FDA 510(k) Premarket Notification

K953503 PRESSUREGUARD IV

CLEARED

Oct 24, 1995
90 days to decision

K953503 · Product code: **FNM** · General Hospital
Source: https://www.510kdatabase.net/k953503/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Mattress, Air Flotation, Alternating Pressure (FNM)
Date received	Jul 26, 1995
Decision date	Oct 24, 1995
Days to decision	90 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Span-America Medical Systems, Inc.**
Location	Mchenry, IL, US
Contact	WANDA TOTTON
510(k) history	19 submissions · 19 cleared · 1980-1998

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k953503/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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