

**K823249 LIFESCOPE II/FOUR**Nov 24, 1982  
23 days to decisionK823249 · Product code: **DRG** · CardiovascularSource: <https://www.510kdatabase.net/k823249/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Transmitters And Receivers, Physiological Signal, Radiofrequency (DRG)
Date received	Nov 1, 1982
Decision date	Nov 24, 1982
Days to decision	23 days
Third-party review	No

**APPLICANT**

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Company	<b>Nihon Kohden America, Inc.</b>
Location	Foothill Ranch, CA, US
510(k) history	166 submissions · 163 cleared · 1979-2012

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

Device record: <https://www.510kdatabase.net/k823249/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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