

**K854505 CARDIMAX FX-406U**Feb 4, 1986  
84 days to decisionK854505 · Product code: **DRS** · Cardiovascular  
Source: <https://www.510kdatabase.net/k854505/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Transducer, Blood-pressure, Extravascular (DRS)
Date received	Nov 12, 1985
Decision date	Feb 4, 1986
Days to decision	84 days
Third-party review	No

**APPLICANT**

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Company	<b>Brentwood Instruments, Inc.</b>
Location	Mchenry, IL, US
Contact	RICHARD ROSENTHAL
510(k) history	21 submissions · 20 cleared · 1981-1990

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k854505/>; 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 26, 2026