

**K911871 CARDIODYNE KINETORR/PC BLOOD PRESSURE MONITORS**Nov 7, 1991  
195 days to decisionK911871 · Product code: **DXN** · Cardiovascular  
Source: <https://www.510kdatabase.net/k911871/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Measurement, Blood-pressure, Non-invasive (DXN)
Date received	Apr 26, 1991
Decision date	Nov 7, 1991
Days to decision	195 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Cardiodyne, Inc.</b>
Location	Mchenry, IL, US
Contact	PATRICK PHILLIPS
510(k) history	3 submissions · 3 cleared · 1977-1991

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k911871/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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