FDA 510(k) Premarket Notification

K864090 SHELF LIFE CHANGE OF STERILE DEVICES

CLEARED

Mar 20, 1987
178 days to decision

K864090 · Product code: **DRS** · Cardiovascular
Source: https://www.510kdatabase.net/k864090/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Transducer, Blood-pressure, Extravascular (DRS)
Date received	Sep 23, 1986
Decision date	Mar 20, 1987
Days to decision	178 days
Third-party review	No

APPLICANT

Company	**Utah Medical Products, Inc.**
Location	Mchenry, IL, US
Contact	EDWIN O GOODMAN
510(k) history	38 submissions · 38 cleared · 1979-2015