FDA 510(k) Premarket Notification

K001802 VENODYNE DVT ADVANTAGE MODEL 610

CLEARED

Jan 9, 2001
209 days to decision

K001802 · Product code: **JOW** · Cardiovascular
Source: https://www.510kdatabase.net/k001802/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Sleeve, Limb, Compressible (JOW)
Date received	Jun 14, 2000
Decision date	Jan 9, 2001
Days to decision	209 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Microtek Medical, Inc.**
Location	Mchenry, IL, US
Contact	THOMAS BONNER
510(k) history	24 submissions · 24 cleared · 1984-2011

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k001802/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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