

**K800500 PULSEOMETER**Apr 8, 1980  
35 days to decisionK800500 · Product code: **DRT** · CardiovascularSource: <https://www.510kdatabase.net/k800500/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Cardiac (incl. Cardiotachometer & Rate Alarm) (DRT)
Date received	Mar 4, 1980
Decision date	Apr 8, 1980
Days to decision	35 days
Third-party review	No

**APPLICANT**

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Company	<b>Kirklands Sporting Goods Co.</b>
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1980-1980

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

Device record: <https://www.510kdatabase.net/k800500/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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