

K925046 PLEUR-EVAC CARDIOTOMY RESERVOIRApr 20, 1993
196 days to decisionK925046 · Product code: **DTN** · Cardiovascular
Source: <https://www.510kdatabase.net/k925046/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Reservoir, Blood, Cardiopulmonary Bypass (DTN)
Date received	Oct 6, 1992
Decision date	Apr 20, 1993
Days to decision	196 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Deknatel, Inc.
Location	Fall River, MA, US
Contact	HARRY SAVARD
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...
