

**K930888 WRISTRECORDER PLUS CARDIAC EVENT  
RECORDING SYSTEM**Jul 21, 1993  
152 days to decisionK930888 · Product code: **DRS** · Cardiovascular  
Source: <https://www.510kdatabase.net/k930888/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Transducer, Blood-pressure, Extravascular (DRS)
Date received	Feb 19, 1993
Decision date	Jul 21, 1993
Days to decision	152 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Ralin Medical, Inc.</b>
Location	Beaverton, OR, US
Contact	DONALD OWEN
510(k) history	2 submissions · 2 cleared · 1993-1994

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k930888/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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