

**K840717 ECS 502**Aug 19, 1985  
549 days to decisionK840717 · Product code: **DRT** · CardiovascularSource: <https://www.510kdatabase.net/k840717/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Cardiac (incl. Cardiotachometer & Rate Alarm) (DRT)
Date received	Feb 17, 1984
Decision date	Aug 19, 1985
Days to decision	549 days
Third-party review	No

**APPLICANT**

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Company	<b>Robert Bosch Corp.</b>
Location	Mchenry, IL, US
510(k) history	21 submissions · 18 cleared · 1979-1986

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

Device record: <https://www.510kdatabase.net/k840717/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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