FDA 510(k) Premarket Notification

K961854 SPR PLUS II OVERLAY SYSTEM (CL250/CL212

CLEARED

Aug 9, 1996
87 days to decision

K961854 · Product code: **FNM** · General Hospital
Source: https://www.510kdatabase.net/k961854/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Mattress, Air Flotation, Alternating Pressure (FNM)
Date received	May 14, 1996
Decision date	Aug 9, 1996
Days to decision	87 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Gaymar Industries, Inc.**
Location	Mchenry, IL, US
Contact	PETER SCOTT
Website	https://www.gaymar.com
510(k) history	27 submissions · 27 cleared · 1976-2011