

K833171 MONITORING KITSep 6, 1984
356 days to decisionK833171 · Product code: **DRS** · CardiovascularSource: <https://www.510kdatabase.net/k833171/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Transducer, Blood-pressure, Extravascular (DRS)
Date received	Sep 16, 1983
Decision date	Sep 6, 1984
Days to decision	356 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/k833171/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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