FDA 510(k) Premarket Notification

K791363 KWIK-AWA VEST

CLEARED

Aug 3, 1979
17 days to decision

K791363 · Product code: **DRG** · Cardiovascular
Source: https://www.510kdatabase.net/k791363/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Transmitters And Receivers, Physiological Signal, Radiofrequency (DRG)
Date received	Jul 17, 1979
Decision date	Aug 3, 1979
Days to decision	17 days
Third-party review	No

APPLICANT

Company	**Medical Specialties, Inc.**
Location	Mchenry, IL, US
510(k) history	40 submissions · 40 cleared · 1979-1995