

K833792 TRANTECSILICONE QUARTZ DISP. TRANSDFeb 10, 1984
102 days to decisionK833792 · Product code: **DRS** · CardiovascularSource: <https://www.510kdatabase.net/k833792/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Transducer, Blood-pressure, Extravascular (DRS)
Date received	Oct 31, 1983
Decision date	Feb 10, 1984
Days to decision	102 days
Third-party review	No

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k833792/> Data sourced from FDA 510(k) public records (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. - Generated May 27, 2026