

**K201341 AcQMap 3D Imaging and Mapping Catheter**Sep 15, 2020  
118 days to decisionK201341 · Product code: **MTD** · CardiovascularSource: <https://www.510kdatabase.net/k201341/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intracardiac Mapping, High-density Array (MTD)
Date received	May 20, 2020
Decision date	Sep 15, 2020
Days to decision	118 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	Karla Schaffner
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/k201341/> 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 June 28, 2026