

**K021325 MODIFICATION OF APEXPRO TELEMETRY SYSTEM**

May 7, 2002  
 11 days to decision

K021325 · Product code: **MHX** · Cardiovascular  
 Source: <https://www.510kdatabase.net/k021325/>

**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Monitor, Physiological, Patient(with Arrhythmia Detection Or Alarms) (MHX)
Date received	Apr 26, 2002
Decision date	May 7, 2002
Days to decision	11 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>General Electric Medical Systems Information Techn</b>
Location	Sugarland, TX, US
Contact	JOELLE NEIDER
510(k) history	33 submissions · 33 cleared · 1999-2002

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

Device record: <https://www.510kdatabase.net/k021325/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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