FDA 510(k) Premarket Notification

K792226 CARDIVAN BLOOD PRESSURE MONITOR

CLEARED

Dec 19, 1979
44 days to decision

K792226 · Product code: **DRT** · Cardiovascular
Source: https://www.510kdatabase.net/k792226/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Cardiac (incl. Cardiotachometer & Rate Alarm) (DRT)
Date received	Nov 5, 1979
Decision date	Dec 19, 1979
Days to decision	44 days
Third-party review	No

APPLICANT

Company	**Paramed Technology, Inc.**
Location	Mchenry, IL, US
510(k) history	3 submissions · 3 cleared · 1979-1989