FDA 510(k) Premarket Notification

K911871 CARDIODYNE KINETORR/PC BLOOD PRESSURE MONITORS

CLEARED

Nov 7, 1991
195 days to decision

K911871 · Product code: **DXN** · Cardiovascular
Source: https://www.510kdatabase.net/k911871/

SUBMISSION DETAILS

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

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