

**K926338 PLEUR-EVAC CARDIOTOMY RESERVOIR # CR-9400 & 9400F**Jul 26, 1994  
585 days to decisionK926338 · Product code: **CAC** · Anesthesiology  
Source: <https://www.510kdatabase.net/k926338/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Apparatus, Autotransfusion (CAC)
Date received	Dec 18, 1992
Decision date	Jul 26, 1994
Days to decision	585 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>Deknatel, Inc.</b>
Location	Fall River, MA, US
Contact	HARRY SAVARD
Website	<a href="https://www.teleflex.com">https://www.teleflex.com</a>
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...

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k926338/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated May 25, 2026