FDA 510(k) Premarket Notification

K851987 FLUSH DEVICE, AUTOMATIC

CLEARED

Aug 16, 1985
101 days to decision

K851987 · Product code: **DRS** · Cardiovascular
Source: https://www.510kdatabase.net/k851987/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Transducer, Blood-pressure, Extravascular (DRS)
Date received	May 7, 1985
Decision date	Aug 16, 1985
Days to decision	101 days
Third-party review	No

APPLICANT

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