

K833964 PEGASUS MATTRESSJan 30, 1984
75 days to decisionK833964 · 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

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Device record: <https://www.510kdatabase.net/k833964/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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