

K962307 RHYDER DIAGNOSTIC CATHETERSep 17, 1996
92 days to decisionK962307 · Product code: **DRF** · Cardiovascular
Source: <https://www.510kdatabase.net/k962307/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Electrode Recording, Or Probe, Electrode Recording (DRF)
Date received	Jun 17, 1996
Decision date	Sep 17, 1996
Days to decision	92 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Fidus Medical Technology Corp.
Location	Fremont, CA, US
Contact	TED ORMSBY
510(k) history	1 submissions · 1 cleared · 1996-1996

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