

K881252 PLEUR-EVAC(R) A6000 ADULT/PEDIATRIC CHEST DRAINAGEJun 17, 1988
86 days to decisionK881252 · Product code: **GCX** · General Hospital
Source: <https://www.510kdatabase.net/k881252/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Apparatus, Suction, Operating-room, Wall Vacuum Powered (GCX)
Date received	Mar 23, 1988
Decision date	Jun 17, 1988
Days to decision	86 days
Third-party review	No

APPLICANT

Company	Deknatel, Inc.
Location	Fall River, MA, US
Contact	OROFINO
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...
