FDA 510(k) Premarket Notification

CLEARED

K962123 AKROTECH 4000T KINETIC TURNING LOW AIR LOSS SYSTEM

Jul 18, 1996
48 days to decision

K962123 · Product code: **FNM** · General Hospital
Source: https://www.510kdatabase.net/k962123/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Mattress, Air Flotation, Alternating Pressure (FNM)
Date received	May 31, 1996
Decision date	Jul 18, 1996
Days to decision	48 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Lumex, Inc.**
Location	Bay Shore, NY, US
Contact	JOE Z ANETTI
510(k) history	8 submissions · 8 cleared · 1990-1997