

K884320 PROPAQ 106(TM)Dec 27, 1988
75 days to decisionK884320 · Product code: **DRT** · Cardiovascular
Source: <https://www.510kdatabase.net/k884320/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Cardiac (incl. Cardiotachometer & Rate Alarm) (DRT)
Date received	Oct 13, 1988
Decision date	Dec 27, 1988
Days to decision	75 days
Third-party review	No

APPLICANT

Company	Protocol Systems, Inc.
Location	Beaverton, OR, US
Contact	JAMES W SANDBERG
510(k) history	15 submissions · 14 cleared · 1988-2000

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