

**K820444 HEART-ALERT**Mar 12, 1982  
22 days to decisionK820444 · Product code: **KHX** · Physical MedicineSource: <https://www.510kdatabase.net/k820444/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Platform, Force-measuring (KHX)
Date received	Feb 18, 1982
Decision date	Mar 12, 1982
Days to decision	22 days
Third-party review	No

**APPLICANT**

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Company	<b>Medical Equipment Devices, Inc.</b>
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1982-1982

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

Device record: <https://www.510kdatabase.net/k820444/>; Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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