FDA 510(k) Premarket Notification

K833792 TRANTECSILICONE QUARTZ DISP. TRANSD

CLEARED

Feb 10, 1984
102 days to decision

K833792 · Product code: **DRS** · Cardiovascular
Source: 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