

**K851987 FLUSH DEVICE, AUTOMATIC**Aug 16, 1985  
101 days to decisionK851987 · Product code: **DRS** · CardiovascularSource: <https://www.510kdatabase.net/k851987/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Transducer, Blood-pressure, Extravascular (DRS)
Date received	May 7, 1985
Decision date	Aug 16, 1985
Days to decision	101 days
Third-party review	No

**APPLICANT**

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Company	<b>Transamerica Delaval, Inc.</b>
Location	Walker, MI, US
Contact	RICHARD G EKSTROM
510(k) history	7 submissions · 7 cleared · 1984-1986

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k851987/>; 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 June 29, 2026