FDA 510(k) Premarket Notification

K833964 PEGASUS MATTRESS

CLEARED

Jan 30, 1984
75 days to decision

K833964 · Product code: **FNM** · General Hospital
Source: https://www.510kdatabase.net/k833964/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Mattress, Air Flotation, Alternating Pressure (FNM)
Date received	Nov 16, 1983
Decision date	Jan 30, 1984
Days to decision	75 days
Third-party review	No

APPLICANT

Company	**Dermalex Co.**
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1984-1984