FDA 510(k) Premarket Notification

K170948 AcQMap High Resolution Imaging and Mapping System

CLEARED

Oct 16, 2017
200 days to decision

K170948 · Product code: **DQK** · Cardiovascular
Source: https://www.510kdatabase.net/k170948/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Computer, Diagnostic, Programmable (DQK)
Date received	Mar 30, 2017
Decision date	Oct 16, 2017
Days to decision	200 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Acutus Medical, Inc.**
Location	Carlsbad, CA, US
Contact	Brenda Clay
510(k) history	24 submissions · 24 cleared · 2017-2023