

K915640 VOYAGR SERIES OF DEFLECTABLE DIAGNOSTIC EP CATHETERMar 9, 1992
83 days to decisionK915640 · Product code: **DRF** · Cardiovascular
Source: <https://www.510kdatabase.net/k915640/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Electrode Recording, Or Probe, Electrode Recording (DRF)
Date received	Dec 17, 1991
Decision date	Mar 9, 1992
Days to decision	83 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Arrhythmia Technologies, Inc.
Location	San Jose, CA, US
Contact	GEORGE M SAVAGE
510(k) history	2 submissions · 2 cleared · 1992-1992

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k915640/>. Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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