

**K220784 AcQMap High Resolution Imaging and Mapping System**Jul 1, 2022  
106 days to decisionK220784 · Product code: **DQK** · Cardiovascular  
Source: <https://www.510kdatabase.net/k220784/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Computer, Diagnostic, Programmable (DQK)
Date received	Mar 17, 2022
Decision date	Jul 1, 2022
Days to decision	106 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Acutus Medical, Inc.</b>
Location	Carlsbad, CA, US
Contact	Sindhu Sridhar
510(k) history	24 submissions · 24 cleared · 2017-2023

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**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k220784/>. 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 14, 2026