FDA 510(k) Premarket Notification

K910882 PROPAQ 102EL, 104EL AND 106EL

CLEARED

May 8, 1991
68 days to decision

K910882 · Product code: **DRT** · Cardiovascular
Source: https://www.510kdatabase.net/k910882/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Cardiac (incl. Cardiotachometer & Rate Alarm) (DRT)
Date received	Mar 1, 1991
Decision date	May 8, 1991
Days to decision	68 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