

**K896697 PLEUR-EVAC(R) A-6020--INFANT SINGLE USE CHEST
DRAI**Feb 8, 1990
72 days to decisionK896697 · Product code: **KDQ** · General Hospital
Source: <https://www.510kdatabase.net/k896697/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Bottle, Collection, Vacuum (KDQ)
Date received	Nov 28, 1989
Decision date	Feb 8, 1990
Days to decision	72 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...

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