FDA 510(k) Premarket Notification

K890198 PARAMED MODEL 9350 BLOOD PRESSURE MONITOR (9350)

CLEARED

Jun 12, 1989
145 days to decision

K890198 · Product code: **DXN** · Cardiovascular
Source: https://www.510kdatabase.net/k890198/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Measurement, Blood-pressure, Non-invasive (DXN)
Date received	Jan 18, 1989
Decision date	Jun 12, 1989
Days to decision	145 days
Third-party review	No

APPLICANT

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

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k890198/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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