

K892915 PLEUR-EVAC A-8002--ADULT/PEDIATRIC DUAL COLLECTIONJun 21, 1989
62 days to decisionK892915 · Product code: **KDQ** · General Hospital
Source: <https://www.510kdatabase.net/k892915/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Bottle, Collection, Vacuum (KDQ)
Date received	Apr 20, 1989
Decision date	Jun 21, 1989
Days to decision	62 days
Third-party review	No

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...

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k892915/> Data sourced from FDA 510(k) public records (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. - Generated May 25, 2026