

**K792226 CARDIVAN BLOOD PRESSURE MONITOR**Dec 19, 1979  
44 days to decisionK792226 · Product code: **DRT** · CardiovascularSource: <https://www.510kdatabase.net/k792226/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Cardiac (incl. Cardiometer & Rate Alarm) (DRT)
Date received	Nov 5, 1979
Decision date	Dec 19, 1979
Days to decision	44 days
Third-party review	No

**APPLICANT**

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Company	<b>Paramed Technology, Inc.</b>
Location	Mchenry, IL, US
510(k) history	3 submissions · 3 cleared · 1979-1989

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k792226/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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