

K833170 FLUSH DEVICEAug 12, 1984
331 days to decisionK833170 · Product code: **DRS** · CardiovascularSource: <https://www.510kdatabase.net/k833170/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Transducer, Blood-pressure, Extravascular (DRS)
Date received	Sep 16, 1983
Decision date	Aug 12, 1984
Days to decision	331 days
Third-party review	No

APPLICANT

Company	Transamerica Delaval, Inc.
Location	Walker, MI, US
510(k) history	7 submissions · 7 cleared · 1984-1986

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Device record: <https://www.510kdatabase.net/k833170/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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