FDA 510(k) Premarket Notification

K884320 PROPAQ 106(TM)

CLEARED

Dec 27, 1988
75 days to decision

K884320 · 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