FDA 510(k) Premarket Notification

K880572 OHMEDA 2350 FINAPRES(R) BLOOD PRESSURE MONITOR

CLEARED

May 10, 1988
89 days to decision

K880572 · Product code: **DXN** · Cardiovascular
Source: https://www.510kdatabase.net/k880572/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Measurement, Blood-pressure, Non-invasive (DXN)
Date received	Feb 11, 1988
Decision date	May 10, 1988
Days to decision	89 days
Third-party review	No

APPLICANT

Company	**Ohmeda Medical**
Location	Madison, WI, US
Contact	DEANA D DICKERSON
510(k) history	120 submissions · 118 cleared · 1984-2013