

**K041607 QUINTON Q-TE**Jul 16, 2004  
31 days to decisionK041607 · Product code: **MHX** · CardiovascularSource: <https://www.510kdatabase.net/k041607/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Abbreviated
Device classification	Monitor, Physiological, Patient(with Arrhythmia Detection Or Alarms) (MHX)
Date received	Jun 15, 2004
Decision date	Jul 16, 2004
Days to decision	31 days
Third-party review	No
Summary / Statement	Summary
Other names	RMS

**APPLICANT**

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Company	<b>Quinton Cardiology, Inc.</b>
Location	Bothell, WA, US
Contact	KAREN BROWNE
510(k) history	1 submissions · 1 cleared · 2004-2004

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k041607/>; 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 July 3, 2026