FDA 510(k) Premarket Notification

K932835 KENDALL T.E.D./SCD SEQUENTIAL COMPRESSION DEVICE

CLEARED

Aug 6, 1993
57 days to decision

K932835 · Product code: **JOW** · Cardiovascular
Source: https://www.510kdatabase.net/k932835/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Sleeve, Limb, Compressible (JOW)
Date received	Jun 10, 1993
Decision date	Aug 6, 1993
Days to decision	57 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Kendall Healthcare Products Co. Div.Of Tyco Health**
Location	Mansfield, MA, US
Contact	John Vozella
510(k) history	66 submissions · 50 cleared · 1989-1997