FDA 510(k) Premarket Notification

K781680 MONITORING, PRESSURE KIT

CLEARED

Oct 6, 1978
4 days to decision

K781680 · Product code: **DRS** · Cardiovascular
Source: https://www.510kdatabase.net/k781680/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Transducer, Blood-pressure, Extravascular (DRS)
Date received	Oct 2, 1978
Decision date	Oct 6, 1978
Days to decision	4 days
Third-party review	No

APPLICANT

Company	**Pharmaseal Div., Baxter Healthcare Corp.**
Location	Mchenry, IL, US
510(k) history	24 submissions · 24 cleared · 1976-1979