

**K873214 APX CARDIO ECHOCARDIOGRAPHY SYSTEM #H4300**Jun 1, 1988  
292 days to decisionK873214 · Product code: **DPT** · CardiovascularSource: <https://www.510kdatabase.net/k873214/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Probe, Blood-flow, Extravascular (DPT)
Date received	Aug 14, 1987
Decision date	Jun 1, 1988
Days to decision	292 days
Third-party review	No

**APPLICANT**

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Company	<b>General Electric Co.</b>
Location	Mchenry, IL, US
Contact	LARRY A KROGER
510(k) history	254 submissions · 254 cleared · 1976-2011

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k873214/>; Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated May 17, 2026