

K880572 OHMEDA 2350 FINAPRES(R) BLOOD PRESSURE MONITORMay 10, 1988
89 days to decisionK880572 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k880572/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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