

**K802917 MARK III ULTRASONIC SCANNING SYSTEM**Jan 16, 1981  
58 days to decisionK802917 · Product code: **DXK** · CardiovascularSource: <https://www.510kdatabase.net/k802917/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Echocardiograph (DXK)
Date received	Nov 19, 1980
Decision date	Jan 16, 1981
Days to decision	58 days
Third-party review	No

**APPLICANT**

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Company	<b>Advanced Technology Laboratories, Inc.</b>
Location	Mchenry, IL, US
510(k) history	61 submissions · 59 cleared · 1980-2002

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

Device record: <https://www.510kdatabase.net/k802917/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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