FDA 510(k) Premarket Notification

K012451 PROPAQ ENCORE MODELS 202, 204, 206

CLEARED

Aug 20, 2001
19 days to decision

K012451 · Product code: **DRT** · Cardiovascular
Source: https://www.510kdatabase.net/k012451/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Monitor, Cardiac (incl. Cardiotachometer & Rate Alarm) (DRT)
Date received	Aug 1, 2001
Decision date	Aug 20, 2001
Days to decision	19 days
Third-party review	No
Summary / Statement	Summary
Other names	PROPAQ CS MODELS 242, 244, 246

APPLICANT

Company	**Welch Allyn Protocol, Inc.**
Location	Beaverton, OR, US
Contact	DONALD M ABBEY
510(k) history	7 submissions · 7 cleared · 2001-2007