

**K883109 MINIMON 7136B PATIENT MONITOR**Jan 17, 1989  
176 days to decisionK883109 · Product code: **DRT** · Cardiovascular  
Source: <https://www.510kdatabase.net/k883109/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Cardiac (incl. Cardiotachometer & Rate Alarm) (DRT)
Date received	Jul 25, 1988
Decision date	Jan 17, 1989
Days to decision	176 days
Third-party review	No

**APPLICANT**

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Company	<b>Kontron, Inc.</b>
Location	Everett, MA, US
Contact	DAVID CROMWICK
510(k) history	1 submissions · 1 cleared · 1989-1989

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k883109/>; Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated July 7, 2026