

**K932835 KENDALL T.E.D./SCD SEQUENTIAL COMPRESSION
DEVICE**Aug 6, 1993
57 days to decisionK932835 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k932835/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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