

**K841031 OXYHI CRYSTALLOID CARDIOPLEGIA SOLUT**May 23, 1984  
72 days to decisionK841031 · Product code: **DTN** · CardiovascularSource: <https://www.510kdatabase.net/k841031/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Reservoir, Blood, Cardiopulmonary Bypass (DTN)
Date received	Mar 12, 1984
Decision date	May 23, 1984
Days to decision	72 days
Third-party review	No

**APPLICANT**

---

Company	<b>American Bentley</b>
Location	Mchenry, IL, US
510(k) history	59 submissions · 59 cleared · 1983-1988

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k841031/> 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 May 27, 2026