

**K962123 AKROTECH 4000T KINETIC TURNING LOW AIR LOSS SYSTEM**Jul 18, 1996  
48 days to decisionK962123 · 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	<b>Lumex, Inc.</b>
Location	Bay Shore, NY, US
Contact	JOE Z ANETTI
510(k) history	8 submissions · 8 cleared · 1990-1997

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k962123/> 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 June 29, 2026