

K090696 NUVANT CARDIAC EVENT MONITORING SYSTEMJun 19, 2009
95 days to decisionK090696 · Product code: **QYX** · CardiovascularSource: <https://www.510kdatabase.net/k090696/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Outpatient Cardiac Telemetry (QYX)
Date received	Mar 16, 2009
Decision date	Jun 19, 2009
Days to decision	95 days
Third-party review	No
Summary / Statement	Summary
Other names	NUVANT MOBILE CARDIAC TELEMETRY SYSTEM

APPLICANT

Company	Corventis, Inc.
Location	San Jose, CA, US
Contact	MADHURI BHAT
510(k) history	7 submissions · 7 cleared · 2009-2014

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k090696/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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