

**K910882 PROPAQ 102EL, 104EL AND 106EL**May 8, 1991  
68 days to decisionK910882 · Product code: **DRT** · CardiovascularSource: <https://www.510kdatabase.net/k910882/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Cardiac (incl. Cardiometer & Rate Alarm) (DRT)
Date received	Mar 1, 1991
Decision date	May 8, 1991
Days to decision	68 days
Third-party review	No

**APPLICANT**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k910882/>; Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated May 26, 2026