FDA 510(k) Premarket Notification

K890432 DEKNATEL HIGH PRECISION VASCULAR PUNCH(TM)

CLEARED

Apr 13, 1989
76 days to decision

K890432 · Product code: **DWS** · Cardiovascular
Source: https://www.510kdatabase.net/k890432/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Instruments, Surgical, Cardiovascular (DWS)
Date received	Jan 27, 1989
Decision date	Apr 13, 1989
Days to decision	76 days
Third-party review	No

APPLICANT

Company	**Deknatel, Inc.**
Location	Fall River, MA, US
Contact	AMY PETERSON
Website	https://www.teleflex.com
510(k) history	37 submissions · 37 cleared · 1976-1997

Deknatel, Inc. is a medical device manufacturer based in Fall River, US. The company specializes in surgical devices and wound closure solutions. Deknatel received FDA 510(k) clearances from total submissions between 1976 and 1997. The company's cleared devices span multiple surgical specialties, with particular strength in anesthesiology and general surgery. Notable product lines include autotransfusion systems, chest drainage devices, and surgical sutures in various materials and configurations. The company is inactive and represents a historical regulatory record. No F...