FDA 510(k) Premarket Notification

K914576 KELLER CARDIAC MONITOR MEDIL KMS870

CLEARED

Dec 30, 1991
76 days to decision

K914576 · Product code: **DRT** · Cardiovascular
Source: https://www.510kdatabase.net/k914576/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Cardiac (incl. Cardiotachometer & Rate Alarm) (DRT)
Date received	Oct 15, 1991
Decision date	Dec 30, 1991
Days to decision	76 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Keller Medical Specialties Products, Inc.**
Location	Antioch, IL, US
Contact	JAY KELLER
510(k) history	6 submissions · 6 cleared · 1990-1999