FDA 510(k) Premarket Notification

K833170 FLUSH DEVICE

CLEARED

Aug 12, 1984
331 days to decision

K833170 · Product code: **DRS** · Cardiovascular
Source: 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