FDA 510(k) Premarket Notification

K090696 NUVANT CARDIAC EVENT MONITORING SYSTEM

CLEARED

Jun 19, 2009
95 days to decision

K090696 · Product code: **QYX** · Cardiovascular
Source: 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