**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k896671/>. Data sourced from FDA 510(k) public records ([accessdata.fda.gov](https://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](https://accessdata.fda.gov). - Generated June 28, 2026