

**K831274 WLECTROCARDIOGRAPH ECG-7204**Oct 27, 1983  
191 days to decisionK831274 · Product code: **DSI** · Cardiovascular  
Source: <https://www.510kdatabase.net/k831274/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Detector And Alarm, Arrhythmia (DSI)
Date received	Apr 19, 1983
Decision date	Oct 27, 1983
Days to decision	191 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/k831274/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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