

**K864090 SHELF LIFE CHANGE OF STERILE DEVICES**Mar 20, 1987  
178 days to decisionK864090 · Product code: **DRS** · CardiovascularSource: <https://www.510kdatabase.net/k864090/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Transducer, Blood-pressure, Extravascular (DRS)
Date received	Sep 23, 1986
Decision date	Mar 20, 1987
Days to decision	178 days
Third-party review	No

**APPLICANT**

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Company	<b>Utah Medical Products, Inc.</b>
Location	Mchenry, IL, US
Contact	EDWIN O GOODMAN
510(k) history	38 submissions · 38 cleared · 1979-2015

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k864090/>, 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 23, 2026